

O₂ evolution in the Paleoproterozoic: Experimental simulation of the relationships between Fe in paleosols and PO₂

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Ferrous iron is supplied by mineral dissolution for water in a weathering profile. Because Fe²⁺ is soluble and Fe³⁺ is insoluble, Fe remaining in the profile as Fe³⁺ is related to PO₂ according to Fe²⁺ oxidation kinetics,

$$-[\text{Fe}^{2+}]/dt = k[\text{Fe}^{2+}][\text{OH}^-]^2[\text{PO}_2]^x \quad (1)$$

where k is the rate constant and x the variable ($0 \leq x \leq 1$). On the other hand, the mineral dissolution rates and Fe behavior during weathering under low O₂ conditions are poorly known. We have simulated weathering under low O₂ conditions in the laboratory, examined the Fe redistribution, and established the relationships between Fe in the dissolution system and PO₂ levels.

Olivine, Mg_{1.82}Fe_{0.18}SiO₄, was dissolved in a glove box at 0.2, 8.9x10⁻³, 9.4x10⁻⁴ and 8.6x10⁻⁵ atm of PO₂ in a flow-through system. To model the Fe behavior during weathering, olivine was also dissolved in the ambient atmosphere changing pH and temperature. To discuss the effects of PO₂ on the Fe behavior during dissolution or weathering, we define a ratio of the total amount of precipitated Fe³⁺ (remaining in a profile) (Fe_{prec}) to that of Fe²⁺ dissolved from mineral (Fe_{diss}) as $\Phi = \text{Fe}_{\text{prec}}/\text{Fe}_{\text{diss}}$. Fig. 1 shows the relationships between Φ and PO₂ (diamonds, observed values; dashed curves, calculated variations with different x values in equation (1)). The Φ value increases with increase in PO₂ (solid curve). Furthermore, the x value decreases with decrease in PO₂. The O₂ level is considered to have increased from 10⁻⁶ to 10⁻³ atm in the Paleoproterozoic. Therefore, our results under low O₂ conditions indicate that Φ is a good indicator of the oxidation state of paleosols, and more importantly, we can quantitatively estimate PO₂ levels at the time of weathering based on the Fe²⁺ and Fe³⁺ concentrations in paleosols.

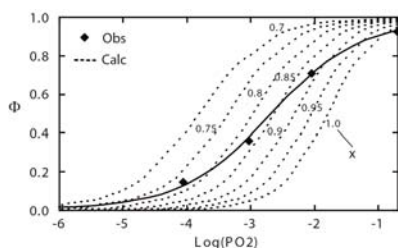


Figure 1: Relationships between Φ and PO₂.

An alternative interpretation of U-Th isochron and estimation of magma residence time

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Mineral U-Th isochron systematics is designed to date the eruption age of young (<350 ka) volcano, but researches provide alternative interpretation of timescales of magma residence and crystallization.

In one case of Tianchi Volcano [1], a reliable mineral U-Th isochron reveals a calculated age of ca. 23ka. Since the eruption age of the volcano is ~1000 aBP, the difference is regarded to reflect the magma residence time. After that, more cases prove that the mineral U-Th isochron provides an approach to estimate the timescale of magma residence and crystallization. However, numerical modelling and theoretical interpretation on this alternative interpretation is still poorly known now, here we report a numerical model on U-Th decay in a prolonged magma crystallization process.

Crystallization in magma chamber can be described by a zoning growth model ($^{230}\text{Th}/^{232}\text{Th}$)_t, and the average isotopic ratio is an integral

$$(1) \quad \left(\frac{^{230}\text{Th}}{^{232}\text{Th}}\right) = \int_0^T \left(\frac{^{230}\text{Th}}{^{232}\text{Th}}\right)_t dv dt$$

is the isotopic ratio in a crystallization zoning belt, and its variation can be described by the normal decay equation,

$$\left(\frac{^{230}\text{Th}}{^{232}\text{Th}}\right)_t = \left(\frac{^{230}\text{Th}}{^{232}\text{Th}}\right)_0 \cdot e^{-\lambda t} + \left(\frac{^{238}\text{U}}{^{232}\text{Th}}\right)_0 \cdot (1 - e^{-\lambda t})$$

Suppose the magma crystallization occurs in a square order of

$$dv = kt \cdot dt$$

we can calculate the equation (1), and get still a linear equation of U-Th isotopes. The linear slope is a function of crystallization time and eruption age

$$m = \frac{1}{\lambda T_m} \cdot (e^{-\lambda(T_2+T_m)} - e^{-\lambda T_2})$$

According to the model, crystallization time of Tianchi Volcano is about 100 ka. This model can also be used on other volcanoes, and can provide more precise estimation on magma crystallization time.

[1] Dunlap CE. 1996. PhD thesis. University of California Santa Cruz. 1-215.