COMMENT

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Open questions on the structures of crystalline water ices

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Water can form a vast number of topological frameworks owing to its hydrogen-bonding ability, with 19 different forms of ice experimentally confirmed at present. Here, the authors comment on open questions and possible future discoveries, covering negative to ultrahigh pressures.

The hydrogen bond as envisioned by Linus Pauling is key to the structures of H₂O ices. A hydrogen bond can be characterised as O–H...O, where the hydrogen atom is linked via a short covalent bond to the donor oxygen and via the longer bond to the acceptor. The one feature that makes ice structures so diverse is that four hydrogen bonds can be formed by a water molecule, although it has only three atoms. Thereby, two atoms may act as H donors, two as H acceptors. In ice, as opposed to water clusters or liquid water, all four hydrogen bonds are developed, resulting in a tetrahedral coordination around each water molecule. This local coordination motif is called the Walrafen pentamer and is found in almost all ice structures¹.

Hydrogen bonding at ultrahigh pressure

Figure 1 summarises the phase diagram for water in a very broad pressure regime, up to pressures as high as encountered in the interior of the ice giants, Uranus and Neptune. Only at ultrahigh pressures, exceeding 50 GPa, the Walrafen pentamer breaks down. In this pressure regime, the hydrogen atom becomes centred between the two oxygen atoms. The molecular nature of H₂O breaks down in two stages. First, the H atoms jump back and forth between the two O atoms (labelled dynamic VII/X in Fig. 1). Second, the H atoms remain static in the centre (labelled atomic solid ice X). Roman numerals are used for ice phases, following the chronological order of the deduction of their crystal structures. Ice X was made in the laboratory in 1984² and represents the 10th polymorph of H₂O. It does not melt when it is heated beyond 2500 K. It experiences a dynamic transition instead: the H atoms start to be mobile in a lattice of O atoms remaining at fixed positions—a property known as superionicity. This ice was discovered only very recently through X-ray experiments on laser-shocked water, and is called ice XVIII since 2019³. Ice XVIII is almost as conductive as metals are. In fact, delocalised H atoms in superionic ice take the same charge carrier role as delocalised electrons in metals. Based on the conditions of the experiments, Uranus

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Apart from the ultralow- and ultrahigh-density ices described so far, all other experimental discoveries pertain to the highly versatile intermediate-pressure range, between about 0.1 and 2.5 GPa as depicted in Fig. 2. In total, 13 out of the currently 19 known ice polymorphs are found in this relatively narrow pressure range. Ices II–IX were discovered prior to the 1970s. Ices XII–XV and β -XV were made in the last two decades. Ices VI and VII in fact exist in the interior of Earth and return to the surface in the form of diamond inclusions¹¹. They have, therefore, been declared as a mineral by the international mineralogic association. At depths of several hundred kilometres, these ices might be of relevance to build friction between cold-subduction slabs, resulting in seismic events one could term “ice quakes.” High-pressure forms of ice are even more abundant in space, where the icy moons are covered in layers of ice up to 1000 km thick so that their own weight suffices to transform the ices to high-pressure polymorphs such as ice III, V, VI, or VII. On Earth pressures within even the deepest glaciers are not high enough to transform hexagonal ice to high-pressure ice forms. The density increase from ice I to ice III to ice V to ice VI and to ice VII (indicated in Fig. 2 through shades of orange/red) is accommodated in the O-atom network in different ways: first, the hydrogen bonds are compressed and bent in ice III. Then the topology and ring sizes change—rather than the six-membered rings in hexagonal ice, ice V contains both smaller and larger rings. Also, ring threading allows for higher densities, as is the case for ice IV. In ices VI and VII, the high densities are reached through two independent H₂O networks that are fully interpenetrating.

Geometrically frustrated ices

The single property that has resulted in the most discoveries of ice phases is hydrogen order. In an ice crystal, both the H atoms and O atoms may occupy lattice positions, resulting in oriented water dipoles (coloured in blue in Fig. 2). Such ices have very low static dielectric constants of about 4. However, in most cases, especially when crystallising the ice from liquid water, only the O atoms are ordered, but not the H atoms (coloured in orange/red) — a property known as geometrical frustration. The water molecules still obey the Bernal–Fowler ice rules¹². However, the water dipoles are oriented randomly causing dielectric constants of above 100. Currently, the six hydrogen disorder-order pairs I_h –XI, III–IX, V–XIII, VI–XV, VII–VIII, and XII–XIV are known. The order-disorder transition temperature varies between 72 K for the ice I_h /XI couple and 273 K for the ice VII/VIII couple. To actually make H-ordered ices and to release the frustration in the H-subnetwork kinetics is the key challenge. It has taken until the 1970s before Kawada achieved to make partly H-ordered ice XI from its disordered parent ice I_h . The ordering process requires waiting times of days at <72 K¹³, but only if extrinsic defects are injected into the hydrogen-bond network through dopants. HCl and KOH have been the most successful dopants, where KOH allows accessing ice XI, and HCl allows accessing the H-ordered high-pressure ices XIII–XV¹⁴. As of now, it is still unknown why HCl accelerates ordering dynamics in high-pressure ices, and KOH in low-pressure ice, but not the other way around.

One specific and unique case that has received much attention lately is ice VI. Not only one, but in fact 45 different types of H order can be imagined that all obey the Bernal–Fowler ice rules. The antiferroelectric ice XV structure discovered in 2009 is only one of the H-ordered pendants of disordered ice VI¹⁵. In our experiments we recently discovered a second one, which we call ice β -XV¹⁶. This represents the 19th polymorph of ice, where elucidation of its crystal structure based on diffraction experiments is ongoing work. While still being contested, this makes the case for the first phase of ice, in which transitions between two differently ordered H-sublattices may be studied—and it is fully open how such a transition progresses. Eighty-five years ago, Pauling considered all H-ordered structures as degenerate, which can be accessed through permutation of H atoms. It is clear now, though, that there are subtle differences in enthalpy, lattice parameters and density. Yet, it is still unclear for all ices known today whether a more stable arrangement of the H atoms exists, and whether this will be realised experimentally someday.

Outlook

Even at ambient pressure ice still holds surprises, as shown through the realisation of a cubic stacking sequence in ice I without hexagonal stacking faults in 2020^{17,18}. That is, it is still far from clear how many condensed phases of water can be distinguished. A recent machine learning study on ice phases actually suggests in total 51 ice polymorphs, more than 30 of which have not yet been reported in experiments¹⁹. If the stable structures that are predicted in simulations can actually be achieved in experiments, we expect to discover more ice phases in the future than have been discovered up to now. Rather than the candidates indicated between the question marks in Figs. 1 and 2, we think that most likely the next ice to be realised in experiments will be in the intermediate pressure range, e.g., the H-disordered pendant to ice II or the H-ordered pendant to ice IV. These are signs for a field that has not reached maturity yet! That is, in spite of a more than hundred year old history of research on ice structures, the field is still young and evolving more than ever.

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T.L. has written the comment. T.M.G. has produced the figures. V.F.L., C.M.T., and T.M.G. have added to the discussion.

Competing interests

The authors declare no competing interests.

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