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Kyoto University
Structural properties and phase transitions in a silica clathrate
Yunfeng Liang, Folorunso O. Ogundare, Caetano R. Miranda, Jamieson K. Christie, and Sandro Scandolo

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Structural properties and phase transitions in a silica clathrate

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Melanophlogite, a low-pressure silica polymorph, has been extensively studied at different tempera-
tures and pressures by molecular dynamics simulations. While the high-temperature form is con-
figured as cubic, the low-temperature phase is found to be slightly distorted, in agreement with ex-
periments. With increasing pressure, the crystalline character is gradually lost. A 8 GPa, the radial
distribution function is consistent with an amorphous state. Like pristine glass, the topology changes,
plastic behavior, and permanent densification appear above ~12 GPa, triggered by Si coordina-
tion number changes. We predict that a partial crystalline and amorphous sample can be obtained
by recovering the sample from a pressure of ~12–16 GPa. © 2011 American Institute of Physics.
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I. INTRODUCTION

The crystalline-to-amorphous transformation under high
pressure and low temperature is currently the subject of in-
tense study, since the discovery of amorphization of H2O
ice Ih under pressure.1 Interest has focused on the pos-
ible occurrence of pressure-induced amorphization in silica
polymorphs;2 the thermodynamic and mechanistic basis of
the process, and the relationship between amorphization by
pressure and by other means.3,4 Amorphization of minerals
(e.g., silica and silicates) is observed in situ with a variety
of structural probes such as Raman and IR spectroscopy and
x-ray diffraction on samples usually compressed in diamond
anvil cells.3,4 One assumes amorphization when diffraction
peaks vanish or when broad, glasslike bands appear in vibra-
tional spectra. The detailed microstructural origins are how-
ever rarely extracted from experiments alone.

Recent work on zeolite collapse has renewed interest in
pressure-induced amorphization,5–7 where crystalline order
can be preserved to some extent in the densified sample.
In particular, Greaves et al. have reported that zeolite Y was
thermally amorphized 20% at 650 °C, 60% at 750 °C, and 100%
at 850 °C. A 90% amorphized specimen was fabricated in
a multianvil press at 8 GPa and room temperature. The de-
gree of amorphization was initially obtained by comparing
the intensity of x-ray diffraction powder patterns with that of
the starting materials5 and then further identified by the bo-
son peaks.6 Further zero-temperature calculations have the-
oretically shown that the different degrees of amorphization

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with the guest molecules $M^{12}$ ($N_2$, $CH_4$) and $M^{14}$ ($CO_2$), respectively. The guest molecules are thought to act as templates during crystallization. Using a mixture of $CO_2$, $CH_4$, and $N_2$ as guest molecules, melanophlogite was first synthesized under hydrothermal conditions from aqueous silica solutions in 1982.14 Melanophlogite is known to distort slightly from cubic symmetry below around $80^\circ C$.15–22 By careful thermal treatment (e.g., heating to above $600^\circ C$), the guest molecules can be removed.9, 10, 12, 18–22 In the past, many studies have been performed in order to understand the guest molecules trapped in the clathrates23, 24 and the role of the guest for the response of the host framework to pressure.25 More recently, crystalline–amorphous transitions have been documented in guest-free melanophlogite through a broadening and drop in the intensity of the x-ray diffraction lines at $\sim 8$ GPa.9, 10 The microstructure information is however still lacking: particularly how the different cages respond to the pressure, whether the partial crystalline structure exists in densified melanophlogite, and whether the topology of the network is preserved as has been suggested in guest-free silicalite.8

In this paper, we present extensive molecular dynamics simulations of guest-free melanophlogite at different temperatures and pressures. Our main findings are that (1) the temperature-induced phase transition can be explained using the same mechanism underlying the $\alpha$–$\beta$ phase transitions in quartz and cristobalite and the zero thermal expansion at high temperature can be explained solely in terms of expansion of the individual cages; (2) the x-ray amorphous material obtained by compressing melanophlogite between 8 and 12 GPa is topologically identical to the crystalline phase; (3) parts of the crystalline structure still exist in densified melanophlogite subjected to pressures of $\sim 12$–16 GPa.

The paper is organized as follows: in Sec. II, we describe the methodology; in Sec. III, we first present the structural changes with temperature, across the $\alpha$–$\beta$ phase transition. There we show that it is possible to analyze the phase transformations by the distortion of the cages. We then present the changes with pressure of the equation of state, x-ray structure factor, radial distribution functions, and the structural changes of the two different cages. In the last part of Sec. III, we discuss the structural properties of the densified sample.

II. METHODOLOGY

In this paper, we present the results of simulations of the pressure-induced amorphization of guest-free melanophlogite. The simulations were carried out using an interatomic force field optimized by best fit on first-principles (density-functional theory) calculations.26 The force field describes the structural and vibrational properties of most $SiO_2$ crystalline polymorphs, liquid, and glass27–31 better than all the available force fields to which it has been compared so far. In particular, it describes the thermodynamic stability of the crystalline polymorphs of silica,31 including the pressure dependence of the lattice constant and the phonon softening across the rutile-to-$CaCl_2$ transition.27, 28 We start our MD simulations from a cell consisting of 368 $SiO_2$ formula units ($2 \times 2 \times 2$ unit cells), which is obtained by first inserting oxygen atoms in the middle of the line connecting two silicon atoms in type I silicon clathrate and then relaxing the structure to 0 GPa and 0 K. The
FIG. 3. Lattice parameters: lengths (a) and angles (b) as a function of temperature obtained as an average over the last 7 ps of the simulation.

initial velocity was set randomly according to the Maxwell distribution. The MD time step was set to 0.72 fs (30 a.u.).

In order to monitor the structural changes (shape and size) of the two cage types ([512] and [51262]) across the transitions induced either by temperature or by pressure, we calculated radial distribution functions between atoms and imaginary points located at the crystallographic center of the cages. We analyzed radial distribution functions using a method introduced recently by Saitta and co-workers to study the short- and intermediate-range order changes of liquid water32 and amorphous ices.33, 34 They ranked the neighbors (to any tagged one) in order of increasing distance and analyzed the (individual or combined) neighbor distance distributions. From this analysis, they drew insights on the structural changes in liquid and amorphous water (ice) in terms of first (short range) and second neighbor shells (intermediate range). By ranking the Si neighbors to the cage center in the [512] cages (cage 1) and taking the first 20 neighbors, we then extract the neighbor list constructed for cage 1 only. Similarly, by ranking the Si neighbors to the cage center in the [51262] cages (cage 2) and taking the first 24 neighbors, we then analyze the distance distribution function with respect to the cage centers to get some insights into the distortion of the two different cages under different temperatures or pressures.

III. RESULTS AND DISCUSSIONS

A. Temperature-induced symmetrization

We first studied the lattice and volume changes at different temperatures (Figs. 2 and 3). It is interesting to note that the volume expands very fast with increasing temperature and then saturates at around 700 K, which is typical of the α–β transition such as that in quartz and cristobalite.27, 28, 35 Like quartz and cristobalite, we find that melanophlogite also shows the dynamic disorder of oxygen atoms above ∼500 K.

While the high-temperature form is found to be cubic, in agreement with experiments, the low-temperature phase is
found to be slightly distorted (Fig. 3), which is also in agreement with experiments.\textsuperscript{10,15–22} The experimental studies have found the low-temperature phase as tetragonal for guest-bearing melanophlogite, whereas a lower symmetry (probably, an orthorhombic phase) was reported for guest-free melanophlogite.\textsuperscript{18} Here, we found that the original cell was not even orthogonal at temperatures below 500 K (see Fig. 3). The calculated lattice parameter length differences at 300 K are within 0.3%, which is again in good agreement with the experiments.\textsuperscript{10,15–22} However, such small differences (e.g., 0.1° in angles or 0.3% in lengths at 300 K) can hardly be resolved by x-ray diffraction patterns as discussed in Ref. 18.

We then investigated the changes of the two different cages by ranking the Si neighbors to the cage centers and calculating the distance distribution functions. At room temperature, the distribution for cage 1 is well peaked at 4.38 Å (shown in Fig. 4), and for cage 2 it is peaked at 4.87 Å with a shoulder at \( \sim 5.0 \) Å. By taking account of the covalent radius of Si atoms, the dimensions of the cages are similar to the values estimated by Gies.\textsuperscript{16} As shown in Fig. 4, cages 1 and 2 in the \( \beta \)-phase (\( > 700 \) K) do not expand with increasing temperature. In particular, the sizes are similar to each other for the samples at 1100 and 1900 K. Since melanophlogite is exclusively made up of two types of the cages, a pentagonal dodecahedron \([5^{12}]\) cage and a tetrakaidecahedra \([5^{12}6^2]\) cage (as shown in Figs. 1(b) and 1(c)), the invariance of the volume of the individual cages in the high-temperature regime is consistent with the zero thermal expansion of the crystal. Similarly, the symmetry distortion of the lattice at low temperature can be understood by analyzing the distortion of the cages. The radial function in cage 1 at 50 K is split into two peaks at 4.30 and 4.47 Å, respectively. In cage 2, the two peaks are located at 4.75 and 5.06 Å. The distortions are as large as 0.31 Å in cage 2 and 0.17 Å in cage 1, respectively. Symmetry distortions of the order of 0.1–0.3 Å for the cages, which become less spherical at low temperature, are consistent with the calculated low-temperature distortions of the lattice parameters.

**B. Pressure-induced amorphization**

Next, we applied a stepwise hydrostatic compression at 300 and 50 K, in pressure increments of 2 GPa, followed, at each pressure, by an equilibration time of 20 ps. No clear volume collapse can be identified in the equation of state, up to a pressure of 28 GPa (Fig. 5). Melanophlogite has higher compressibility than glass and the equation of state at lower pressure lies in between two recent experiments,\textsuperscript{8,10} while the unit volume converges to pristine glass at pressures above 12 GPa. No sharp volume collapse has been found in the whole studied range. We compared the x-ray structure factor at different pressures in Fig. 6. At 0 GPa and room temperature,
FIG. 7. Si–Si radial distribution functions at 0 and 8 GPa.

FIG. 8. Radial distribution functions with respect to the central point of the cages at five different pressures: for pentagonal dodecahedra [512] cage (a) and tetrakaidecahedra [51262] cage (b).

FIG. 9. Angle distributions with respect to the central point of the cages at five different pressures: for pentagonal dodecahedra [512] cage (a) and tetrakaidecahedra [51262] cage (b).

the diffraction pattern is consistent with the experimental data.\textsuperscript{9,10} The relative intensity differs between different experimental reports. Our data show higher intensities in the second and third peaks (i.e., [202] and [222] assuming a tetragonal unit cell\textsuperscript{14} than those of Ref. 9 but similar to those of Ref. 10. The fact that the experimental data differ may result from the fact that the experimental specimens contain different degrees of disorder. With increasing pressure, we found that the strongest peak [423] and other peaks, shown in Fig. 6(b), decrease in intensity, which is in good agreement with experiments.\textsuperscript{9} At 8 GPa, the x-ray structure factor is consistent with an amorphous state. As shown in Fig. 7, the Si–Si pair distribution function is also flat at large distances, at 8 GPa. The results of the radial distribution function analysis for the response of the cages to pressure are shown in Fig. 8. The cages compress mainly through a bending of the Si–O–Si angle and their distortion increases with pressure. We show the angle distribution functions between Si–center–Si,
where “center” is the center of the cages, in Fig. 9. The peak at 42° corresponds to the nearest-neighbor Si atoms in cage 1, the angle at 70° corresponds to the next-nearest neighbors within the same pentagonal rings, and the angles at 110°, 138°, and 177° correspond to the neighbors within the face pentagonal rings. The angle distribution in cage 2 can also be assigned to well-defined cage elements. Under pressure, cage 2 loses the two large-angle features at 8 GPa and three sharp peaks at 12 GPa, while cage 1 still contains all the features at 12 GPa. This suggests that larger cages are more easily distorted under pressure.

The addition of guest molecules can stabilize a clathrate lattice, acting as an organizing center for the formation of the lattice. The interactions between the guests and the host cages hold the cage in shape at much higher pressures than those of guest-free clathrates. Our simulations are for guest-free lattices, but we can explore this hypothesis by looking at how the structure of the cages changes at distances corresponding to the cage diameter with increasing pressure. For clarity, the Si–Si pair separations between atoms on the surface of each cage were divided into four groups, based on their interatomic distance, $r_0$, at zero pressure: $r_0 < 4 \text{ Å}$, $4 \text{ Å} < r_0 < 6 \text{ Å}$, $6 \text{ Å} < r_0 < 8 \text{ Å}$, and $r_0 > 8 \text{ Å}$. (The largest Si–Si separation at zero pressure is 9.1 Å in cage 1 and 10.7 Å in cage 2.) The separations were tracked as pressure was increased, and Fig. 10 shows these distances at 12 and 24 GPa. At 12 GPa, the distortion of the cages has simply broadened the distribution of distances. At 24 GPa, however, this broadening is much larger and there are clear signs of topological changes. In both cages, there are substantial peaks at the Si–O–Si bond length (~2.4 Å) from pairs with $4 \text{ Å} < r_0 < 6 \text{ Å}$. There are also smaller contributions at this distance and substantial contributions at ~4 Å from atoms with $r_0 > 6 \text{ Å}$. This implies that pairs of silicon atoms which were at large separations at zero pressure are forming chemical bonds at 24 GPa, presumably in a Si–O–Si or longer chain, altering the topology and irreversibly amorphizing the structure. It also implies that this amorphization would not occur at this pressure in the presence of guest atoms. Guest atoms would physically obstruct the formation of Si–O–Si bonds starting from pairs of atoms separated by nearly the diameter of the whole cage and hence maintain the structure of the melanophlogite cages to higher pressures, as seen in Ref. 25 for clathrasils.

C. The recovered sample

The recovered sample shows densification above around 12 GPa (Fig. 11), which suggests that the reduction of the

![FIG. 10. Si–Si radial distribution functions on the surface of pentagonal dodecahedra [512] at 12 (a) and 24 GPa (b) and tetrakaidecahedra [51262] at 12 (c) and 24 GPa (d). Pair distances divided into four groups based on their separation, $r_0$, at zero pressure.](image-url)
Bragg peaks below 12 GPa is not a consequence of bonding changes. This is confirmed by the analysis of the coordination changes and ring statistics (Figs. 12 and 13). The coordination increases only above ~12 GPa, and as a consequence, the topology changes to generate a densified glass. The crystalline melanophlogite contains only five-membered, six-membered, and ten-membered rings. The topology of the recovered sample from 12 GPa has two five-membered rings and one six-membered ring less than the starting crystal. The sample recovered from 16 GPa shows some memory of the crystal according to the ring size distribution functions, which in this case contains a maximum for five-membered rings and a second peak for ten-membered rings. At higher pressure, the densification process saturates and generates a ring statistics similar to that of the densified glass. It is noteworthy that at higher subjected peak pressure, the densification is saturated the way pristine glass behaves as reported in Ref. 29.

We remark that the increase of small-membered rings with peak pressure is consistent with Raman experimental data. A recent study of the pressure-induced amorphization of silicalite using RMC modeling has suggested the existence of a topologically ordered amorphous silica phase. We observed a topologically ordered amorphous structure at 8–12 GPa, but this structure reverts back to melanophlogite upon releasing pressure. Our recovered densified amorphous phase is topologically disordered. In other words, irreversible amorphization in melanophlogite is a consequence of the topological changes. We remark that the density of our topologically ordered amorphous phase at 8 GPa coincides with the density of the densified amorphous phase at 0 GPa and also with the density of densified silicalite (2.7 g/cm³), so density alone cannot be used to discriminate between topologically distinct structures. Molecular dynamics simulations similar to the ones reported here for melanophlogite might be able to provide complementary information, with respect to RMC, on the structure of the densified amorphous phase obtained from silicalite.

IV. CONCLUSIONS

To summarize, we have shown that the Tangney–Scandolo potential can reproduce the experimentally reported transition from a low-temperature distorted structure to a high-temperature cubic structure in melanophlogite. The distortion has been analyzed based on the structural changes of the individual cages. The simulation of high-pressure melanophlogite does not show a sharp volume collapse; instead, it gradually transforms into an amorphous phase. Below about 12 GPa, the transition is reversible: there is no significant change in the bonding or topology, as shown by the ring statistics distributions, and analysis of the coordination numbers and structure. Above this pressure, the cages deform sufficiently to alter the bonding and topology, and phases recovered from above this pressure are altered irreversibly. The densified amorphous phase at intermediate pressure...
contains some crystalline features as manifested by the ring statistics distributions. This connection between topological changes and permanent amorphization offers an alternative interpretation of the recent experimental results obtained on silicalite.\(^8\)