**Raman Spectroscopy and Petrology of Antarctic CR chondrites: Comparison with other carbonaceous chondrites.** M. Komatsu<sup>1,2</sup>, T. J. Fagan<sup>2</sup>, A. Yamaguchi<sup>1,3</sup>, T. Mikouchi<sup>4</sup>, M. E. Zolensky<sup>5</sup>, and M. Yasutake<sup>1,3</sup>. <sup>1</sup>Graduate University for Advanced Studies, SOKENDAI (komatsu\_mutsumi@soken.ac.jp), <sup>2</sup>Dept. Earth Sci., Waseda University, <sup>3</sup>National Institute of Polar Research, <sup>4</sup>Dept. Earth Planet. Sci., University of Tokyo, <sup>5</sup>NASA Johnson Space Center, USA.

# **Introduction:**

In Renazzo-type carbonaceous (CR) chondrites, abundant primary Fe,Ni-metal is preserved in chondrules, but the matrix is characterized by fine-grained magnetite and phyllosilicates [1,2]. This combination of reduced Fe in chondrules with oxidized Fe and phyllosilicates in the matrix has been attributed to aqueous alteration of matrix at relatively low temperatures [1,2].

In this study, we use Raman spectroscopy of matrix and petrology of amoeboid olivine aggregates (AOAs) to evaluate secondary alteration processes in a set of Japanese Antarctic CR chondrites. Raman spectra from organic matter in chondrites can be used as indicators of metamorphic temperature [3,4], and AOAs are sensitive to fluid-assisted metamorphism [5,6]. Our results indicate minimal thermal metamorphic effects, particularly in comparison to metamorphism of the CV3 Allende.

### Methods:

Raman and petrologic data were obtained from polished thin sections (PTS) of CR chondrites A-881828, Y-791498, and Y982495. and carbonaceous chondrite A-881595 in this project. A-811595 was originally classified as a CR chondrite [7], but reclassification as an ungrouped C3 has been suggested [8,9]. Nonetheless, we include A-881595 in this study. In addition to the CRs, some CO and CV chondrites were studied for comparison.

Imaging, mineral identification and EDS analyses were performed using a JEOL JSM-7100F FE-SEM at National Institute of Polar Research (NIPR), and Hitachi S-4500 SEM at University of Tokyo. Raman spectra were collected using a JASCO Raman Spectrometer at NIPR. Analytical parameters are similar to those in [4].

# **Results and Discussion:**

# Raman spectra of polished thin sections

The main tracers of thermal maturity of organic matter (OM) in chondrites are: (1) width of the D-band (band associated with defects; full-width at half peak maximum = FWHM-D) and (2) ratio of peak intensities of the D-band vs. G-band (associated with graphite) =  $I_d/I_g$ . With increasing thermal maturity, FWHM-D decreases and I<sub>d</sub>/I<sub>g</sub> increases [3]. Bonal et al. [4] use these parameters to show variations in metamorphic grade in the CV3 chondrites, with Efremovka near 3.1-3.4 and Allende ≥ 3.6. Their study is based on raw samples, whereas our work is based on spectra from PTS. In our previous work [10], we showed that our Raman

spectra from Efremovka and Allende PTS yielded parameters similar to the results of [4], indicating that reliable OM maturity can be also obtained from PTS.

Raman spectra were collected on randomly selected matrix areas. Raman spectra from CR chondrites in this study exhibit first-order carbon Dand G-bands, at  $\sim 1350$  cm<sup>-1</sup> and  $\sim 1600$  cm<sup>-1</sup> respectively (Fig. 1). They appear similar to those of Y-81020 (CO3.05) and Y980145 (CV3, unclassified petrologic subtype). They are distinct from Allende CV3, which experienced aqueous and thermal alteration (CV>3.6).

Bonal et al. [4] showed that  $I_d/I_g$  increases in petrologic type 3.0 to 3.7, consistent with our results from CO and CV chondrites (Fig. 1). Because the organic composition difference between CR and CV/CO is minimal, Raman parameters can be used for interclass comparison to compare their metamorphic grades [4, 13]. The spectra collected for this study from CR chondrites, all show relatively low I<sub>d</sub>/I<sub>g</sub>, suggesting low thermal maturity, particularly in comparison with Allende (Fig. 1).

# Petrology of AOAs in Antarctic CR chondrites

AOAs in CR chondrites in this study are less abundant than CV and CO chondrites; only a few AOAs are found in each thin section. AOAs in the CR chondrites are composed of forsterite, Al-diopside, anorthite, spinel, Fe,Ni-metal and FeS (Fig. 2a-d). As described in previous studies [2, 11, CR AOAs do not contain secondary 12], the nepheline or fayalitic olivine. In A-881595, some anorthite is replaced by Mg,Al-silicates (Fig.2h), that are similar to phyllosilicates in AOAs in aqueously altered chondrites such as Kaba [6] and Murchison [14].

# Matrix mineralogy

Matrices of CR chondrites are composed of fine-grained minerals such as framboidal and platelet magnetite, sulfides, calcite and phyllosilicates [1]. The abundance of magnetite varies among the CR chondrites, and lath-shaped phyllosilicates (<3 µm) are present in A-881595.

The matrix olivine grains of the CO and CV chondrites examined in this study show a general correlation between grain size and porosity vs. metamorphic subtype (Fig. 3;[6]). Also, the formation of fayalitic olivine appears to have resulted from aqueous alteration. For the CR chondrites, matrix textures show little porosity, similar to weakly metamorphosed CVs and COs (Fig. 3). The presence of lath-shaped phyllosilicates in A-881595 may

indicate heavy aqueous alteration of this sample. **References:** 

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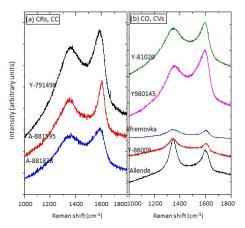


Fig.1. Representative Raman spectra of matrices of carbonaceous chondrites in this study. Spectra are vertically shifted for clarity. (a) CR chondrite Y-791498 and A-881828, and A-881595. (b) Y-81020 (CO3.05), Y980145 (CV3), Efremovka (CV3.1-3.4), Y-86009 (CV3), and Allende (CV>3.6). In each chondrite the first order carbon D- and G-bands are present.

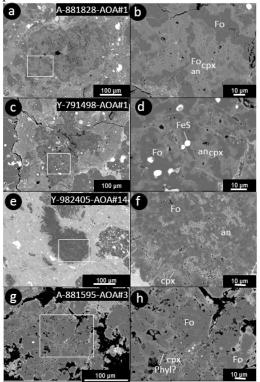


Fig.2. BSE images of AOAs from CR chondrites and A-881595. (a,b) A-881828 (c,d) Y-791498 (e,f) Y982405 (g,h) ungrouped C3 chondrite A-881595.

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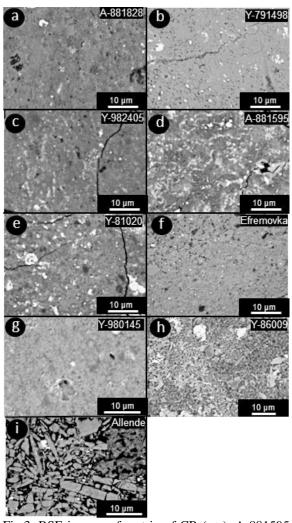


Fig.3. BSE images of matrix of CR (a-c), A-881595 (d), Y-81020 primitive CO (e), and CV chondrites (f-h).