An oceanic oxidation event coincident with the Shuram carbonate-carbon isotope excursion

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The origin of the Ediacaran Shuram negative carbonatecarbon isotope ($\delta^{13}C_{carb}$) excursion (<-10 ‰) remains hotly debated due to its large magnitude and variability in the absence of a coupled excursion in organic carbon ($\delta^{13}C_{ore}$). Here we present the first measurements of carbonate iodine-tocalcium ratios (I/Ca) across this $\delta^{13}C_{\text{carb}}$ excursion, providing evidence for an oxidation event and against a diagenetic origin¹. The oxidized iodine species, iodate, is restricted to oxic waters and is the exclusive iodine species incorporated during carbonate precipitation², meaning changes in I/Ca can record relative fluctuations in dissolved O2 concentrations. Further, controls on the total marine iodine reservoir, and hence I/Ca, include the burial and remineralization of organic matter (OM), iodine's biggest sink². I/Ca within the Khufai of Oman and Doushantuo of South China both increase distinctly in phase with the shift to negative $\delta^{13}C_{\text{carb}}$ values. Our studies of diagenetic alterations to primary I/Ca ratios of modern and recent carbonates reveal that, if anything, primary I/Ca ratios decrease during carbonate diagenesis. The globally extensive increase in I/Ca is best explained by an increase in the total marine iodine reservoir linked to an oxidation event promoting enhanced OM remineralization, thus supporting previous models for the magnitude of the $\delta^{13}C_{\mbox{\tiny carb}}$ excursion and lack of coupled $\delta^{13}C_{\mbox{\tiny org}}$ behavior as a result of oxidation of a fossil organic source³. However, a decrease in pre-Shuram I/Ca values at the end of the $\delta^{13}C_{carb}$ excursion infers a return to reducing conditions, suggesting that the Shuram does not represent a terminal oxidation event.

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