

# $\delta^{142}\text{Ce}$ minus $\delta^{146}\text{Nd}$ value as a redox indicator in Earth's surface environments

JIANGHAO BAI, ZHIBING WANG, JINLONG MA AND  
GANGJIAN WEI

Guangzhou Institute of Geochemistry, Chinese Academy of Sciences

Presenting Author: baijianghao@gig.ac.cn

Stable Ce isotope ratios are considered potential indicators for tracing the redox state of Earth's surface; however, their quantitative relationship remains undefined. Here, we investigated the influences of both redox and non-redox processes on stable Ce isotope fractionation in a terrestrial environment by analyzing the weathering products, selective soil chemical extracts, and parent materials of a typical basaltic profile formed in Hainan Island, South China. During redox processes, the preferential enrichment of the heavier  $^{142}\text{Ce}$  in the tetravalent Ce and/or the inheritance of the heavier isotope signature from the soluble led to the crystalline Fe-Mn (hydro)oxide phases having the highest  $\delta^{142/140}\text{Ce}$  values (up to  $+0.184 \pm 0.040\%$ ). This phenomenon is likely governed by the oxidative adsorption of Mn oxides, while Fe-oxides appear to be less important. In addition, distinct Mn concentrations are important in the  $\tau_{\text{Th,Ce}}$  and  $\delta^{142/140}\text{Ce}$  values but did not correlate with Ce anomalies, indicating that the Mn (hydro)oxides  $\delta^{142/140}\text{Ce}$  values may be more reliable than Ce/Ce\* in quantitatively tracing redox conditions. Regarding non-redox processes, we discussed the impact mechanisms, including external input, pH, organic matter complexation, adsorption/co-precipitation of the poorly crystalline Fe-Mn oxide, and the formation of clay minerals on stable Ce isotopic fractionation. Results indicated that Fe-Mn-Al oxides preferentially enriched the heavier  $^{142}\text{Ce}$ , while the formation of kaolinite preferentially incorporated the lighter  $^{140}\text{Ce}$ . However, the impact of external input, pH, organic matter complexation, and clay mineral adsorption on stable Ce isotopes appeared to be limited. Furthermore,  $\delta^{142/140}\text{Ce}$  values displayed a positive correlation with  $\delta^{146/144}\text{Nd}$  in both the  $0.5 \text{ mol L}^{-1}$  HCl leachates and residual phases, suggesting that stable Ce and Nd isotopes behave similarly during non-redox processes. To isolate the signal of stable Ce isotopes in response to redox conditions, we proposed that  $\delta^{142/140}\text{Ce}$  minus  $\delta^{146/144}\text{Nd}$  could serve as a quantitative proxy for tracing redox fluctuations in terrestrial environments.