The heating rate controls chert $\delta^{18}O$

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The oxygen isotope composition of chert reveals an increasing trend throughout Earth's history. This trend has been explained by decreasing seawater temperatures [e.g. 1], increasing seawater δ^{18} O [e.g. 2], or by an overprinting of the rock record [3]. These hypotheses are called into question by recent evidence for constant seawater δ^{18} O [e.g. 4] and seawater temperatures <40°C in the Archean [5, 6].

To re-evaluate the utility of chert $\delta^{18}O$ in the framework of detritus-controlled rates of silica diagenesis [7], we have analyzed $\delta^{18}O_{VSMOW}$ at the bulk- and μ m- scale of pure cherts and siliceous shales of Precambrian-Cambrian age from SE China. From bulk rock $\delta^{18}O$ we have calculated the $\delta^{18}O$ of the authigenic silica and measured $\delta^{18}O$ of silica in detritusrich samples by SIMS.

Authigenic silica ranges from 12.6 to 24.9 ‰ δ^{18} O and correlates with detritus concentrations, an attribute of temperature-dependent oxygen isotope fractionation during opal-CT to quartz transformation. Peak diagenetic temperatures of *circa* 260°C, as determined by Raman spectrometry of carbonaceous material, show that chert reliably records the diagenetic δ^{18} O signature set during the final diagenetic silica dissolution-reprecipitation step.

Our work shows that δ^{18} O depends on the crustal thermal gradient, because the rate of silica polymorph transformation predominantly depends on temperature [e.g. 8] and because temperature exerts major control over δ^{18} O. We calculate how thermal gradient and burial rate affect δ^{18} O of authigenic quartz. Our results suggest that at least a part of the long-term secular increase in δ^{18} O of chert is caused by decreasing geothermal gradients due to the cooling of the solid Earth.

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