



## **Metasomatic alteration of zircon in high pH fluids under high-grade conditions**

**D.E. Harlov**

GeoForschungsZentrum Potsdam, Telegrafenberg, D-14473 Potsdam, Germany  
(dharlov@gfz-potsdam.de)

In nature zircon is one of the principle accessory minerals used for the dating of geologic processes. As a consequence, the stability of zircon in the presence of various possible metamorphic fluids under a range of P-T conditions and its subsequent instability with respect to some of these fluids has begun to be explored experimentally as well as speculated upon in a series of studies of metasomatised zircons in nature (see review in Geisler et al., 2007). Natural alteration of zircon takes place via one of two metasomatically induced processes: diffusion-reaction or coupled dissolution-reprecipitation. Both processes result in the zircon being either partially or totally replaced by either a new re-equilibrated zircon composition or a totally new zircon-related phase. In this study, fragments (50 - 200  $\mu\text{m}$ ) from a large, inclusion-free, clear, light brown, euhedral zircon collected from a nepheline syenite pegmatite (Seiland magmatic province, northern Norway) are experimentally reacted in 20 mg batches with a series of high pH fluids plus a Th source (5 mg  $\text{ThO}_2$  +  $\text{ThSiO}_2$ ) in sealed Pt capsules at 900 °C and 1000 MPa for 8 days in the piston cylinder press using a  $\text{CaF}_2$  setup. Fluids included 5 mg 2 N NaOH, 5mg 2 N KOH, 10 mg  $\text{Na}_2\text{Si}_2\text{O}_5$  + 5 mg  $\text{H}_2\text{O}$ , and 5 mg  $\text{Ca}(\text{OH})_2$  + 5 mg  $\text{H}_2\text{O}$ . The results of these experiments indicate that only the fluid containing  $\text{Ca}(\text{OH})_2$  reacted with the zircon. This reaction took the form of partial replacement, via dissolution-reprecipitation, of the majority of the zircon grains with baddeleyite ( $\text{ZrO}_2$ ). The rims of the remaining zircon grains are partially replaced by zircon enriched in Th. The replaced rims are characterized by a sharp compositional boundary between the altered and original zircon as well as by a micro-porosity. The preliminary results from these experiments suggest that in nature,

one possible high pH fluid responsible for the alteration of zircon under high-grade conditions could be one with a substantial  $\text{Ca}(\text{OH})_2$  component.

Geisler, T., Schaltegger, U., and Tomaschek, F. (2007) Re-equilibration of zircon in aqueous fluids and melts. *Elements* 3, 43-50.