

Isotope fractionation due to vaporization in tektites: An experimental study

C.A. MACRIS^{1*}, P. NI², E.A. DARLING¹, A. SHAHAR²,
AND E.D. YOUNG³

¹Indiana University – Purdue University Indianapolis, IN, USA
(*correspondence: camacris@iupui.edu)

²Geophysical Lab, Carnegie Institution for Science,
Washington, D.C., USA

³University of California, Los Angeles, CA, USA

Tektites are natural glasses formed as quenched impact melt ejecta. Because they experienced extreme heating while entrained in a hot impact vapor plume, tektites allow insight into these ephemeral events, which play a critical role in planetary accretion and evolution. The isotopic composition of tektite parent materials may be modified by vapor/liquid fractionation at high T in the plume due to preferential loss of light isotopes from the melt through evaporation. Trends from tektite O isotope studies reveal a dichotomy: tektite $\delta^{18}\text{O}$ values are $\sim 4.0\text{-}4.5\%$ lower than their protoliths[1] opposite in direction to a vaporization induced fractionation; yet increasing $\delta^{18}\text{O}$ with decreasing SiO_2 in tektites[2] is consistent with vapor fractionation. Copper isotope studies show that tektite $\delta^{65}\text{Cu}$ values are $1.98\text{-}6.99\%$ higher than protoliths[3], with increases in $\delta^{65}\text{Cu}$ showing a negative correlation with Cu concentration; trends consistent with evaporation induced fractionation.

To understand how volatilization fractionates O and Cu isotopes, we conducted vaporization experiments at high T in an aerodynamic levitation laser furnace. O isotope fractionation experiments used natural tektite material heated to $2150\text{-}2200$ °C for $50\text{-}90$ s while levitated in Ar. Mass losses were from 23 to 26%, reflecting evaporation of Si and O from the melt. The starting tektite had a $\delta^{18}\text{O}$ of 10.06% and the residues ranged from 13.14% to 14.30% . Cu isotope fractionation experiments used synthetic basaltic glass doped with Cu_2O and heated to $2000\text{-}2150$ °C for 30 to 180 s while levitated in Ar. The starting material had a $\delta^{65}\text{Cu}$ of 3.69% , and the residues ranged from 20.26% to 21.51% after 30 s. Based on the $\delta^{65}\text{Cu}$ to Cu concentration relationship, an empirical fractionation factor (α) of 0.9970 to 0.9972 was calculated for the evaporative loss of Cu. These results provide valuable insights into how evaporation fractionates O and Cu isotopes at high temperatures on Earth and other planetary bodies.

[1]Luft et al., *Geochim Cosmochim Acta* **51** (1987). [2]Taylor & Epstein, *J Geophys Res* **74** (1969). [3]Moynier et al., *Geochim Cosmochim Acta* **74** (2010).