

## Micro-Raman spectroscopic study on Ararki (L5) chondrite

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We present here the Micro Raman spectroscopic characterization of Ararki (L5) Chondrite. The mineralogical and chemical composition data of the Kaprada meteorite have been discussed elsewhere [1]. Laser-Raman spectra of chromites from Ararki chondrites show characteristic peaks of natural chromites in the region 400 to 800  $\text{cm}^{-1}$  with A1g mode at 700  $\text{cm}^{-1}$  yielding the composition Cr# of about 87, which in excellent agreement with the EPMA data. There is a well known correlation occurs between the A1g mode and the composition of the chromites. Viz., the highest frequency band shifts from 685  $\text{cm}^{-1}$  for high chromium content to 770  $\text{cm}^{-1}$  for Al-rich end in natural chromites. The equilibration temperature is determined by the composition of the few grains of chromite by adopting the method described by Wlotzka [2]. Ararki meteorite showed the chromite composition with  $\text{Fe} \# 100 \times \{\text{Fe}/(\text{Fe}+\text{Mg})\} = 87$  to 89  $\text{Cr} \# 100 \times \{\text{Cr}/(\text{Cr}+\text{Al})\} = 86$  to 87, which yielded the equilibration temperature of be about 730 to 750  $^{\circ}\text{C}$ . Olivine in L6 chondrite is used to determine the residual stress present in the meteorite [3] The residual stress in the olivine is estimated by using the Raman line at 820  $\text{cm}^{-1}$ , and is found to be about 300 MPa for the Ararki meteorite.

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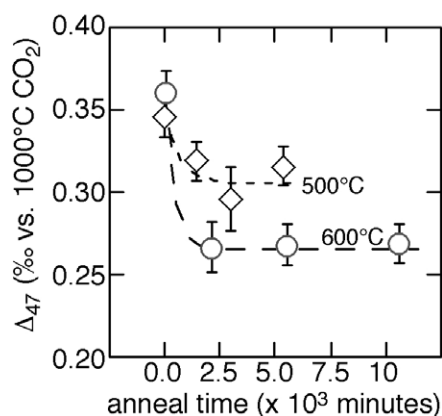
[1] Bhandari N, *et al.*, 2008, *Meteoritics and Planetary Science* v **43**, 761-770. [2] Wlotzka F, 2005, *Meteoritics and Planetary Science* **40**, 1673. [3] Miyamoto M, 1995, *Geophys. Res. Lett* **22**, 437-440.

## Experimental study of solid-state $^{13}\text{C}$ - $^{18}\text{O}$ bond reordering in calcite

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The carbonate clumped isotope thermometer is based on the relative abundance of  $^{13}\text{C}$ - $^{18}\text{O}$  bonds in carbonate minerals, and thus it is important to understand the temperatures and conditions where solid state  $^{13}\text{C}$ - $^{18}\text{O}$  bond reordering occurs on geologically short timescales. This is relevant to the preservation of primary signals in paleoclimate applications, as well as applications in the intermediate temperature realm (ca. 50 – 300 $^{\circ}\text{C}$ ) relevant to diagenesis, hydrothermal mineralization, and basin evolution [1]. I present results from heating experiments designed to deduce Arrhenius parameters and ‘closure temperatures’ for the  $^{13}\text{C}$ - $^{18}\text{O}$  order/disorder process in calcite. Carrara marble heated at 500 $^{\circ}\text{C}$  and 600 $^{\circ}\text{C}$  under dry conditions is measurably changed in ‘clumped isotope’ distribution on timescales of hours to days (Figure 1). The  $\Delta_{47}$  values of  $\text{CO}_2$  extracted from carbonate fail to reach equilibrium values predicted by the theoretical temperature scale of Guo *et al.* [1] (0.26 to 0.25‰), possibly indicating incomplete approach to equilibrium, a non first-order rate law, or other factors. Continuing work will address these questions with additional experiments over an extended the range of temperatures and time durations.



**Figure 1:**  $\Delta_{47}$  values of Carrara marble samples heated at 500 and 600 $^{\circ}\text{C}$  in the presence of dry  $\text{CO}_2$  gas in sealed quartz tubes. Dashed lines are first order reaction progress models. The mean value for unheated Carrara marble is 0.352‰.

[1] Eiler *et al.* 2009 *GCA* **73**, A3222. [2] Guo *et al.* 2009 *GCA* **73**, 7203-7225.