



Since 1959

Can Water Microdroplets Make Soil?

A path to sustainable nanotechnology



Matter in confinement for sustainability

Co-founder

InnoNano Research Pvt. Ltd.
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VayuJAL Technologies Pvt. Ltd.
Aqueasy Innovations Pvt. Ltd.
Hydromaterials Pvt. Ltd.
EyeNetAqua Pvt. Ltd.
Deepspectrum Analytics Pvt. Ltd.

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Professor-in-charge



International Centre for Clean Water



REMARKS

Spontaneous weathering of natural minerals in charged water microdroplets forms nanomaterials

[illegible]

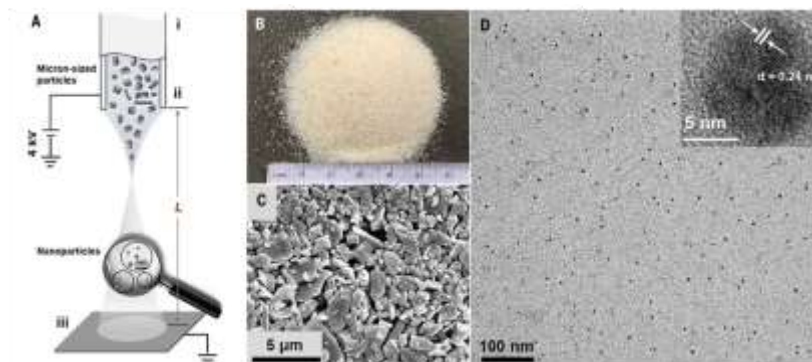
For our applications, we proposed alternative models of the form $f(\mathbf{x}) = \mathbf{w}^T \phi(\mathbf{x})$, where $\phi(\mathbf{x})$ is a vector of features. For example, we can use the features $\phi(\mathbf{x}) = [1, x_1, x_2, \dots, x_n]^T$, where x_i is the i -th component of \mathbf{x} . This is a linear model in the space of features.

After the delay, the results of the second study were published. It reported that the α -synuclein-deficient mice performed better than the control mice on all six measures of motor function. Interestingly, the control mice that were fed the α -synuclein-deficient mice had a 50% reduction in the amount of α -synuclein in the brain. This suggests that the α -synuclein-deficient mice may have a protective effect on the brain.

with 100 mM of 10 mM sodium chloride at 20 mM. Low levels of 0.5 M of 10 mM sodium chloride in the dialysis buffer (Fig. 2) were obtained. The concentration of disaccharide 15.5 mM was low in the dialysis buffer, which resulted in a higher disaccharide concentration of 10 mM, consistent with disaccharide concentrations [2, 4]. This product was then obtained as a homogeneous disaccharide disaccharide (TDS) and had a 5.0 to 10.0 mM disaccharide concentration (Fig. 2). The product was then obtained as a homogeneous disaccharide disaccharide (TDS) and had a 5.0 to 10.0 mM disaccharide concentration (Fig. 2). The product was then obtained as a homogeneous disaccharide disaccharide (TDS) and had a 5.0 to 10.0 mM disaccharide concentration (Fig. 2).

To determine what our future child development study might entail, we interviewed a few people who were involved in the program. We asked them what they thought the program should be like, and what they thought the program should be able to do. We also asked them what they thought the program should be able to do for the children. We asked them what they thought the program should be able to do for the children. We asked them what they thought the program should be able to do for the children.

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A scale of 1000

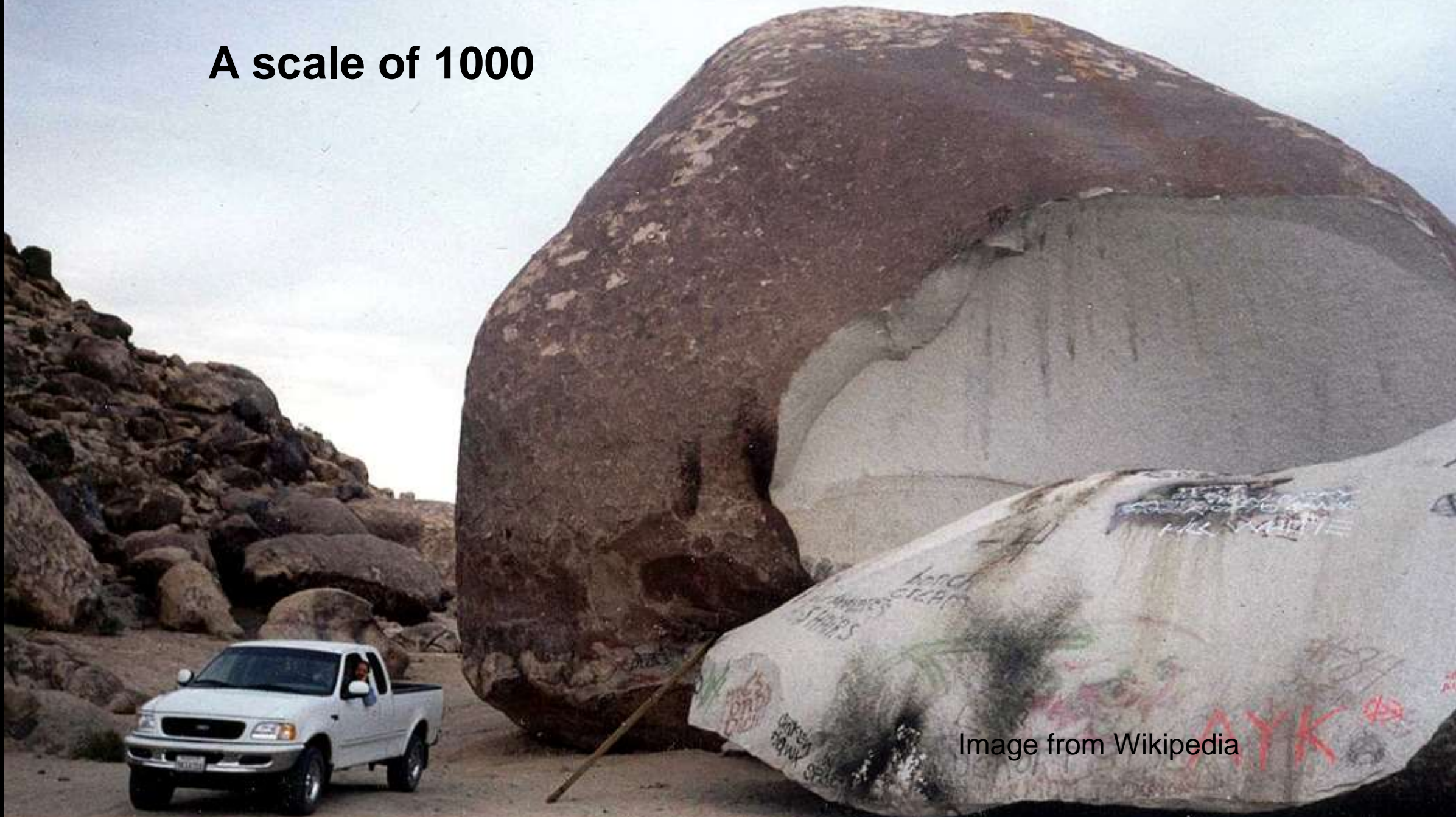
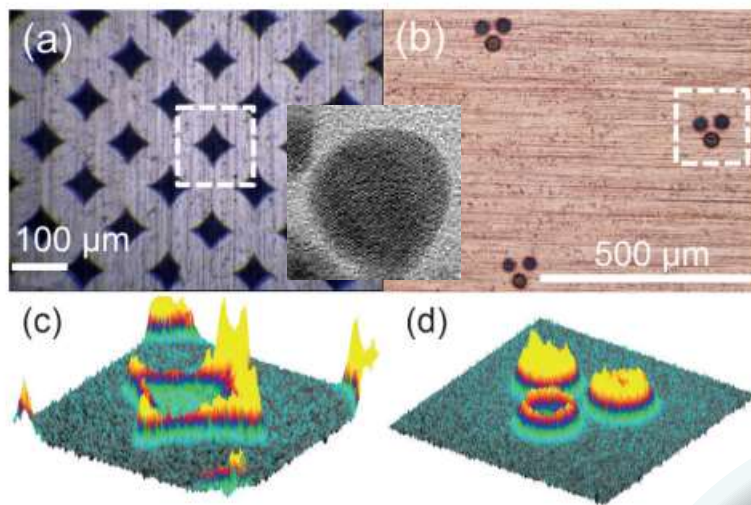
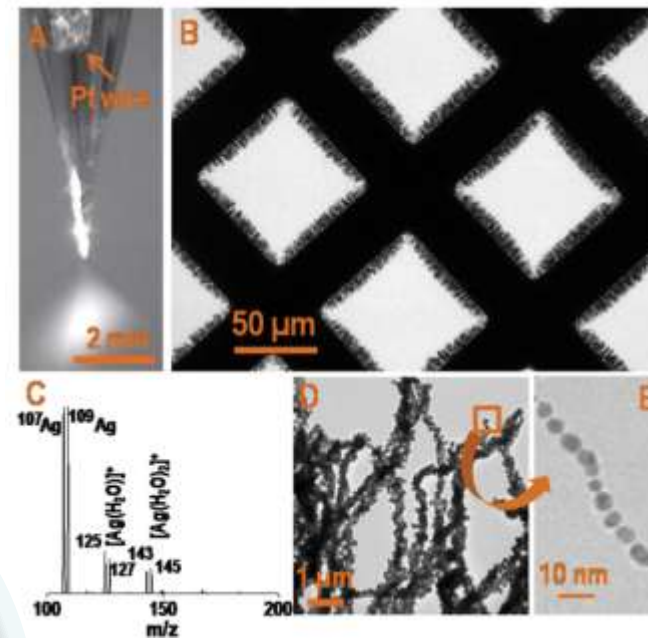


Image from Wikipedia

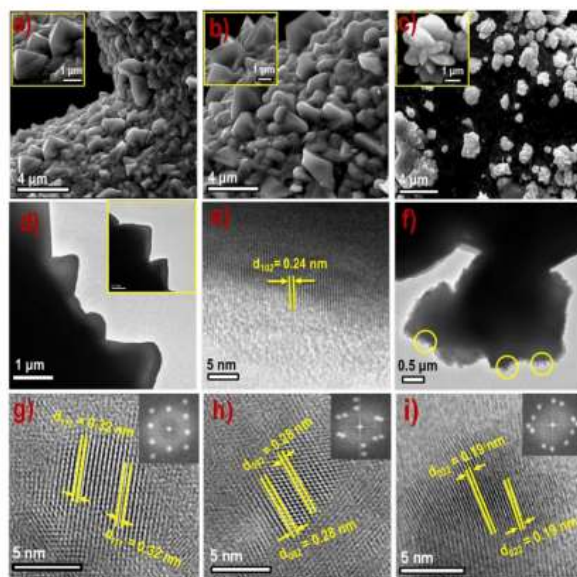
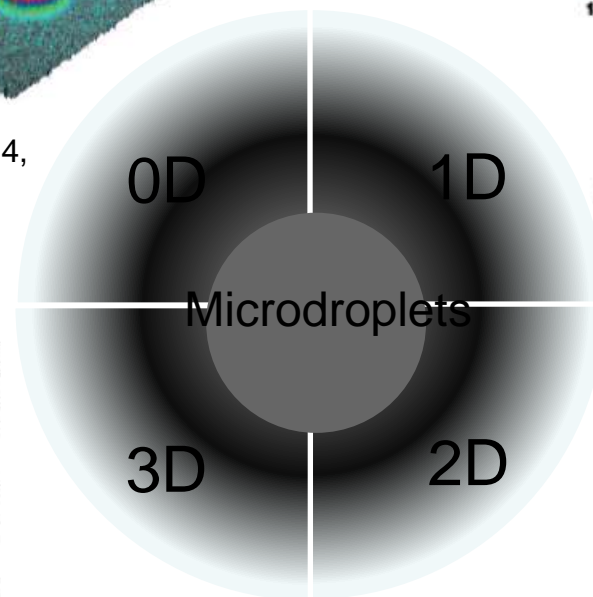
Functional Nanomaterials



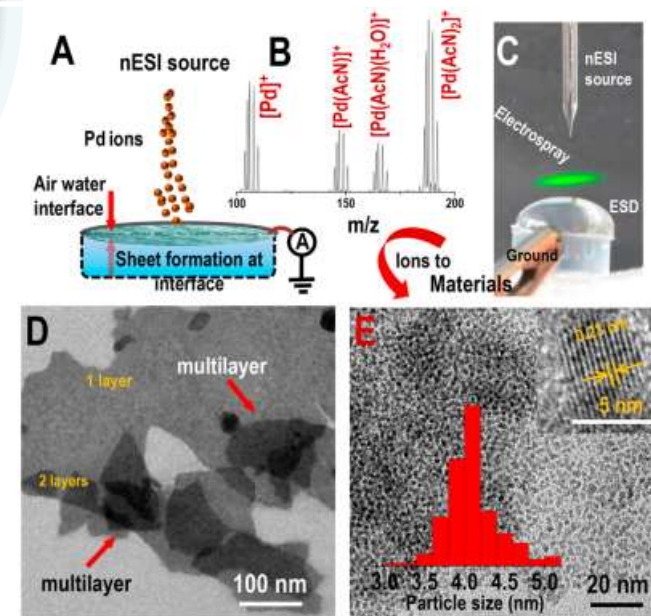
Anyin Li, et. al., *Angew. Chem. Int. Ed.* 2014, 53, 12528–12531.



Depanjan Sarkar et. al., *Adv. Mater.* 2016, 28, 2223–2228.



Arijit Jana et. al., *J. Mater. Chem. A*, 2019, 7, 6387–6394.



Depanjan Sarkar, et. al., *J. Phys. Chem. C* 2018, 122, 17777–17783.

Chemical Science

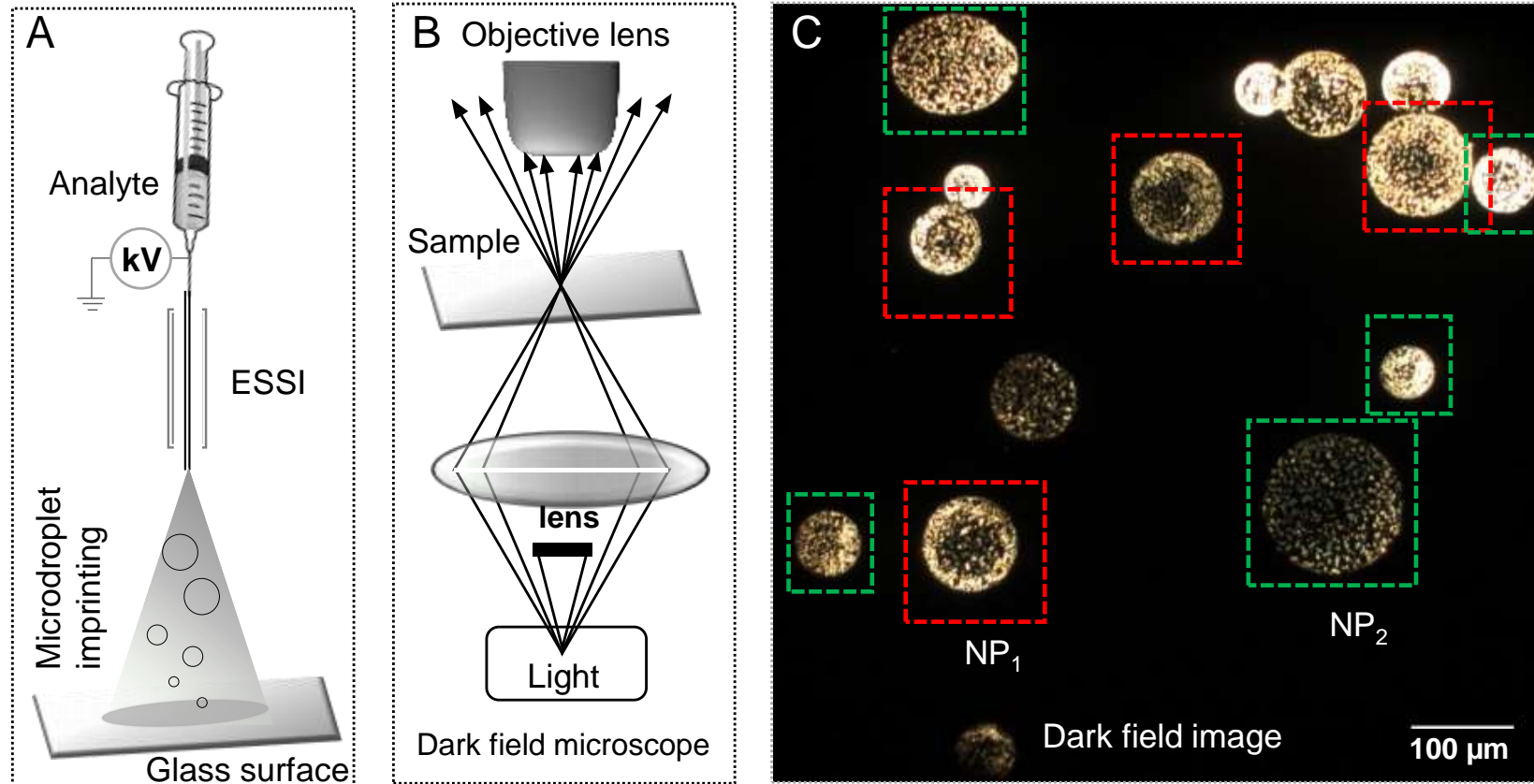
Volume 13
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7 December 2022
Pages 13251–13634

rsc.li/chemical-science



ISSN 2041-6539

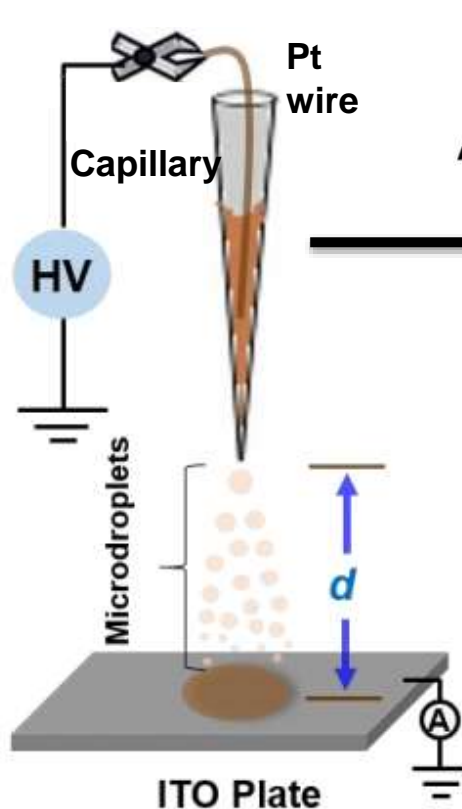
Understanding Microdroplets



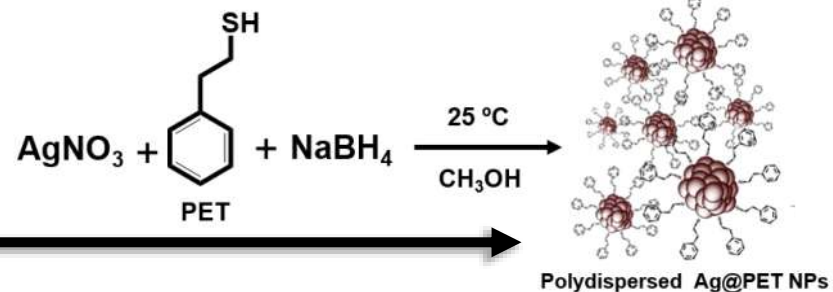
Transformation of Materials in Microdroplets

Ambient Microdroplet Annealing of Nanoparticles

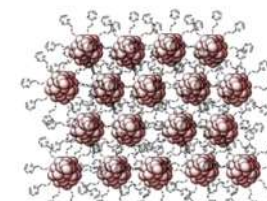
Experimental set-up



Synthesis of polydisperse NPs

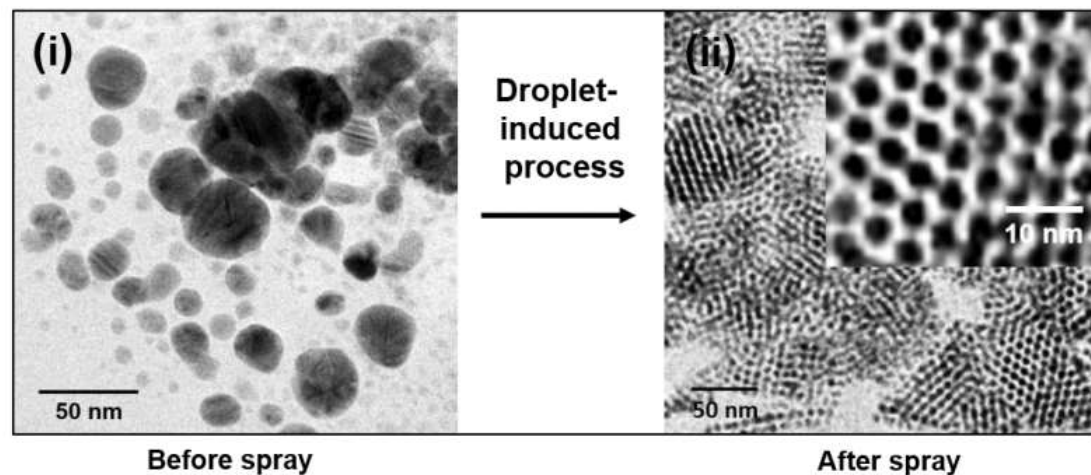


Ambient
microdroplet
annealing



Assembly of uniform nanocrystals

Transformation process





Thanks to ChatGPT

Weathering in Nature

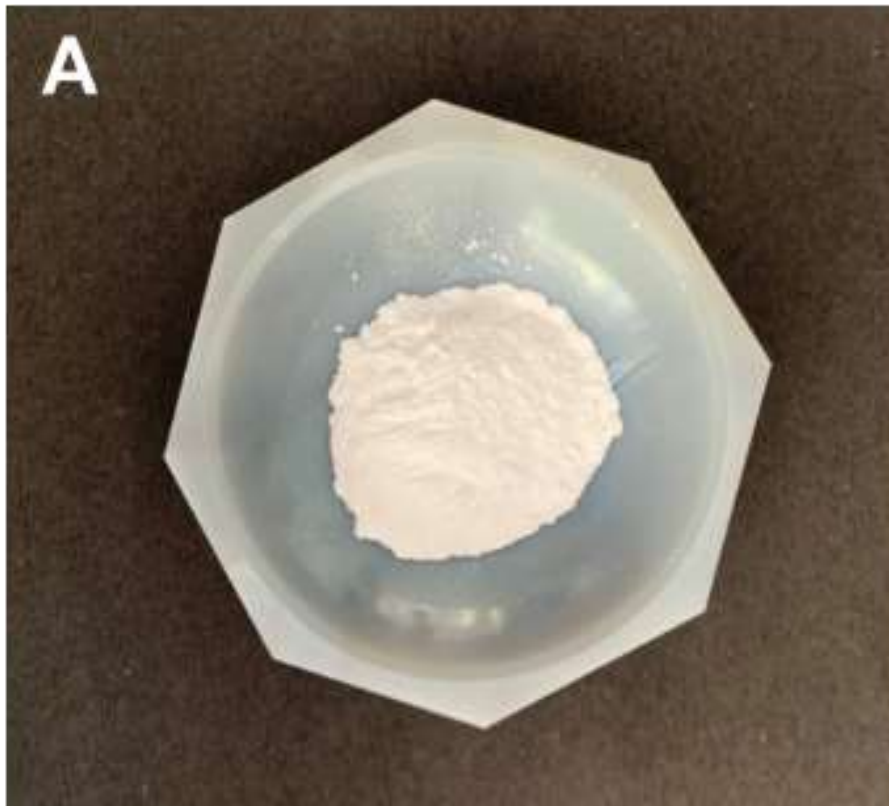


Sand, the Ubiquitous Material

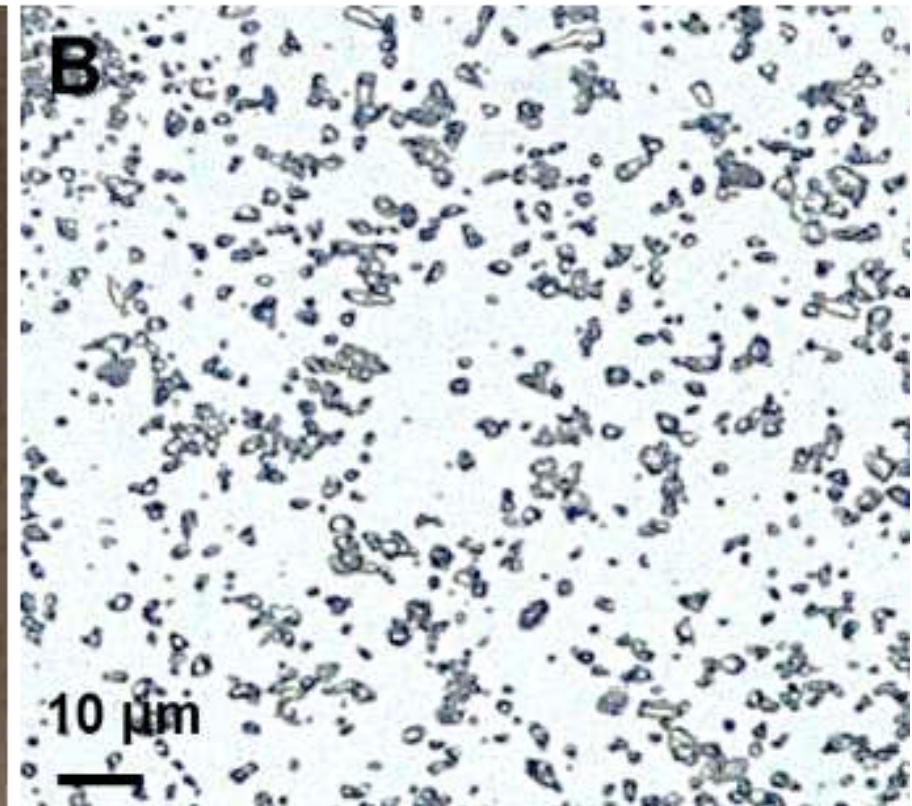


Images from Wikipedia



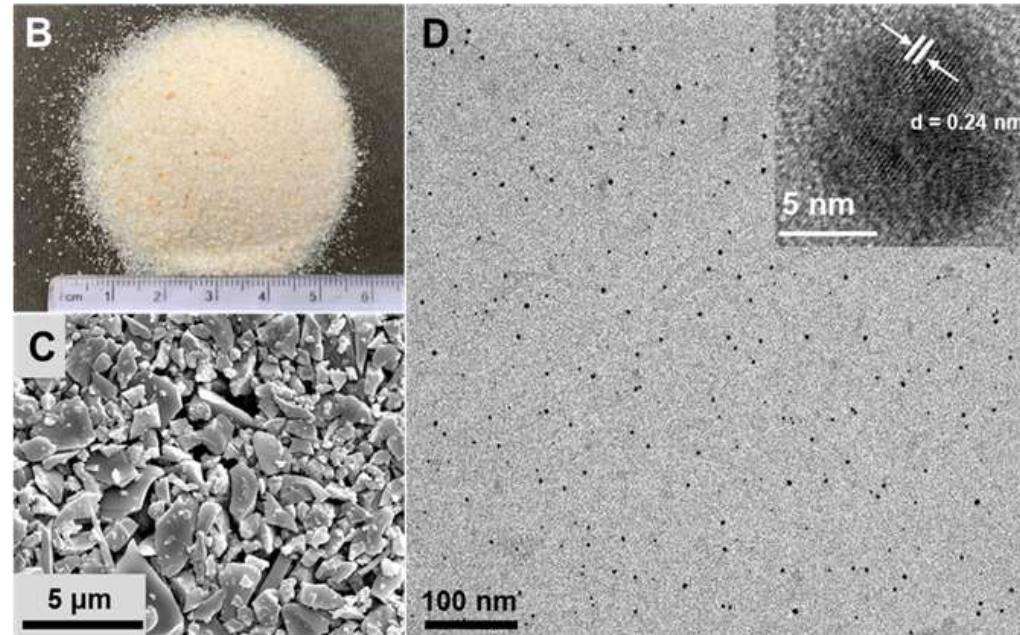
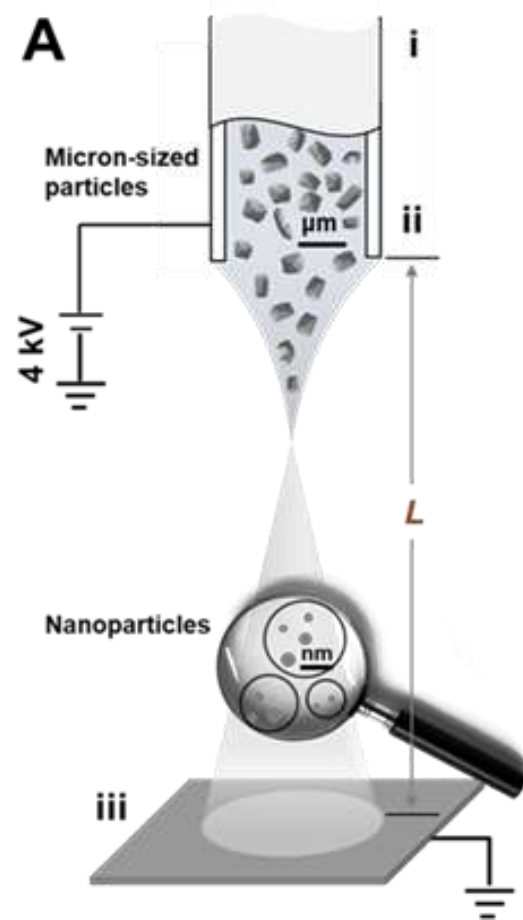


Ground silica

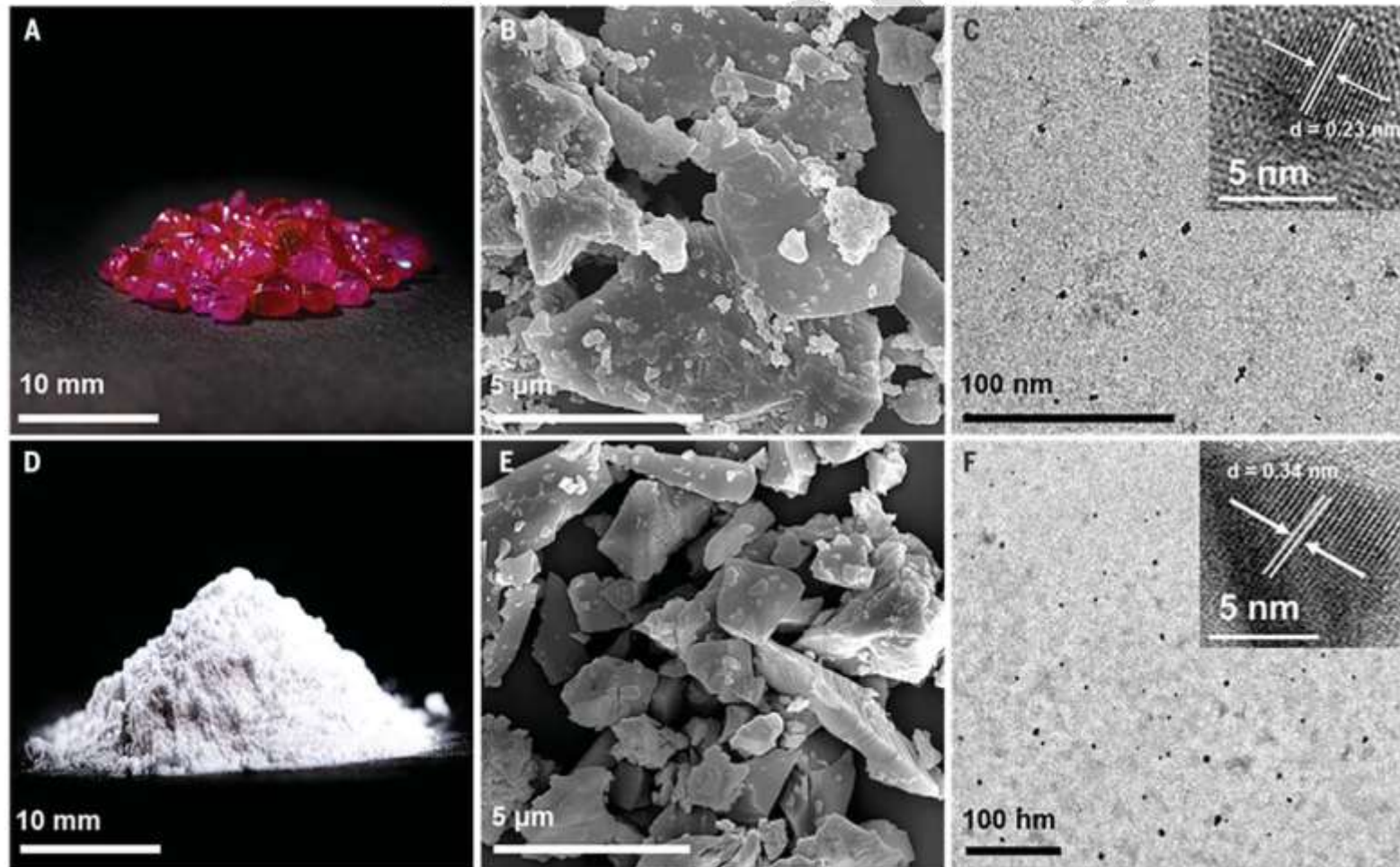


Optical image of silica

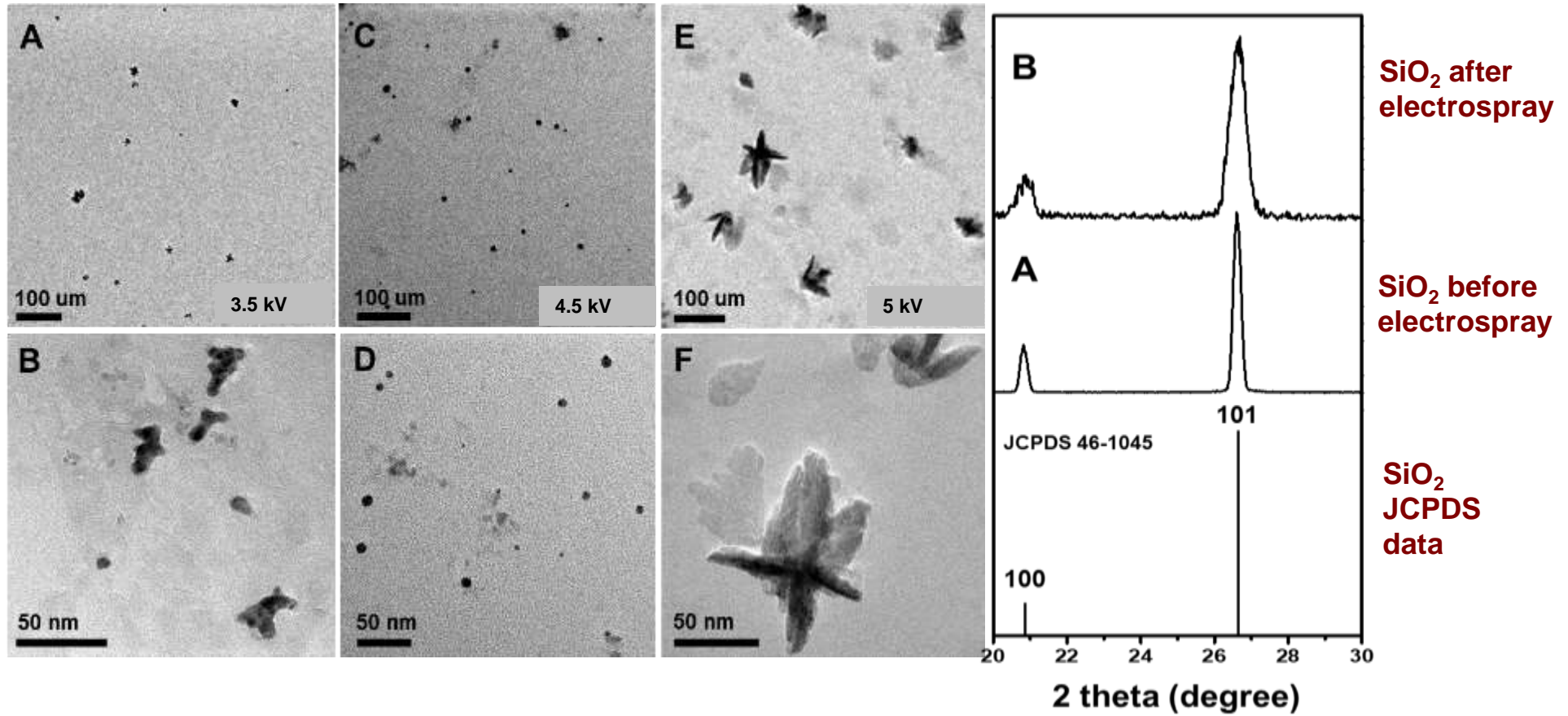
Weathering of Minerals in Microdroplets



Ruby, Fused Alumina

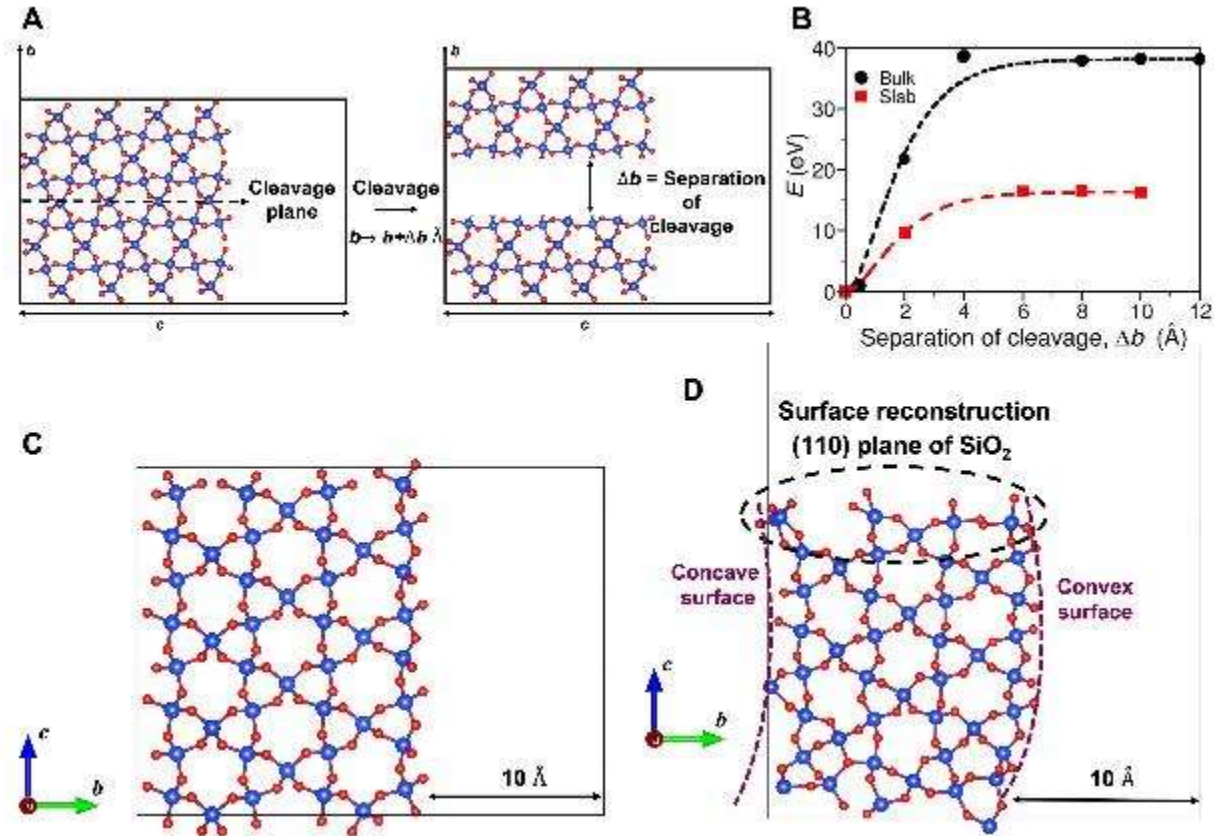
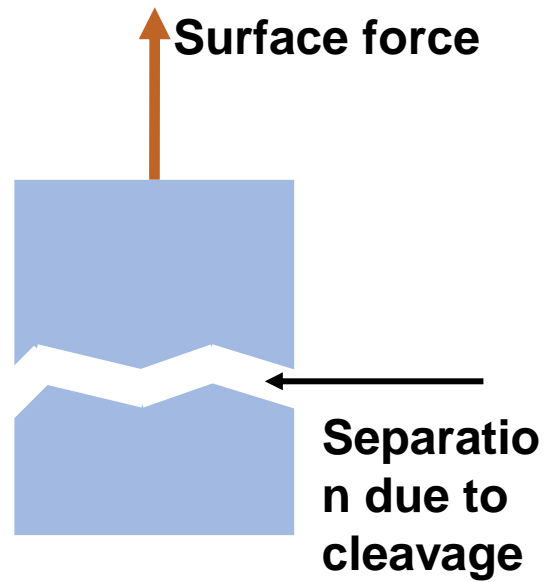


Fragmentation of Silica – Varying Conditions

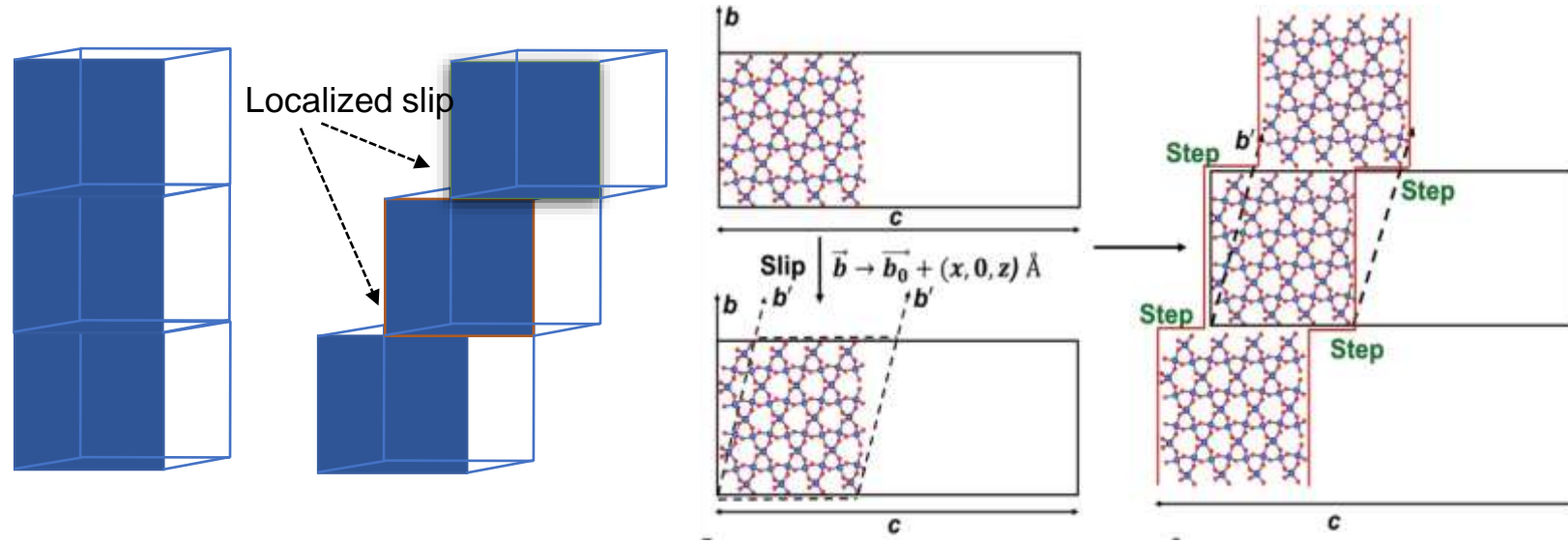


Mechanism: Cleavage

The process of cleavage and surface reconstruction visualized with first-principles simulations



Mechanism: Slip



This instability leads to the formation of a stacking fault on the (010) plane, achieved with slip localized at (010) plane

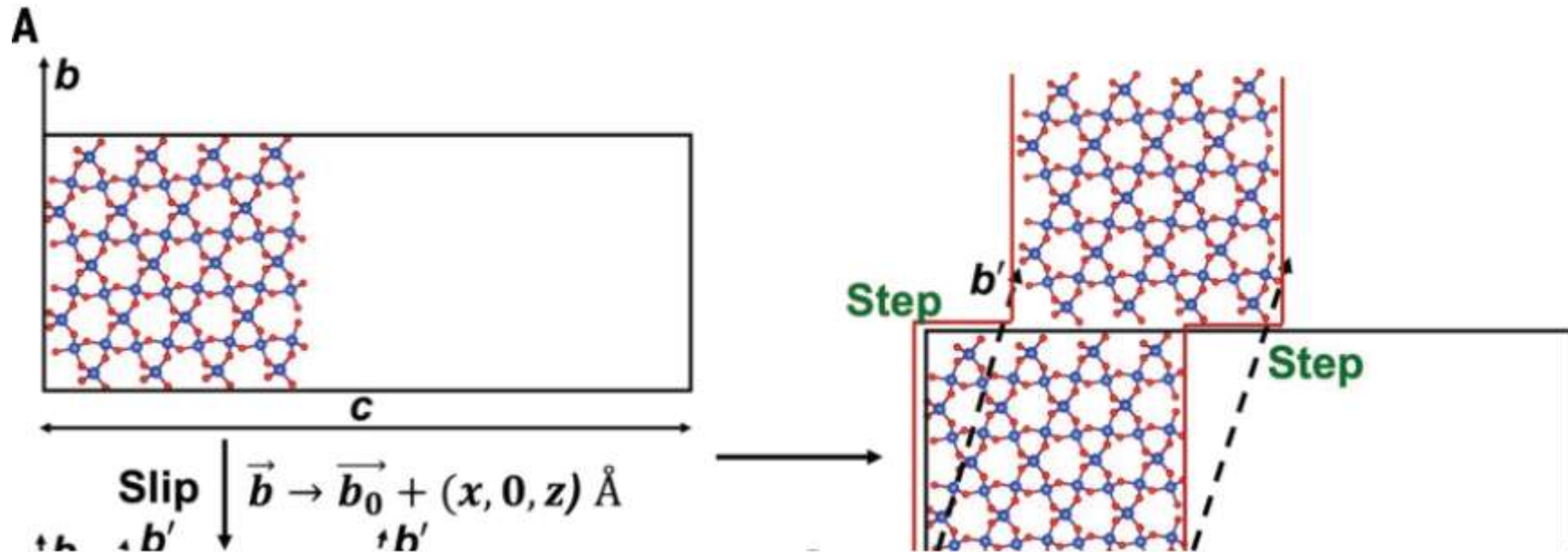
Stacking fault

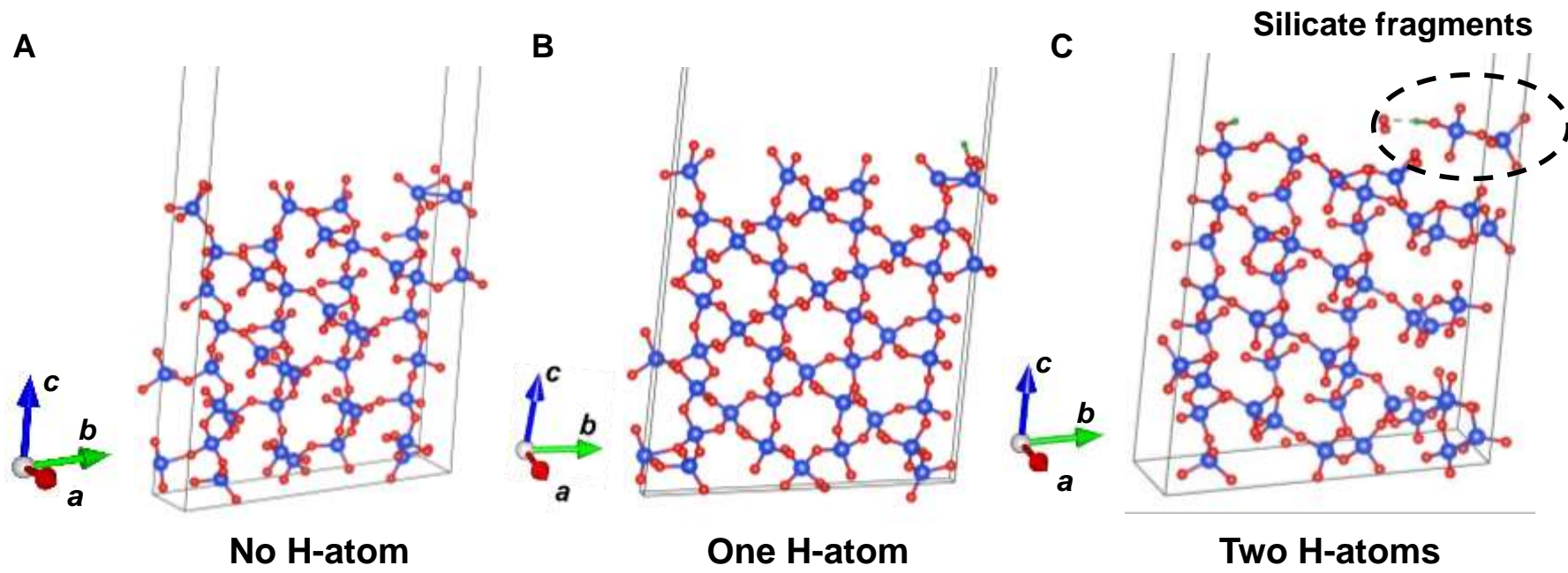
$$\vec{b} \rightarrow \vec{b}_0 + (x, 0, z),$$

$(x, z \in [0, 1])$ - fractional coordinates

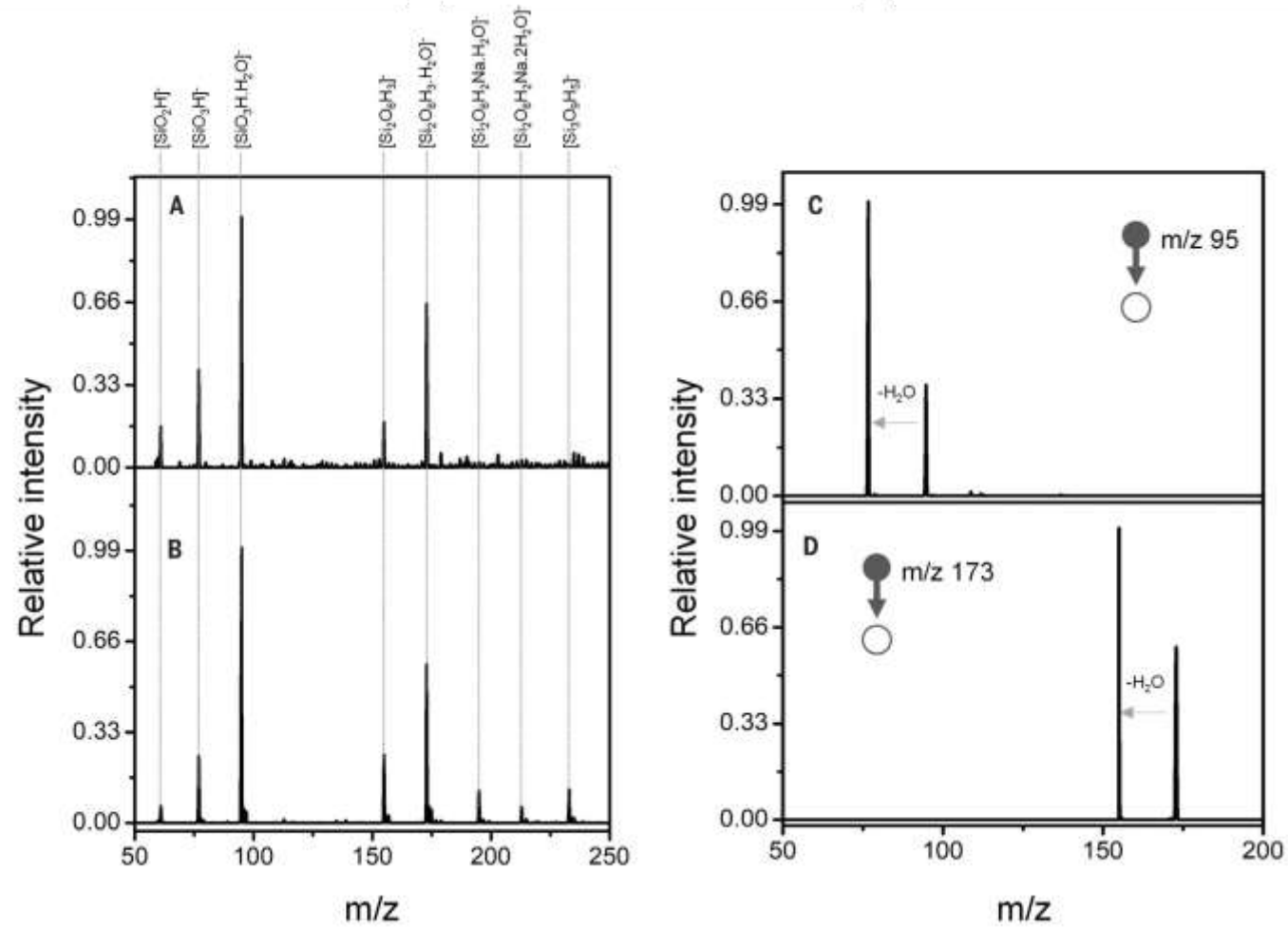
SFEs of (010) direction with (0, 0), (0, 0.5), (0.5, 0) and (0.5, 0.5) slip configurations on the (110) plane of SiO_2

SFE (J/m^2)	Slab					
	x	z	w/o H-atom	1 H-atom	2 H-atoms	E
	0.0	0.0	0	0	0	0
	0.5	0.5	-1.21	-0.93	-0.88	-1.20
	0.5	0.0	1.20	1.18	0.90	1.12
	0.0	0.5	-0.07	0.89	-0.83	-0.09

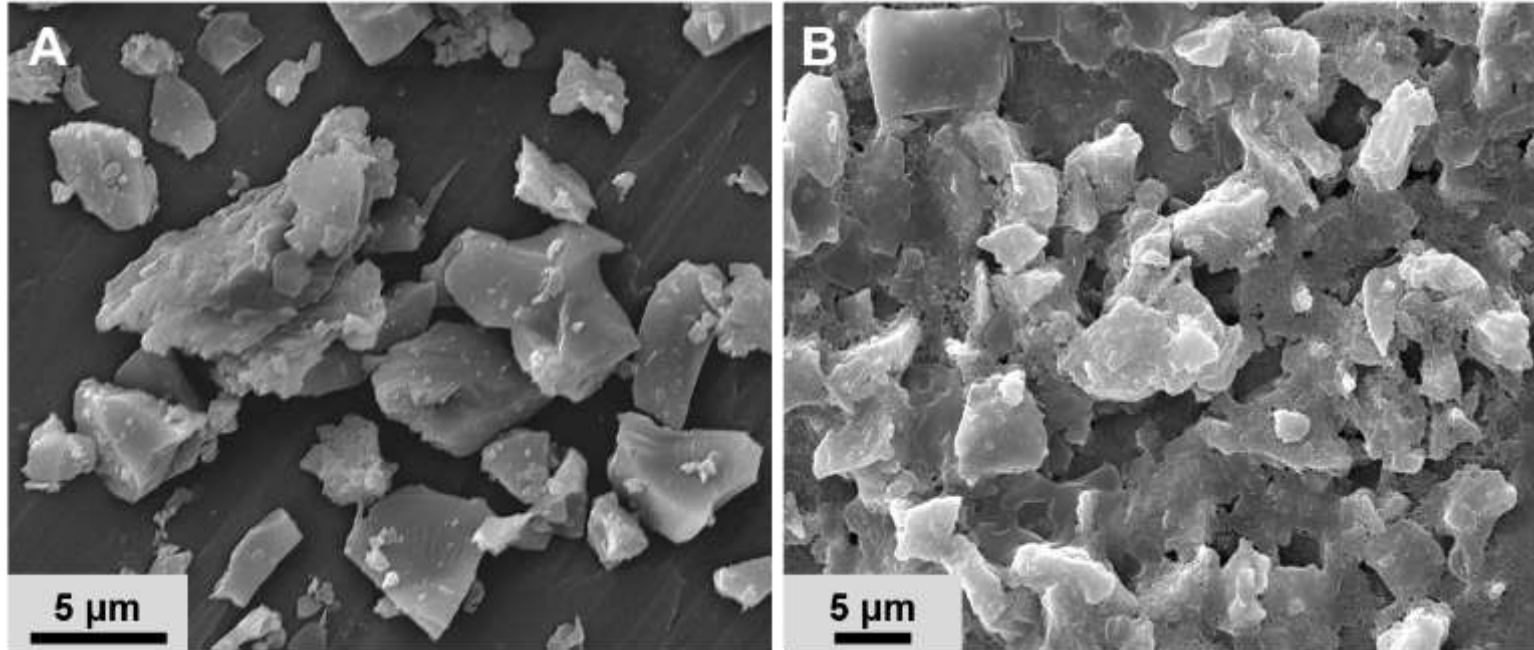




Mass Spectrometry of the Fragments

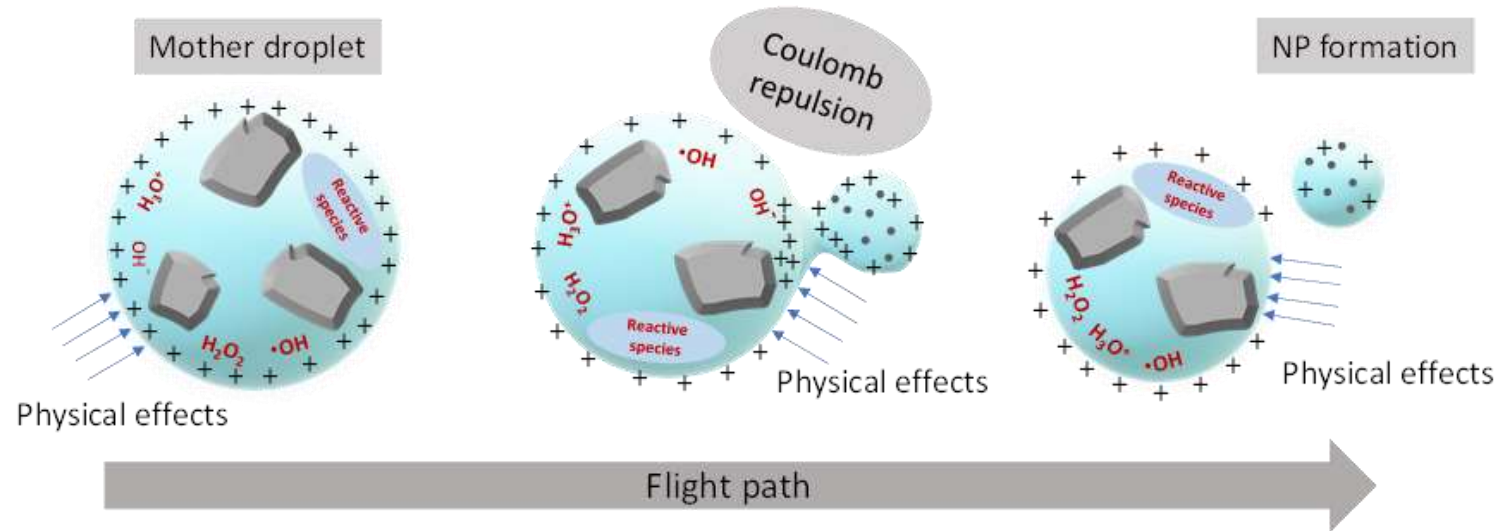


Effect of charged microdroplets on quartz



Increased surface roughness after the spray

Mechanism of nanoparticle formation



Rayleigh, On the
equilibrium of liquid
conducting masses
charged with electricity,
Philosophical Magazine,
1882

$$Q = 8\pi (\epsilon_0 \gamma R^3)^{1/2}$$

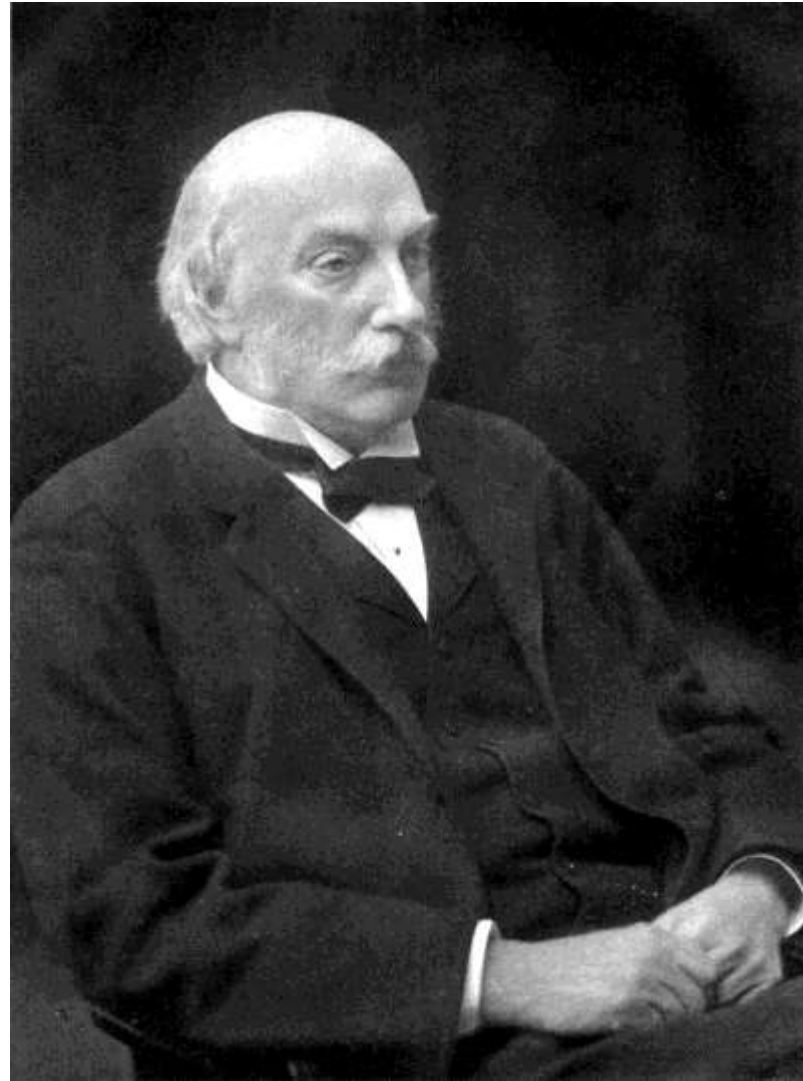
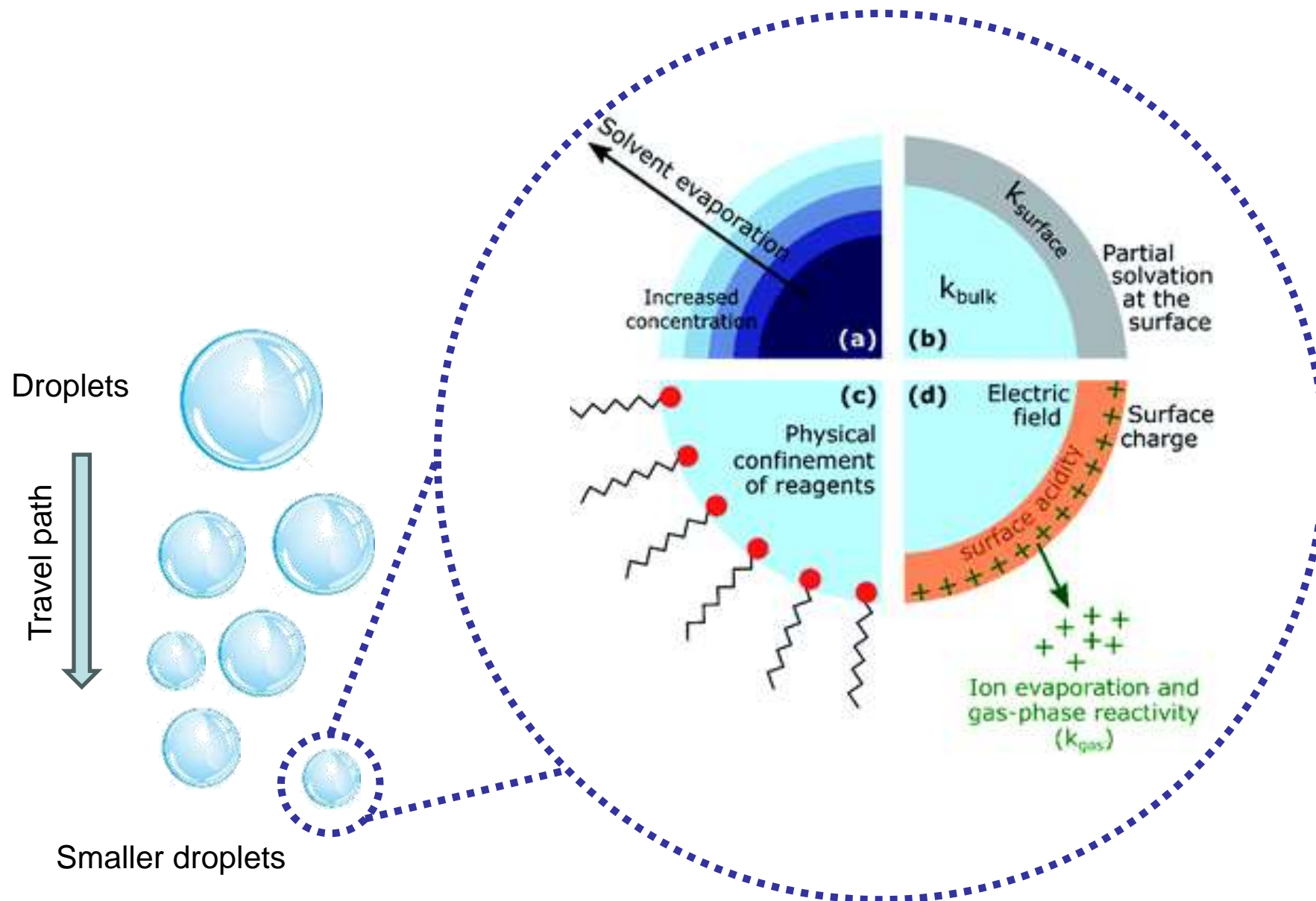


Image from Wikipedia

Understanding Microdroplets



PERSPECTIVES

CHEMISTRY

Breaking down microdroplet chemistry

Charged microdroplets accelerate mineral disintegration

By R. Graham Cooks and Dylan T. Holden

Charged microdroplets are commonly observed in clouds, sea spray, and other natural aerosols. The chemistry that occurs at the air-water interface of these droplets is often distinct from what is observed in bulk solution, which is of considerable interest because chemical reactions can be accelerated at this boundary (1, 2). This may have implications for environmental processes such as the weathering of rocks, which contributes to soil formation. On page 1012 of this issue, Spoorthi *et al.* (3) report that micrometer-scale mineral particles can rapidly break down into nanoparticles when in charged aqueous microdroplets (see the figure). This points to a potential role for atmospheric water droplets in the natural disintegration of minerals.

To examine material degradation, Spoorthi *et al.* borrowed methodology used to accelerate bond-forming chemical reactions. By spraying an aqueous suspension of microparticles of natural minerals, the authors produced nanoparticles of minerals in high yield. Specifically, Spoorthi *et al.* used an electrospray device to emit a jet of liquid droplets (by applying high voltage) containing mineral particles of natural quartz, ruby, or synthetic alumina that ranged in size from 1 to 5 μm in diameter. The authors observed the production of nanoparticles that were 5 to 10 nm in diameter. Moreover, the fragmentation occurred in approximately 10 ms.

Such material degradation and chemical synthesis experiments are united by the extremes of chemical reactivity that occur at the air-water interface, where reagents are partially solvated (4). Whether formed through nebulization, splashing from a surface, or other means, microdroplet populations will include droplets with nonzero net charges. The small radius of curvature in a microdroplet produces a very strong electric field (5) that can support a double layer of electric charge at the air-water interface. The change in geometry (radius of curvature)

converts a two-dimensional air-water interface with limited electric field into a sphere with an electric field of a strength approaching the order of chemical bond energies (3 to 4.5 eV/Å). Coulombic fission (the splitting of charged microdroplets due to excess charge overcoming the surface tension) and evaporative processes further increase the surface area, reduce the radius of curvature, and augment the surface electric field of the droplet.

The unusual chemical nature of the air-water interface results in much remarkable chemistry. For example, amino acids in water undergo dehydration to form peptides in this environment (6), whereas bulk water simply solvates amino acids. The superacidic interface activates amino acids and removes water to yield peptides. In addition to such acid-base reactions, redox chemistry results from the formation of strong oxidants and reductants from water at the interface. For example, a high hydronium ion (H_3O^+) concentration at the interface derived from fleetingly charged surface water molecules ($\text{H}_2\text{O}^+/\text{H}_2\text{O}^-$) coexists with oxidative species such as hydrogen peroxide (H_2O_2) and OH^\bullet . These redox species enable a variety of spontaneous chemical trans-

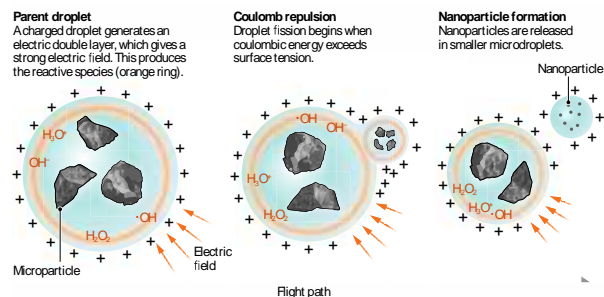
formations, including carbon-oxygen (C-O) bond cleavage in phosphonates, which yields the corresponding phosphonic acid (7), and in the Baeyer-Villiger oxidation of aryl ketones to give esters (8). These considerations thereby enable simultaneous acid-base and oxidation-reduction chemistry in a single population of droplets (7).

Through their study, Spoorthi *et al.* have added natural weathering to a list of processes in which accelerated interfacial microdroplet reactions play an important role. Other processes include those in the atmosphere, both natural and anthropogenic, the latter typified by pollution that involves nitrate photochemistry (9). A substantial number of accelerated catalyst-free microdroplet reactions form the basis for chemical syntheses that generate a variety of small molecules (10), including the facile and high-throughput functionalization of drugs. This latter approach can be scaled up so that microdroplet reactions produce substantial small-molecule products. Prebiotic chemistry, including peptide and nucleotide formation, is another process that is accelerated at the microdroplet air-water interface (11).

The millisecond timescale of quartz degradation reported by Spoorthi *et al.* matches the known microsecond-to-millisecond timescale for accelerated bond-formation and bond-cleavage chemical reactions in microdroplets (1). This reinforces the conclusion that the chemical basis for accelerated weathering lies in the powerful acidic and hydrolytic nature of the air-water interface. The authors further suggest a role for the superacid interface in inducing slippage at crystal plane boundaries in quartz and ruby fragmentation. Their simulations show that individual protons inserted into the slip configuration mineral

Micro-to-nano transitions in minerals at the air-water interface

Reactions that promote mineral disintegration are accelerated at the air-water interface of microdroplets. Key reactive species are the result of the effects of a high electric field at the surface of the water droplets.



Department of Chemistry, Purdue University, West Lafayette, IN, USA. Email: cooks@purdue.edu

Water structure and electric fields at the interface of oil droplets

<https://doi.org/10.1038/s41586-025-08702-y>

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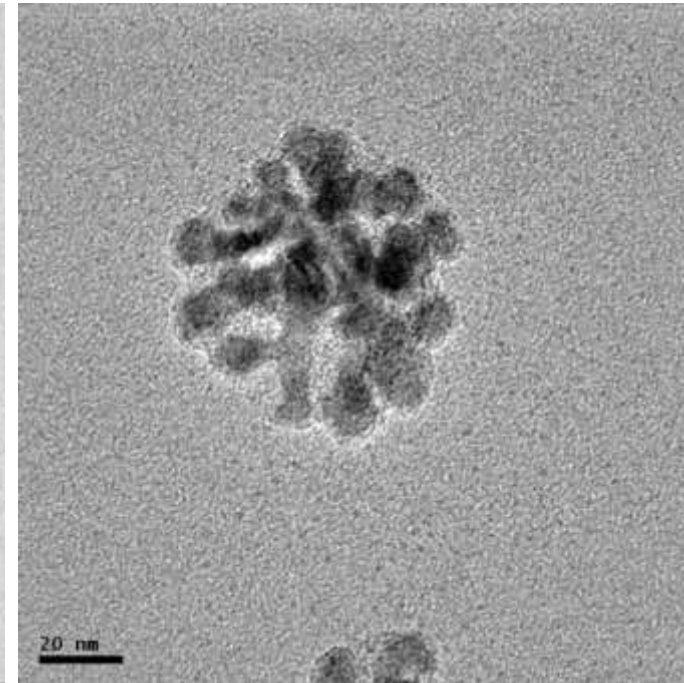
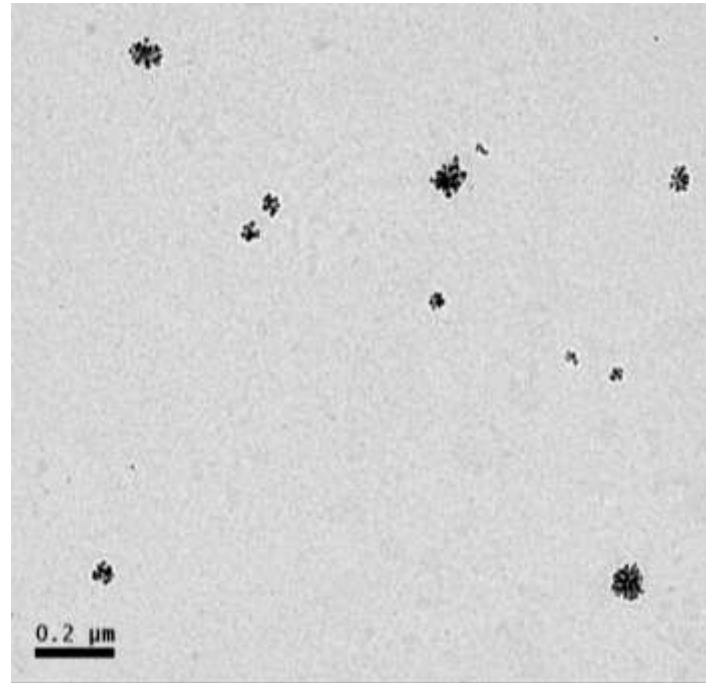
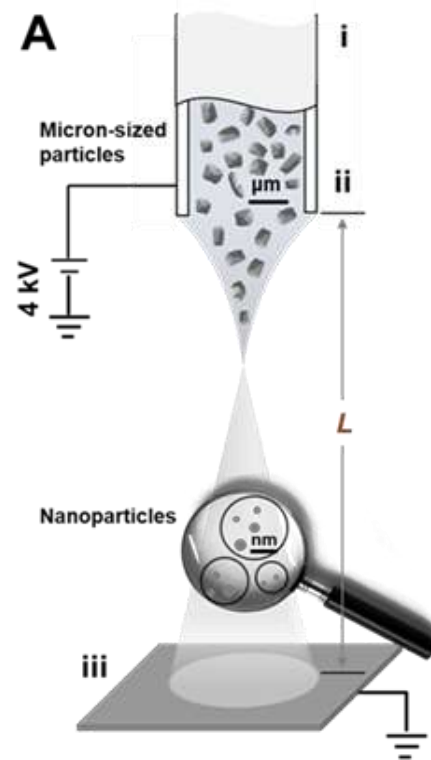
Published online: 19 March 2025



Lixue Shi^{1,5}, R. Allen LaCour^{2,3,5}, Naixin Qian¹, Joseph P. Heindel^{2,3}, Xiaoqi Lang¹, Ruoqi Zhao^{2,3}, Teresa Head-Gordon^{2,3,4} & Wei Min¹

Interfacial water exhibits rich and complex behaviour¹, playing an important part in chemistry, biology, geology and engineering. However, there is still much debate on the fundamental properties of water at hydrophobic interfaces, such as orientational ordering, the concentration of hydronium and hydroxide, improper hydrogen bonds and the presence of large electric fields^{2–5}. This controversy arises from the challenges in measuring interfacial systems, even with the most advanced experimental techniques and theoretical approaches available. Here we report on an in-solution, interface-selective Raman spectroscopy method using multivariate curve resolution^{6,7} to probe hexadecane-in-water emulsions, aided by a monomer-field theoretical model for Raman spectroscopy⁸. Our results indicate that oil–water emulsion interfaces can exhibit reduced tetrahedral order and weaker hydrogen bonding, along with a substantial population of free hydroxyl groups that experience about 95 cm^{−1} redshift in their stretching mode compared with planar oil–water interfaces. Given the known electrostatic zeta potential characteristic of oil droplets⁹, we propose the existence of a strong electric field (about 50–90 MV cm^{−1}) emanating from the oil phase. This field is inferred indirectly but supported by control experiments and theoretical estimates. These observations are either absent or opposite in the molecular hydrophobic interface formed by small solutes or at planar oil–water interfaces. Instead, water structural disorder and enhanced electric fields emerge as unique features of the mesoscale interface in oil–water emulsions, potentially contributing to the accelerated chemical reactivity observed at hydrophobic–water interfaces^{10–13}.

How do they form?

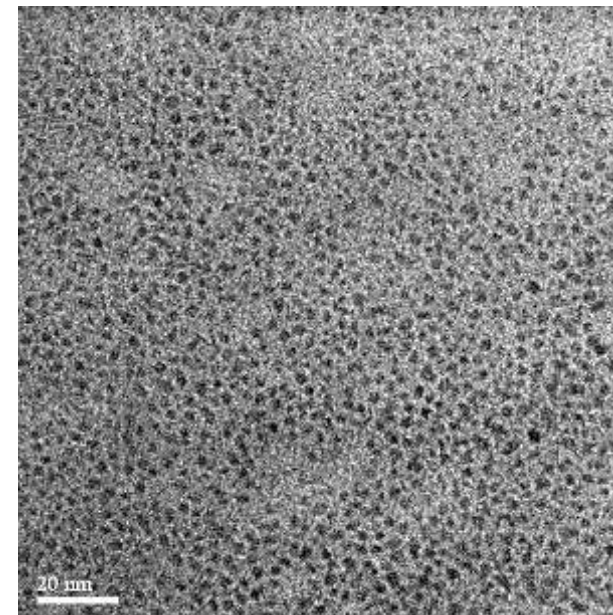
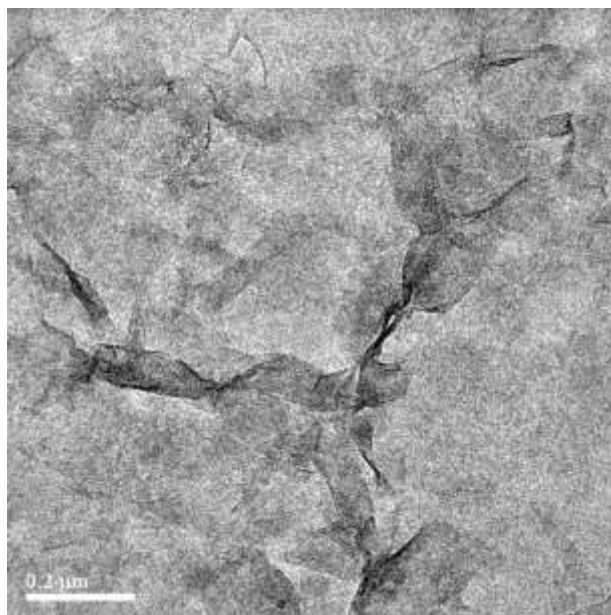
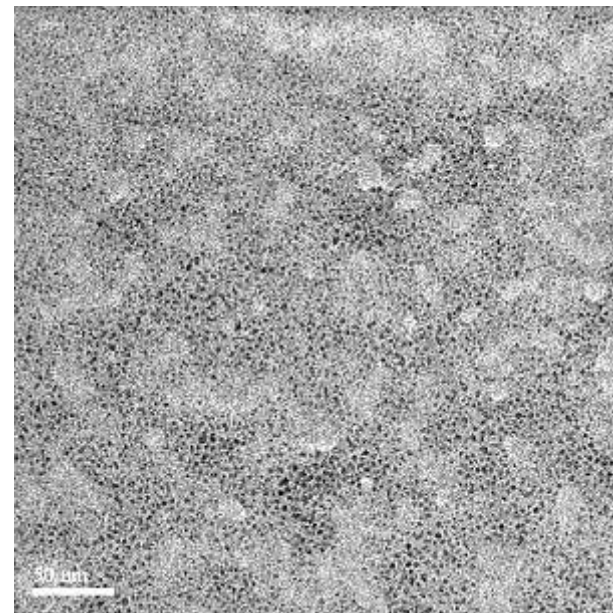
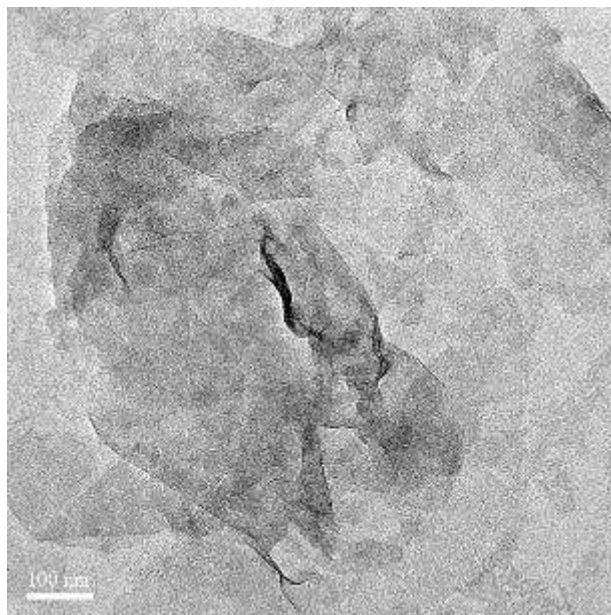
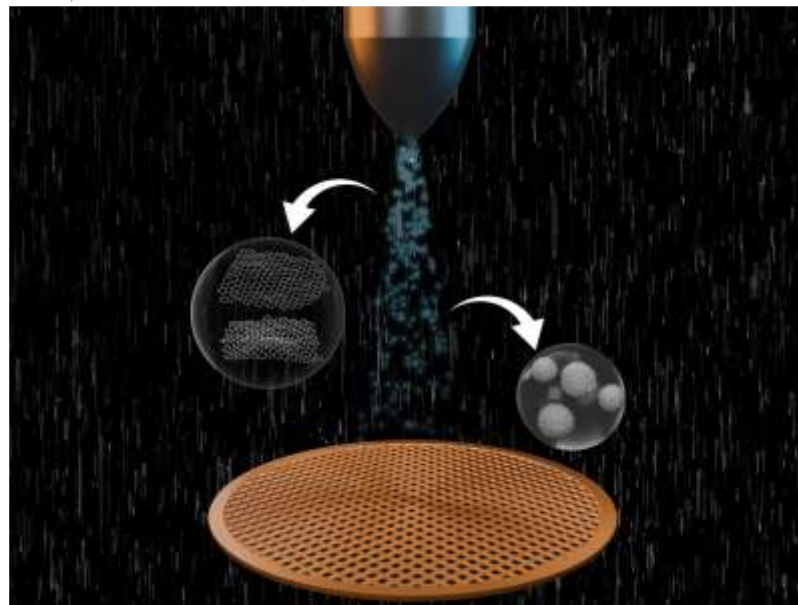


MoS₂ Nanosheets

ChemComm

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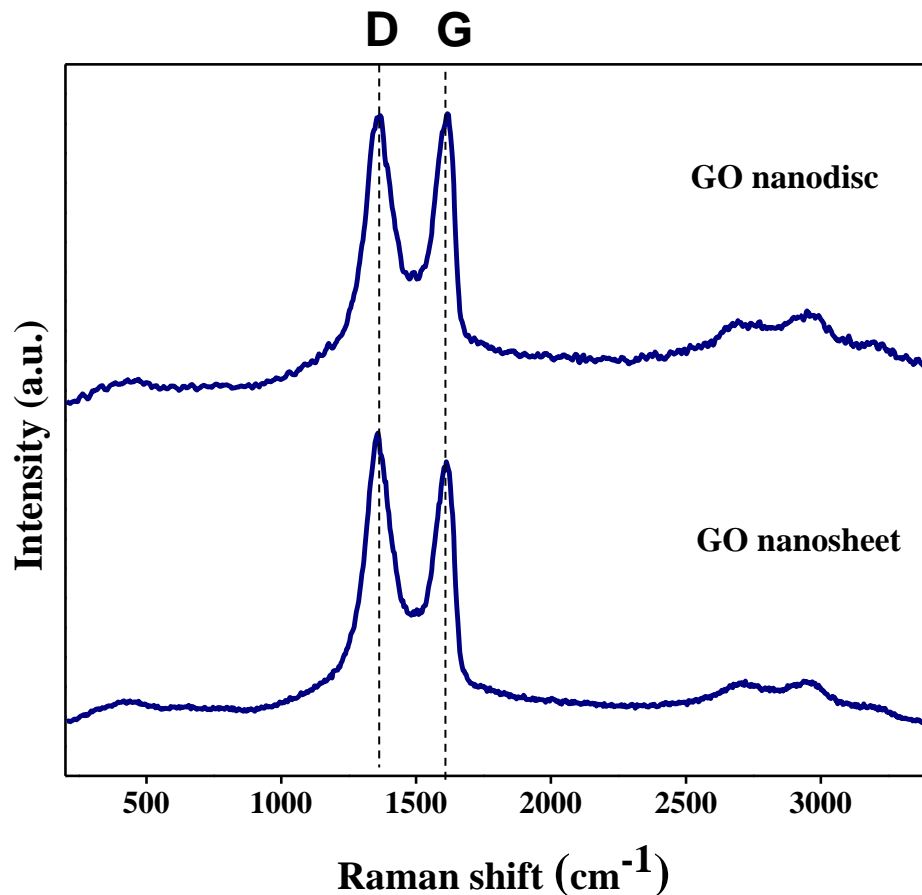
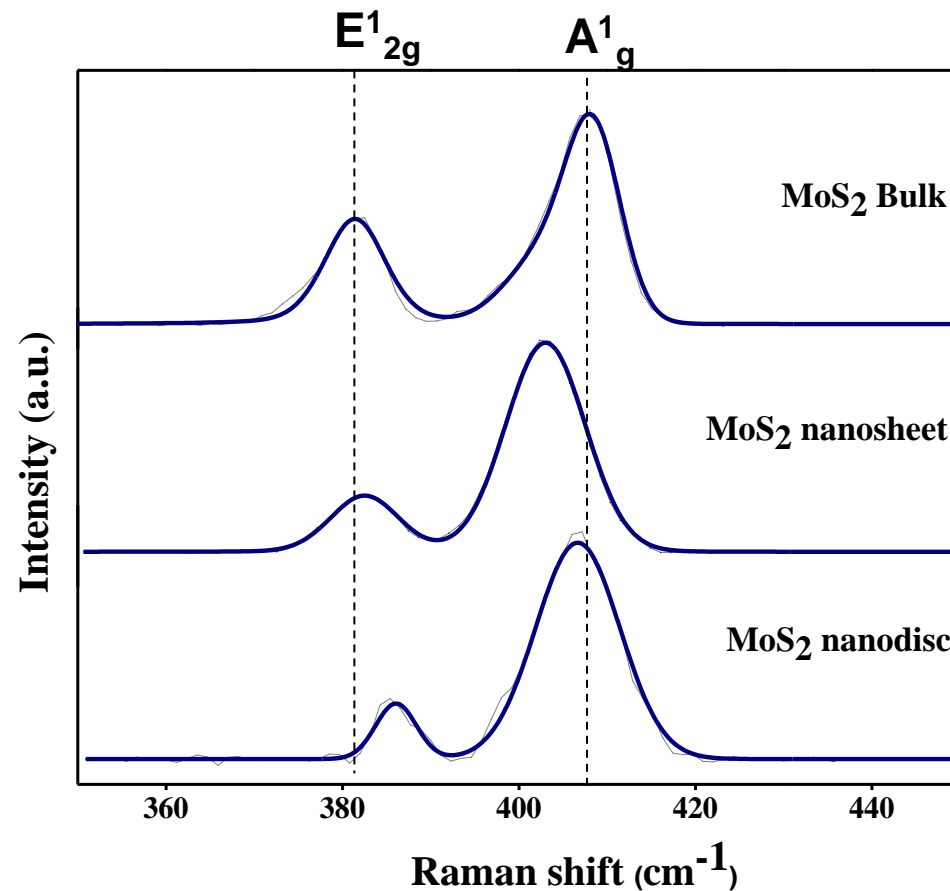
Volume 61
Number 30
18 April 2025
Pages 5529–5676



MoS₂ Nanosheet

MoS₂ Nanoparticles

Raman Spectra of MoS₂ and Graphene Oxide Nanosheets

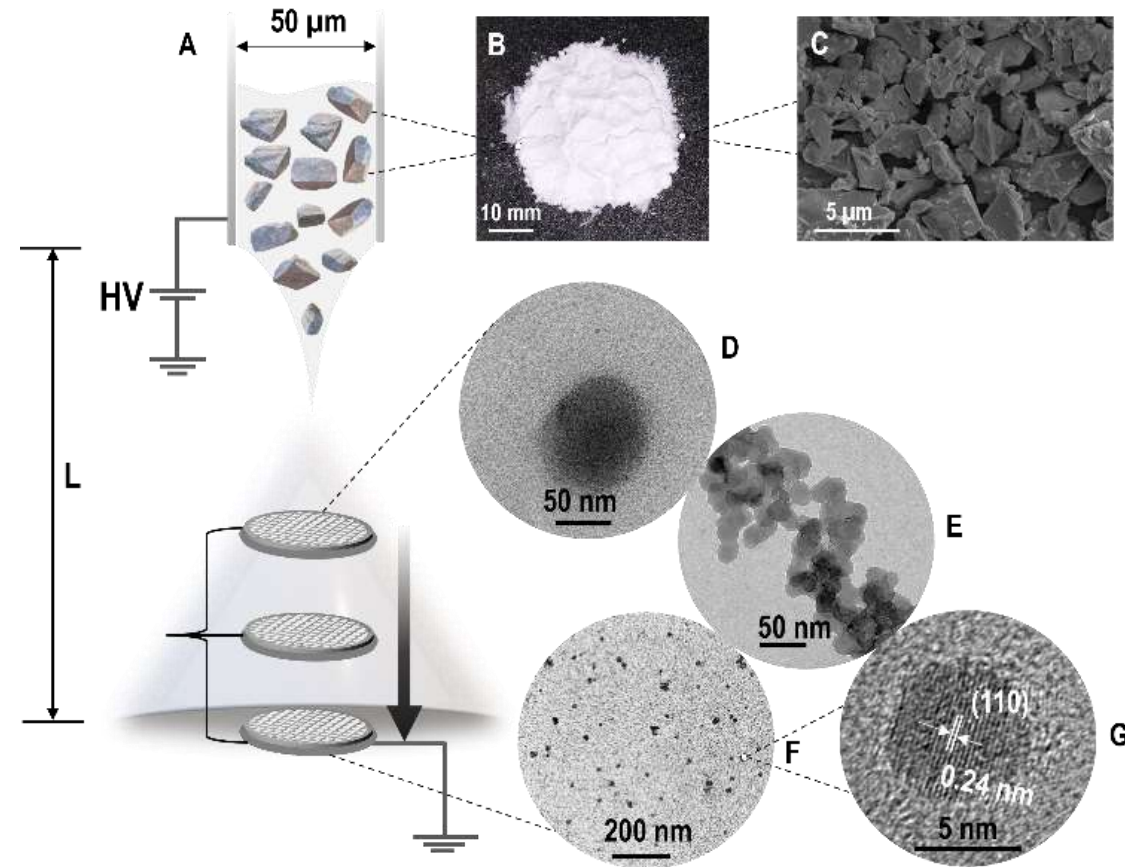


	E^1_{2g} (cm ⁻¹)	A^1_g (cm ⁻¹)
Bulk	381.34	407.67
NS	382.88	402.95
ND	386.01	406.67

Relative peak intensity

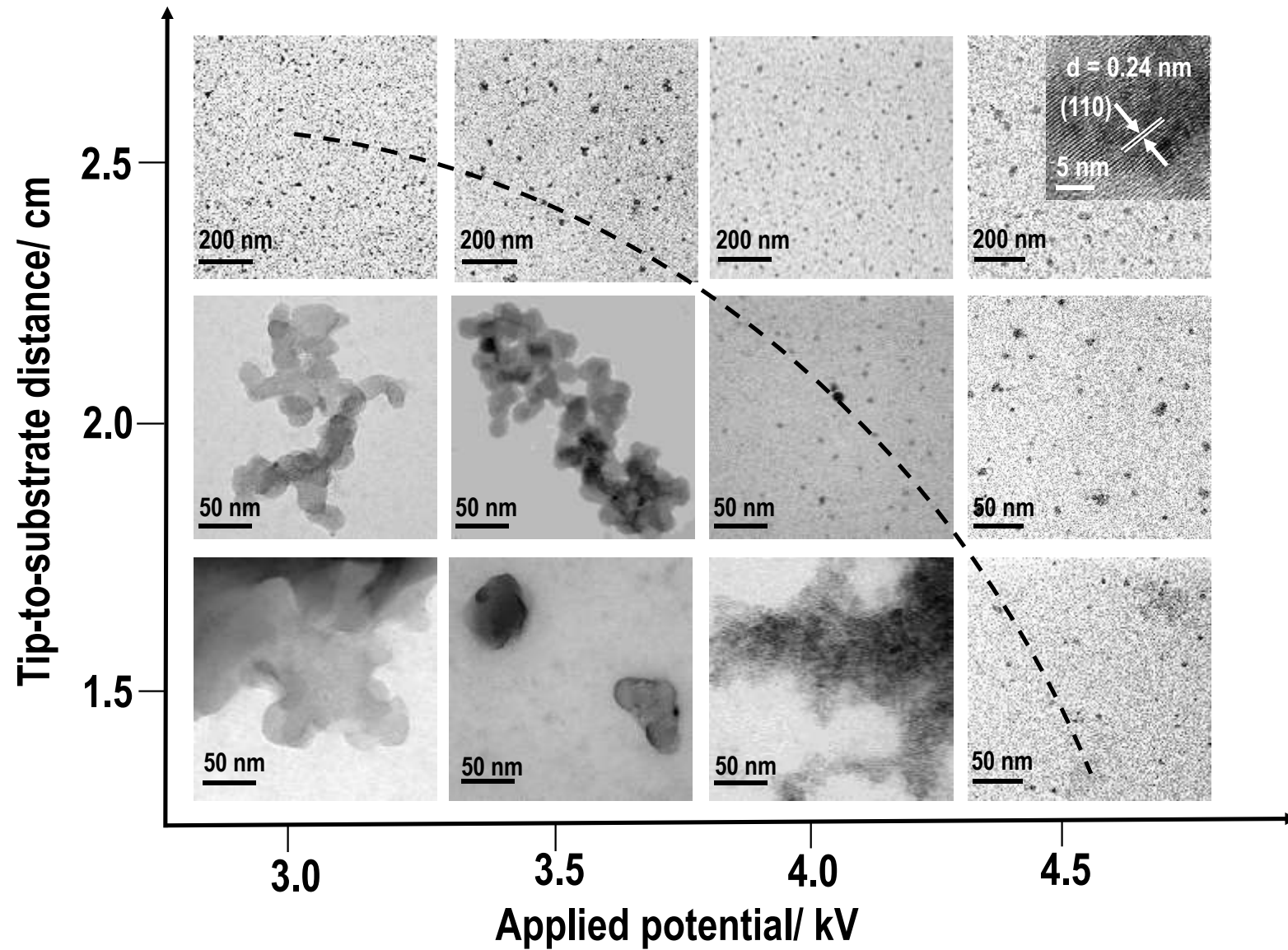
Spoorthi et al. Chem. Comm. 2025

Unveiling steps in the weathering of minerals

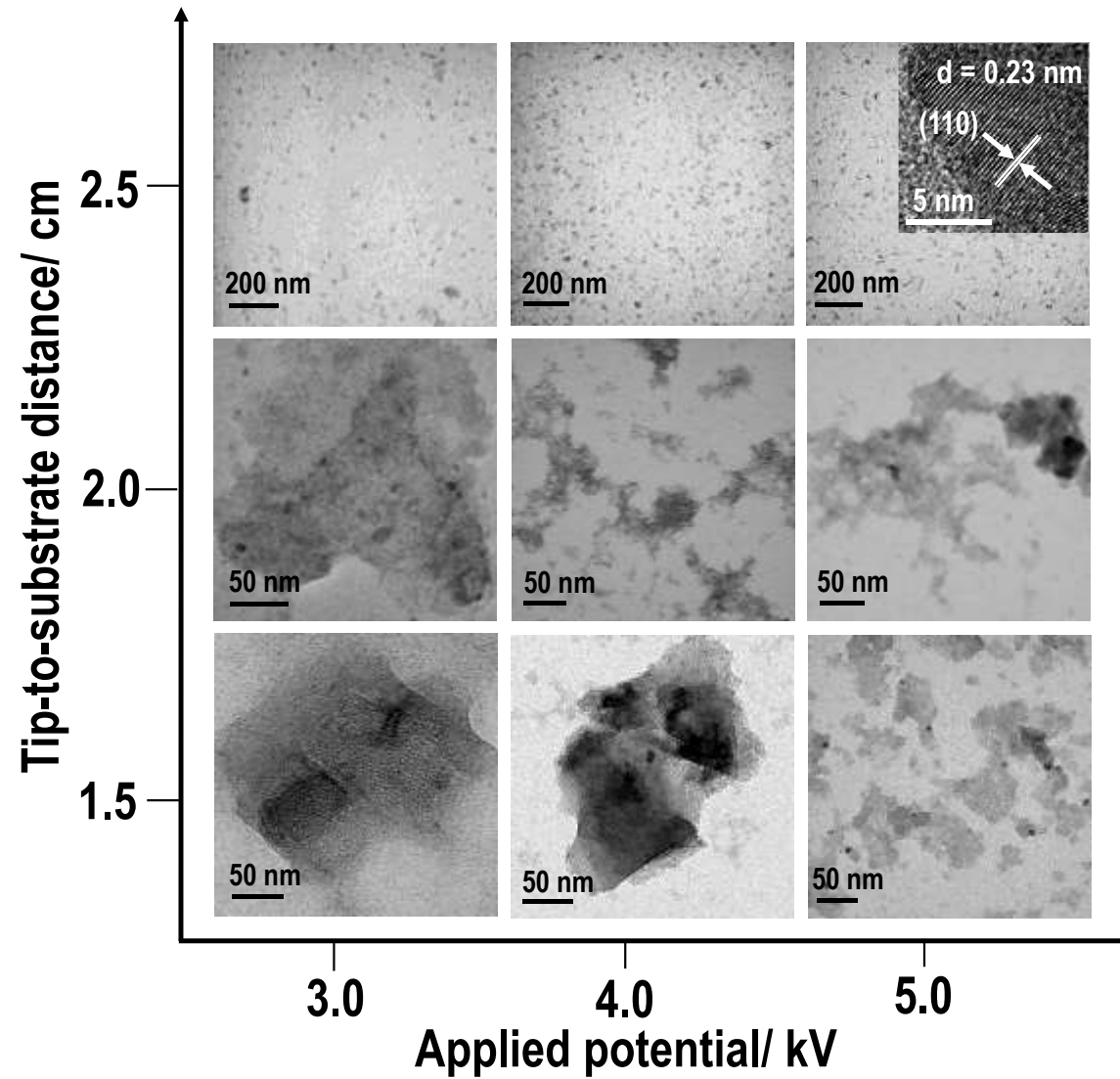


Anubhav Mahapatra et al, Submitted

Disintegration of quartz



Disintegration of ruby



Technology and implementation

PNAS PNAS PNAS

Biopolymer-reinforced synthetic granular nanocomposites for affordable point-of-use water purification

Mohan Udhaya Sankar¹, Sahaja Aigal¹, Shihabudheen M. Maliyekkal¹, Amrita Chaudhary, Anshup, Avula Anil Kumar, Kamalesh Chaudhari, and Thalappil Pradeep²

Unit of Nanoscience and Thematic Unit of E

Edited by Eric Hoek, University of California

Creation of affordable materials for clean drinking water is one of the most promising ways to provide drinking water for all. Combining the composites to scavenge toxic species and other contaminants along with the ability to provide affordable, all-inclusive drinking water without electricity. The critical problem is the synthesis of stable materials that can be used in the presence of complex contaminants in drinking water that deposit and cause health issues. Here we show that such composites can be synthesized in a simple and effective way without the use of electrical power. The nanocomposites have sand-like properties, such as higher shear strength and stability. These materials have been used as a water purifier to deliver clean drinking water. The ability to prepare nanostructures at ambient temperature has wide relevance for water purification.

hybrid | green | appropriate technology | frugal science | developing world



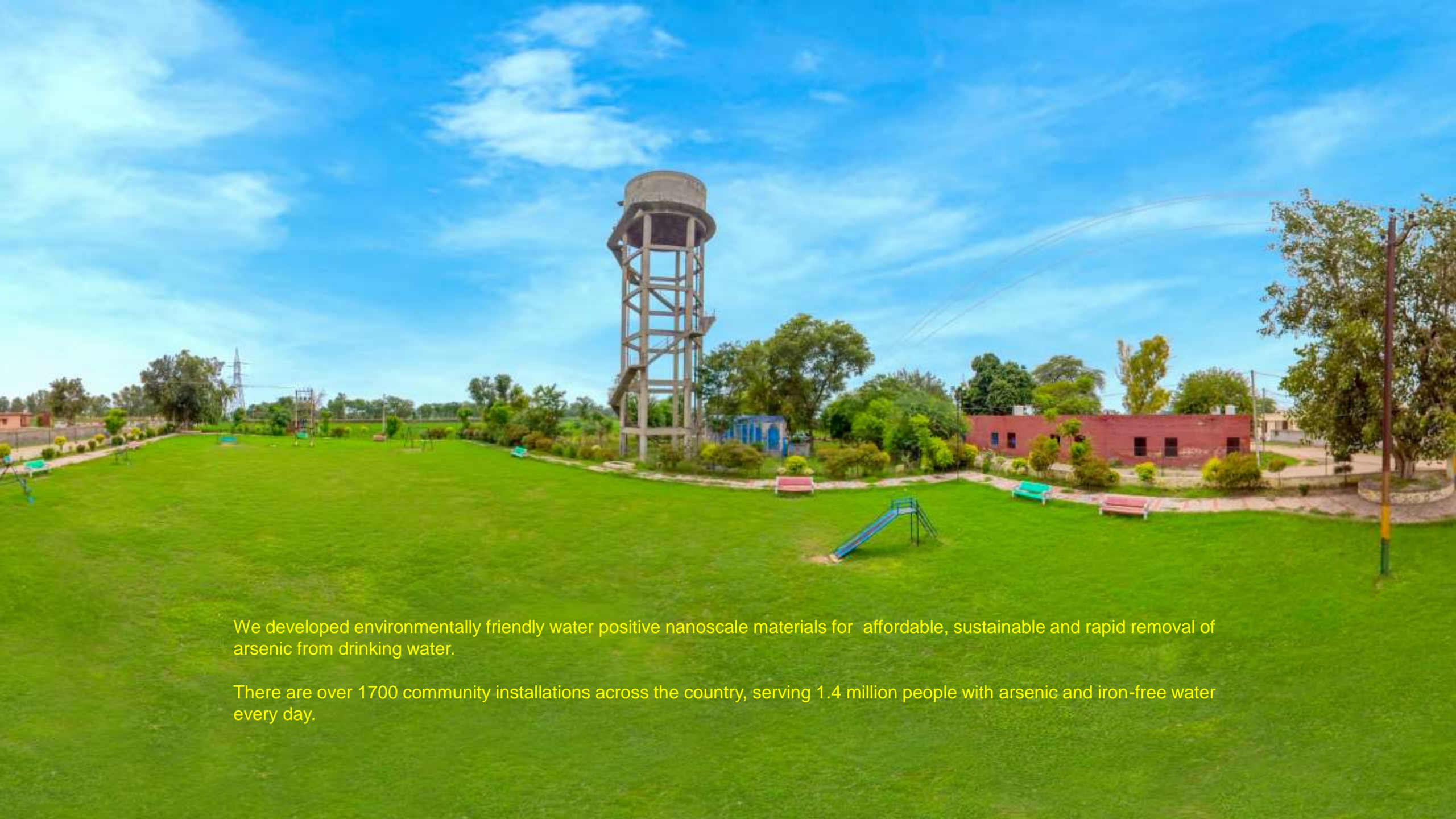
ology Madras, Chennai 600 036, India

, 2013 (received for review November 21, 2012)

not available; and (c) continued retention of silver in the matrix is difficult. We demonstrate a unique family of nanocrystalline chitosan granular composite materials prepared at room temperature through an aqueous route. The granular composition is attributed to abundant -OH groups on chitosan, which help in the crystallization of silver hydroxide and also ensure strong covalent bonding between the particles and the matrix. X-ray photoelectron spectroscopy (XPS) confirms that the composition is rich in silver. Using hyperspectral imaging, the silver leaching in the water was confirmed. We demonstrate a simple way to reactivate the silver nanoparticle composites for continual antimicrobial activity in drinking water. These composites have been developed that can remove contaminants in water. We demonstrate an affordable water purification device based on such composites designed for rural areas and undergoing field trials in India, as a potential solution for widespread eradication of the waterborne disease burden.

a potential solution for widespread eradication of the waterborne disease burden.

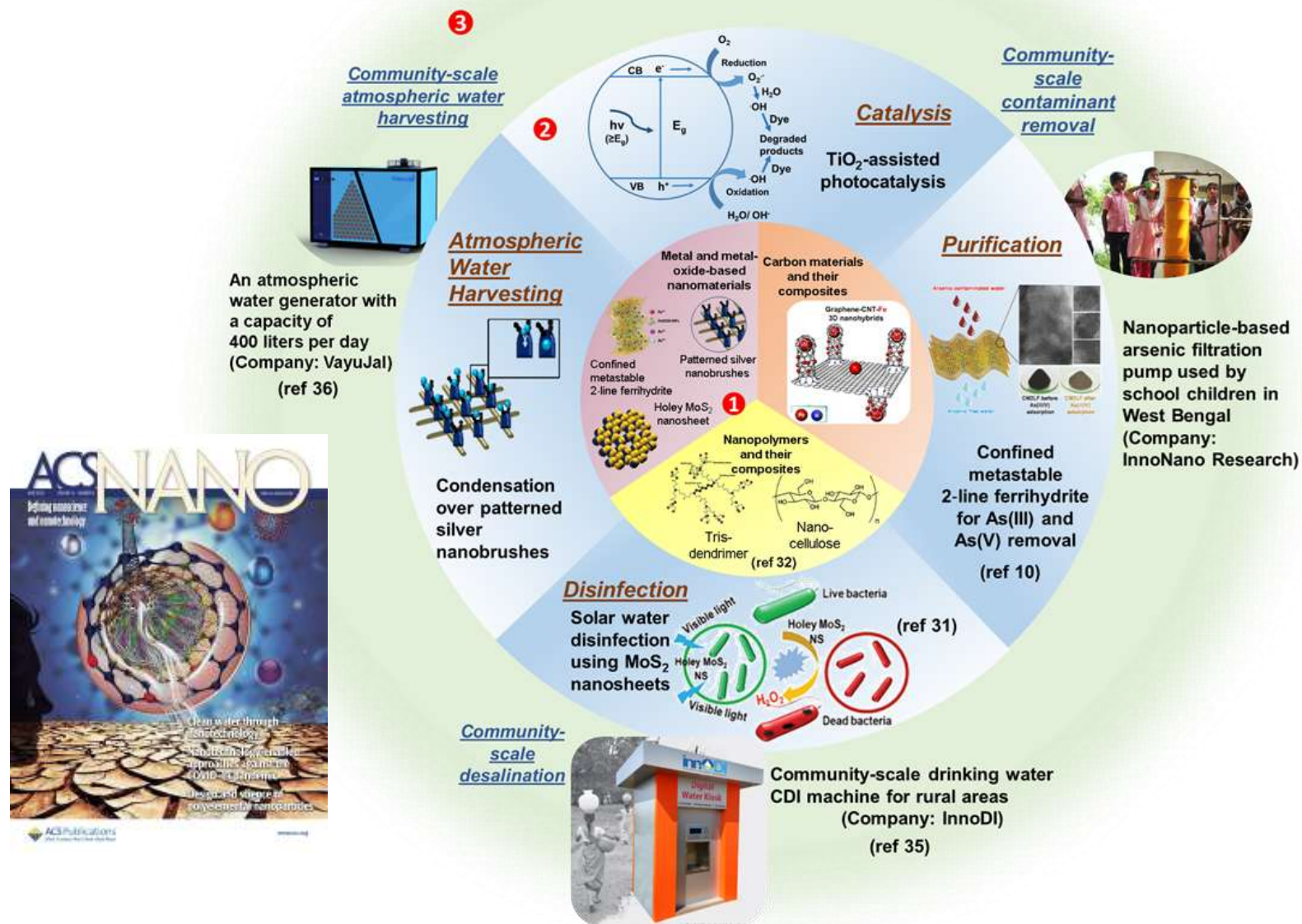
Results and Discussion



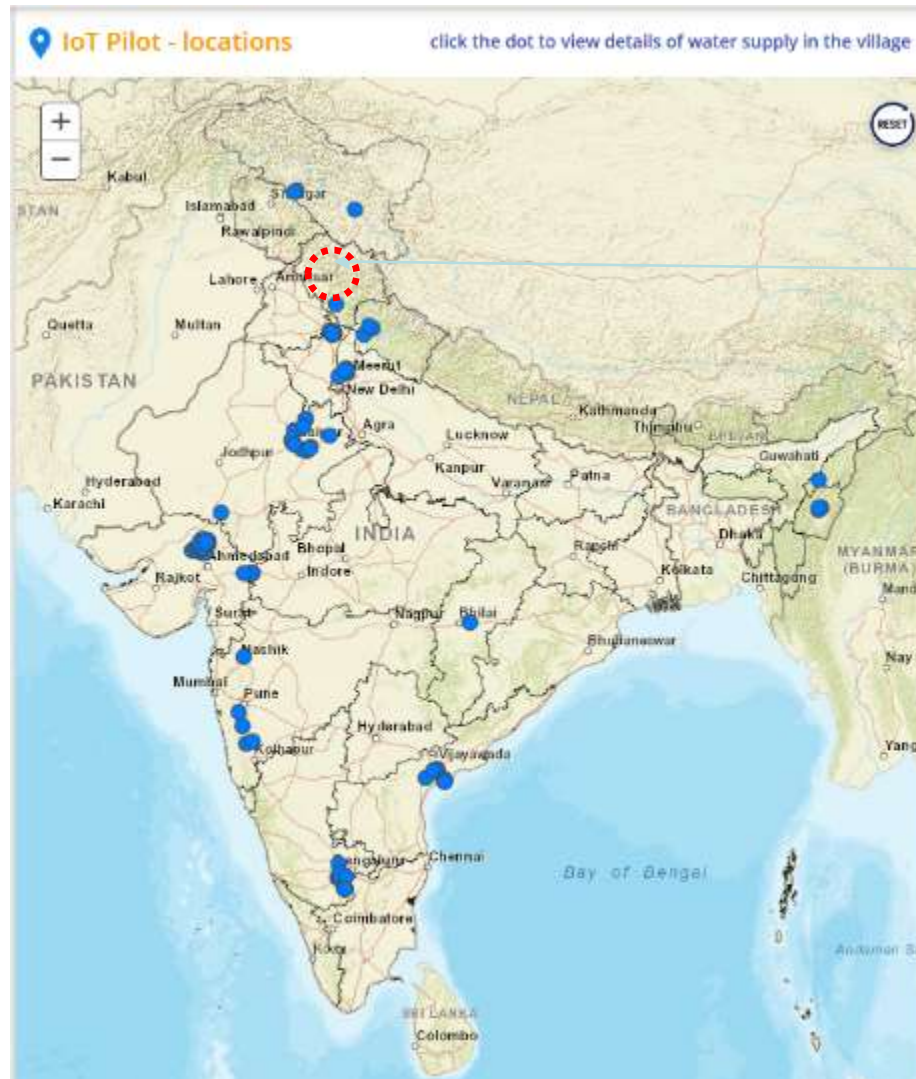
We developed environmentally friendly water positive nanoscale materials for affordable, sustainable and rapid removal of arsenic from drinking water.

There are over 1700 community installations across the country, serving 1.4 million people with arsenic and iron-free water every day.

Evolution of materials to products

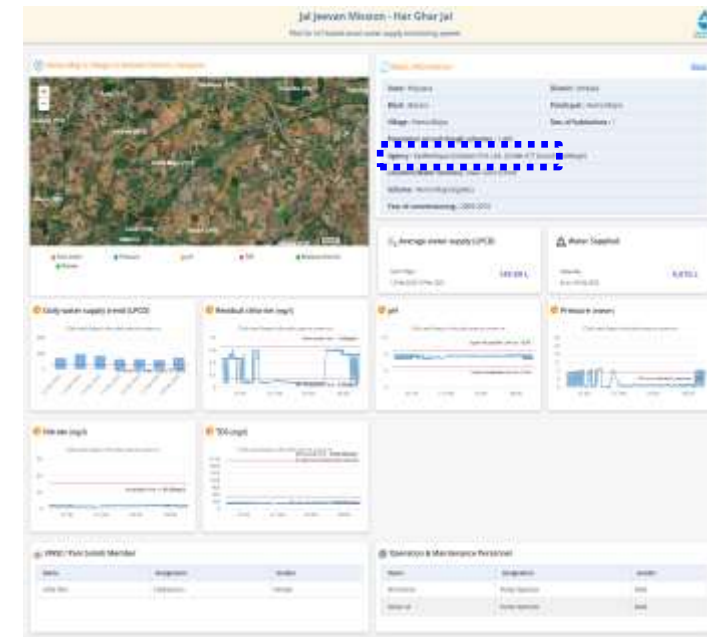


India's water is being monitored



IITM/IISc

Installations made by four companies

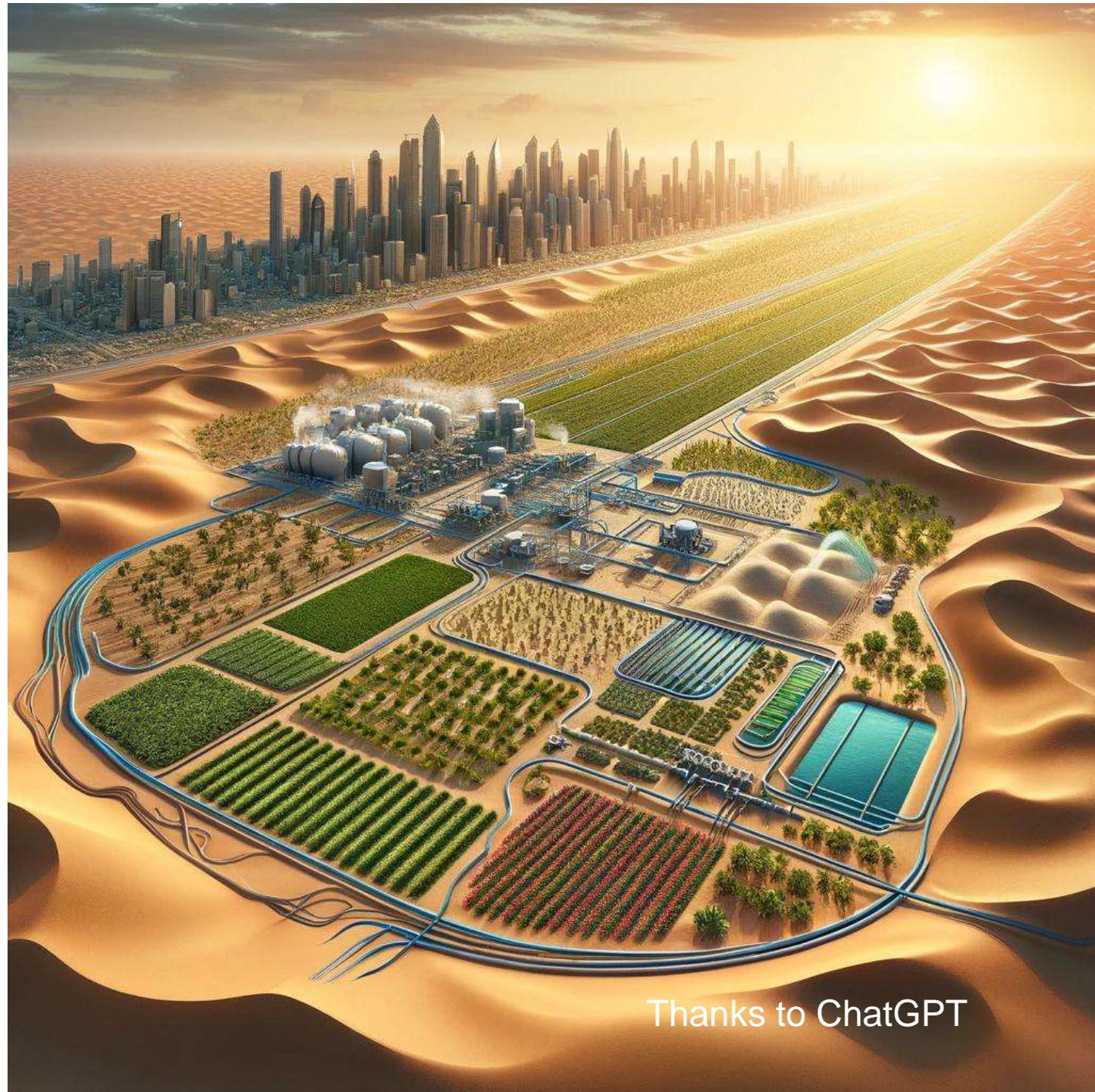




Thanks to ChatGPT

Vision

Make soil using
processed wastewater
and make deserts
bloom.



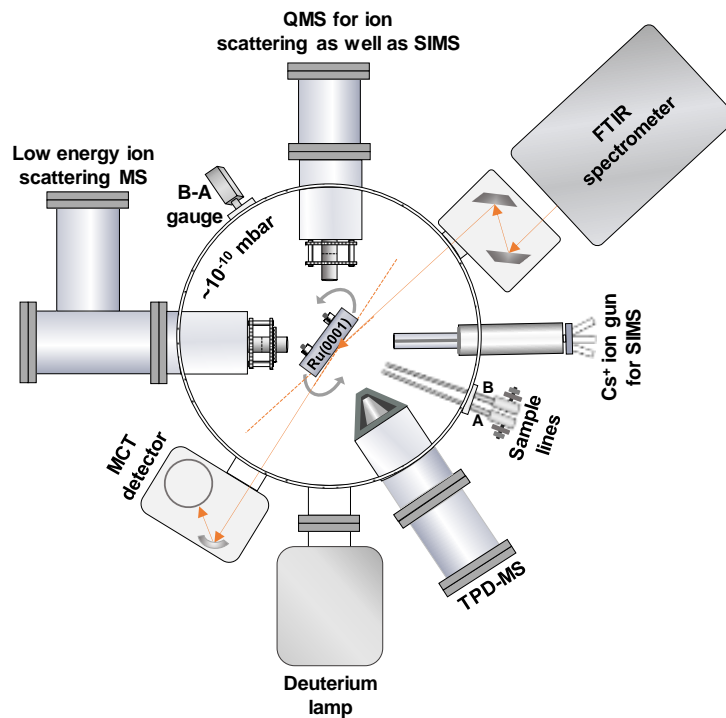
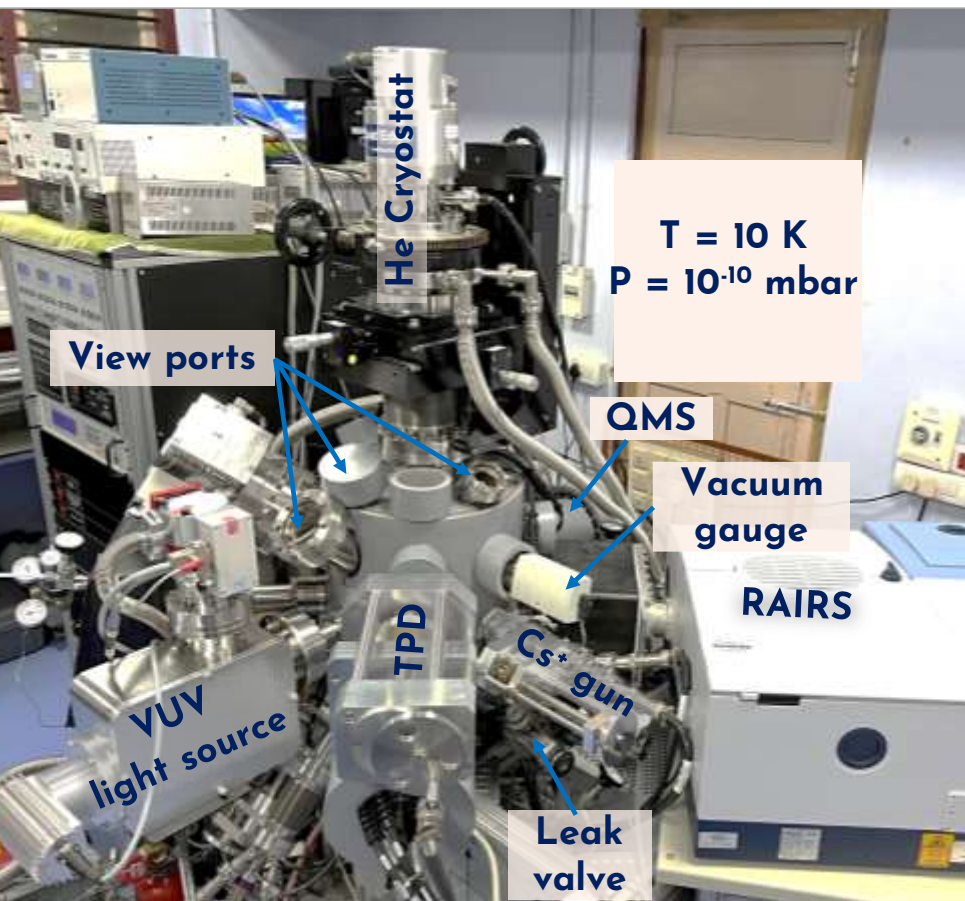
Thanks to ChatGPT



Can Clathrate Hydrates Exist in Space?

Exploring astrobiology

Instrumentation

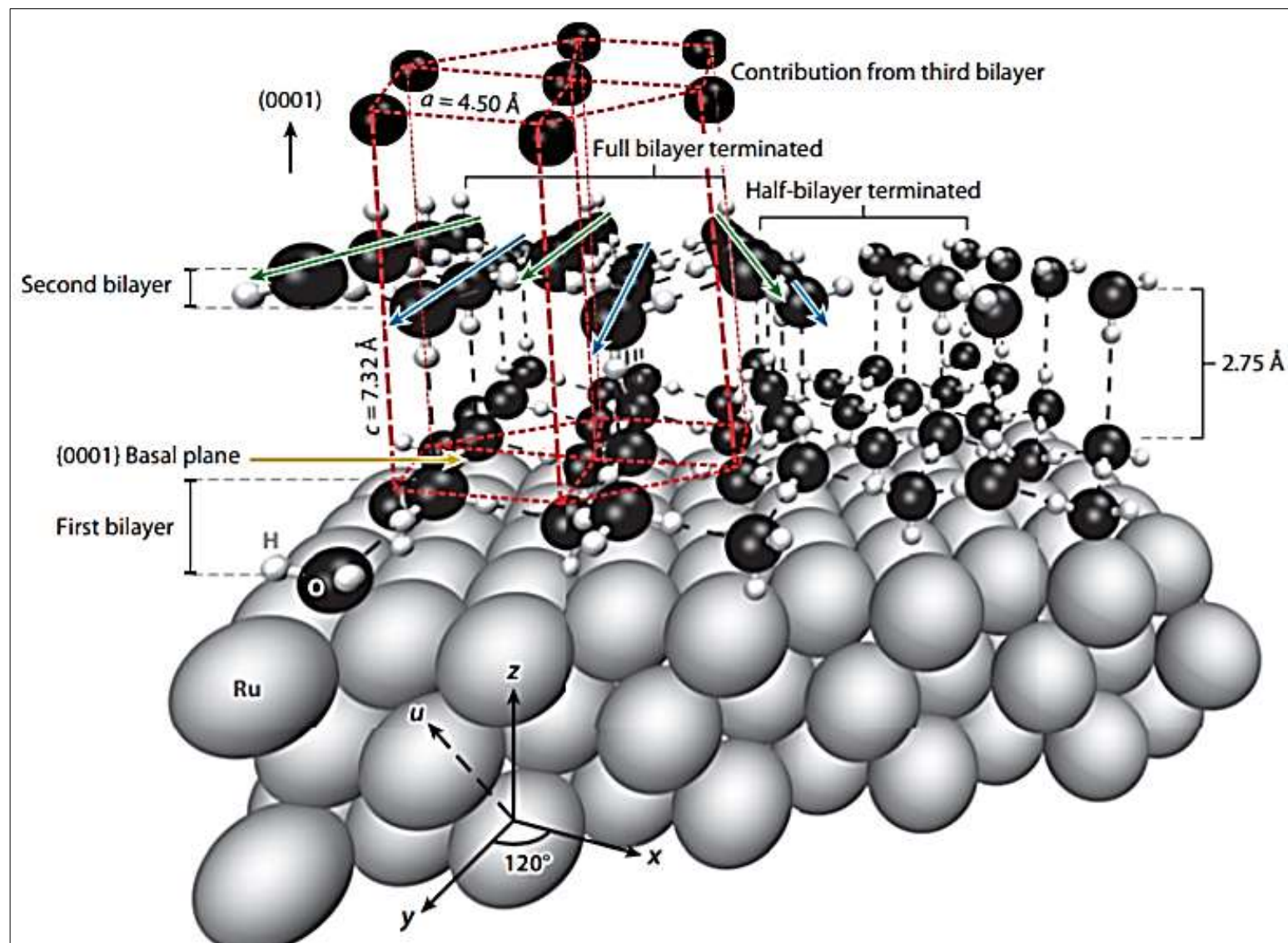


115-400 nm



Bag, S. et al., *Rev. Sci. Instrum.* **2014**, 85, 014103/1-014103/7

Viswakarma, G. et al., *J. Phys. Chem. Lett.*, **2023**, 14, 2823–2829

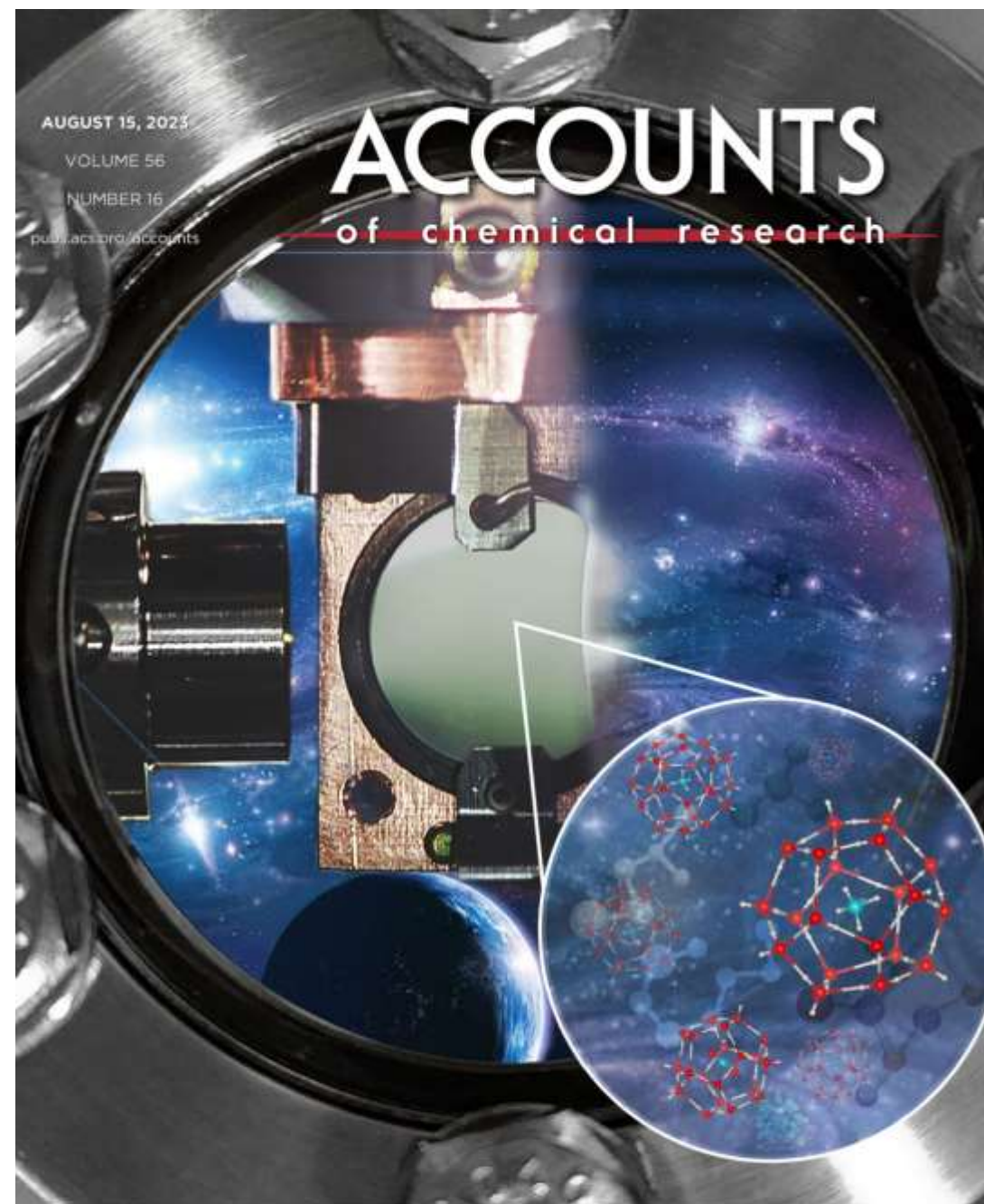


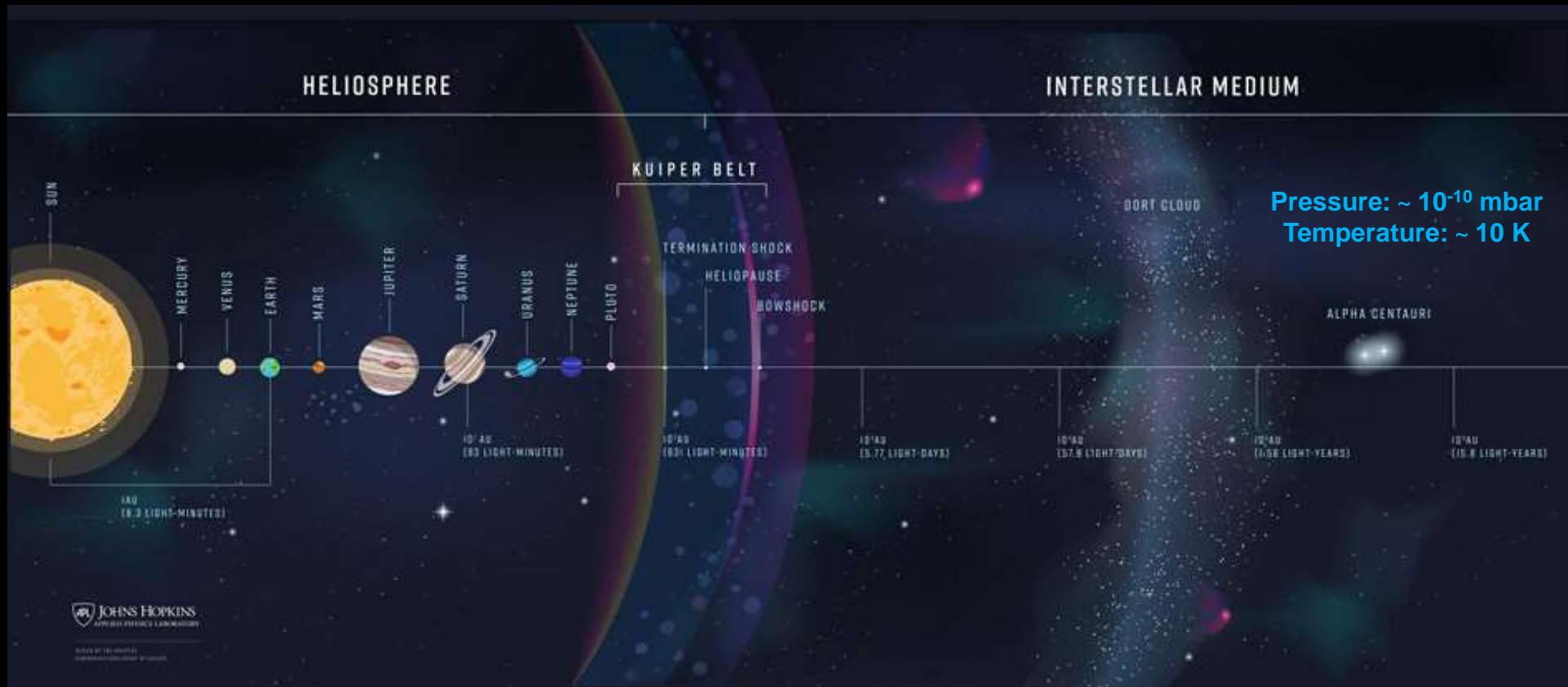
Formation and Transformation of Clathrate Hydrates under Interstellar Conditions

Jyotirmoy Ghosh, Gaurav Vishwakarma, Rajnish Kumar,* and Thalappil Pradeep*

Cite This: <https://doi.org/10.1021/acs.accounts.3c00317>

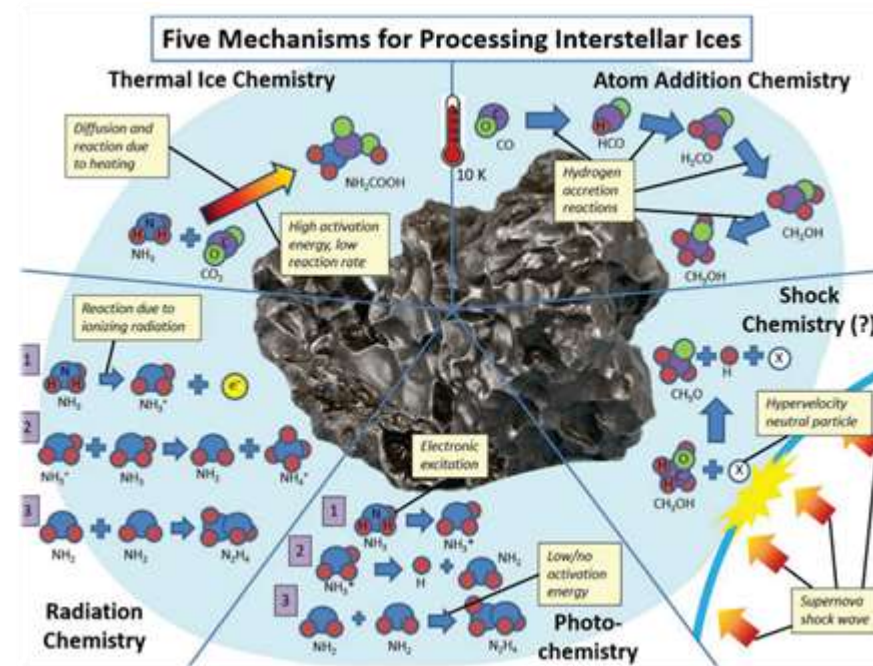
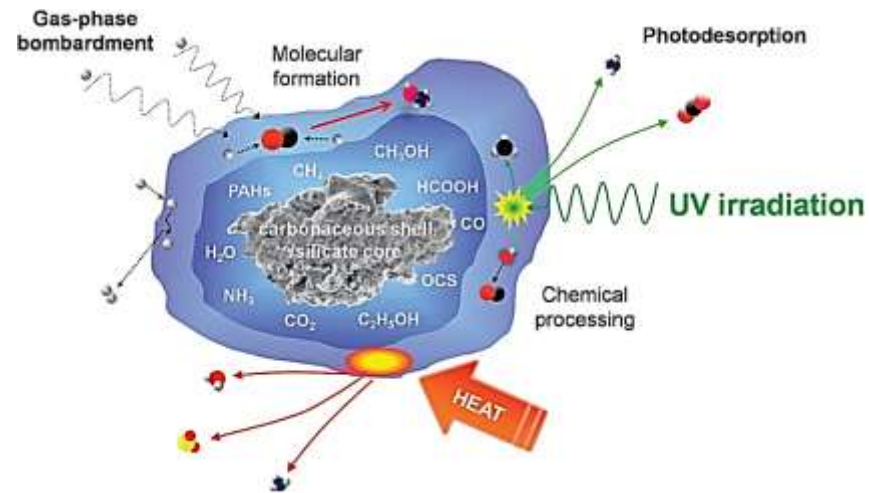
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Diffuse clouds: $T \sim 100$ K, $n \sim 100$ molecules per cm^3
 Dense clouds: $T \sim 10$ - 100 K, $n \sim 10^4$ - 10^8 molecules per cm^3
 On Earth sea level: $T \sim 300$ K, $n \sim 3 \times 10^{19}$ molecules per cm^3

Interstellar ices



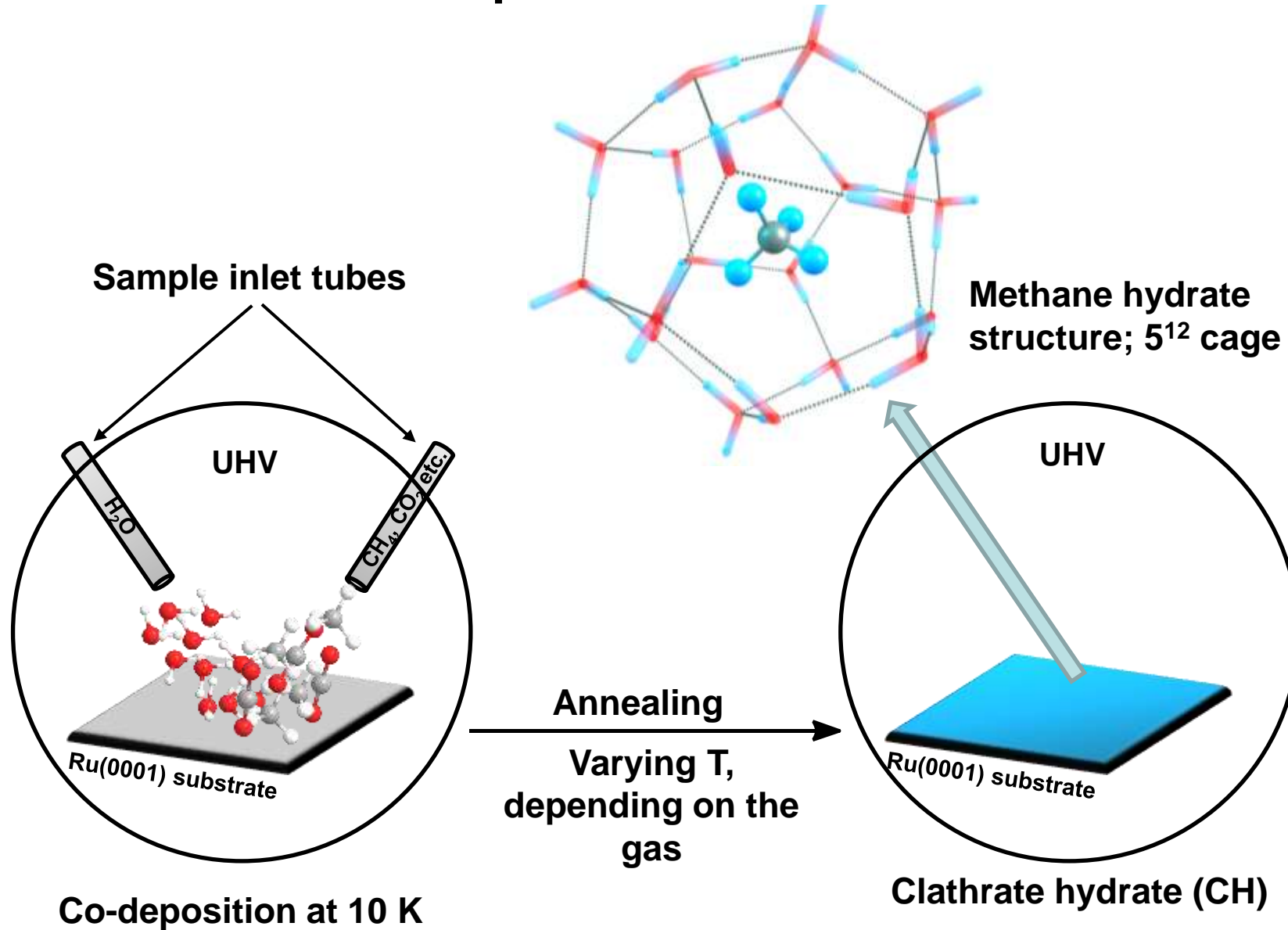
Silicates and carbonaceous material – 0.01-0.5 μm

Arumainayagam, C. R. *et al.*, *Chem. Soc. Rev.*, **2019**, 48, 2293–2314

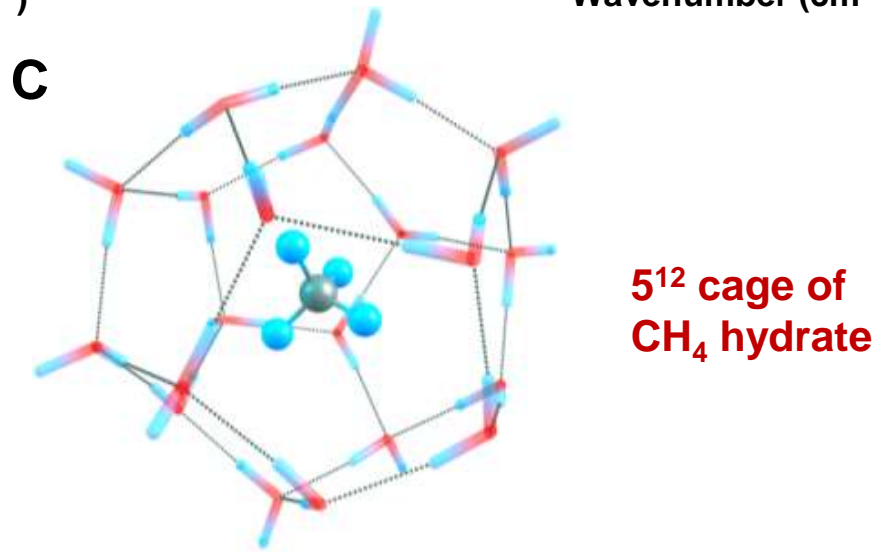
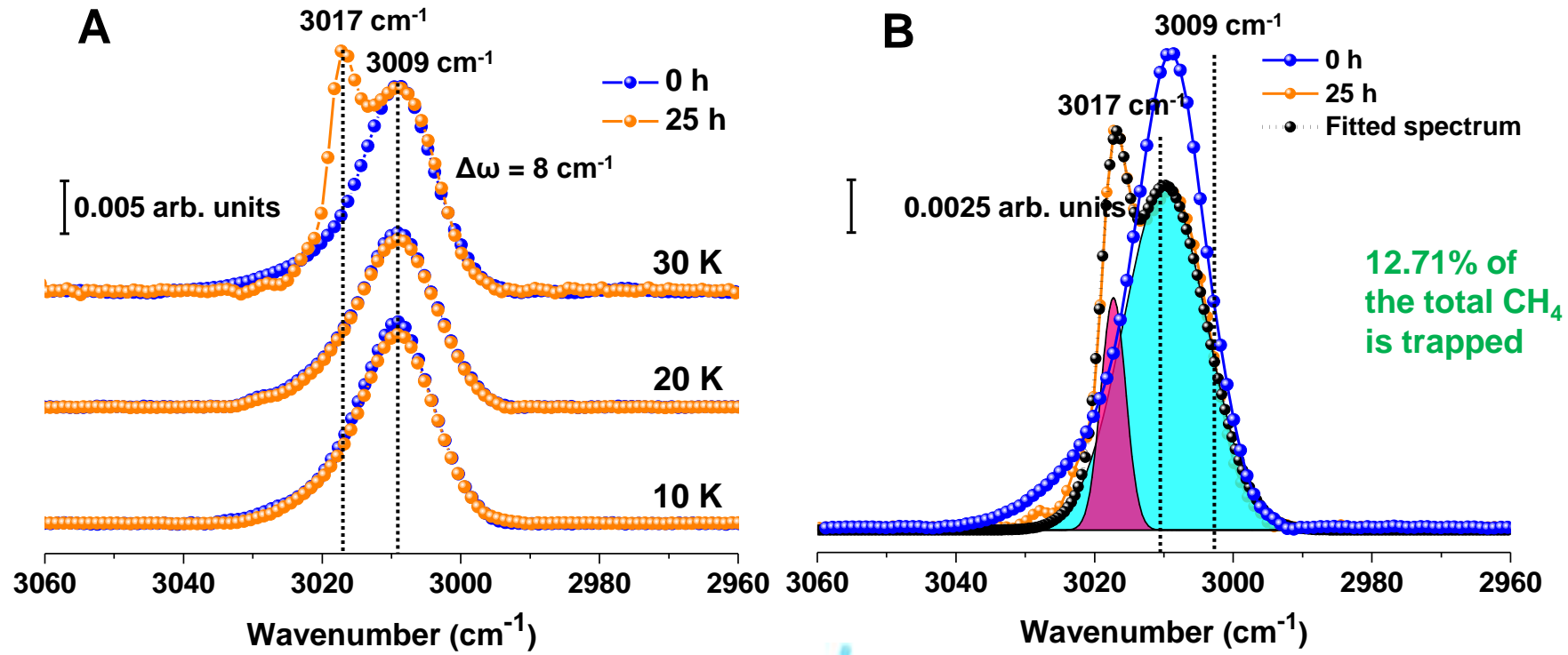
Clathrate hydrates in interstellar environment

Ghosh, J. *et al.*, *Proc. Natl. Acad. Sci. U.S.A.*, **2019**, 116, 1526-1531

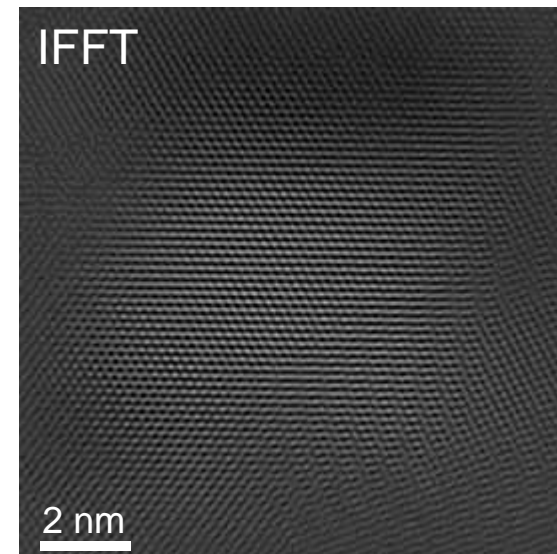
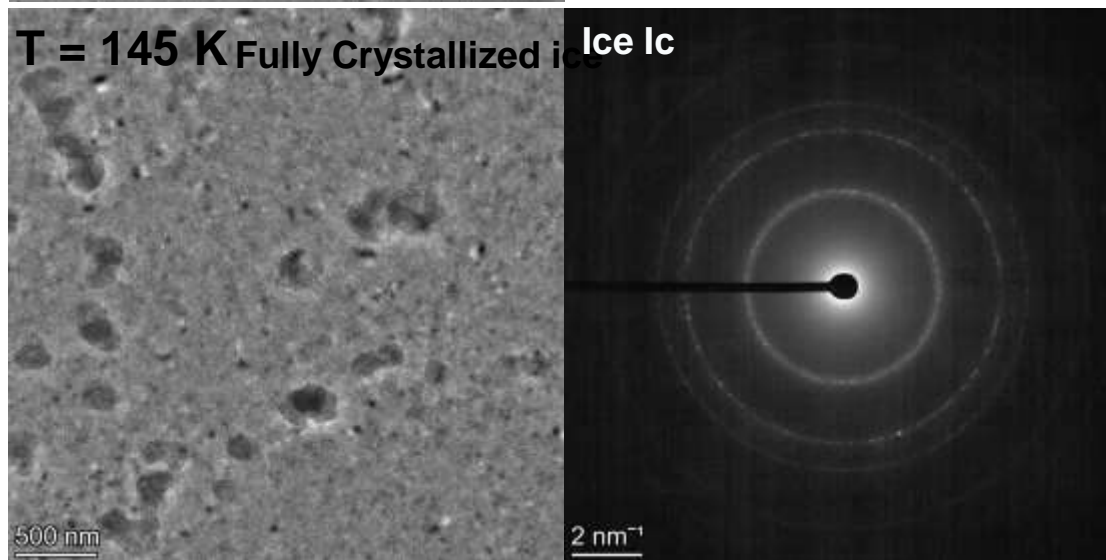
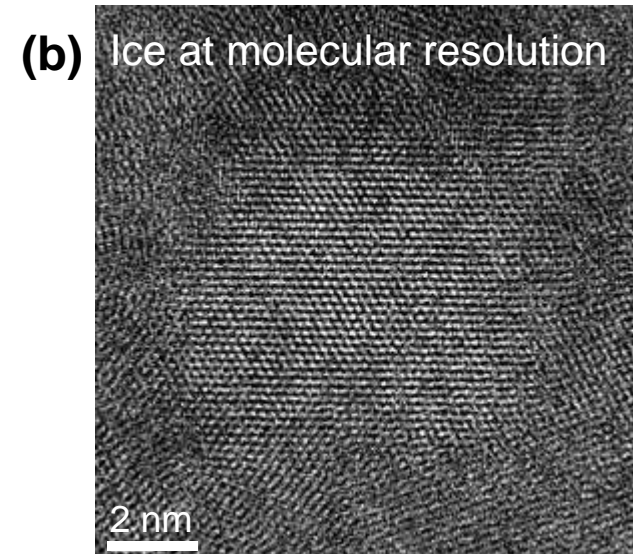
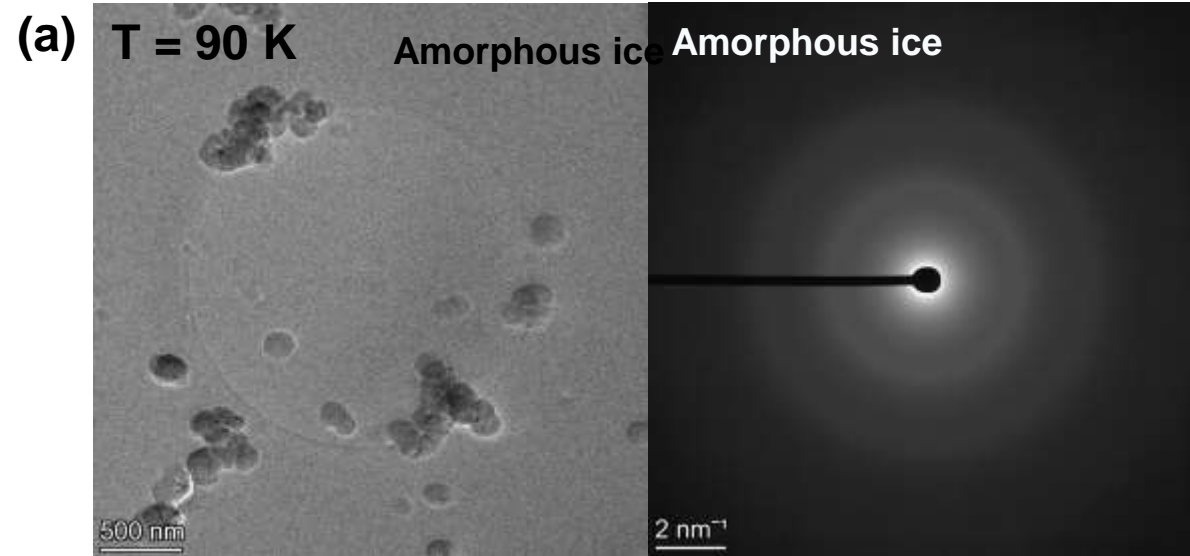
Experimental method



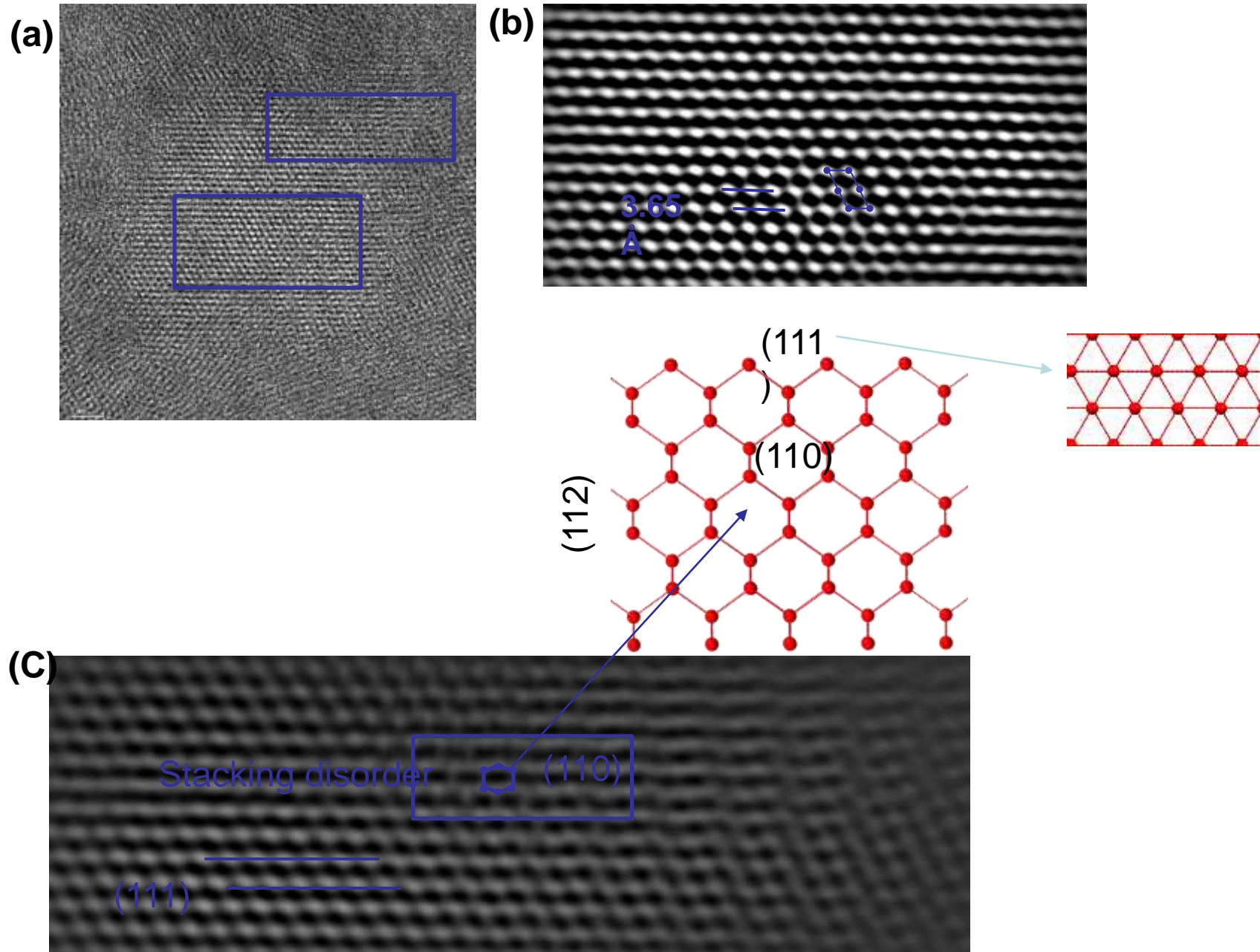
Clathrate hydrates in interstellar environment



Observing growth of crystalline ice from amorphous ice



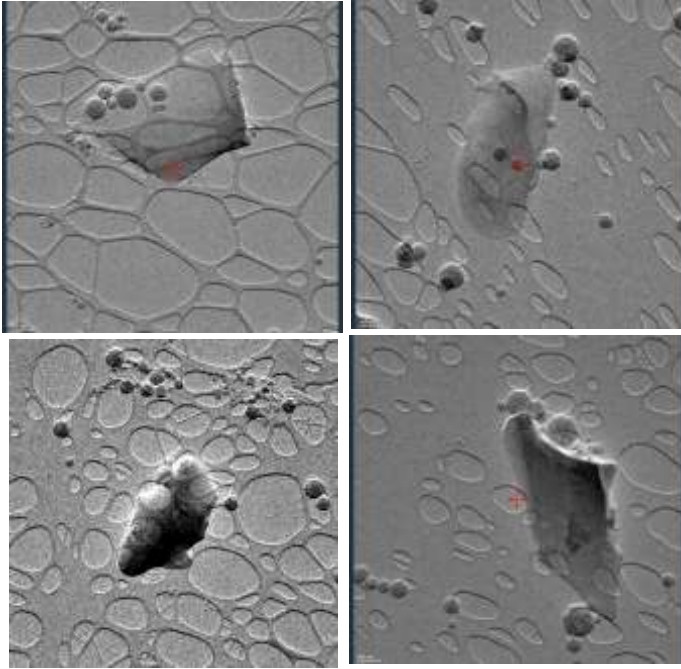
Imaging cubic ice at molecular resolution



Electron diffraction of nanometer-scale crystals of clathrate hydrate

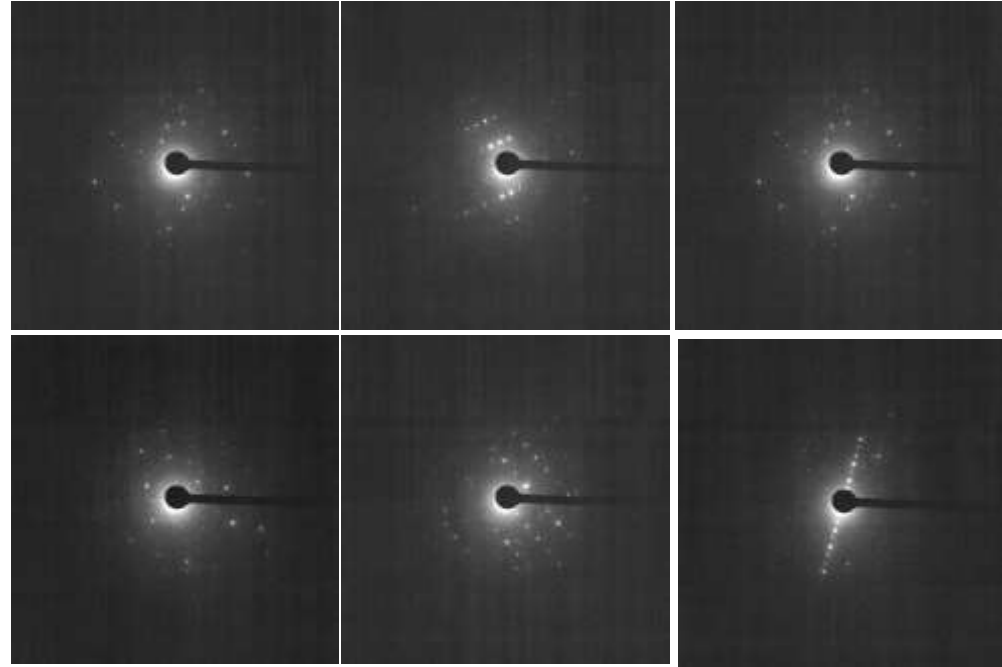
(a)

THF-CH₄ Clathrate hydrate crystal

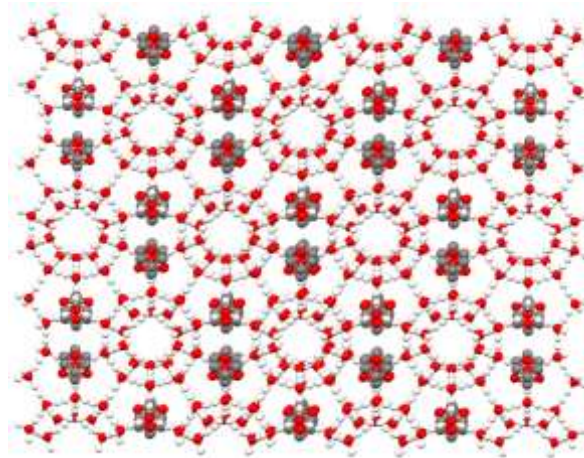
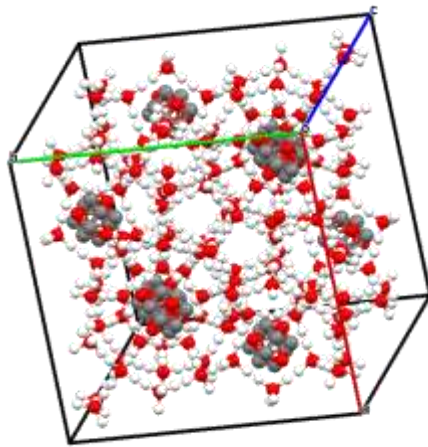


(b)

3 D electron diffraction



(c)

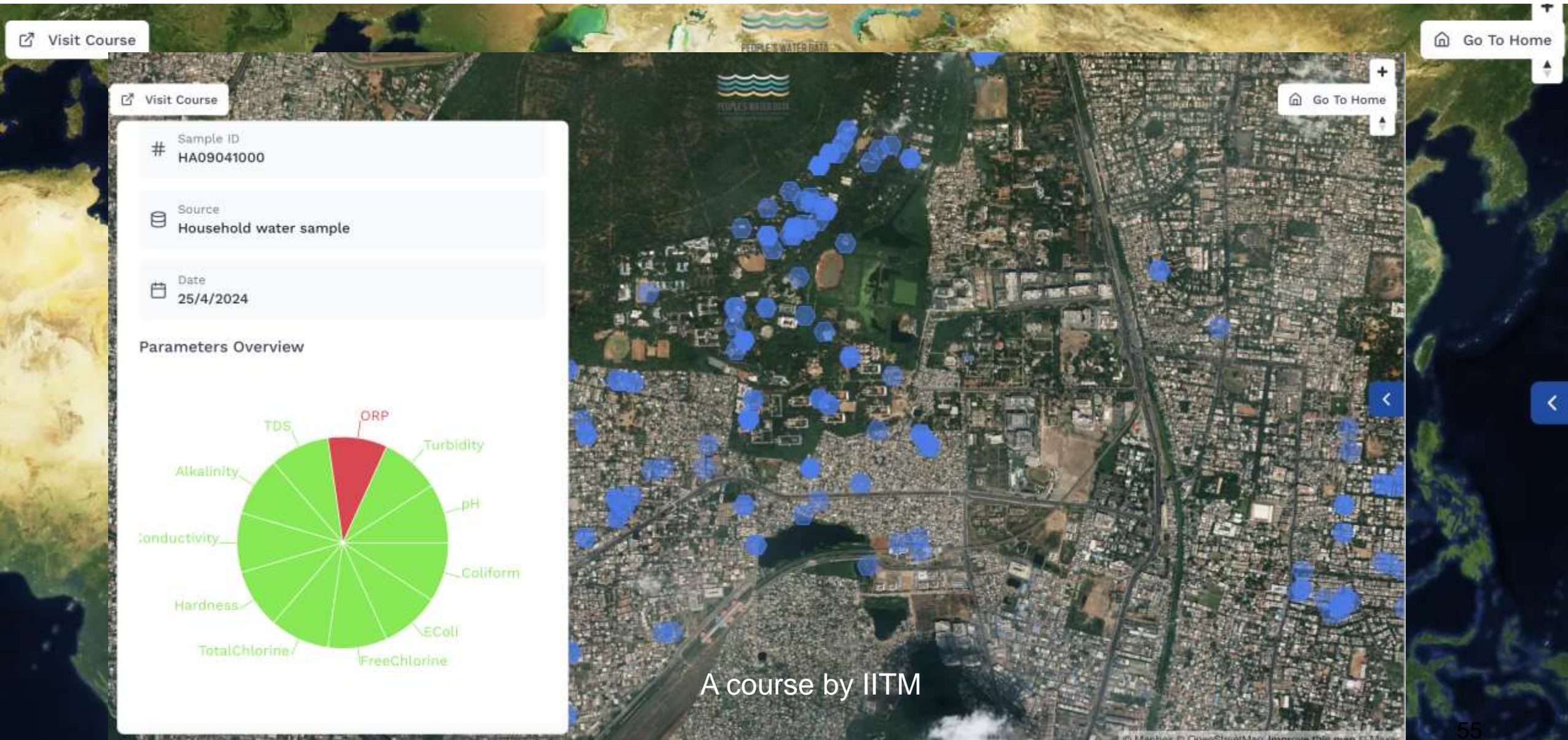


Crystal structure

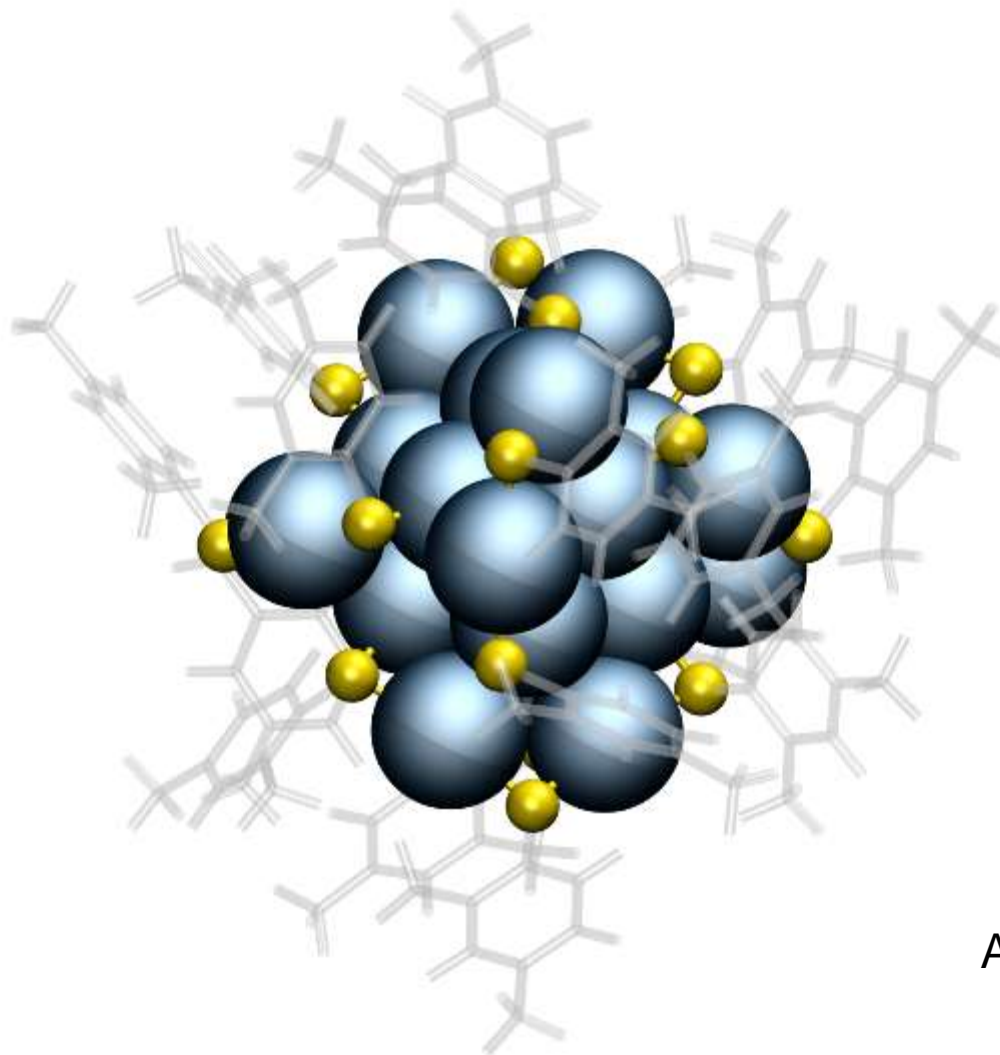


International Centre for Clean Water

People's Water Data

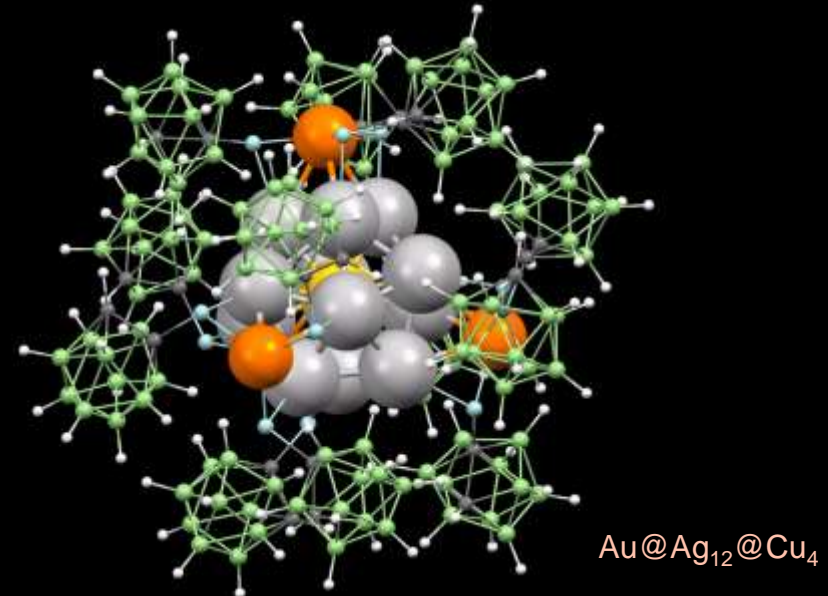
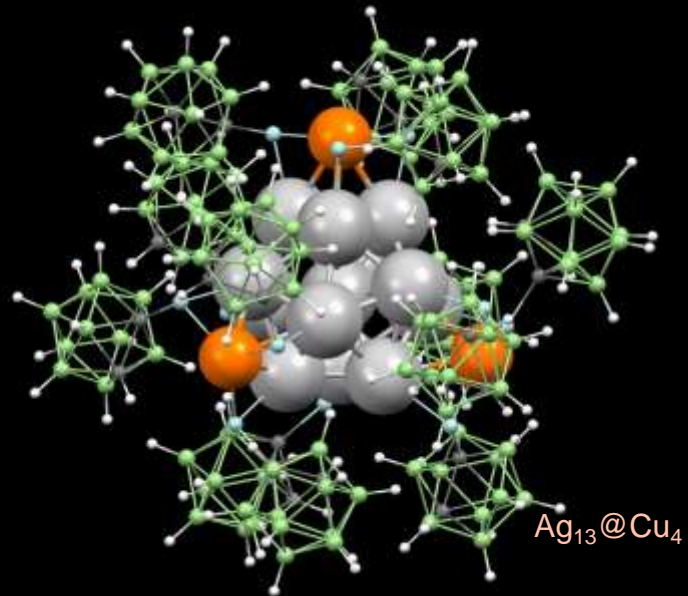
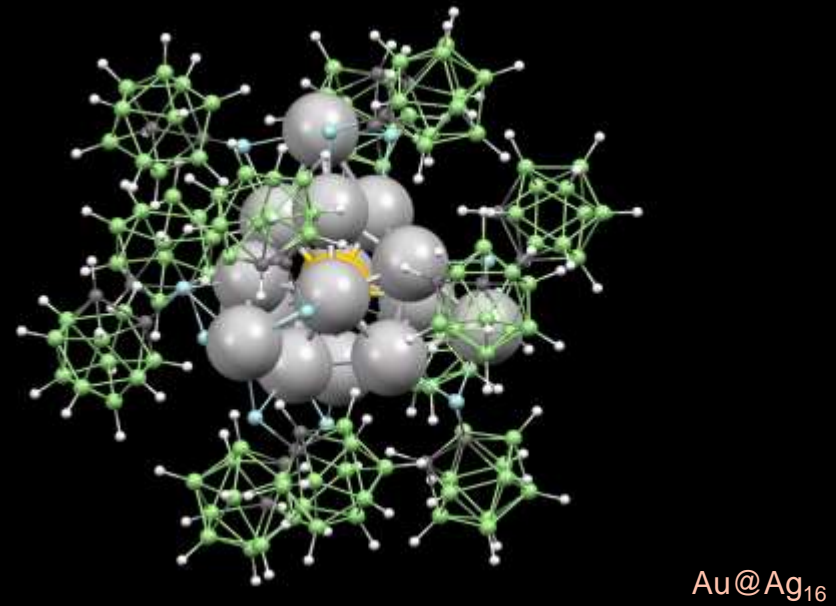
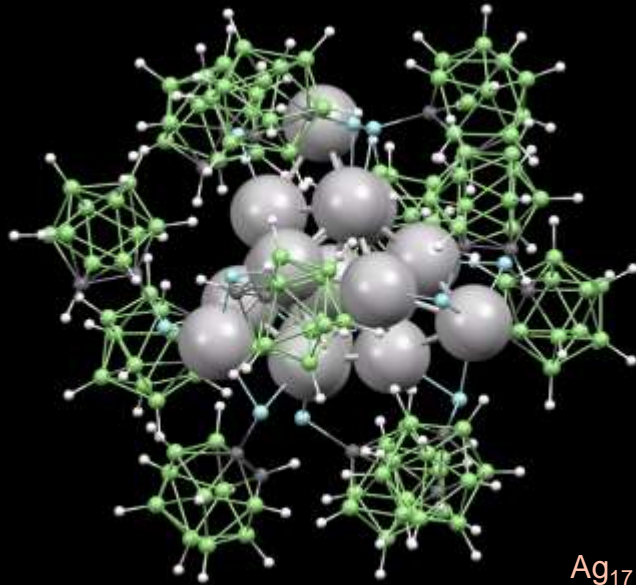


New molecules



$\text{Au}_{25}, \text{Ag}_{25}, \text{Ag}_{29}$

Structure of M_{17} Nanoclusters



Conclusions

Natural minerals break spontaneously in charged water microdroplets

It occurs only in water... so far

Studies on a variety of materials

Facile due to proton-induced slip

Detailed investigations are essential to know more

Implications to the production of specific nanomaterials and soil in general

Other collaborators



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Robin Ras



Manfred Kappes



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Olli Ikkala



Tomas Base



Horst Hahn



Biswarup Pathak



K. V. Adarsh



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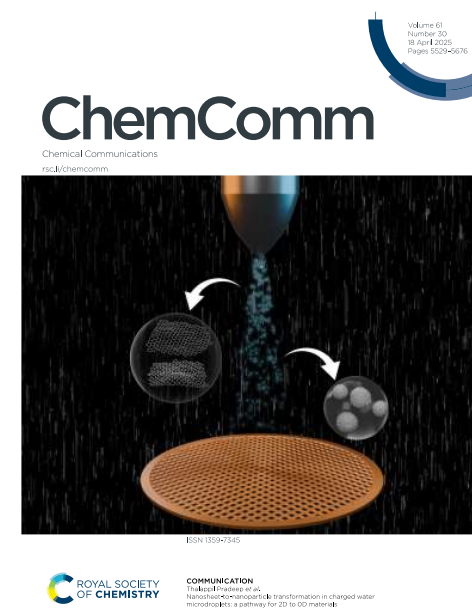
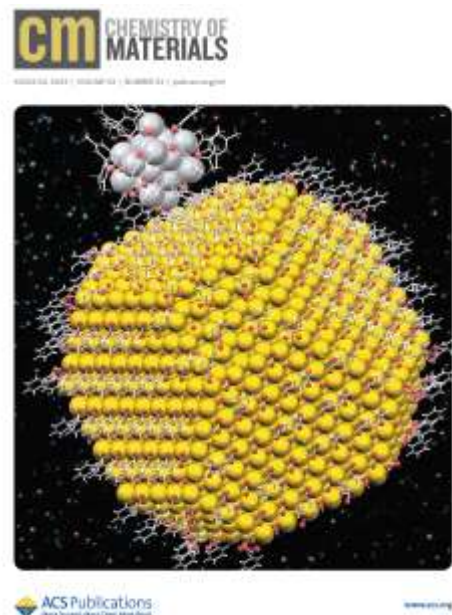
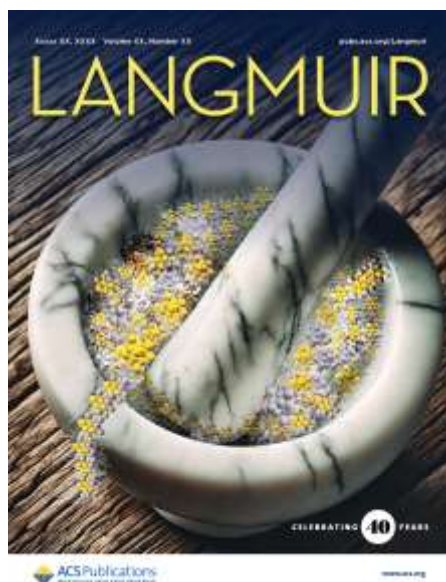


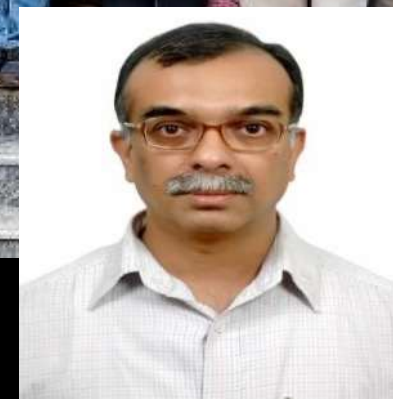
Vivek Polshettiwar

Department of Science and Technology

Institute of Eminence

Many Outstanding Individuals







Indian Institute of Technology Madras



Bhaskar Ramamurthi/V. Kamakoti

Thank you all

