Chondrite volatility trends revisited

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At the Lunar and Planetary Science Conference, we showed that there is complementarity between EH and CK chondrites for certain elements in bulk meteorite patterns [1,2]. In search for more elements showing similar behaviour, we plotted bulk chondrite data (Fig. 1) from [3]. Data are normalized to "volatile-free" CI abundances considering 20% H_2O and 3.5% C [4,1].

Fig. 1: Volatility trend for EH and CV/CK chondrites.



A general complementary "mirror-like" trend is observed for elements condensing above 1060 K (Au). CV and CK chondrites are enriched in refractory elements, whereas EH (and EL chondrites, not shown) are depleted. Between 1400 and 1300 K the patterns cross over with EH chondrites becoming enriched over CV/CK. This relative enrichment of EH continues with decreasing condensation temperature.

The temperature interval at which the element patterns cross over corresponds to the transition between refractory inclusion and chondrule formation. EH and EL contain <<1 vol.% refractory inclusions. Removal of a refractory component from the enstatite chondrite forming region has been suggested previously [e.g. 5]. Our observations indicate that this refractory component could have been transported into the CV/CK chondrite forming region before chondrite parent bodies accreted.

References

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Spatial and temporal (30 yr.) variations of lead geochemical signature in a macrotidal highly polluted estuary

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Two dated sediment cores (²¹⁰Pb) were sampled in 2005 on tidal flats of the Scheldt estuary at the vicinity of the heavily industrialized harbour city of Antwerp. Coring was close to the estuarine turbidity maximum and in the reactive 0-10 psu zone. The cores were compared with suspended particulate matter (SPM) and surface sediments (SS), sampled between 2002 and 2006 from the Scheldt river to the southern bay of the North Sea. Pb isotopic compositions measured by MC-ICP-MS were compiled regarding to grain-sized distributions as well as sedimentological, mineralogical and geochemical data. The present study aims at characterizing the sources and the evolution of Pb inputs during last 30 yr. in that complex environment.

As a whole, Pb isotopic data for SPM, SS and cores from the Antwerp area show a narrow range of variations. ²⁰⁶Pb/²⁰⁷Pb data vary of 1.9 ‰ (RSD) from the mean value of 1.1629 (with 1.1630 for SS and SPM, and 1.1628 for cores) suggesting that ~70% of particulate Pb has an anthropogenic origin. A close look at the Pb isotopic ratios highlights systematic variations reflecting different controlling mecanisms. In the SPM and SS samples, the larger Pb isotopic variations (from $1.1582 \pm 0,000010$ -2sd- to $1.1697 \pm$ 0,000017) are related to daily tidal cycles, where the relative contribution of fluvial and marine waters controls the isotopic signature. In the cores, the relative lower variation range (from 1.1608 ± 0.000011 to 1.1646 ± 0.000007) may result from a larger time-series integration and homogenization of the surface layers by bioturbation. Pb isotopic variations are correlated with grain-sized distributions, that are controlled by changes in deposition modes, mainly related to neap tide/spring tide cycles. In contrast, the decrease in Pb concentration, recorded from the 75-80's period to 2005, does not imply any significant isotopic answer. Therefore, Pb sources in the Scheldt estuary may remain unchanged, in component and relative contribution, over 30 yr. The water quality improvement, in terms of Pb contents, is not related to the removal of the leaded gasoline, but only to industrial emission control policies and establishment of water treatment plants.