High pressure and high temperature in situ X-ray observation of hydrous wadsleyite, $Mg_{1.75}SiO_4H_{0.5}$ under the condition of the mantle transition zone

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Preliminary high pressure and high temperature in situ X-ray observations of hydrous wadsleyite, Mg_{1.75}SiO₄H_{0.5} were carried out using a combination of the synchrotron radiation and a large volume multianvil high pressure device (SPEED-1500). We made one run during this beam time.

Hydrous wadsleyite was, in advance, synthesized from the mixture of MgO, Mg(OH), and SiO, powders as a composition of Mg/Si=1.75 and H₂O=5 wt % using a multi-anvil high pressure apparatus (ORENGE-2000) in Ehime University. The sample was sealed in Pt capsule to avoid H₂O loss during the experiment. The synthesis condition was 15 GPa, 1200°C and 30 min. The recovered sample was characterized by a conventional x-ray diffraction and an EPMA. Almost crystals were hydrous wadsleyite, and small amount of clinopyroxene was coexisted in the charge. The lattice parameters a=8.252(1), b=11.550(3)c=5.680(1)Å, respectively, and the chemical composition was ~1.8 of the Mg/Si and the deficit of EPMA weight total was ~ 3 wt%, which corresponds to the H₂O content.

In the in situ experiment, pressures were applied first at a ram load of 500 ton (\sim 16GPa), and then temperatures were increased to 800°C to release the deviatric stress (Fig.

1). In this experiment, NaCl was used as a pressure marker, and the pressures were calculated by Decker scale. After releasing the deviatric stress, the X-ray diffraction patterns of the sample were collected every $100\,^{\circ}$ C from 800°C to room temperature for 300 seconds. Then we tried to release the pressure at 600 $^{\circ}$ C by decreasing the ram load from 500 to 250 ton (\sim 15GPa), and collected the diffraction patterns at 800, 400, 200 and 25 $^{\circ}$ C. In this beam time, two beam dumps were occurred (total 5 hours), so we can not collect enough data to determine the equation of state of hydrous wadsleyite. We will continue this type of experiments in the next beam time.

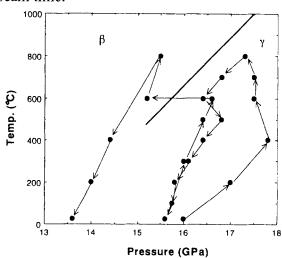


Fig. 1 P-T path in the present experiment. A bold line represents the β - γ phase boundary in Mg_2SiO_4 from Akaogi et al. (1989).