

## **Hematite and magnetite precipitates in olivine from the Sulu peridotite: A result of dehydrogenation-oxidation reaction of mantle olivine?**

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### **ABSTRACT**

Analytical electron microscopic observations have been carried out on a garnet peridotite from the Maobei area, Sulu ultrahigh-pressure terrane. The results showed that olivine in this garnet peridotite (5.3–6.6 GPa; 853–957 °C), contains precipitates of chromian magnetite and chromian-titanian hematite at dislocations and (001) faults. Specific crystallographic relationships were determined between these precipitates and the olivine host, viz.  $[101]_{\text{Mt}}//[001]_{\text{Ol}}$ ,  $[110]_{\text{Mt}}//[0\bar{1}1]_{\text{Ol}}$ , and  $[0\bar{1}1]_{\text{Mt}}//[011]_{\text{Ol}}$ ; and  $[0001]_{\text{Hm}}//[100]_{\text{Ol}}$  and  $[10\bar{1}0]_{\text{Hm}}//[001]_{\text{Ol}}$ . These oriented oxides are not associated with silicate/silica phases and therefore cannot be accounted for by the mechanism of olivine oxidation. It is postulated that these magnetite and hematite precipitates most likely have resulted from dehydrogenation-oxidation of nominally anhydrous mantle olivine during rock exhumation. In view of the contrasting diffusion rates of H and Fe in the olivine lattice, it is suggested that the formation process might actually take place in steps. Hydrogen diffusion with concomitant quantitative oxidation of  $\text{Fe}^{2+}$  to  $\text{Fe}^{3+}$  in olivine occurred early during initial rock exhumation and was followed by slow Fe diffusion forming magnetite/hematite at stacking faults and dislocations within the olivine lattice. Two requirements are essential under such a scenario: an ample amount of H content of the olivine, and an appropriate exhumation rate, probably in the range of 6–11 mm/year, of the host rock. It is also noted that such dehydrogenation-oxidation processes may hamper a correct estimate of the actual *P-T* conditions and mantle oxidation state based on mineral chemistries present in mantle eclogite/peridotite. The present study demonstrates that oriented mineral inclusions may not necessarily form through exsolution processes *sensu stricto*, but may form through a series of more complicated reaction mechanisms.

**Keywords:** Magnetite, hematite, olivine, dehydrogenation-oxidation, UHP peridotite