The origin of sulfur in Canary Island magmas and its implications for Earth's deep sulfur cycle

ZOLTÁN TARACSÁK¹, MARGARET E. HARTLEY², RAY BURGESS², MARIE EDMONDS¹, MARC-ANTOINE LONGPRÉ³, BRIAN DAVID MONTELEONE⁴, ROMAIN TARTÈSE² AND ALEXANDRA (SASHA) V. TURCHYN¹

Oxygen fugacity (fO_2) controls whether sulfur is present as sulfide or sulfate in silicic magmas: increasing fO_2 can lead to an order of magnitude increase in sulfur solubility [1]. This is reflected in the sulfur content of comparatively reduced and oxidised melts erupting at mid-ocean ridges (0.1-0.15 wt%) and at subduction-related volcanic arcs (up to 0.7 wt%), respectively [2]. Some ocean islands basalts (OIBs), including those erupted at El Hierro in the Canary Islands, reach S contents similar to subduction-related magmas [3]. While at subduction zones elevated sulfur contents and fO_2 is explained by slab-mantle interaction [1], the cause of sulfur enrichment and oxidation at ocean islands is more enigmatic.

We use sulfur isotope (δ^{34} S) microanalyses of melt inclusions and glasses alongside bulk δ^{34} S analyses of lavas from El Hierro to investigate the origin of observed sulfur enrichment. Melt inclusions have $\delta^{34}S$ values between -1.6% and +4.1% (V-CDT), and S contents between 200 and 5300 µg/g, while lavas have δ^{34} S between -15.5 and +1.4‰ and S contents <400 µg/g. Using degassing and melting models we estimate undegassed melt $\delta^{34}S$ (between 0±2%) and mantle source S content (310±120 μg/g) for El Hierro magmas. On average, 60% of the sulfur in the mantle source is of recycled origin. To satisfy radiogenic isotope constrains on the fraction of recycled material in the mantle source (<10% [5]) we estimate that the recycled component should contain ≥1800 µg/g sulfur. A silica-rich partial melt of recycled oceanic crust could reach these modelled sulfur contents if it was partially oxidised (Fe³⁺/ Σ Fe of >0.15). Based on our results we propose that recycling of Fe³⁺ in silicates is the source of elevated fO2 in OIBs, while silicate melts that contain S⁶⁺ offer an effective pathway for oxidation in the upper mantle via melt-rock reactions.

References:

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¹University of Cambridge

²University of Manchester

³City University of New York

⁴Woods Hole Oceanographic Institution