Geophysical Research Abstracts, Vol. 7, 02690, 2005 SRef-ID: 1607-7962/gra/EGU05-A-02690 © European Geosciences Union 2005



## Hydrogen incorporation in a ringwood ite analogue: $\mathbf{Mg}_{2}\mathbf{GeO}_{4}$ spinel

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Single-crystals of Mg<sub>2</sub>GeO<sub>4</sub> spinel, a stable analogue to mantle ringwoodite at atmospheric pressure up to 800°C, were synthesized from Mg<sub>2</sub>GeO<sub>4</sub> olivine in hydrous conditions at 1.9 GPa and  $\sim 1000$ °C. Mg<sub>2</sub>GeO<sub>4</sub> spinel has been extensively used for the study of phase transformations [1] and rheological properties [2]. However, the hydrous phase and hydrogen incorporation has not yet been considered.

IR spectra show sharp O-H stretching peaks at 3531 and 3502 cm<sup>-1</sup>. Quantification of IR spectra reveals hydrogen in the order of 5 to 10 ppm wt H<sub>2</sub>O. A complete hydrogen/deuterium exchange was achieved at 700 °C leading to a diffusion coefficient of deuterium of about  $3 \cdot 10^{-14}$  m<sup>2</sup>/s. This value is of the same order than diffusion coefficients observed in other mantle minerals, such as garnet or diopside.

The O-O distance of ~ 2.9 Å calculated from the spectra of Mg<sub>2</sub>GeO<sub>4</sub> spinel is consistent with the GeO<sub>4</sub> tetrahedra edge length known from crystal structure data. In IR spectra of hydrous mantle ringwoodites the O-H peaks appear at 3105 cm<sup>-1</sup> [3], correlating to an O-O distance of ~ 2.7 Å [4] which is the SiO<sub>4</sub> tetrahedra edge length in the ringwoodite structure. Whereas the hydration mechanism and the site occupancies involved are still in discussion for mantle spinel [3,5], the IR measurements on Mg<sub>2</sub>GeO<sub>4</sub> confirm the idea of protonation at the tetrahedral edges [6].

This study was supported by the EU through a Marie Curie fellowship HPMF-CT-2002-2098.

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