

Poster

(*P, T*)-behavior of an intermediate scapolite of unusual *I4/m* symmetry

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Scapolites are usually formed under metamorphic conditions in the presence of fluids, but are also reported as constituents of upper mantle xenoliths [1,2]. The scapolite series of minerals represents a complex non-binary solid solution, whose end members are marialite [Na₄Al₃Si₉O₂₄Cl], meionite [Ca₄Al₆Si₆O₂₄CO₃] and silvialite [Ca₄Al₆Si₆O₂₄SO₄]. The members whose composition falls on the marialite-meionite joint appears to be the most common in natural occurrences [3,4]. The representatives close to marialite, on one side, and to meionite, on the other side, are usually reported to crystallize in the tetragonal *I4/m* space group, while intermediate scapolites are usually found in the primitive space group *P4₂/n*. In this study, we report a scapolite sample from Madagascar, which composition falls between those of the end-members marialite and meionite: (Na_{1.86}Ca_{1.86}K_{0.23}Fe_{0.01})(Al_{4.36}Si_{7.64})O₂₄[Cl_{0.48}(CO₃)_{0.48}(SO₄)_{0.01}]. Based on both X-ray and neutron single-crystal diffraction data, an anomalous *I*-centered lattice (*I4/m* space group) is observed. This unusual symmetry, for an intermediate scapolite, may be assigned to the presence of anti-phase domains too small to be detected by diffraction techniques. Experimental data based on *in situ* single-crystal X-ray diffraction show that the unusual *I*-centered lattice is always preserved at varying pressure and temperature up to 1000°C, whereas a phase transition towards a triclinic polymorph, that can be described according to a non-conventional *I*-1 symmetry, was found to occur at 9-10 GPa, with a minor influence from temperature. These results suggest that intermediate scapolites may crystallize with an *I*-centered crystal structure, whose preservation at ambient conditions may provide information on the thermal history (fast cooling) of the sample.

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