

Equation of state and phase transition in KAlSi_3O_8 hollandite at high pressure

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ABSTRACT

The tetragonal hollandite structure (KAlSi_3O_8 hollandite) has been studied up to 32 GPa at room temperature using high-pressure in-situ X-ray diffraction techniques. A phase transformation from tetragonal $I4/m$ phase to a new phase was found to occur at about 20 GPa. This transition is reversible on release of pressure without noticeable hysteresis and hence this new high-pressure phase is unquenchable to ambient conditions. The volume change associated with the transition is found to be small (not measurable), suggesting a second order transition. The diffraction pattern of the high-pressure phase can be indexed in a monoclinic unit cell (space group $I2/m$), which is isostructural with $\text{BaMn}_8\text{O}_{16}$ hollandite. The γ angle of the monoclinic unit cell increases continuously above the transition. A Birch-Murnaghan equation of state fit to pressure-volume data obtained for KAlSi_3O_8 hollandite yields a bulk modulus $K_0 = 201.4$ (7) GPa with $K'_0 = 4.0$.