

Permanent Dissipative Structures in Water: The Matrix of Life?

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PROMETE SRL – CNR SPIN OFF, NAPOLI, ITALY



promete

Meeting in memory of Mae-Wan Ho

15-16 June, 2017

London Mathematical Society,

De Morgan House, 57-58 Russell Square,

London WC1B 4HS

Permanent Dissipative Structures in Water: The Matrix of Life?

There is now clear evidence that various kinds of low energy physical perturbations can induce the spontaneous formation of dissipative structures in liquid water.

It has recently been discovered that these structures are remarkably persistent even in the solid phase.

Using easily reproducible experimental methods, we have observed large quantities of supramolecular aggregates of water.

Are such structures the matrix of life itself?



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**_environment &
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3

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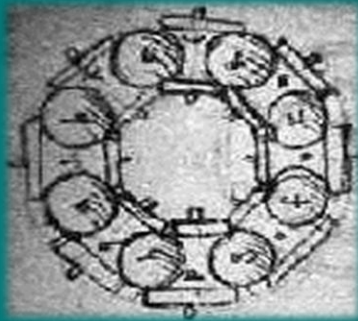
**_technology
transfer**

**CORE
BUSINESS AREAS
OF GROWTH**

An example of missing Technology Transfer ...



1500



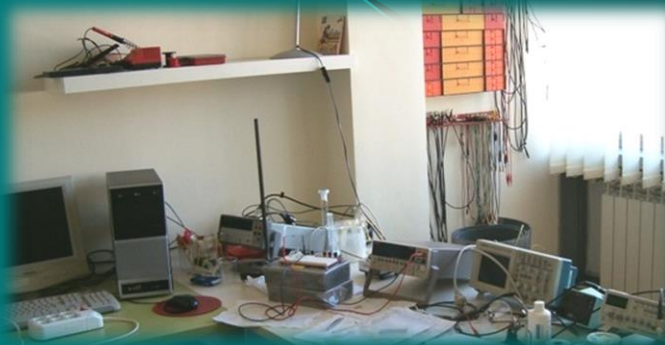
400 years ...

1900



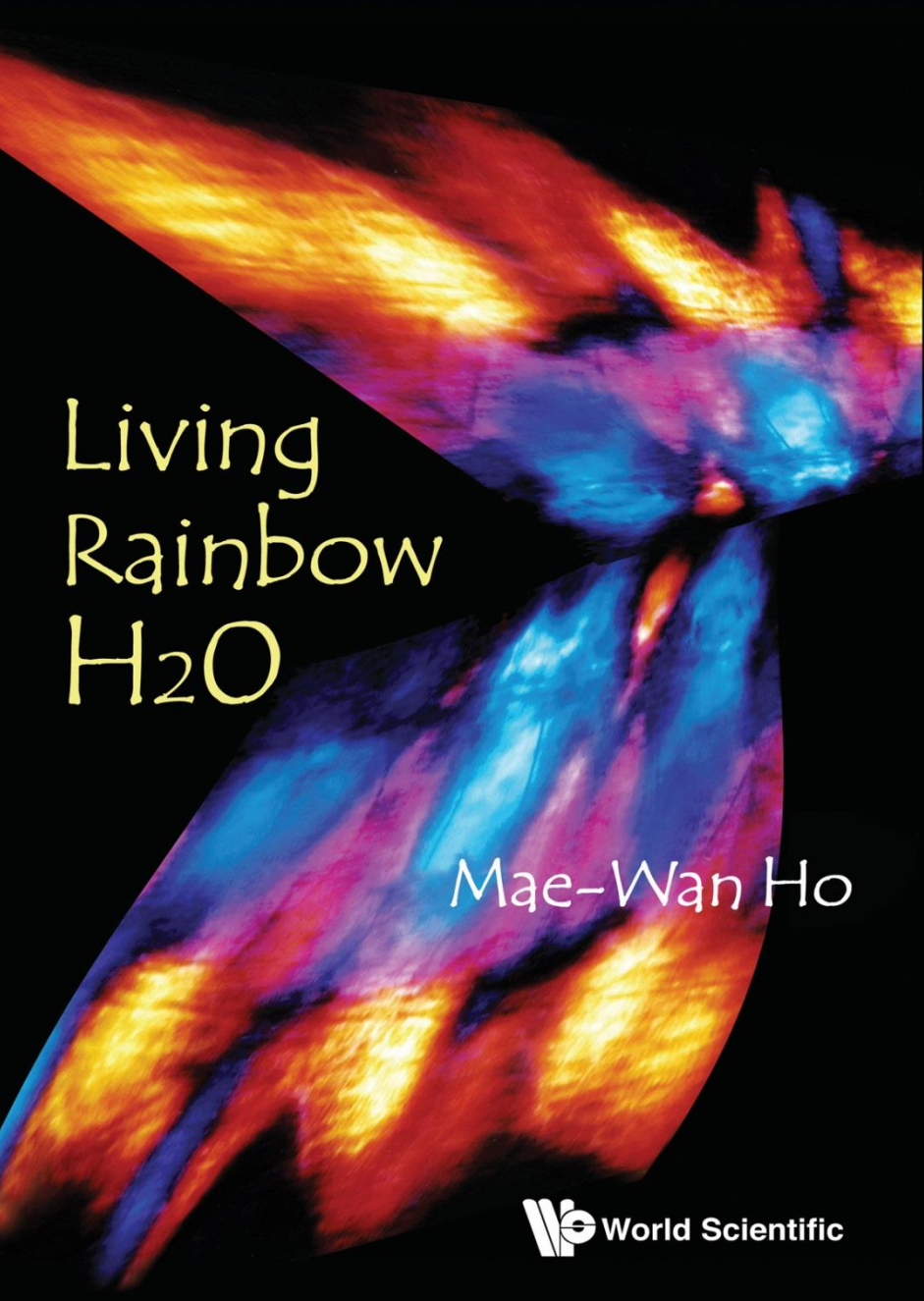


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Living Rainbow H₂O

Mae-Wan Ho

*“Water is the means,
medium and message of life,
the rainbow within that
mirrors the one in the sky”*

*A unique synthesis of the latest
quantum physics and chemistry
of water that tells you why it is
so fit for life*

*Sequel to The Rainbow Worm
in the author’s quest for the
meaning of life*

 World Scientific

le connes
inattese sioni

LA FRONTIERA, L'IGNOTO, L'ERESIA

Napoli, Nov 2011

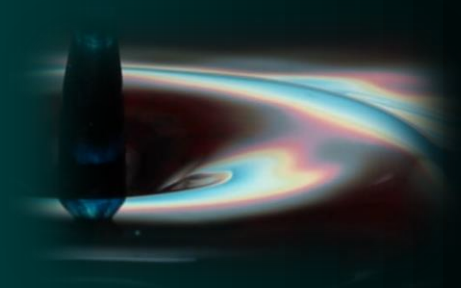
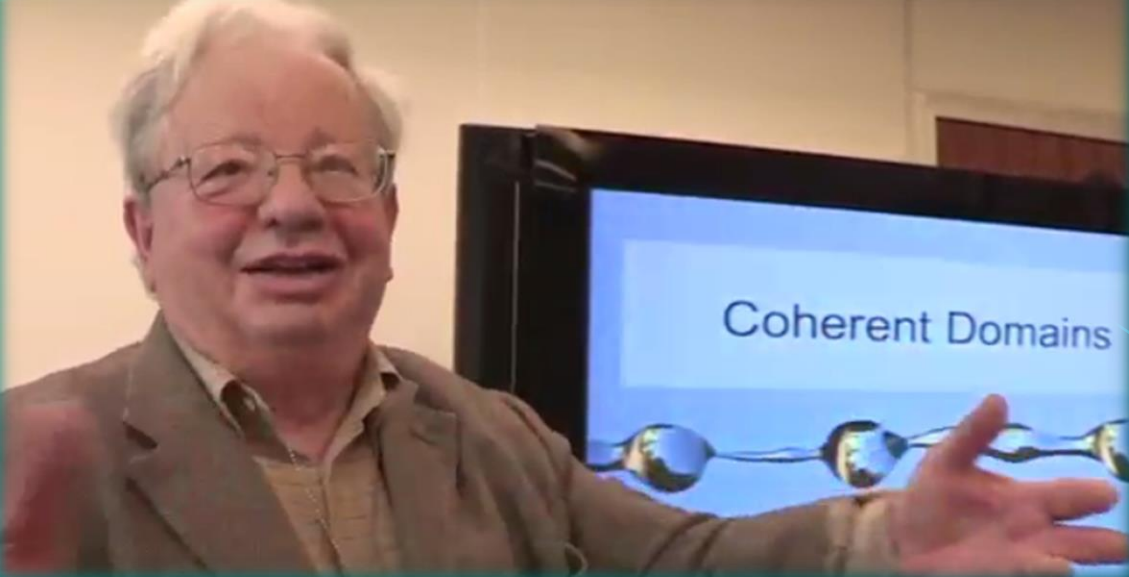
**Istituto Italiano
per gli Studi Filosofici**



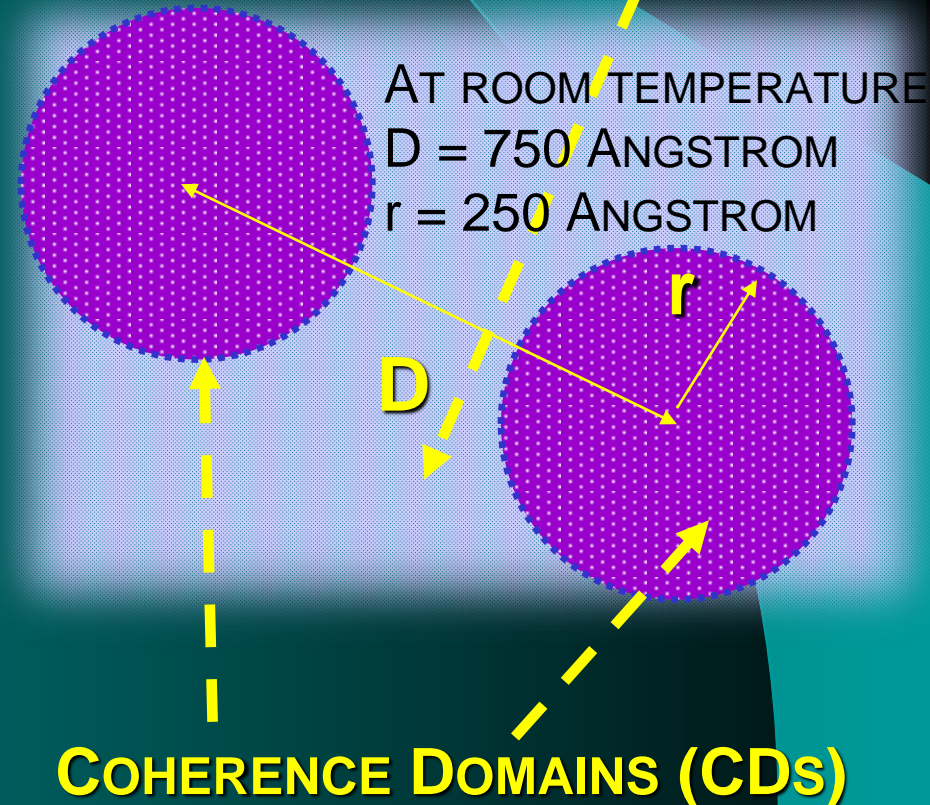
Roberto Germano (primo a destra) e Mae-Wan Ho (terza da destra)







NON-COHERENT PHASE











Chiral micron-sized H₂O aggregates in water: Circular dichroism of supramolecular H₂O architectures created by perturbing pure water

Vittorio Elia^{a,2}, Tamar A. Yinnon^b, Rosario Oliva^a, Elena Napoli^a,
Roberto Germano^c, Fabrizio Bobba^d, Angela Amoresano^a

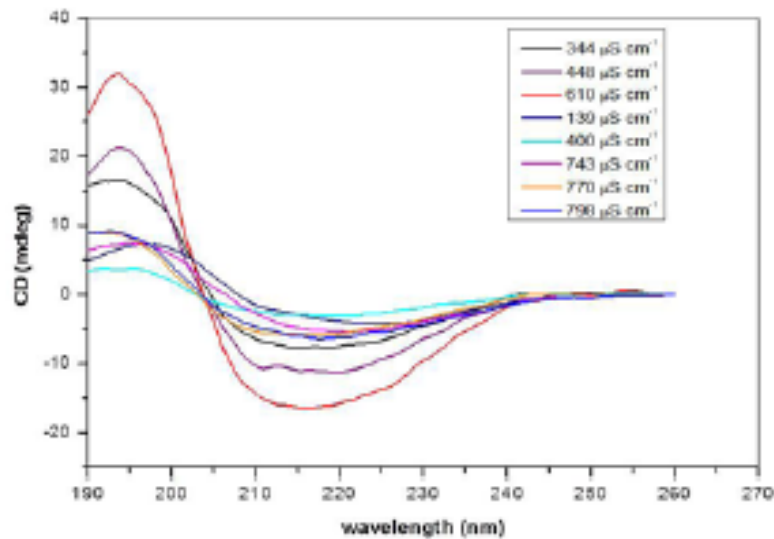


Figure 4. CD spectra of INW samples with different electric conductivity. The CD in mdeg is plotted as a function of the wavelength in nm for samples with electric conductivity as specified in the inset. The temperature of the samples is 20 °C.

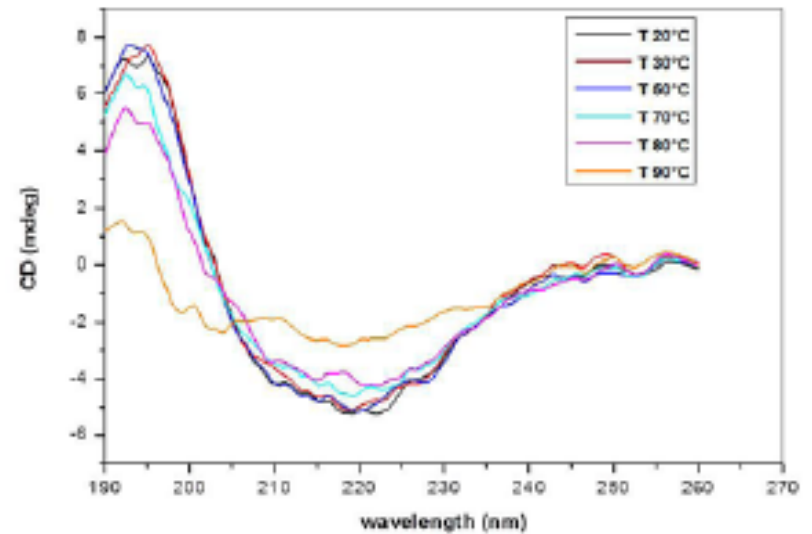
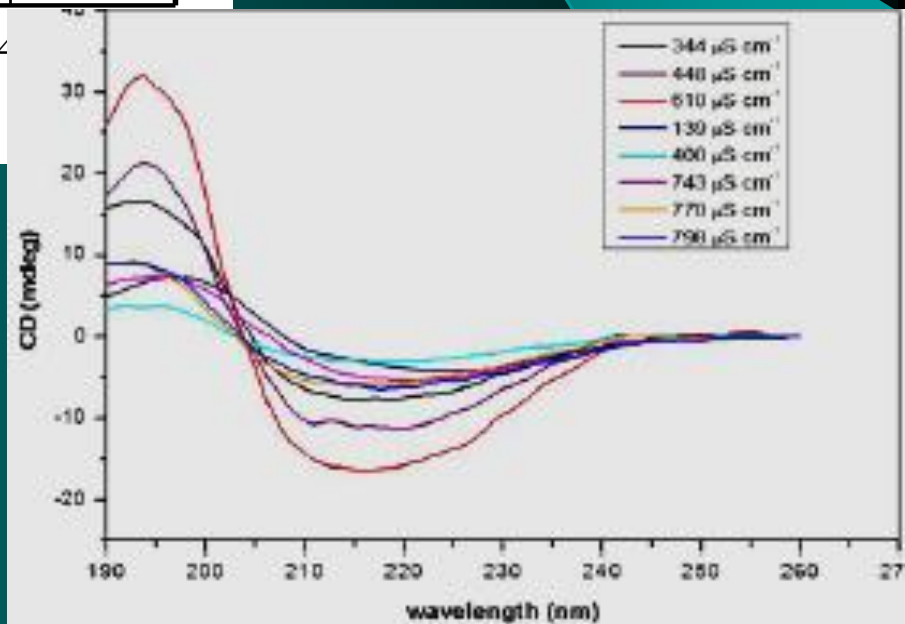
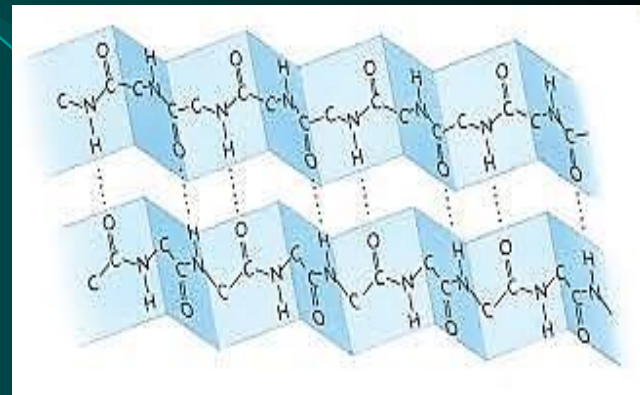
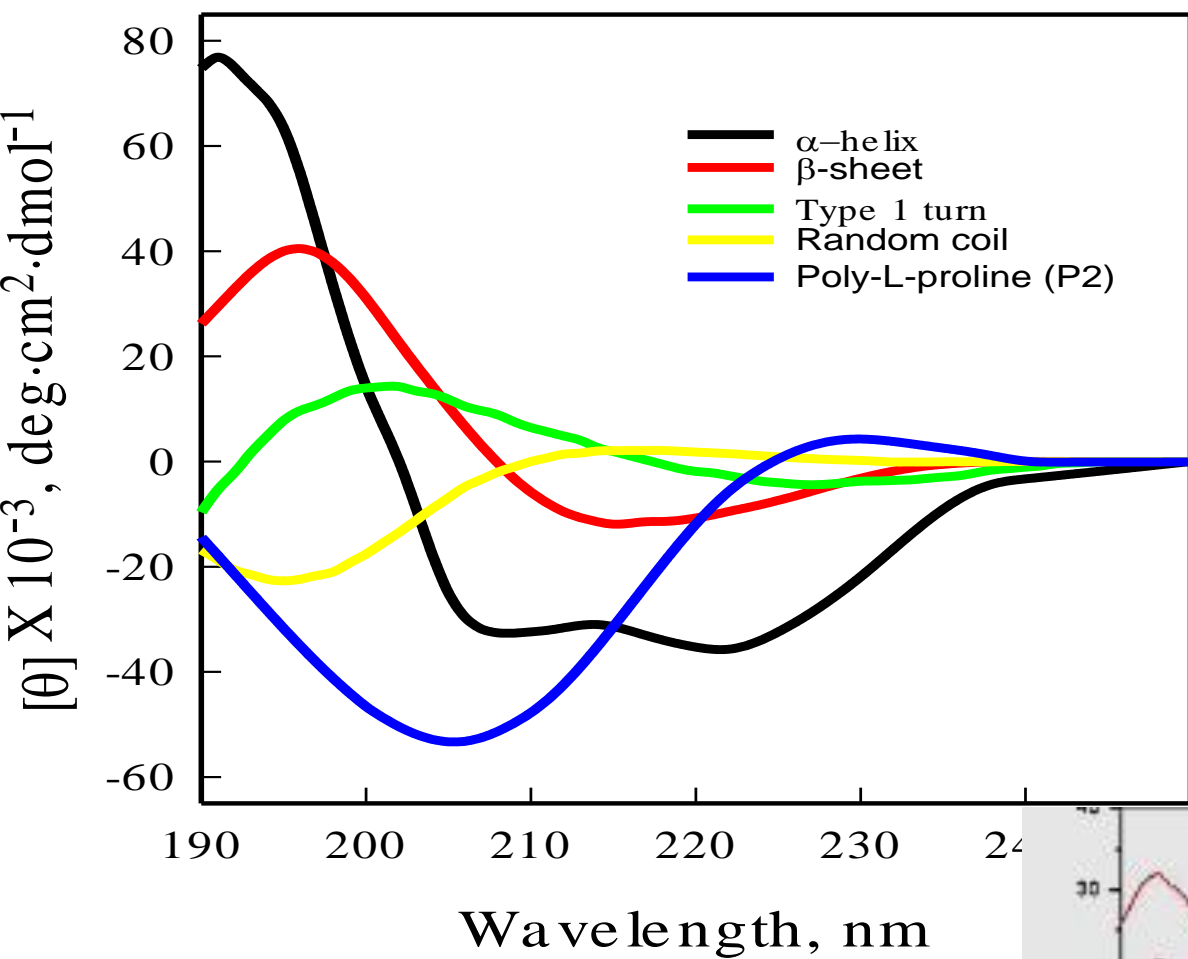


Figure 5. CD spectra of an INW sample at different temperatures. The CD in mdeg is plotted as a function of the wavelength in nm for a sample



UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H₂O ARCHITECTURES GENERATED BY PERTURBING PURE WATER

Authors

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Elena Napoli^a, Roberto Germano^c, Fabrizio Bobba^d,
Angela Amoresano^a

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**UV-FLUORESCENCE BEHAVIOR OF
SUPRAMOLECULAR H_2O ARCHITECTURES
GENERATED BY PERTURBING PURE WATER**

MICRON-SIZED SUPRAMOLECULAR H_2O AGGREGATES ARE PRODUCIBLE BY PERTURBING PURE WATER, I.E., IMMERSING A HYDROPHILIC MEMBRANE IN MILLI-Q WATER, STIRRING THE LIQUID, REMOVING AND DRYING THE MEMBRANE, AND THEN REPEATING THESE STEPS (ITERATIVELY NAFIONATED WATER).

INW SHOWS A FLUORESCENCE SPECTRA CONSISTING OF A BROAD BAND SPANNING FROM 290 TO 550 NM.

ON DILUTION OF THE SAMPLE, THE FLUORESCENCE INTENSITY DIMINISHES, AND THIS EMISSION IS ABSENT IN MILLI-Q WATER. THE ABSENCE OF ANY SHARP EMISSION PEAKS AND THE SMEARED OUT CHARACTER OF THE SPECTRA INDICATE PRESENCE OF EXCIMERS.

UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H₂O ARCHITECTURES GENERATED BY PERTURBING PURE WATER

TO ENSURE THAT IMPURITIES (MOLECULES RELEASED BY THE MEMBRANE, ORGANIC- OR BIO-CONTAMINANTS) DO NOT UNDERLIE AFOREMENTIONED PHENOMENA, WE EMPLOYED STATE-OF-THE-ART ANALYTIC TECHNIQUES:

MATRIX-ASSISTED LASER DESORPTION/IONIZATION TIME OF FLIGHT AND GAS CHROMATOGRAPHY BOTH COUPLED TO MASS SPECTROSCOPY, AND ION CHROMATOGRAPHY.

THE PERTURBED WATER ONLY CONTAINS 10⁻⁶ M FLUORINE AND SULFATE IONS RELEASED BY THE MEMBRANE AND MINUSCULE AMOUNTS OF CONTAMINANTS, BUT LYOPHILIZING THE INW PERTURBED WATER PRODUCES A PONDERAL SOLID RESIDUE!!

PURE WATER (BIDISTILLED) →

PHYSICAL PERTURBATIONS →

→ LARGE CHANGES IN PHYSICO-CHEMICAL PARAMETERS !!!
(pH, HEAT OF MIXING, ELECTRICAL CONDUCTIVITY, VISCOSITY...)

→ ANALYSES

→ IT IS STILL ...PURE WATER!!



BUT

→ BUT

→ CIRCULAR DICHROISM: MACROMOLECULES!!!

AND

→ UV FLUORESCENCE : ELECTRONIC ENERGETIC TRANSITION
(280 NM)



from a 'non-bonding' (lone-pair) n orbital to

an 'antibonding' π orbital designated as π^* .

NON EXISTENT IN H₂O MOLECULES!!!!!!

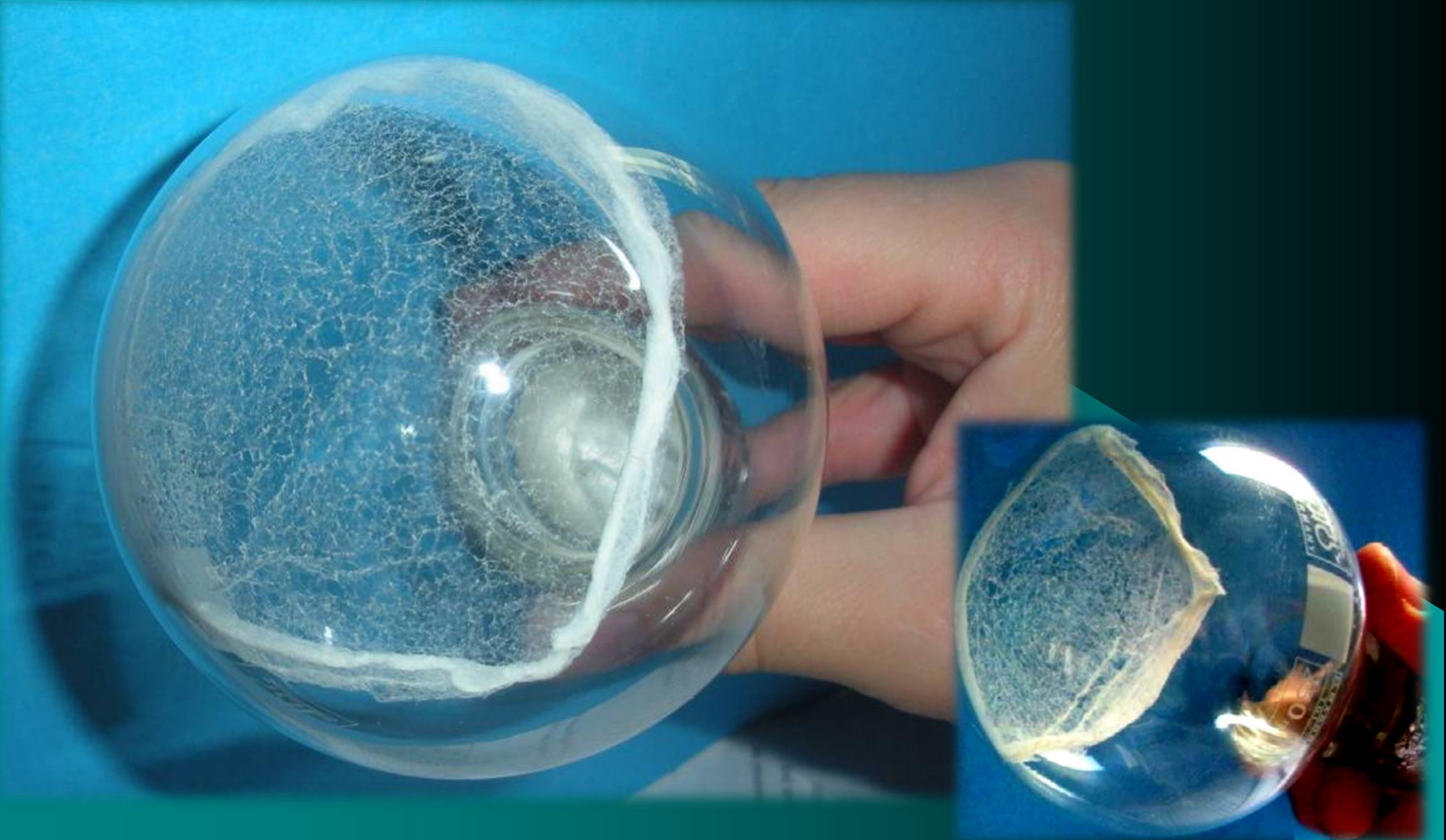
MOREOVER...

MOREOVER

IF DRYING (OR FREEZEDRYING) →
A SOLID !!!!

**STABLE AT AMBIENT TEMPERATURE
AND PRESSURE**





APPEARANCE OF THE EVIDENTLY PONDERAL SOLID RESIDUES: SOME TEN OF MILLIGRAMS COMING FROM LYOPHILIZATION OF ONLY 250 ML OF ITERATIVELY NAFIONATED WATER (IFW) → VITTORIO ELIA

Water with a Nafion membrane

I. Wash membrane in Milli-Q water.

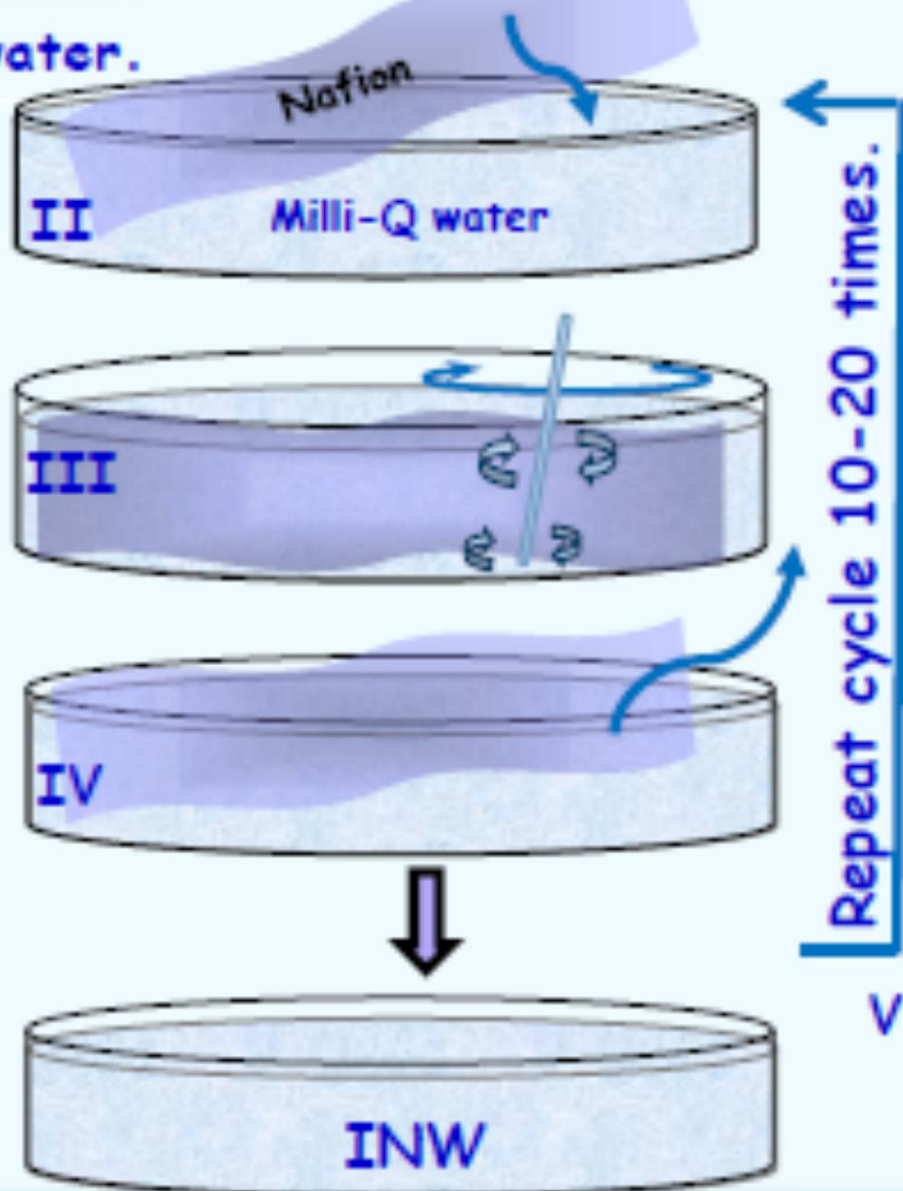
II. Immerse membrane.

III. Gently stir liquid
and turn over membrane.

IV. Take out membrane
and dry it in air.

V. Repeat steps II-IV
10-20 times.

Iterative Nafionized water
(INW)

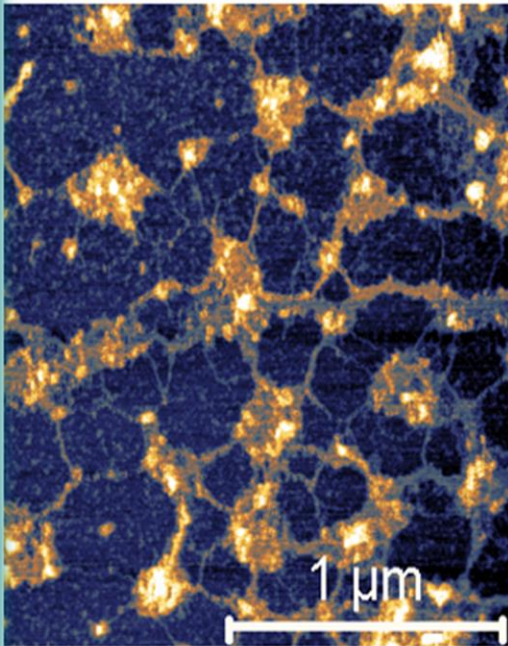


XEROSYDRYLE

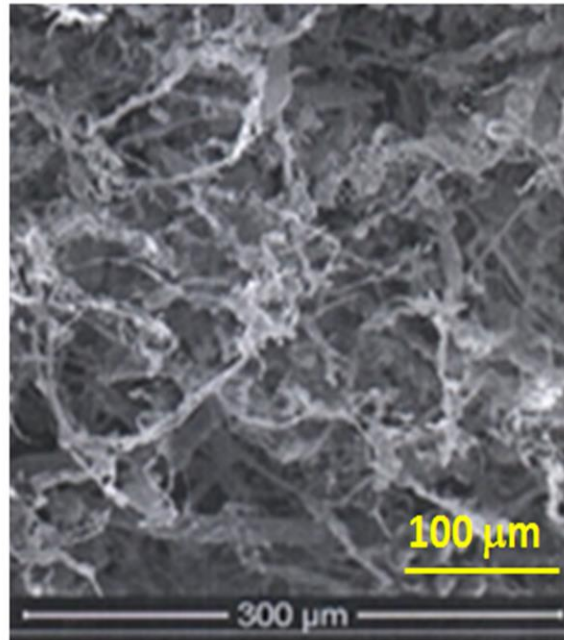
FROM ANCIENT GREEK:

XEROS (DRY) - IDRO (WATER) - YLE (MATTER)

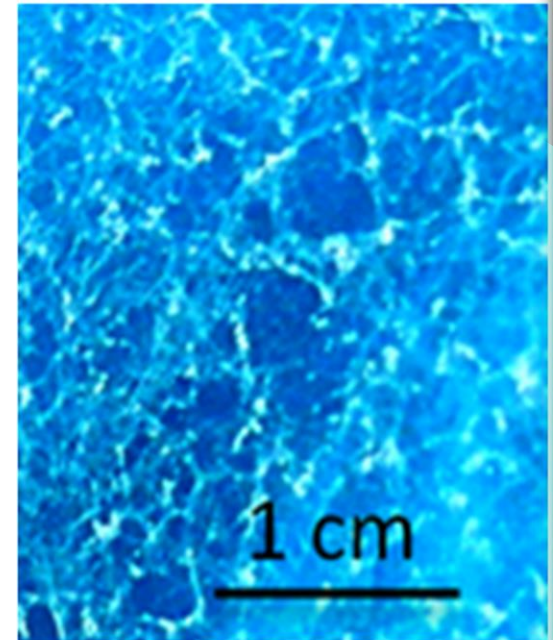
XEROSYDRYLE FRACTAL APPEARANCE



10^{-6} m

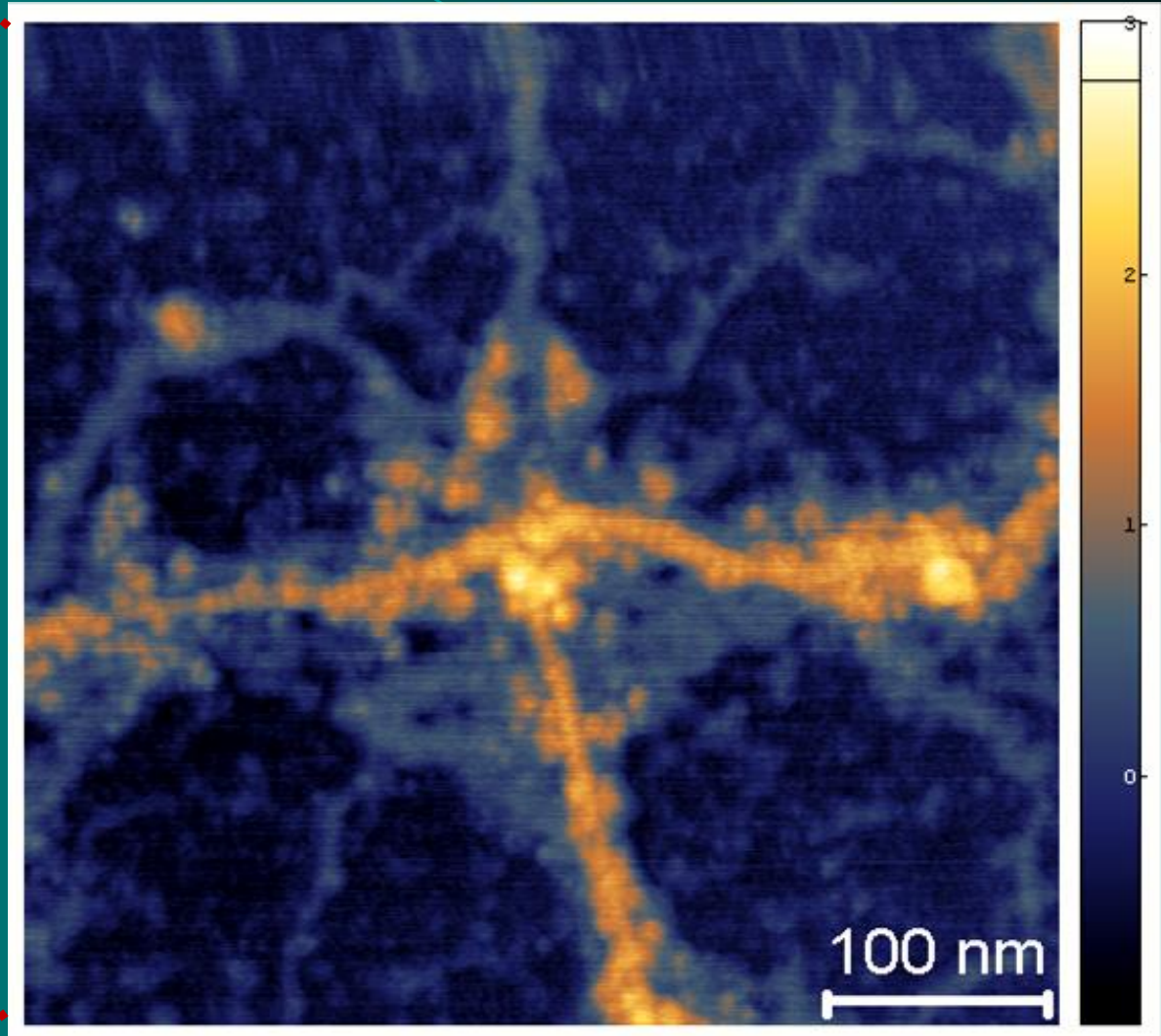
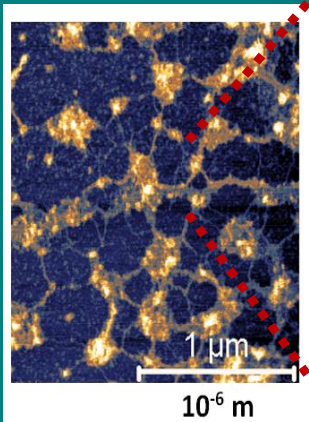


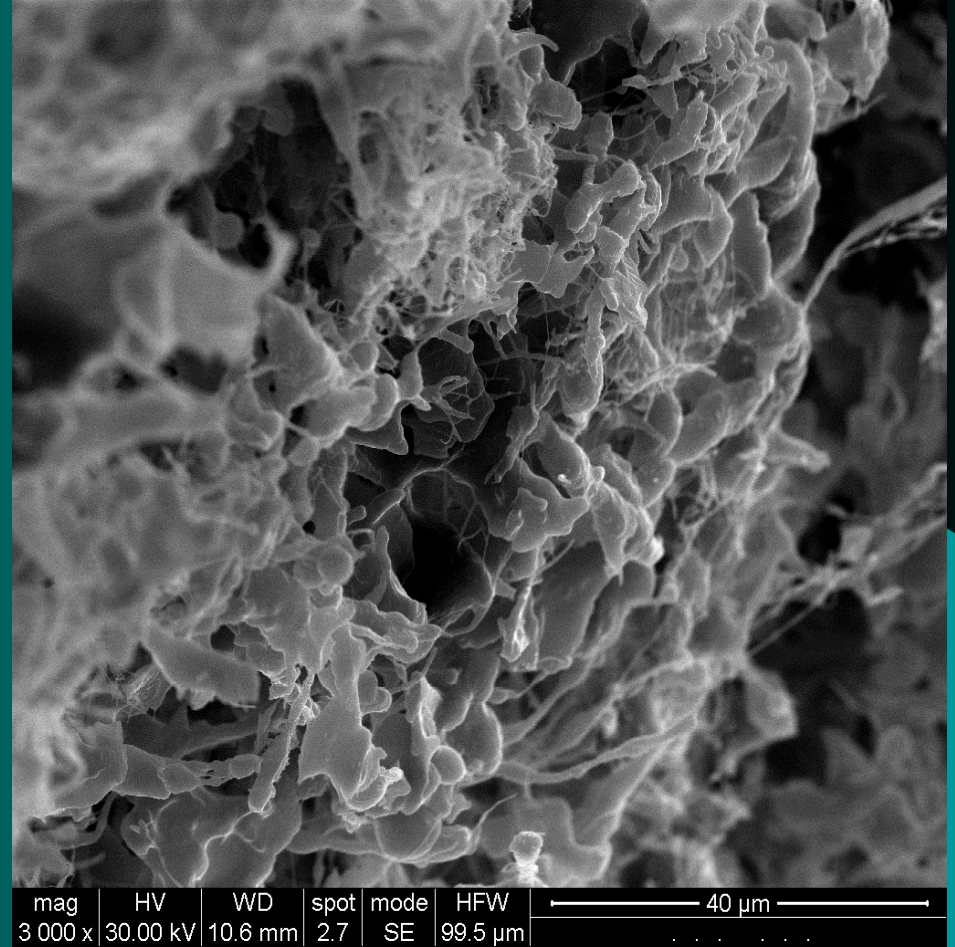
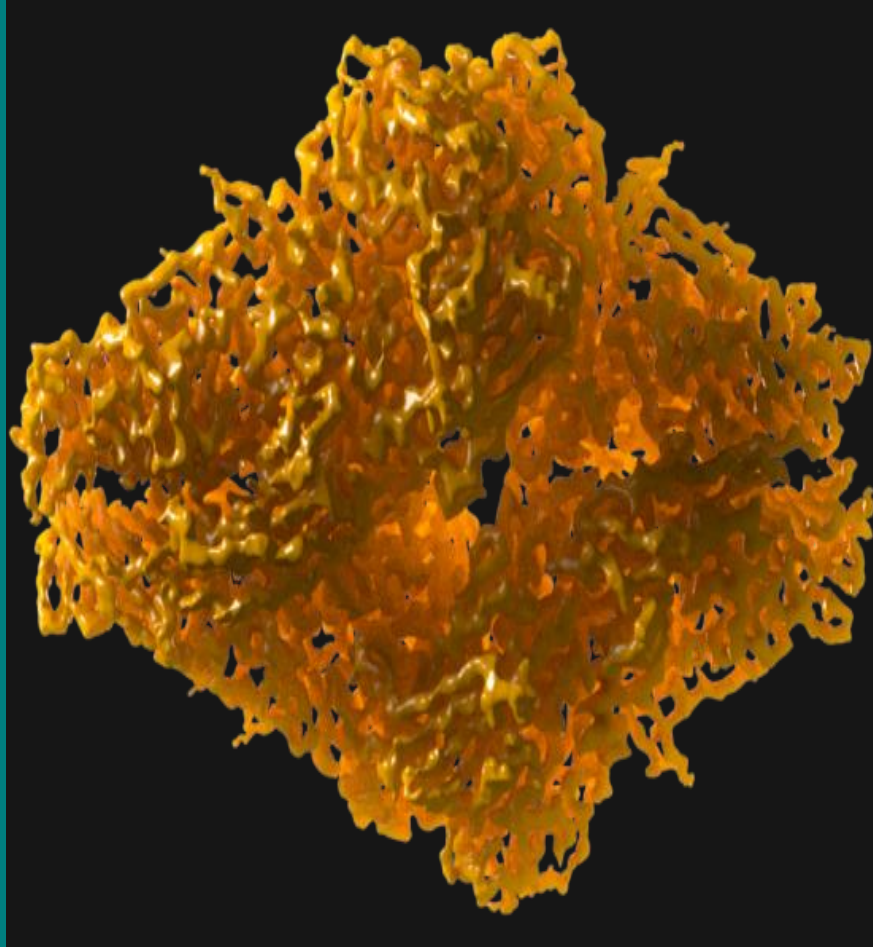
10^{-4} m



10^{-2} m

**(AFM) THE MINIMUM MEASURED HEIGHT OF FIBRILS IS ABOUT 1NM,
WHEREAS THEIR LENGTHS ARE OF HUNDREDS OF NANOMETRES**



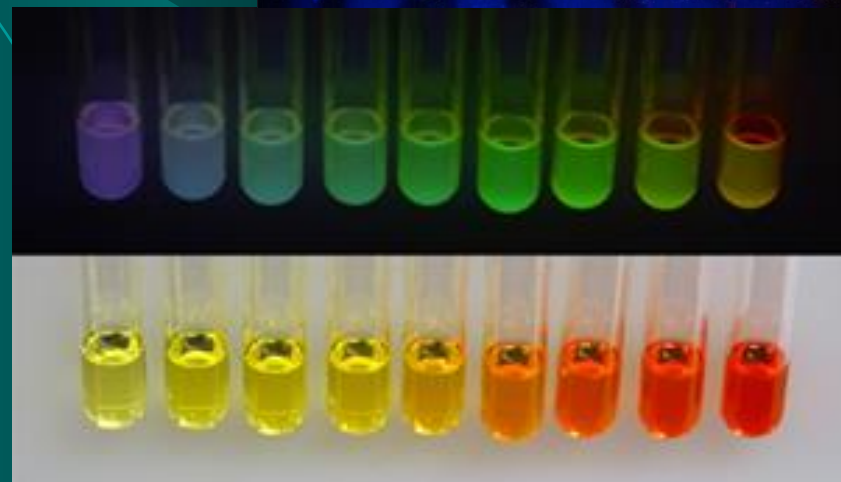
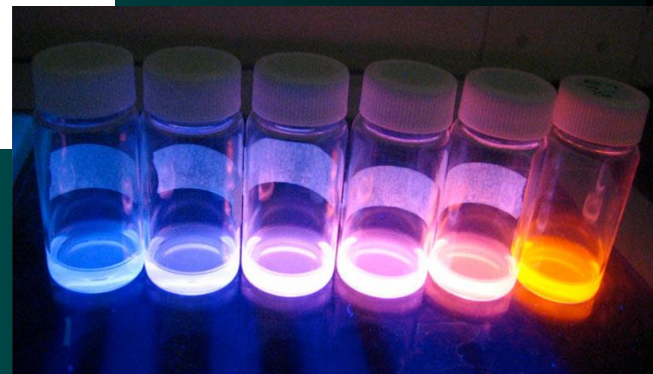
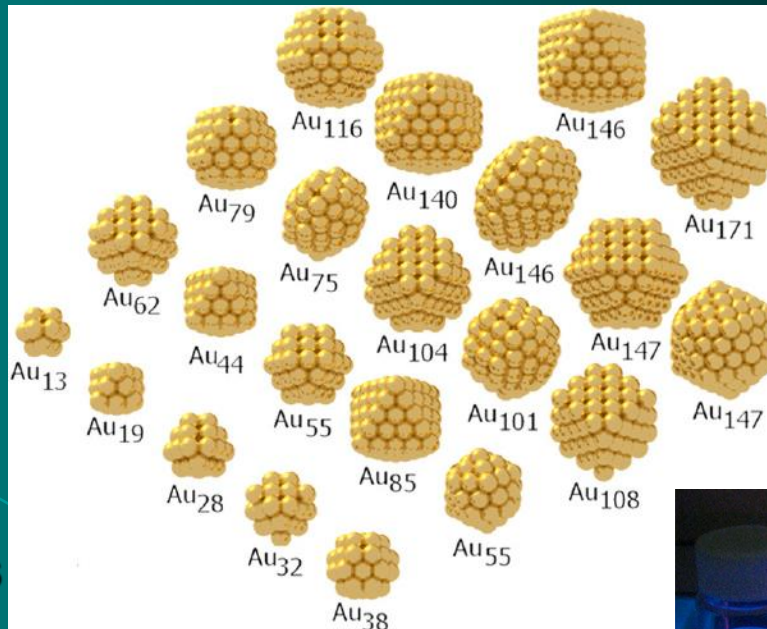


NO WONDER IF ...

THE GOLD NANOPARTICLES

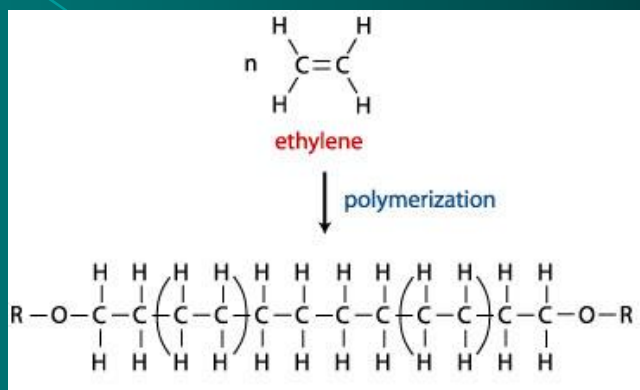
BEHAVE DIFFERENTLY ... EVEN FLUORESCENT...

FROM BULK GOLD



NO WONDER IF...

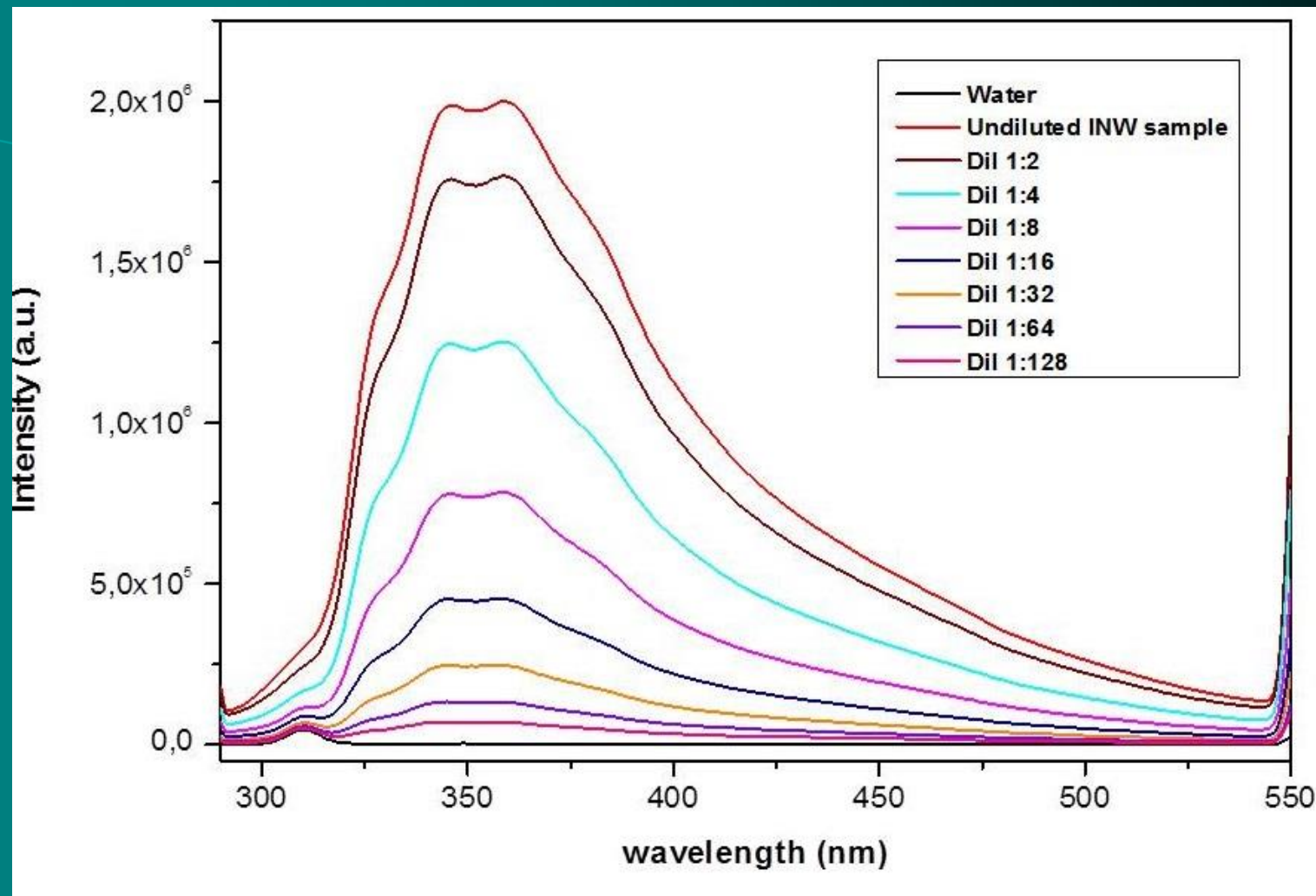
GAS ETHYLENE →



POLYMERIC SOLID POLYETHYLENE !!!!

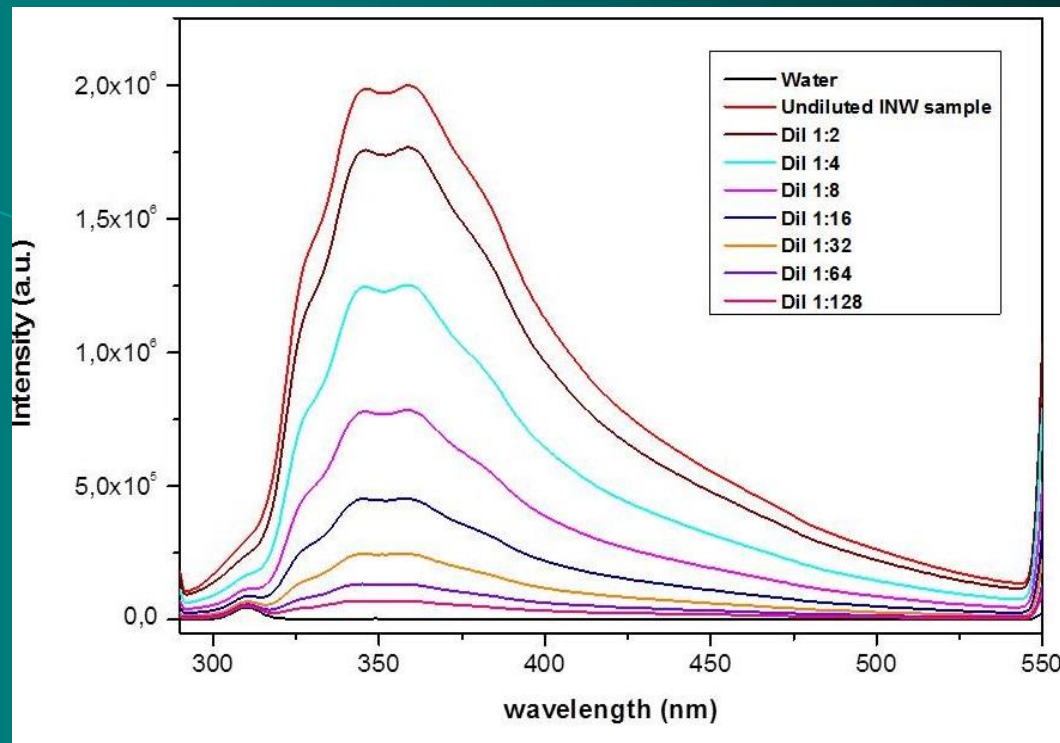


UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H_2O ARCHITECTURES GENERATED BY PERTURBING PURE WATER



THE FIGURE PRESENTS THE UV-VISIBLE (UV-VIS) FLUORESCENCE SPECTRA OF AN INW SAMPLE, OF ITS SERIAL DILUTED LIQUIDS AND OF THE CONTROL MILLI-Q WATER. THE SPECTRA RESULT FROM EXCITATION OF THE LIQUIDS WITH 280 NM RADIATION.

UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H_2O ARCHITECTURES GENERATED BY PERTURBING PURE WATER



THE FLUORESCENCE INTENSITIES ARE PLOTTED AS FUNCTION OF WAVELENGTH IN nm FOR SAMPLES KEPT AT 25 °C.

THE RED LINE PRESENTS THE FLUORESCENCE OF AN INW SAMPLE WITH $X_{INW}=1300 \mu\text{Scm}^{-1}$

THE COLORED LINES PRESENT THE FLUORESCENCE OF SAMPLES OBTAINED BY SUCCESSIVE DILUTIONS OF THIS $X_{INW}=1300 \mu\text{Scm}^{-1}$ SAMPLE.

THE SAMPLE WAS DILUTED WITH MILLI-Q WATER.

THE DILUTION RATIO (DIL) IS SPECIFIED IN THE INSET.

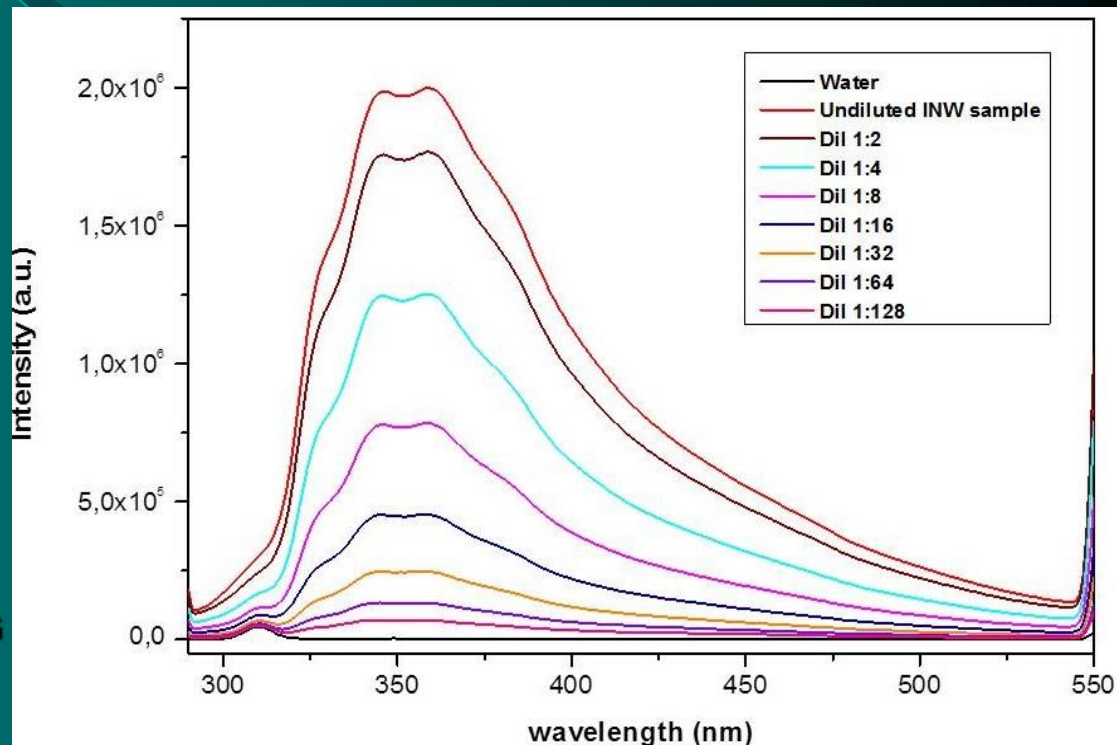
THE BLACK LINE PRESENTS THE FLUORESCENCE OF THE CONTROL, I.E., MILLI-Q WATER.

UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H₂O ARCHITECTURES GENERATED BY PERTURBING PURE WATER

**INW HAS A DISTINCTIVE ABSORPTION PEAK IN THE 250 - 320 nm RANGE
WITH MAXIMUM AT 280 nm.**

THE PEAK IS ABSENT IN BULK WATER.

**INW'S FLUORESCENCE SPECTRA
CONSIST OF A BROAD BAND SPANNING
FROM 290 TO 550 nm.**



UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H_2O ARCHITECTURES GENERATED BY PERTURBING PURE WATER

THE UV ABSORBANCE AND FLUORESCENCE SPECTRA OF INW RESEMBLE THOSE OF INTERFACIAL WATER ADJACENT TO NAFION MEMBRANES (WAN) (POLLACK)

THE UV ABSORBANCE OF INW MOST CLOSELY RESEMBLES THAT OF WAN AT A DISTANCE OF ABOUT 450 MICRON FROM THE NAFION MEMBRANE (WAN-450 nm).

IN PARTICULAR, INW AND WAN-450 HAVE IN COMMON THE STRONG ABSORBANCE ENHANCEMENT WHEN THE WAVELENGTH DECREASES FROM 240 TO 200 nm.

EXCIMER-TYPE UV-VIS FLUORESCENCE BANDS HAVE BEEN OBSERVED FOR WATER-ETHANOL MIXTURES AND VARIOUS AQUEOUS SOLUTIONS, E.G., SOLUTIONS OF LITHIUM CHLORIDE OR SUCROSE.

THE UV ABSORBANCE IN THE 240 - 200 nm RANGE HAS NOT YET BEEN CLARIFIED FOR WAN OR OTHER STRUCTURED WATERS.

ANALOGOUS ABSORBANCE HAS BEEN OBSERVED FOR MAGNETIZED WATER SUGGESTING THAT IT IS RELATED TO THE DYNAMICS OF ELECTRONS IN WATER.

UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H₂O ARCHITECTURES GENERATED BY PERTURBING PURE WATER

ANALYSES (I)

WE SHOWED THAT PERTURBING PURE WATER WITH A NAFION MEMBRANE AND STIRRING LEADS TO UV ABSORBANCE AND FLUORESCENCE SPECTRA ATTRIBUTABLE TO MICRON-SIZED SUPRAMOLECULAR H₂O AGGREGATES.

THESE AGGREGATES FORMED DURING THE LIQUID-LIQUID TRANSITION TRIGGERED BY THE PERTURBATIONS.

TO SPOT CONTAMINANTS AND ELIMINATE SUSPICION THAT THESE UNDERLIE OUR FINDINGS, WE CARRIED OUT EXTENSIVE ANALYSES.

OUR MASS SPECTROSCOPY ANALYSES EVINCE THAT THE AGGREGATES ARE COMPOSED OF H₂O.

UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H₂O ARCHITECTURES GENERATED BY PERTURBING PURE WATER

ANALYSES (II)

OUR MALDI-TOF/MASS SPECTROSCOPY

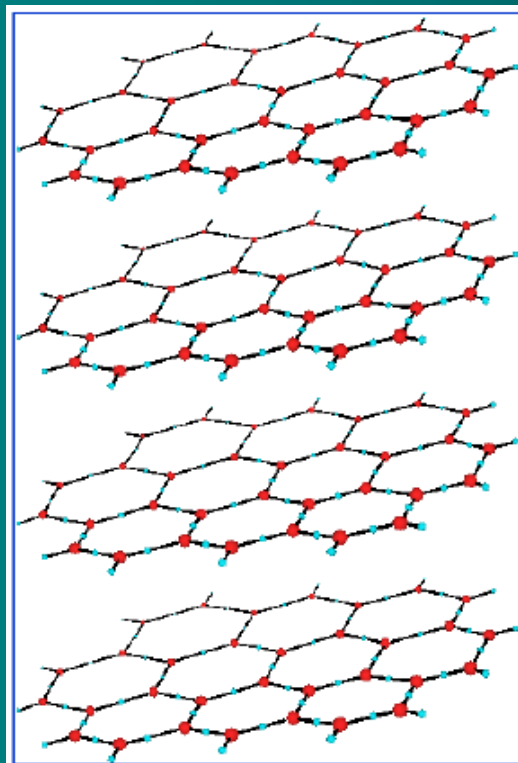
MATRIX-ASSISTED LASER DESORPTION/IONIZATION – TIME OF FLIGHT

AND GAS CHROMATOGRAPHY/MASS SPECTROSCOPY

ANALYSES SHOW THAT ONLY MINISCULE AMOUNTS OF ORGANIC OR BIOLOGICAL CONTAMINANTS ARE PRESENT IN THE PERTURBED WATER. THESE CONTAMINANTS CANNOT ACCOUNT FOR FORMATION OF THE MICRON-SIZED AGGREGATES.

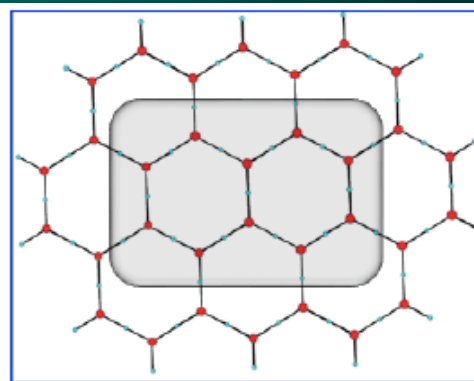
OUR ION CHROMATOGRAPHY ANALYSES SHOW THAT THE SOLE INORGANIC MOLECULES RELEASED BY THE NAFION MEMBRANES ARE FLUORINE AND SULFATE IONS. IN THE PERTURBED WATER, THESE IONS ARE PRESENT AT CONCENTRATIONS OF ABOUT 10^{-6} MOL / L. THESE IONS CANNOT ACCOUNT FOR FORMATION OF ITS MICRON-SIZED AGGREGATES, ITS HIGH ELECTRIC CONDUCTIVITY OR LOW PH.

**(A) SCHEMATIC REPRESENTATION OF THE
MULTILAYER HONEYCOMB ICE-LIKE MODEL.**

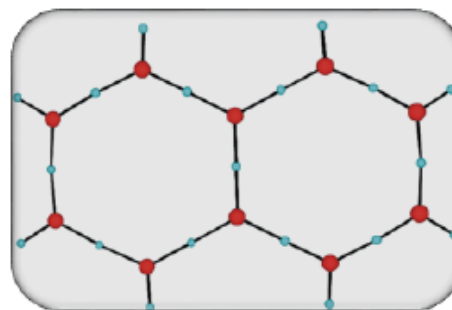


(a)

(B) A TEN-HEXAGON LAYER.



(b)



(c)

**(C) THE $H_{19}O_{10}$ SYSTEM COMPOSED BY TWO FUSED WATER HEXAGONS IS THE MONOMER
USED IN THE THEORETICAL STUDY:**

SEGARRA-MARTÍ J, ROCA-SANJUÁN D, MERCHÁN M (2014)

**CAN THE HEXAGONAL ICE-LIKE MODEL RENDER THE SPECTROSCOPIC
FINGERPRINTS OF STRUCTURED WATER? FEEDBACK FROM QUANTUM-
CHEMICAL COMPUTATIONS. *ENTROPY* 16 (7) : 4101-4120.**

WHEN TWO $[\text{H}_{19}\text{O}_{10}]^-$ ANIONIC SYSTEMS ARE SITUATED IN A PARALLEL ORIENTATION AT 2.4 Å WITHIN THE CONSTRAINTS OF $\text{D}_{2\text{H}}$ SYMMETRY (INPUT), GEOMETRY OPTIMIZATION OF THE RESULTING GLOBAL SYSTEM $[\text{H}_{38}\text{O}_{20}]^{2-}$ LEADS, AFTER A LONG OPTIMIZATION PROCESS WHERE THE ORIGINAL SPATIAL SYMMETRY IS LOST AND BREAKING OF THE WAVE FUNCTION OCCURS, TO THE BULK-TYPE WATER AGGREGATE DISPLAYED IN OUTPUT.

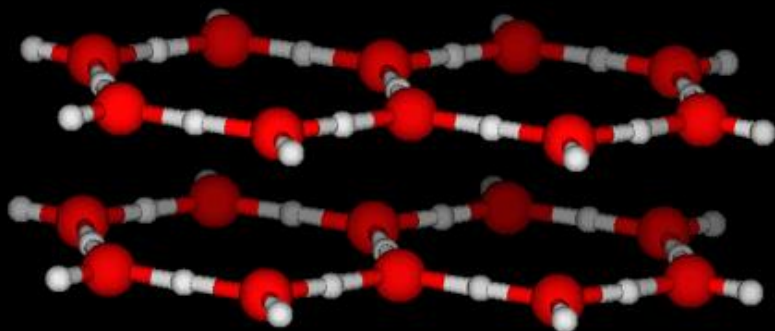
Segarra-Martí J, Roca-Sanjuán D, Merchán M (2014)

Can the hexagonal ice-like model render the **spectroscopic fingerprints of structured water?**

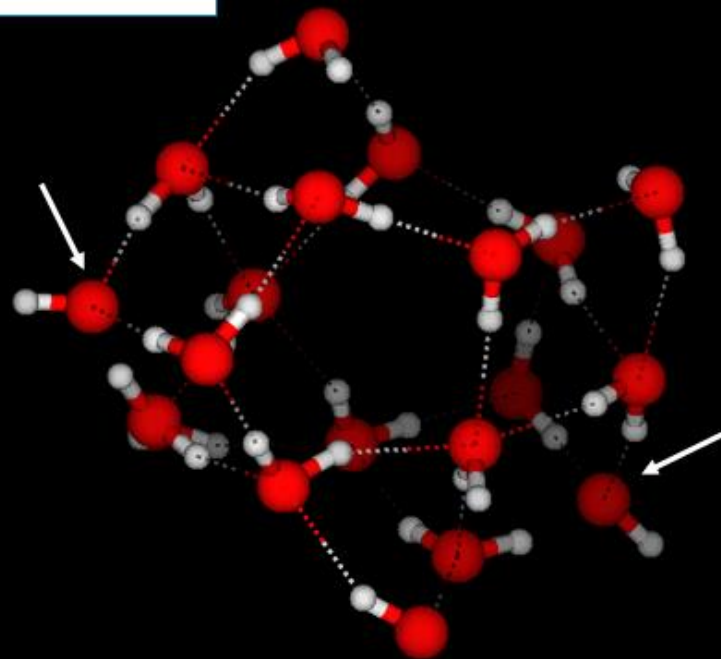
Feedback from quantum-chemical computations.

Entropy 16(7):4101-4120.

INPUT



OUTPUT

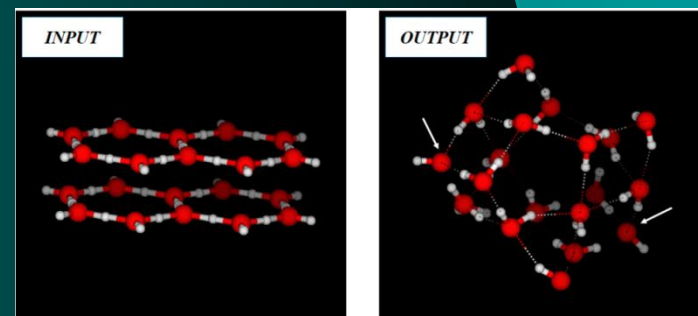


FROM: SEGARRA-MARTÍ J, ROCA-SANJUÁN D, MERCHÁN M (2014)
CAN THE HEXAGONAL ICE-LIKE MODEL RENDER THE SPECTROSCOPIC FINGERPRINTS OF STRUCTURED WATER? FEEDBACK FROM QUANTUM-CHEMICAL COMPUTATIONS.
ENTROPY 16 (7) : 4101-4120.

THE SPECTROSCOPIC FEATURES RECORDED FOR STRUCTURED WATER ARE FULLY CONSISTENT WITH THOSE COMPUTED FOR THE NEUTRAL MOLECULAR MODELS USED TO DESCRIBE THE ESSENTIALS OF THE MULTILAYER HONEYCOMB ARRANGEMENTS.

THE NEUTRAL π -STACKED DIMER $[H_{38}O_{20}]$ HAS BEEN REVEALED TO BE PARTICULARLY SUITED FOR THIS PURPOSE,

BOTH FOR THE ABSORPTION AND FLUORESCENCE EVENTS.

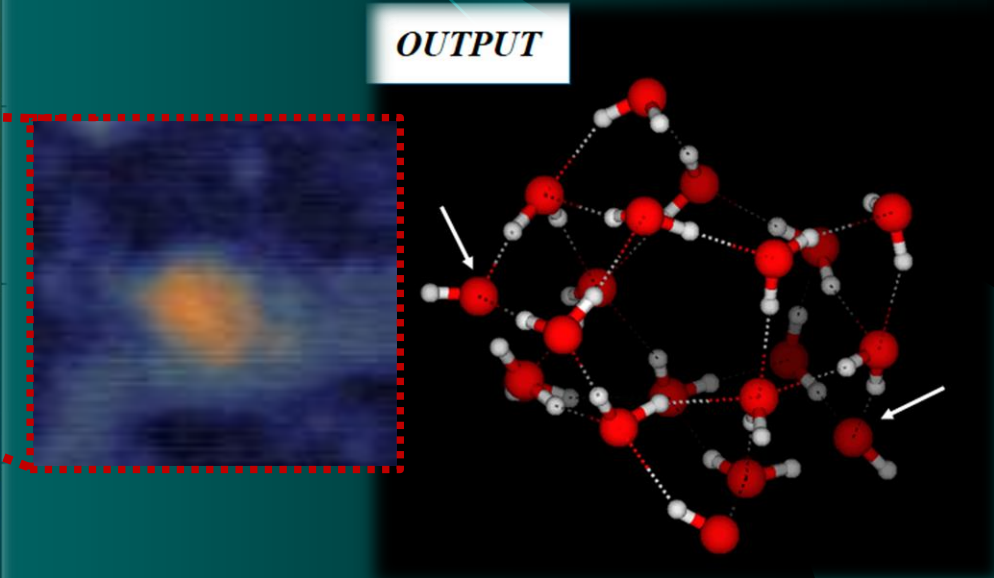
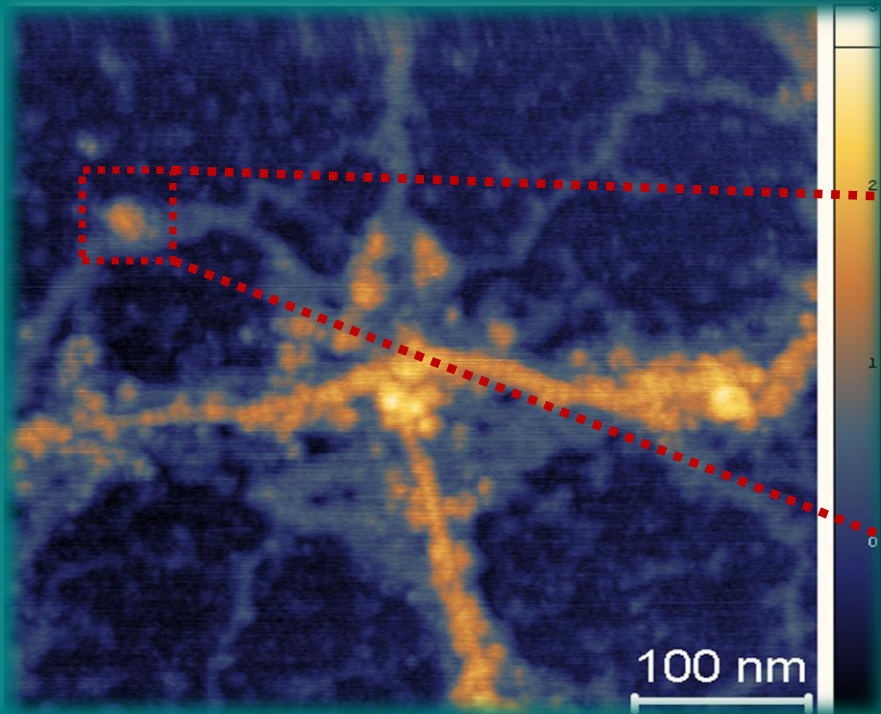
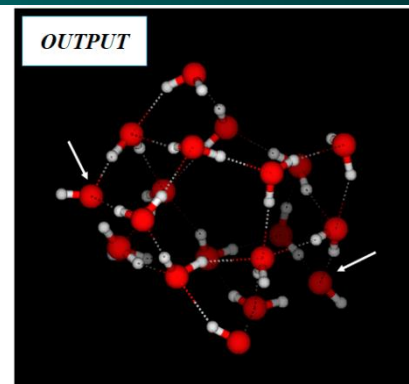
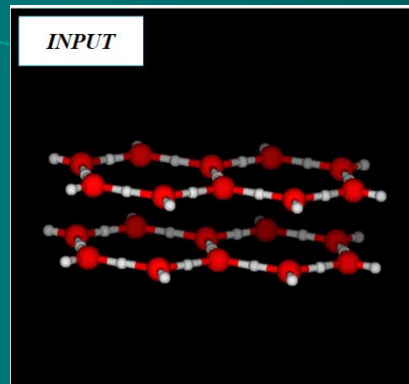


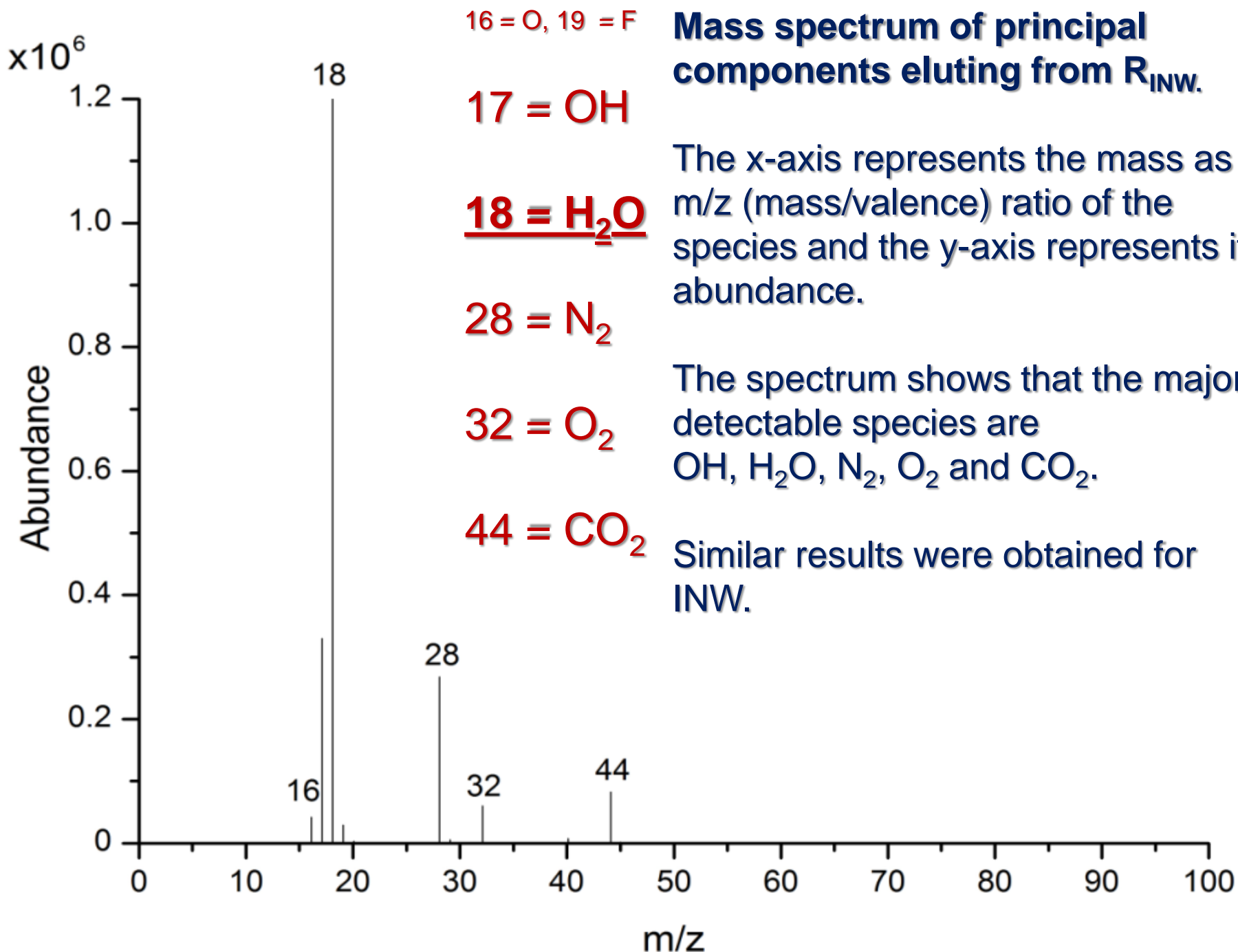
FROM: SEGARRA-MARTÍ J, ROCA-SANJUÁN D, MERCHÁN M (2014) **CAN THE HEXAGONAL ICE-LIKE MODEL RENDER THE SPECTROSCOPIC FINGERPRINTS OF STRUCTURED WATER? FEEDBACK FROM QUANTUM-CHEMICAL COMPUTATIONS.** *ENTROPY* 16 (7) : 4101-4120.

THEREFORE, THE RESULTS PROVIDED FULLY SUPPORT THE SCENARIO ENVISIONED IN THE REALM OF BIOLOGY BY ALBERT SZENT-GYÖRGYI. IN THE CONCLUSION SECTION OF THE BOOK ENTITLED “BIOENERGETICS”, PUBLISHED IN 1957, HE STATED:

“LUCRETIAN BIOCHEMISTRY INVOLVES THE ASSUMPTION THAT NO INTERACTION CAN TAKE PLACE BETWEEN MOLECULES WITHOUT THEIR TOUCHING ONE ANOTHER. SUPPORT IS GIVEN IN THIS BOOK TO THE IDEA THAT MANIFOLD INTERACTIONS CAN TAKE PLACE WITHOUT SUCH BODILY CONTACT, EITHER THROUGH ENERGY BANDS OR THROUGH THE ELECTROMAGNETIC FIELD, WHICH THUS APPEARS WITH WATER AND ITS STRUCTURES AS THE MATRIX OF BIOLOGICAL REACTIONS.”

QUANTUM CHEMISTRY SHOWS INDEED THAT WATER MAY FORM STRUCTURES THAT ABSORB AND TRANSMIT ENERGY PRECISELY IN THE SAME ENERGY RANGE AS THAT ARISING FROM AROMATIC AMINO ACIDS AND DNA/RNA NUCLEOBASES.





**PROBABLY, IT MAY IMPORTANT TO HIGHLIGHT THAT
UP TO NOW THE PAPERS PUBLISHED BY US ON THESE VERY STABLE
AGGREGATE OF PURE WATER ARE THE FOLLOWING:**

- 1) V. Elia, E.Napoli, Dissipative Structures in Extremely Diluted Solutions of Homeopathic Medicines. A Molecular Model based on Physico-Chemical and Gravimetric evidences, *International Journal of Design and Nature*, 5 (1), 39-48, (2010)**
- 2) V. Elia, G. Ausanio, A. De Ninno, F. Gentile, R. Germano, E. Napoli, M. Niccoli, Experimental evidence of stable aggregates of water at room temperature and normal pressure after iterative contact with Nafion polymer membrane, *WATER*, 5, 16-26 (2013)**
- 3) V. Elia, G. Ausanio, F. Gentile, R. Germano, E. Napoli and M. Niccoli, Experimental evidence of Stable Water Nanostructures in Extremely Diluted Solutions, at Standard Pressure and Temperature, *Homeopathy*, 103, 1, 44-50 (2014)**
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Large Supramolecular Water Clusters Caught on Camera - A Review

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UV-FLUORESCENCE BEHAVIOR OF SUPRAMOLECULAR H_2O ARCHITECTURES GENERATED BY PERTURBING PURE WATER

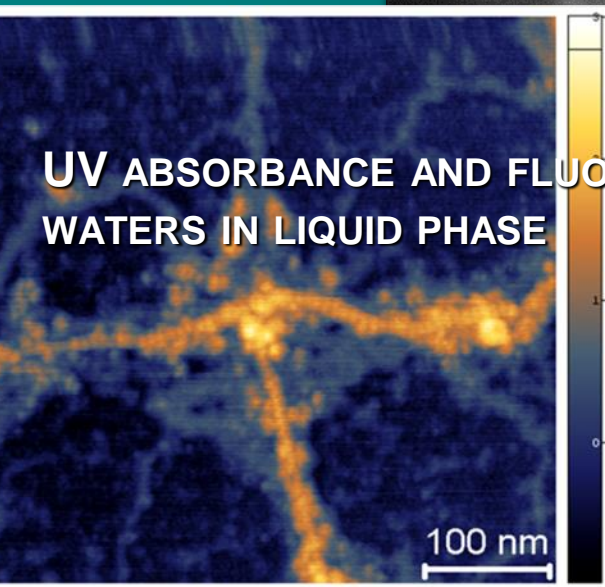
CONCLUSIONS

UV ABSORBANCE AND FLUORESCENCE SPECTRA RESEMBLE THOSE OF OTHER STRUCTURED
WATERS IN LIQUID PHASE

BUT

WE ISOLATED A NEW SOLID MATERIAL
AT NORMAL PRESSURE AND TEMPERATURE
CONSTITUTED BY SOVRAMOLECULAR STRUCTURES OF H_2O :

XEROSYDRYLE



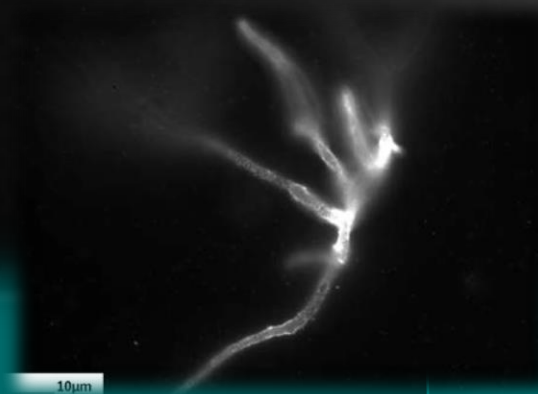
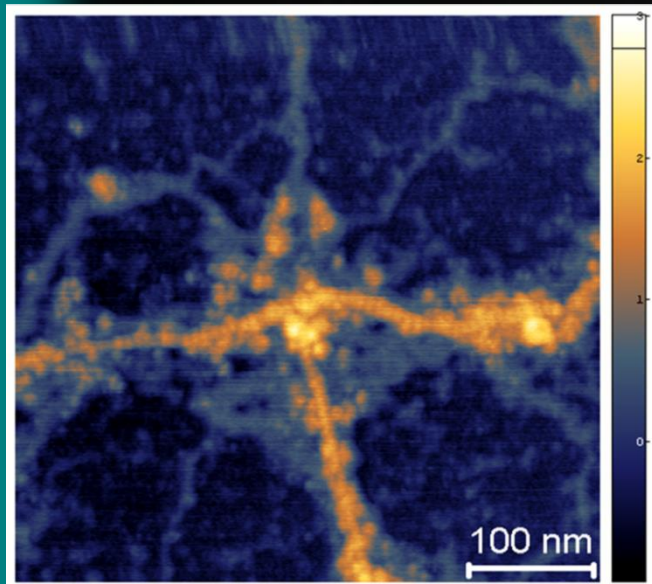
mag	HV	WD	spot	mode	HFW
3 000 x	30.00 kV	10.6 mm	2.7	SE	99.5 μ m

40 μ m

**IT IS POSSIBLE TO BEGIN TO INVESTIGATE THE NATURAL HYPOTHESIS
THAT THESE SPONTANEOUS DISSIPATIVE SYSTEMS
FORMING IN PURE LIQUID WATER (MADE BY H₂O)**

**ABLE TO “CONDENSE” IN CERTAIN CONDITIONS
AND BECOMING SOLID AT AMBIENT TEMPERATURE AND PRESSURE,**

MAY BE THE MATRIX OF LIFE ITSELF



**I DEDICATE THIS TALK TO THE LIVING MEMORY
OF
MAE WAN HO**

THANK YOU FOR YOUR ATTENTION !!

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