

**LETTER**

**Very low solubility of rutile in H<sub>2</sub>O at high pressure and temperature, and its implications for Ti mobility in subduction zones**

**PETER TROPPER<sup>1,2</sup> AND CRAIG E. MANNING<sup>2,\*</sup>**

<sup>1</sup>Institute of Mineralogy and Petrography, University of Innsbruck, Innrain 52, A-6020 Innsbruck, Austria

<sup>2</sup>Department of Earth and Space Sciences, University of California, Los Angeles, California 90095-1567, U.S.A.

**ABSTRACT**

The solubility of rutile in H<sub>2</sub>O has been measured at 1000–1100 °C, 1–2 GPa. The data indicate that solubility is very low over the investigated range, with a maximum of 4.7 millimol/kg H<sub>2</sub>O at 1100 °C, 2 GPa. The data were fit with the equation  $\log m_{\text{Ti}} = 4.892 - 10470/T + 0.1923P$ , where  $m_{\text{Ti}}$  is Ti molality,  $T$  is in Kelvins, and  $P$  in GPa. When compared to previous results, the new data indicate substantially lower solubility, opposite pressure dependence, and thermodynamic properties of the reaction  $\text{rutile} = \text{TiO}_{2,\text{aq}}$  that are now consistent with other oxide hydrolysis reactions. Calculations of Ti transport during mantle metasomatism by H<sub>2</sub>O in subduction zone environments predict much lower Ti mobility at all conditions. These results offer strong support for models of Ti retention in eclogites during slab devolatilization, and require that examples of significant Ti mass transfer be explained by complexing agents in solution, most likely aluminosilicate complexes.