

Pressure-induced structural phase transition of the iron end-member of ringwoodite (γ -Fe₂SiO₄) investigated by X-ray diffraction and Mössbauer spectroscopy

**ERAN GREENBERG,^{1,4,*} LEONID S. DUBROVINSKY,¹ CATHERINE MCCAMMON,¹
JEROME ROUQUETTE,² INNOKENTY KANTOR,^{3,5} VITALI PRAKAPENKA,³ GREGORY KH. ROZENBERG,⁴
AND MOSHE P. PASTERNAK⁴**

¹Bayerisches Geoinstitut, University of Bayreuth, Bayreuth 95447, Germany

²Institut Charles Gerhardt Montpellier, UMR CNRS 5253, équipe C2M, Université Montpellier II, 34095 Montpellier Cedex 5, France

³Consortium for Advanced Radiation Sources, University of Chicago, 9700 South Cass Avenue, Argonne, Illinois 60439, U.S.A.

⁴School of Physics and Astronomy, Tel-Aviv University, Tel-Aviv 69978, Israel

⁵European Synchrotron Radiation Facility, 6 rue Jules Horowitz, Grenoble 38043, France

ABSTRACT

We have carried out X-ray diffraction and Mössbauer spectroscopy measurements on the spinel phase γ -Fe₂SiO₄ (ringwoodite) at ambient temperature and pressures up to 66 GPa using diamond anvil cells. At pressures above 30 GPa, a previously unknown structural phase transition to a rhombohedrally distorted spinel phase has been observed (space group $R\bar{3}mR$). Mössbauer spectroscopy measurements reveal two different Fe²⁺ sites at high pressure with an abundance ratio of 3:1, in agreement with the two crystallographic sites occupied by the iron in this distorted spinel structure. The unit-cell volume of the low-pressure spinel phase as a function of pressure results in a bulk modulus of $K_0 = 197(3)$ GPa using the second-order Birch-Murnaghan equation of state, and $K_0 = 201(8)$ GPa and $K' = 3.7(7)$ when using a third-order equation of state. The pressure evolution of the unit-cell volume and the Mössbauer hyperfine parameters are in good agreement with previous studies, which were limited to a lower pressure range.

Keywords: High pressure, phase transition, ringwoodite, XRD, Mössbauer