



Deposited via The University of Leeds.

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/id/eprint/128093/>

Version: Accepted Version

---

**Proceedings Paper:**

Vollmer, C, Leitner, J, Busemann, H et al. (2017) Nitrogen functional chemistry of organic grains in CR Chondrites and IDPs. In: Meteoritics and Planetary Science. 80th Annual Meeting of the Meteoritical Society (2017), 23-28 Jul 2017, Santa Fe, NM, USA. Wiley. Article no: A372, p. 6206. ISSN: 1086-9379. EISSN: 1945-5100.

<https://doi.org/10.1111/maps.12934>

---

© 2017 The Meteoritical Society. This is the peer reviewed version of the following article: (2017), 80th Annual Meeting of the Meteoritical Society (2017). Meteorit Planet Sci, 52: 1. which has been published in final form at <https://doi.org/doi:10.1111/maps.12938>. This article may be used for non-commercial purposes in accordance with the Wiley Terms and Conditions for Self-Archiving.

**Reuse**

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

**Takedown**

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing [eprints@whiterose.ac.uk](mailto:eprints@whiterose.ac.uk) including the URL of the record and the reason for the withdrawal request.

## NITROGEN FUNCTIONAL CHEMISTRY OF ORGANIC GRAINS IN CR CHONDRITES AND IDPs.

C. Vollmer<sup>1</sup>, J. Leitner<sup>2</sup>, H. Busemann<sup>3</sup>, N. H. Spring<sup>4</sup>, D. Kepaptsoglou<sup>5</sup>, Q. M. Ramasse<sup>5</sup>, and P. Hoppe<sup>2</sup>

<sup>1</sup>Universität Münster, Inst. f. Mineralogie, Corrensstr. 24, 48149 Münster, Germany, christian.vollmer@wwu.de,

<sup>2</sup>Max-Planck-Institut für Chemie, Hahn-Meitner-Weg 1, 55128 Mainz, Germany, <sup>3</sup>ETH Zürich, Institut für Geochemie und Petrologie, Clausiusstr. 25, Zürich, CH, <sup>4</sup>University of Alberta, Earth and Atmospheric Sciences, Edmonton, Canada, <sup>5</sup>SuperSTEM Laboratory, Keckwick Lane, Daresbury, UK.

**Introduction:** The provenance of the most pristine biomolecules, e.g., amino acids or nucleobases, on the early Earth is still a matter of debate [e.g., 1]. Starting from very simple interstellar CHON-bearing species such as CO<sub>2</sub>, H<sub>2</sub>CO (formaldehyde), HCN, or NH<sub>3</sub>, complex polymerization reactions can either occur in the interstellar medium or the solar nebula, e.g., by irradiation of coated dust grains, on cometary or asteroidal parent bodies by aqueous alteration, or by reactions on Earth in the atmosphere or at mineral interfaces in hydrothermal systems. Disentangling these complex modification processes is important to understand the evolution of the first pristine relevant biomolecules [see, e.g., 2, for a recent review].

**Samples and Methods:** Extraterrestrial samples that have not suffered severe thermal overprint such as petrologic type 2-3 carbonaceous chondrites or interplanetary dust particles (IDPs) are suitable candidates to search for pristine biomolecules. These samples contain on average several wt.% of organic C mainly in the form of kerogen-like molecules (“insoluble organic matter”, IOM) that can be isotopically anomalous, most notably in H or N. The origins of these anomalies are usually attributed to fractionation reactions in cold environments [e.g., 3], but recent ideas suggest that they might also be explained by high temperature solar nebula processes [4] or hydrothermal reactions on asteroidal bodies [5] in the case of nitrogen. The aim of this study is to locate organic matter in these samples “in-situ” by electron microscopy (SEM) and NanoSIMS techniques and to characterize the functional chemistry, specifically of nitrogen, and mineralogy by low voltage (60 kV) high resolution scanning transmission electron microscopy (STEM) combined with electron energy loss spectroscopy (EELS). This approach is less representative than bulk analyses on extracted organic matter performed since decades, but provides important information on unprocessed, single grains within their petrographic context.

**Results and Discussion:** Results on organic grains from CR chondrites Renazzo and Elephant Moraine (EET) 92161 and several IDPs from the L2006 collector confirm our previous findings that the polymerization of aromatic rings in the CR grains is increased compared to the IDP grains, probably due to subtle fluid-induced reactions on the CR parent body [6]. These polycyclic aromatic hydrocarbons are not directly relevant to early lifeforms, but could have served as scaffolds for the first primitive cell membranes [7]. Other functional groups observed by EELS at the C-K edge include several types of C-O bonding environments (ketones, aldehydes, carboxylic acids) that may act as precursors for  $\alpha$ -amino acids via Strecker synthesis reactions [8] or for complex sugars via formose reactions [9]. Carbonate bonding is prevalent in the CR as well as in the IDP samples, not always as discrete calcite grains, but dispersed within the organic matter, which underlines the importance of fluids for organic synthesis reactions. Furthermore, we could document strong nitrogen functionality (C-N nitrile/imine or imidazole and NH<sub>x</sub> amine bonding) in the CR grains, but so far not in the IDP organics. This may indicate the importance of fluids for advancing polymerization and stabilization reactions of N-compounds or the presence of functionally different primary nitrogen reservoirs sampled by cometary and asteroidal samples. Nitrogen functionality is of particular importance, as for example nitrile/imine and amine bonding environments are precursors of amino acids, whereas N-heterocycles such as imidazole are important constituents of nucleobases. Recent analyses by N-XANES on bulk samples of Tagish Lake organics have demonstrated as well that N-K edge functionality can be fitted by different amounts of amine and imine/nitrile or imidazole functional groups [10]. This supports recent ideas that aqueous complexation reactions starting from NH<sub>2</sub>-bonding environments were catalyzed on meteorites parent bodies to form biorelevant molecules such as certain amino acids and nucleobases [11].

*The DFG is acknowledged for funding this project in the course of the SPP 1833 “Building a Habitable Earth”.*

**References:** [1] Dalai P. et al. (2016), *Elements* 12, 401. [2] Alexander C. M. O'D. et al. (2017), *Chemie der Erde*, <https://doi.org/10.1016/j.chemer.2017.01.007>. [3] Busemann H. et al. (2006), *Science* 312, 727. [4] Kuga M. et al. (2015), *Proceedings of the National Academy of Sciences* 112, 7129. [5] Pizzarello S. and Bose M. (2015), *The Astrophysical Journal* 814, 107. [6] Vollmer C. et al. (2014), *Proceedings of the National Academy of Sciences* 111, 15338. [7] Ehrenfreund P. et al. (2006), *Astrobiology* 6, 490. [8] Simkus D. N. et al. (2016), *Meteoritical Society Meeting* (Berlin), abstr. #6215. [9] Cody G. D. et al. (2011), *Proceedings of the National Academy of Sciences* 108, 19171. [10] Cody G. D. and Alexander C. M. O'D. (2017), *Lunar and Planetary Science Conference XLVIII*, abstr. #2747. [11] Rotelli L. et al. (2016), *Nature Scientific Reports* 6, 38888.