

Geobarometric constraints to low-P evolution of lithospheric mantle

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During tectonic exhumation at extensional settings (mid ocean ridge and passive continental margin), mantle peridotites can recrystallize from spinel- to plagioclase-facies conditions [1, 2]. This subsolidus decompressional evolution is documented in peridotites from orogenic and ophiolitic massifs. Recent subsolidus experiments on fertile and depleted lherzolites in the Na₂O-CaO-FeO-MgO-Al₂O₃-SiO₂-Cr₂O₃-TiO₂ system, at P=0.27-1.0 GPa, T=900-1200°C have evidenced systematic mineral compositional variations within the plagioclase facies [3]. Experimental results indicate that Ca-Na partitioning between plagioclase and clinopyroxene strictly depends on P; in particular, plagioclase records a marked anorthite content increase (An=59-83) within a rather narrow range of P. Remarkably, plagioclase composition is similar in both fertile and depleted lherzolites, which have quite different bulk X_{Cr} but comparable bulk Na₂O/CaO ratios. This indicates that the composition of plagioclase does not depend on bulk X_{Cr} and reasonably can be used as a marker of recrystallization pressure in ultramafic rocks with fixed Na₂O/CaO ratios. In order to constrain the P-T slope of plagioclase An-isopleths, and make them applicable to natural occurrences, a new set of piston cylinder experiments on fertile lherzolite have been performed between 0.5-0.8 GPa, 1050-1150°C. Experimental results are then applied to plagioclase-facies recrystallized peridotites. We consider as case study the peridotites and associated pyroxenites from Northern Apennine (External Liguride ophiolitic Unit, Italy). They show textural evidence of plagioclase-facies re-equilibration and systematic compositional changes in minerals [4], consistent with experimental results. Detailed microstructural and mineral chemistry investigations have revealed An-reverse zoning in plagioclase of both peridotites and pyroxenites (An=57-80 and An=62-84 respectively), likely related to different stages of low P re-equilibration. Here we show that An content in plagioclase represents a potential geobarometer to reconstruct the subsolidus decompressional evolution of lithospheric mantle at extensional settings.

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Space - time variability of ²¹⁰Pb fluxes in the Arabian Sea

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Time series samples of settling sediment particles were collected during 1986-'87 from three stations in a east to west transect along 15°N and one station at 10°N during 1993-'94 from ~1000m (shallow) and ~3000m (deep) waters of the Arabian Sea.[1] The traps deployed at these four locations provide unique environmental settings to examine the application of ²¹⁰Pb as a tracer for water column transport as the magnitude of the particle flux is laterally variable at these locations. The objectives of these studies are to measure the settling fluxes of ²¹⁰Pb, to understand their spatial and seasonal variation and their relation to the water column scavenging processes.

The ²¹⁰Pb concentration is in the range of 10 - 600 (dpm g⁻¹) and the corresponding fluxes are 0.2 - 0.4 (dpm.cm⁻² y⁻¹) in the shallow and 0.4 - 0.7 (dpm.cm⁻² y⁻¹) in the deep traps. The ²¹⁰Pb concentration as well as the fluxes showed more pronounced seasonal variation than the spatial as well as the vertical variation.

The overall geochemical mass balance scenario of ²¹⁰Pb in water, settling particles and underneath sediment revealed a large advective export of ²¹⁰Pb to the continental margins and its subsequent enhanced deposition fluxes indicative of 'boundary scavenging' mechanism.[2, 3].

The ²¹⁰Pb Flux (F) / ²¹⁰Pb production (P) ratio – (F/P) and organic Carbon (C_{org}) as a proxy for C_{org} content prediction concept [4] pose basic question of applicability for Arabian Sea water column features.

[1] Ramswamy *et al.* (1991) *Deep-Sea Research I* **38**, 169-184. [2] Borole (2002) *Deep-Sea Research I* **49**, 1055-1069. [3] Scholten *et al.* 2005 *EPSL* **230**, 319-337. [4] Moore & Dymond (1998) *Nature* **331**, 339-341.