

# Carbon Isotope Composition of Basalts from Kama'ehuakanaloa (Loihi Seamount) : Primordial vs. Recycled Carbon in the Hawaiian Mantle Plume

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We analyzed the C isotope composition of vesicle CO<sub>2</sub>, plus He isotopes and He and CO<sub>2</sub> concentrations in the vesicle and glass phase of 37 submarine basalts from the summit, north and south rifts, and east flank of Loihi Seamount. Tholeiites and transitional basalts lie in a narrow range of vesicle  $\delta^{13}\text{C} = -0.9$  to  $-4.6\text{‰}$ , while alkali basalts range from  $-2.1$  to  $-7.2\text{‰}$  (Fig. 1). Calculated total (vesicle+glass)  $\delta^{13}\text{C}$  in tholeiites and transitional basalts range from  $-2.5$  to  $-6.0\text{‰}$  for  $\Delta = \delta_{\text{vapor}} - \delta_{\text{melt}} = +2$  to  $+4\text{‰}$ . Although the vapor-melt isotope fractionation factor ( $\Delta$ ) for Loihi basalts is uncertain, there is a broad trend of vesicle  $\delta^{13}\text{C}$  with the proportion of total CO<sub>2</sub> trapped in vesicles. This trend suggests that  $\Delta$  may be  $\leq +2\text{‰}$  for Loihi magma, and implies total  $\delta^{13}\text{C} = -3$  to  $-5\text{‰}$  for most lavas. This range resembles mantle source values deduced from gas-rich MORBs and Iceland basalts, and for Kilauea deduced from its fumarole gas. However, this similarity is a conundrum because all Loihi basalts extensively degassed their initial CO<sub>2</sub>. CO<sub>2</sub>/Ba systematics and crystal fractionation modeling reveal that Loihi primary magmas (MgO=18 wt.%) lost >97% of their initial CO<sub>2</sub> (0.6 to 1.8 wt.%). Total  $\delta^{13}\text{C}$  vs. total CO<sub>2</sub> in most Loihi tholeiitic and transitional basalts approximates a closed-system degassing trajectory. Extrapolating this trend to initial (pre-degassing) CO<sub>2</sub> concentrations implies  $\delta^{13}\text{C} = -0.5$  to  $-3.0\text{‰}$  in Loihi primary magmas having the highest <sup>3</sup>He/<sup>4</sup>He ratios (>30 R<sub>A</sub>). This primary magma estimate is based on  $\Delta=+2$  during closed-system degassing; larger values of  $\Delta$ , as measured experimentally, and/or open-system degassing, lead to even higher estimates of  $\delta^{13}\text{C}$  for primary magmas. The Hawaiian plume source therefore seems to be characterized by  $\delta^{13}\text{C}$  that is markedly higher than the values of  $-4$  to  $-6\text{‰}$  commonly found throughout Earth's mantle. This higher value of  $\delta^{13}\text{C}$  could imply some tectonic recycling of inorganic C (limestone) to the Loihi mantle source. However, a mantle source  $\delta^{13}\text{C}$  of  $-0.5$  to  $-3.0\text{‰}$  is also within the possible range of Earth's primordial carbon given the large range of  $\delta^{13}\text{C}$  in chondritic meteorites.

