Pressure Dependence of Fragile-to-Strong Transition and a Possible Second Critical Point in Supercooled Confined Water

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By confining water in nano-pores of silica glass, we can bypass the crystallization and study the pressure effect on the dynamical behavior in deeply supercooled state using neutron scattering. We observe a clear evidence of a cusp-like fragile-to-strong (F-S) dynamic transition. Here we show that the transition temperature decreases steadily with an increasing pressure, until it intersects the homogenous nucleation temperature line of bulk water at a pressure of 1600 bar. Above this pressure, it is no longer possible to discern the characteristic feature of the F-S transition. Identification of this end point with the possible second critical point is discussed.

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Water is a continuing source of fascination to scientists because of its many counterintuitive low-temperature properties. Although the stable form of water at sufficiently low temperature is inevitably crystalline, liquid water can also exist in a metastable supercooled state far below the melting point. One of the most intriguing questions related to the unusual properties of supercooled water is whether two critical points may exist in a single component liquid.

It has been known since 1970s that a number of thermodynamic response functions of supercooled water, notably the isothermal compressibility \( K_T \) and the constant-pressure specific heat \( C_P \), show a power-law divergence behavior at a singular temperature, experimentally determined to be 228 K at the ambient pressure. Concurrently, the transport properties, such as the shear viscosity \( \eta \) and the inverse self-diffusion constant \( D \), diverge according to power-laws toward the same singular temperature. The anomalies of the thermodynamic quantities become plausible if one postulates the existence of a second low-temperature critical point at about 228 K and at somewhat elevated pressure. On the other hand, the transport coefficient anomalies are reminiscent of the dynamical behavior of a supercooled liquid near the so-called kinetic glass transition temperature, predicted by Mode-Coupling theory (MCT).

Search for the predicted first-order liquid-liquid transition line and its end point, the second low-temperature critical point in water, has been hampered by intervention of the homogenous nucleation process, which takes place at 235 K at the ambient pressure. However, by confining water in nano-pores of mesoporous silica MCM-41-S with cylindrical pores of 14 Å diameter, we have been able to study the dynamical behavior of water in a temperature range down to 160 K, without crystallization. Using high-resolution Quasi-Elastic Neutron Scattering (QENS) method and Relaxing-Cage Model (RCM) for the analysis, we determine the temperature and pressure dependences of the average translational relaxation time \( \langle \tau_T \rangle \) for the confined supercooled water.

Micellar templated mesoporous silica matrices MCM-41-S, which have 1-D cylindrical pores arranged in 2-D hexagonal arrays, were synthesized by following a similar method for synthesizing MCM-48-S previously. Using the short chain cationic surfactants, \( C_{12} \)TMAB, as templates and adding \( \beta \)-type zeolite seeds as silica source, we obtain MCM-41-S (S denotes seed) with smaller pore sizes and stronger silica walls than the traditional MCM-41. In this pressure experiment, we chose the silica with a pore diameter of 14 Å because for the fully hydrated sample the differential scanning calorimetry (DSC) data shows no freezing peak down to 160 K. The sample is then hydrated by exposing to water vapor in a closed chamber until it reaches the full hydration level of 0.5 gram \( H_2O \)/1 gram Silica.

High-resolution QENS spectroscopy method is used to determine the temperature and pressure dependences of \( \langle \tau_T \rangle \) for the confined water. Because neutrons can easily penetrate the thick-wall high-pressure cell and because it is predominantly scattered by hydrogen atoms in water, rather than by the matrices containing it, incoherent QENS is an appropriate tool for our study. Using two separate high-resolution QENS spectrometers, we are able to measure the translational-relaxation time from 0.2 ps to 10,000 ps over the temperature and pressure range. The high-pressure experiments were performed at both the High-Flux Backscattering (HFBS) and the Disc-Chopper Time-of-Flight (DCS) spectrometers in the NIST Center for Neutron Research (NIST NCNR). The two spectrometers used to measure the spectra have two widely different dynamic ranges (for the chosen experimental setup), one with an energy resolution of 0.8 \( \mu eV \) (HFBS) and a dynamic range of \( \pm 11 \mu eV \), and the other with an energy resolution of 20 \( \mu eV \) (DCS) and a

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dynamic range of $\pm 0.5$ meV in order to be able to extract the broad range of relaxation times from the measured spectra. The same high pressure system, including specially designed aluminium pressure cell, was used on the two instruments. Helium gas, the pressure-supplying medium, fills the whole sample cell, and applies pressure to the fully hydrated sample. The experiment at each pressure was done with a series of temperatures, covering both below and above the transition temperature. Altogether, 1000 spectra were collected, spanning 9 pressures: ambient, 100, 200, 400, 800, 1200, 1600, 2000, and 2400 bars.

QENS experiments measure the Fourier transform of the Intermediate Scattering Function (ISF) of the hydrogen atoms, $F_H(Q, t)$, of water molecules in the pores of the silica matrix. Molecular Dynamics (MD) simulations have shown that the ISF of both bulk and confined water can be accurately described as a two-step relaxation: a short-time Gaussian-like (in-cage vibrational) relaxation followed by a plateau. This function is fairly insensitive to temperature and confined water through MD and experimental data. By considering only the spectra with wave vector transfer $Q < 1.1$ Å$^{-1}$, we can safely neglect the contribution from the rotational motion of water molecule in ISF. The RCM describes the translational dynamics of water at supercooled temperature in terms of the product of two functions:

$$F_H(Q, t) \approx F_T(Q, t) = F^S(Q, t) \exp \left[ -\left( t/\tau_T(Q) \right)^\beta \right],$$

$$\tau_T(Q) = \tau_0 (0.5Q)^{-\gamma}, \langle \tau_T \rangle = \tau_0 \Gamma(1/\beta)/\beta,$$  

(1)

where the first factor, $F^S(Q, t)$, represents the short-time vibrational dynamics of the water molecule in the cage. This function is fairly insensitive to temperature variation, and thus can be calculated from MD simulation. The second factor, the $\alpha$-relaxation term, contains the stretch exponent $\beta$, and the $Q$-dependent translational relaxation time $\tau_T(Q)$, which is a strong function of temperature. The latter quantity is further specified by two phenomenological parameters $\tau_0$ and $\gamma$, the exponent controlling the power-law $Q$-dependence of $\tau_T(Q)$. $\langle \tau_T \rangle$ is a $Q$-independent quantity where $\Gamma$ is the gamma function. It essentially gives a measure of the structural relaxation time of the hydrogen-bond cage surrounding a typical water molecule. The temperature dependence of the translational relaxation time at each pressure is then calculated from three fitted parameters, $\tau_0$, $\beta$, and $\gamma$, by analyzing a group of quasi-elastic peaks at different $Q$ values simultaneously. For this analysis, we chose seven spectra for data taken at HFBS and eleven spectra for data taken at DCS, at each temperature.

We show in Fig.1 as an example, two complete sets (temperature series) of QENS area-normalized spectra. The broadening of the quasi-elastic peaks becomes more and more noticeable as temperature increases. In Fig.1A, we may notice, from shoulders of these spectral lines, that two groups of curves, 231-250 K and 198-209 K, are separated by the curve at a temperature of 220 K. This visual information reinforces the result of the detail analysis shown in Fig.2 that there is an abrupt dynamical transition at $T_L = 216$ K. Fig.1B shows the RCM analysis of the spectrum taken at $T = 220$ K, close to the transition temperature. On the other hand in Fig.1C, the spectra at pressure 1600 bar show a rather smooth variation with temperature, indicating that there is no sharp transition. This pressure corresponds to the end point of the line of $F-S$ transitions shown in Fig.2. Fig.1D is, again, an RCM analysis of the spectrum taken at $T = 214$ K of this pressure. RCM, as one can see, reproduces the experimental spectral line shapes of confined water well. The broadening of the experimental data over the resolution function, shown in Figs.1B and 1D, leaves enough dynamic information to be extracted by RCM.

The behavior of shear viscosity $\eta$ or equivalently the structural relaxation time $\tau$ of a supercooled liquid approaching its glass transition temperature is called ‘fragile’ when it varies according to the so-called Vogel-Fulcher-Tammann (VFT) law; and the behavior is called ‘strong’ when $\eta$ or $\tau$ obeys Arrhenius law. For water, a fragile liquid at room temperature and at moderately supercooled temperatures, Itô and coworkers proposed that a ‘fragile-to-strong’ transition would occur at around 228 K, based on a thermodynamic argument. The $F-S$ transition in a molecular liquid like water may...
be interpreted as a variant of kinetic glass transition predicted by the ideal MCT, where the real structural arrest transition is avoided by an activated hoping mechanism below the transition.

In Fig. 2 we report the temperature variation of $\langle \tau_T \rangle$ for water molecules as a function of pressure. It is seen that Figs. 2A, 2B, 2C and 2D show clearly a transition from a VFT law: $\langle \tau_T \rangle = \tau_0 \exp \left[ D(T_0/(T - T_0)) \right]$, where $D$ is a constant providing the measure of fragility and $T_0$, the ideal glass transition temperature, to an Arrhenius law: $\langle \tau_T \rangle = \tau_1 \exp \left[ E_A/RT \right]$, where $E_A$ is the activation energy for the relaxation process and $R$, the gas constant. This transition of VFT to Arrhenius behavior, previously observed at ambient pressure, is the signature of a F-S dynamic transition predicted by Ito et al. In this paper, we show the extension of this transition into finite pressures. The transition temperature, $T_L$, as the crossing point of the VFT law and Arrhenius law, is calculated by $1/T_L = 1/T_0 - (Dk_B)/E_A$. However, in Figs. 2E and 2F, the cusp-like transition becomes rounded off and there is no clear-cut way of defining the F-S transition temperature. Note that in Fig. 2F, there is still a hint of fragile behavior at high enough temperature.

Summarizing all the results, we show in a P-T plane, in Fig. 3 the observed pressure dependence of $T_L$ and its estimated continuation, denoted by a dash line, in the pressure region where no clear-cut F-S transition is observed. One should note that the $T_L$ line has a negative slope, parallel to TMD line, indicating a lower density liquid on the lower temperature side. This $T_L$ line also approximately tracks the $T_H$ line, and terminates in the upper end when intersecting the $T_H$ line at 1600 bar and 200 K, at which point the character of the dynamic transition changes. We shall discuss the significance of this point later on. A special feature of the $T_L$ line at the lower end should be noted as well. The line essentially becomes vertical after around 200 bar and the transition temperature approaches a constant value of $\sim 225$ K.

Since $T_L$ determined experimentally is a dynamic transition temperature, it is natural to question whether the system is in a liquid state on both sides of the $T_L$, and if so, what would the nature of the high-temperature and low-temperature liquids be? Sastry and Angell have recently shown by a MD simulation that at a temperature $T \approx 1060$ K (at zero pressure), below the freezing point 1685 K, the supercooled liquid silicon undergoes a first-order liquid-liquid phase transition, from a fragile, dense liquid to a strong, low-density liquid with nearly tetrahedral local coordination. Prompted by this finding, we may like to relate, in some way, our observed $T_L$ line to the liquid-liquid transition line, predicted by MD simulations of water and speculating on the possible location of the low-temperature critical point.

According to our separate inelastic neutron scattering experiments, which measure the librational density of states of water contained in 18 Å pore size MCM-41-S, water remains in disordered liquid state both above and below the F-S transition at ambient pressure. Furthermore, our analysis of the F-S transition for the case of ambient pressure indicates that the activation energy...
barrier for initiating the local structural relaxation is $E_A = 4.89$ Kcal/mol for the low-temperature strong liquid. Yet, previous inelastic scattering experiments of stretch vibrational band of water indicate that the effective activation energy of breaking a hydrogen bond at 258 K (high-temperature fragile liquid) is 3.2 Kcal/mol. Therefore, it is reasonable to conclude that the high-temperature liquid corresponds to the high-density liquid (HDL) where the locally tetrahedrally coordinated hydrogen bond network is not fully developed, while the low-temperature liquid corresponds to the low-density liquid (LDL) where the more open, locally ice-like hydrogen bond network is fully developed.

It is appropriate now to address the possible location of the second critical point. Above the critical temperature $T_C$ and below the critical pressure $p_C$, we expect to find a one-phase liquid with a density $\rho$, which is constrained to satisfy an equation of state: $\rho = f(p,T)$. If an experiment is done by varying temperature $T$ at a constant pressure $p < p_C$, $\rho$ will change from a high-density value (corresponding to HDL) at sufficiently high temperature to a low-density value (corresponding to LDL) at sufficiently low temperature. Since the fragile behavior is associated with HDL and the strong behavior with LDL, we should expect to see a clear F-S transition as we lower the temperature at this constant $p$. Therefore, the cusp-like F-S transition we observed should then occur when we cross the so-called Widom line in the one-phase region. On the other hand, if the experiment is performed in a pressure range $p > p_C$, corresponding to the two-phase region and crossing the Liquid-Liquid (L-L) coexistence line, the system will be consisting of mixture of different proportions of HDL and LDL as one varies $T$. In this latter case, $\langle \tau_T \rangle$ vs. $1/T$ plot will not show a clear-cut F-S transition (the transition will be washed out) because the system is in a mixed state. The above picture would then explain the dynamical behavior we showed in Fig. In Fig a clear F-S transition is observed up to 1200 bar and beyond 1600 bar the transition is rounded off. From this observation, the reasonable location of the L-L critical point is estimated to be at $p_C = 1600 \pm 400$ bar and $T_C = 200 \pm 10$ K, shown by a star in Fig.

Additionally, in a recent MD simulation using Jagla Model Potential by Xu et al, a small peak was found in the specific heat $C_P$ when crossing the Widom line at a constant $p$. Meanwhile, Maruyama et al conducted an experiment on adiabatic calorimetry of water confined within nano-pores of silica gel. It was found that water within 30 Å pores was well prevented from crystallization, and also showed a small $C_P$ peak at 227 K at ambient pressure. This experimental result further supports that the F-S transition we observed at 225 K at ambient pressure is caused by the crossing of the Widom line in the one-phase region above the critical point.

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