

Chemical substitutions, paragenetic relations, and physical conditions of formation of hōgbomite in the Sittampundi layered anorthosite complex, South India

PULAK SENGUPTA,^{1,*} UTTAM K. BHUI,¹ INGO BRAUN,² UPAMA DUTTA,¹ AND D. MUKHOPADHYAY³

¹Department of Geological Sciences, Jadavpur University, Kolkata 700032, India

²Mineralogisch-Petrologisches Institut, Universität Bonn, Poppelsdorfer Schloss, 53115 Bonn, Germany

³Department of Geology, University of Calcutta, 29, Ballygunge Circular Road, Kolkata 700019, India

ABSTRACT

Layered anorthosite, chromiferous pyroxenite, and spatially associated mafic and felsic rocks of the 2.9 Ga Sittampundi layered complex (SLC), South India, underwent high-grade metamorphism at ca. 2.5 Ga and were subjected to amphibolite-facies metamorphism accompanied by intrusion of granitoid plutons during late-Neoproterozoic tectonothermal activity (0.72–0.45 Ga). During the latter event, anorthosite developed millimeter- to centimeter-thick compositional layers rich in clinopyroxene+amphibole+clinozoisite+chlorite±chromite and corundum+spinel+chlorite. Tiny grains of hōgbomite replaced only the grains of corundum and spinel and are in textural equilibrium with chlorite. The studied hōgbomite contains appreciable TiO₂ (>4.4 wt%) but insignificant NiO and ZnO. Cr₂O₃ content reaches up to 0.35 wt% only in chromite-bearing samples. Systematic partitioning of Fe and Mg between hōgbomite and associated Fe-Mg minerals demonstrate attainment of chemical equilibrium among these phases. Integrating textural relations and algebraic analyses of the phase compositions, several reactions were constructed involving the associated oxide phases (spinel, corundum, hōgbomite), calcite, and silicates (amphibole, chlorite, anorthite, clinozoisite). Interpretation of the reaction reveals that (1) Mg²⁺ and Ti⁴⁺ were mobile for more than 2 cm during the formation of hōgbomite and chlorite, and (2) chloritization of amphibole in the clinopyroxene+amphibole-bearing layers released Ti that was transported to the spinel+corundum-bearing layers to develop hōgbomite. Stability fields of some critical mineral assemblages in P - X_{CO_2} and T - X_{CO_2} space combined with geothermobarometry in the associated rocks tightly constrain the growth of hōgbomite in the presence of aqueous fluids ($X_{\text{CO}_2} < 0.15$) to the P - T range of 7 ± 1 kbar, 650 ± 50 °C. These aqueous fluids, presumably derived from the Pan-African granitoid batholiths, chloritized amphibole grains, and transported the released Ti⁴⁺ to the spinel+corundum-bearing layers to develop hōgbomite. Topological relations in isothermal-isobaric fugacity diagrams ($\log f_{\text{O}_2}$ - $\log f_{\text{S}_2}$ and $\log f_{\text{O}_2}$ - $\log f_{\text{H}_2\text{O}}$) in the system FeO-Al₂O₃-TiO₂-O₂-S₂-H₂O-CO₂ (+MgO, Cr₂O₃) indicate that the stability and compositional characteristics of natural hōgbomite are strongly influenced by f_{O_2} , f_{S_2} , $f_{\text{H}_2\text{O}}$, and concentrations of other soluble species (Ti, Mg, Cr, etc.) in the metamorphic fluids.

Keywords: Hōgbomite, corundum, spinel, anorthosite, Ti-mobility, Sittampundi, South India