## Zircon solubility in supercritical fluids and zirconium mobilization in subduction zones

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Supercritical fluids may serve as effective agents for the transfer of Zr and other high field strength elements (HFSE) in subduction zone, yet zircon solubility investigation in such solute-rich fluids is rare. Here, zircon solubility (ZrO<sub>2</sub> content at zircon saturation) in solute-rich supercritical fluids in KAlSi<sub>3</sub>O<sub>8</sub> ( $\pm$  K<sub>2</sub>O  $\pm$  Al<sub>2</sub>O<sub>3</sub>) – H<sub>2</sub>O systems was investigated at 2.0-6.0 GPa and 800-1000°C, close to the top-slab conditions at sub-arc depths. All the supercritical fluids were quenched to hydrous glasses and thus can be analyzed using electron probe.

ZrO<sub>2</sub> content in the supercritical fluids increases with temperature and solute content and alkalinity (molar K/Al ratio) but decreases with pressure. At 2.0 GPa, ZrO<sub>2</sub> content in KAlSi<sub>3</sub>O<sub>8</sub> – H<sub>2</sub>O fluids (K/Al = 1.0) increases from <100 ppm at 800-900°C and solute content < 50 wt% to 500-700 ppm at 1000°C and solute content >70 wt%. At 2.0 GPa and 1000 °C, addition of K<sub>2</sub>O to the KAlSi<sub>3</sub>O<sub>8</sub> - H<sub>2</sub>O system strongly enhances ZrO<sub>2</sub> content to ~10000 ppm at K/Al of 1.67. Therefore, solute content and alkalinity exert primary controls on zircon solubility. At 4.0 and 6.0 GPa, the crystallization of muscovite and kyanite results in K/Al higher than 1.0 in the KAlSi<sub>3</sub> $O_8$  – H<sub>2</sub>O fluids and the negative effect of pressure on zircon solubility was offset and even reversed due to the enrichment of K2O. Comparison with literature data shows that zircon solubility in solute-rich supercritical fluids is at least 10 times higher than in dilute aqueous fluids. During subduction, the fluid released from slab will become higher in solute and alkali/Al due to increasing P-T and formation of Al-rich residual phases. Zircon solubility and element carrying capacity of supercritical fluid thus increase with subduction depth. The experimental results not only explain zircon dissolutionredeposition in metamorphic rocks, but also zircon presence in wedge peridotites and Zr enrichment of primitive arc basalts relative to ocean-ridge basalts.