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"Hot" Zircons from the Merlin Kimberlite Field, Northern Territory, Australia

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Introduction

Kimberlites and lamproites are volatile-rich, potassic ultramafic magmas that form at depths in excess of 150 km. They ascend to the surface rapidly (time spans of hours to days), entraining mantle and crustal wall rock fragments along the way. If diamondiferous portions of the mantle are transgressed, diamonds will be transported to the Earth's surface. Similarly, zircon and other U-bearing minerals will be entrained in the erupting magma during transit through the crust, and will also be transported to the surface.

Current diamond exploration strategies include the search for kimberlite indicator minerals (KIMS) such as chromite, garnet and Cr-diopside in stream sediments and soil samples. KIMS are, however, prone to chemical dissolution in tropical weathering conditions. Therfore, alternative approaches involving the geochemical analysis of resistant minerals such as zircon would enhance exploration success in tropical regions such as Australia, India and Brazil.

Background

Zircons begin to retain Pb when they cool below about 900°C so U/Pb ages generally record the age of formation of the grain. However, because helium readily diffuses out of zircon at temperatures above 180°C (Reiners et al., 2002), zircon that originates from mid- to lower-crustal levels will not retain the radiogenic helium produced during the radioactive decay of naturally occurring U and Th:

This means that mid- to lower-crustal zircons will have young (U-Th)/He ages (Figure 1). In contrast, upper crustal zircons will have very high contents of radiogenic helium (and old (U-Th)/He ages) because all of the ⁴He atoms produced over time have been locked into this "cooler" zircon structure (T<180°C).



The result is that kimberlitic and lamproitic zircons from the mid- to lower-crust will have a markedly different U-Th-Pb-He age to zircons eroded from the upper crust. In fact, the zircon (U-Th)/He age populations should be bimodal where zircons are sampled from all crustal depths (Figure 1). This work reports the results of U-Th-Pb-He dating of samples obtained from the Merlin Kimberlite, Northern Territory, Australia.



Figure 1. Helium retention in zircon is temperature dependent and controlled primarily by the Earth's geothermal gradient. Zircon grains from the lower- to mid-crustal regions entrained within a kimberlite eruption ("hot" zircon) will have ages equivalent to the eruption age of the kimberlite, whereas upper crustal zircons ("cold" zircon) will have cratonic ages. In this hypothetical example where a kimberlite eruption occurred today in a Proterozoic craton, the U-He age distribution of zircon xenocrysts would be either 0 for "hot" zircons transported from great depth or 1700 Myr for "cold" zircons transported from relatively shallow levels.

Samples and Methods

The Merlin kimberlite deposits are hosted within the Early Cambrian Bukalara Sandstone (Muir, 1980). We separated zircon from: (i) a bulk sample of the Sacramore kimberlite pipe in the Merlin kimberlite field and (ii) a core sample of unmineralized Bukalara Sandstone located approximately 10 km from the kimberlite. Individual zircons were mounted in epoxy and polished to expose a surface suitable for SHRIMP analysis in the John de Laeter Centre of Mass Spectrometry, Curtin University of Technology, Perth, Australia. The zircon standard BR266 (559.1Ma, 206 Pb/ 238 U = 0.09059; Stern 2001) was used to calibrate U-Th-Pb isotopic ratios and elemental abundances. Corresponding 206 Pb/ 238 U ages were corrected for common Pb using the 208 method (eg., Stern 1997).

Helium was thermally extracted from single crystals, heated using a 1064 nm Nd-YAG laser. ⁴He abundances were determined by isotope dilution using a pure ³He spike, calibrated daily against an independent ⁴He standard. The uncertainty in the sample ⁴He measurement is <1%. The U and Th content of degassed apatite were determined by isotope dilution using ²³⁵U and ²³⁰Th spikes. Zircon was digested in Parr bombs using HF. Standard solutions containing the same spike amounts as samples were treated identically as were a series of unspiked reagent blanks. For single crystals digested in small volumes (0.3-0.5 ml), U and Th isotope ratios were measured to a precision of < 2%. Overall the (U-Th)/He thermochronology method at CSIRO has a precision of 2.5% and more detailed description of U and Th analysis can be found in Evans et al. (2005). The zircon (U-Th)/He and U/Pb ages were plotted using Isoplot/Ex 2.2 (Ludwig 2000).



Figure 2. Zircon (U-Th)/He and U/Pb ages for Merlin kimberlite displayed as a function of relative probability.

Results





2), consistent with the Mesoproterozoic formation of the North Australian Craton and indicating that the zircons in the mineral separate obtained from the kimberlite are clearly of xenocrystic origin. That the mean U/Pb age of zircon from the Bukalara Sandstone country rock is statistically indistinguishable from that of the Merlin kimberlite zircon (Figure 3) provides even more definitive evidence.

The Merlin kimberlitic zircon (U-Th)/He thermochronometry (n=33) yielded a weighted average age of 368±18 Ma (Figure 2), within error of previously determined Rb/Sr (367±4 Ma) and ⁴⁰Ar/³⁹Ar (376-380±4 Ma) dating of phlogopite model ages for the Merlin field (Lee et al., 1998; Hell et al., 2003). The (U-Th)/He results indicate that over 95% of the zircon grains from the Merlin kimberlite have helium ages equivalent to that of the age of emplacement of the diamondiferous kimberlites (367-380 Ma: Lee et This result suggests that the zircon al., 1998). xenocrysts could only have originated from mid- to deep-crustal protoliths where ambient temperatures exceeded the helium closure temperature for zircon (~180°C) at the time of eruption.

Because the majority of zircon grains from the Merlin kimberlite are "hot" zircons, with (U-Th)/He ages indicative of a deep crustal origin, helium thermochronology of zircon can potentially be used as a geochemical exploration tool for kimberlite deposits in tropical environments where chemical weathering destroys traditional KIM phases.



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