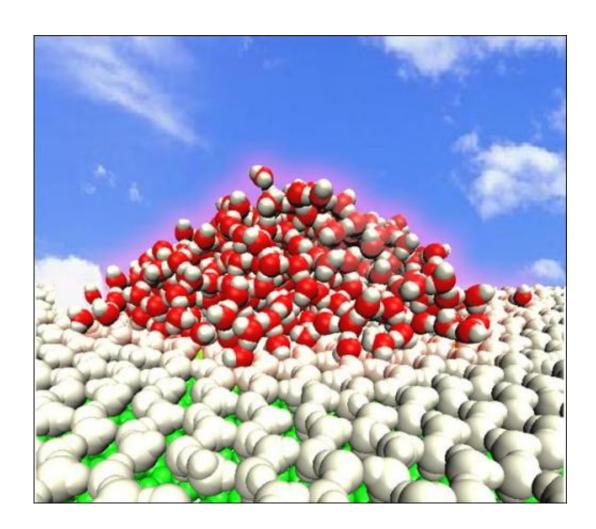


Scientists challenge classical phenomenon that water always completely wets water

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A liquid water droplet on a water monolayer on a polar model surface has been observed at room temperature, and has been termed an 'ordered water monolayer that does not completely wet water.' The blue sky and white cloud represent room temperature; green spheres represent solid surfaces; silver spheres represent an ordered water monolayer; and red and white spheres represent oxygen and hydrogen atoms of the liquid water molecules. Credit: Science China Press



Viewed macroscopically at room temperature, when water contacts other water, it will spread out and finally both mix together. In 2009, Chinese researchers first reported observing the formation of a liquid water droplet on a water monolayer at room temperature, and termed the phenomenon an ordered water monolayer that does not completely wet water. Since then, this phenomenon has been observed on many real surfaces by numerical simulations and experiments by research groups worldwide.

The molecular scale behavior of <u>water</u> at a solid/liquid interface holds fundamental significance in a diverse set of technical and scientific contexts, ranging from the efficiency of oil mining to the activity of biological molecules. Recently, it has become recognized that both the physical interactions and the <u>surface</u> morphology have significant impact on the behavior of interfacial water, including the water structures and wetting properties of the surface.

In a new review, Chunlei Wang, Yizhou Yang and Haiping Fang of the Shanghai Institute of Applied Physics report recent advances in atomlevel understanding of interfacial water that exhibits an ordered character on various solid surfaces at room or cryogenic temperature.

In recent decades, they report, "with the development of computer simulation tools and experimental tools, it is possible to obtain atomic level pictures of the water structure and the complex hydrogen bond network near the interfaces."

"Utilizing these tools, researchers have revealed that water structure may show appealing ordered structures depending on the surface morphology both at cryogenic temperatures and <u>room temperature</u>," they explain. "These ordered structures of water not only exhibit novel pictures but



also affect the various surface properties, such as the surface electrochemical property, surface wetting behavior and surface frictions, etc."

"Special focus has been devoted to the wetting phenomenon of an "ordered water monolayer that does not completely wet water," along with the underlying mechanism, on model and real solid surfaces at room temperature," they report in the Beijing-based journal *SCIENCE CHINA Physics, Mechanics & Astronomy*.

The three scholars likewise outline possible applications of this phenomenon.

According to classical understanding, when water contacts other water, it will spread out and finally both mix together, i.e., water always completely wets water due to the hydrogen bonds formed among water molecules. In 2005, an ice monolayer that did not completely wet ice on metal Pt(111) surfaces was observed at extremely low temperatures.

It is well known that <u>liquid water</u> is essential for almost every biological process. Thus, if this exceptional behavior of liquid water exists, it might be crucial in understanding many biological activities.

Until relatively recently, this phenomenon had not been expected to be observed at room temperature. Like the superconductor and Bose–Einstein condensation, results obtained at cryogenic temperatures are usually not transferable to those observed at room temperature unless a new mechanism is introduced.

At room temperature, thermal fluctuations usually break the hydrogen bond networks in a water monolayer and lead to more hydrogen bonds formed between the monolayer and other water molecules. Thus, the water monolayer at room temperature is expected to completely wet



water.

However, in 2009, Haiping Fang and co-researchers numerically observed a liquid water droplet on a water monolayer on a polar model surface at room temperature, and termed the phenomenon an "ordered water monolayer that does not completely wet water" (Phys. Rev. Lett. 2009, 103, 137801).

This began to challenge the classical picture that a water monolayer always completely wets water. This work expanded understanding of hydrophobicity and hydrophilicity. The related mechanism lies in ordered water structures that reduce the possibilities of hydrogen bonds formed between the monolayer and the contacting water molecules above the monolayer. It should be noted that there remained a considerable number of dangling OH bonds in this room temperature water monolayer, in contrast with previous studies that did not report dangling OH bonds in the ordered water monolayer on a Pt(111) surface and a mica surface.

Over the past several years, more scientists have observed the phenomenon of a liquid water droplet on a water monolayer, both experimentally and numerically, on real surfaces including a self-assembled monolayer (SAM) surface with the terminal of –COOH, a BSA(bovine serum albumin)-Na2CO3 membrane, and an anatase and rutile TiO2 surface, a talc surface, a Pt(100) surface, a hydroxylated Al2O3 surface, a hydroxylated SiO2 surface.

More information: C L Wang, YZ Yang, HP Fang, "Recent advances on an ordered water monolayer that does not completely wet water at room temperature." Sci China-Phys Mech Astron, 2014, 57: 802-809, DOI: 10.1007/s11433-014-5415-3 . link.springer.com/article/10.1 ... 07/s11433-014-5415-3



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