



## Are light $^{13}\text{C}$ diamonds derived from preserved primordial heterogeneity or subducted organic carbon? Using numerical modelling of multi-component mass balanced mixing of stable isotopes

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During the subduction of oceanic crust light volatile elements such as S, C and H are recycled into the upper mantle wedge via slab dehydration and partial melting of oceanic lithosphere. This is evident as arc magmas have higher concentrations of  $\text{SO}_2$ ,  $\text{CO}_2$  and  $\text{H}_2\text{O}$  than mid-ocean ridge basalts (Wallace, 2005). It is also calculated that 50% of the carbon and >70% of the sulphur subducted is returned to the earth's deep mantle (Wallace, 2005).

This work is testing the notion that the subducted organic carbon is a possible source of growth medium for diamonds. Mantle materials display an interesting bimodality in carbon isotopes with a large peak demonstrating the mean mantle value of  $\sim -5\text{‰}$  and a smaller peak consistent with organic carbon at  $\sim -25\text{‰}$  (Deines, 2001). The source of the bimodality remains unresolved with the main theories being; subducted organic carbon, preserved primordial heterogeneity and the existence of a HPHT fractionation process (for a review see Cartigny, 2005).

To test the idea that such organic values of  $\text{d}^{13}\text{C}$  in diamond (ranging from  $-11$  to  $-37\text{‰}$ ) are derived from subducted organic carbon it is essential to compare the  $\text{d}^{13}\text{C}$  values in diamond to other isotopic systems, such as the values for  $\text{d}^{15}\text{N}$  in diamond, as well as values for  $\text{d}^{34}\text{S}$  and  $\text{d}^{18}\text{O}$  in associated syngenetic mineral inclusions.

We have calculated the percentage of organic C-O-N-S in sediments relative to mean mantle values for  $\text{d}^{13}\text{C}$ ,  $\text{d}^{15}\text{N}$ ,  $\text{d}^{34}\text{S}$  and  $\text{d}^{18}\text{O}$  required to produce the observed isotopic ratios found in natural diamonds and syngenetic mineral inclusions. This was done by way of multi-component mass balanced mixing of stable isotopes between sedimentary, organic and mantle materials of varying measured isotope compositions.

### References:

Cartigny, P. 2005. Elements 1, 79-84  
Deines, P. 2001. Earth Science Reviews 58, 247-278  
Wallace, P.J. 2005. Journal of Volcanology and Geothermal Research 140, 217– 240