

(1986), argued that the spherules were carbon-saturated at 1200–1225 °C and therefore that the silicate liquid must have contained graphite. However, in the Fe-C system the stable graphite liquidus is much steeper than the metastable cohenite liquidus, and although these alloys were cohenite-saturated, they were not graphite-saturated. Hence, the silicate magma probably did not contain graphite and carbon was not the dominant control on $f(\text{O}_2)$. Thus, it may be possible to reconcile the main constraints on ureilite petrogenesis without high pressures.

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U-Pb study of the Acapulco meteorite. Ch. Göpel, G. Manhès and C. J. Allègre. Lab. Géochimie et Cosmochimie, I.P.G., 4, Place Jussieu, 75252 Paris, Cedex 05, France.

The Acapulco meteorite is currently studied by a consortium in order to elucidate the origin and history of its parent body. This meteorite is characterized by an approximately chondritic bulk chemistry; its major element composition is close to that of average H-chondrites. Volatile elements and noble gases are however enriched nearly to the level of C1 chondrites, and U and Th show an even higher enrichment (Palme *et al.*, 1981). The texture of Acapulco is equigranular; chondrules are absent. Previous work with long-lived isotopes indicated some unexpected results considering the recrystallized texture and the estimated high equilibration temperature (1350 °K): a K/Ar age of 4.7 ± 0.3 Ga, an internal Sm-Nd isochron indicating a recrystallization age of 4.60 ± 0.03 Ga and recently an Ar/Ar age of 4.51 Ga (Palme *et al.*, 1981; Prinzhofer *et al.*, 1992).

We present U and Pb concentrations and Pb isotopic compositions measured in bulk samples and mineral separates from Acapulco.

Small bulk samples show slightly radiogenic Pb isotopic compositions ($^{206}\text{Pb}/^{204}\text{Pb} = 22\text{--}36$); similar to the majority of meteorites, they appear to have more radiogenic Pb than could have been derived from U-decay. The U concentrations range from 64 to 96 ppb. Compared to C1, Acapulco is enriched in U by a factor of 5–10. ^{204}Pb concentrations of the fragments (~6 ppb) are lower than C1 level (47 ppb); they lie in the range of H 4–6 chondrites (0.2 up to 7 ppb) (Tatsumoto *et al.*, 1976).

Three different mineral phases were separated and analyzed: plagioclase, troilite and phosphate. The phosphate separate shows a high U-concentration. Using the modal abundance of the minerals, phosphates account for half of the U in the bulk rock. The Acapulco phosphate contains more ^{204}Pb (17.1 ppb) than phosphates from L-, H- and LL-chondrites, and its resulting Pb isotopic composition is less radiogenic. The Pb/Pb model age of the phosphate relative to primordial lead is 4.557 ± 0.002 Ga.

The plagioclase is the main carrier of the ^{204}Pb in the bulk meteorite (80%). Different separates show low U concentrations; the Pb isotopic compositions are close to the primordial lead, but distinct.

The Pb isotopic composition of the troilite is intermediate; U and Pb concentrations are similar to those measured by Unruh in troilites from L-chondrites (Unruh, 1982).

TABLE 1. U-Pb data of some Acapulco samples.

	$^{206}\text{Pb}/$ ^{204}Pb	$^{207}\text{Pb}/$ ^{204}Pb	$^{208}\text{Pb}/$ ^{204}Pb	Pb (ppb)	U (ppb)	$^{238}\text{U}/$ ^{204}Pb
WR 28	22.801 ± 0.0019	18.688 ± 0.020	32.198 ± 0.043	507.8	89.6	11.385 ± 0.055
WR 37	26.220 ± 0.022	20.802 ± 0.022	32.627 ± 0.043	455.5	95.7	14.64 ± 0.21
Troi	13.00 ± 0.68	12.53 ± 0.41	31.98 ± 0.81	109.0	3.7	1.72 ± 0.25
Plag 28	9.345 ± 0.009	10.348 ± 0.013	29.411 ± 0.043	1912.0	0.01	0.0022 ± 0.0004
Phos	148.09 ± 0.18	96.42 ± 0.14	48.68 ± 0.08	5107.0	2759.7	137.0 ± 1.1

WR = small fragment, troi = troilite, plag = feldspar, phos = phosphate. All data are corrected for mass discrimination and analytical blank.

The U-Pb systematics clearly indicate that this meteorite has been affected by two distinct processes: the plagioclase data indicate an old perturbation or reequilibration process, whereas the isotopic composition of the troilite and the bulk fragments translates a recent addition of radiogenic lead. We will discuss these data and evaluate different models concerning the history of Acapulco.

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The continuing search for the location of ^{15}N -enriched nitrogen in Acfer 182. M. M. Grady,¹ C. T. Pillinger² and J. W. Arden.³ ¹Dept. of Mineralogy, Natural History Museum, Cromwell Road, London SW7 5BD, UK. ²Dept. of Earth Sciences, The Open University, Walton Hall, Milton Keynes MK7 6AA, UK. ³Dept. of Earth Sciences, Univ. of Oxford, Park Road, Oxford, UK.

Acfer 182 is an unusual chondrite, with abundant small chondrules and CAIs (mean diameter *ca.* 100 μm), and rich in metal (*ca.* 15 vol%). It is closely related to ALH 85085, and, like that meteorite, is highly enriched in ^{15}N (bulk $\delta^{15}\text{N}$ *ca.* +600‰; $\delta^{15}\text{N}_{\text{max}} = +1584\%$ at 900 °C; Grady and Pillinger, 1992). Stepped combustion of Acfer 182 (see Fig. 1) releases ^{15}N over a wide temperature range, indicating that its carriers must be dispersed throughout the meteorite, possibly occurring in carbonaceous material, fine-grained matrix, clasts and metal. The highest relative abundance of ^{15}N is found in phase “N_c,” so far unidentified mineralogically, with a C/N *ca.* 10, which releases its nitrogen on combustion of the whole-rock at 850–950 °C. N_c is more apparent in Acfer 182 than ALH 85085, accounting for *ca.* 8 ppm of the total nitrogen inventory of 85.4 ppm.

An attempt to isolate N_c by physical means proved unsuccessful (Grady and Pillinger, 1992); therefore, chemical treatments were tried: an HF/HCl-resistant residue was prepared from 9 g of fragments. Examination of the remaining material confirmed that it was dominantly composed of Mg-Al spinels, chromite, hibonite and Cr-rich sulphides. Approximately two thirds of the original amount of nitrogen in the sample has been lost on dissolution (see Fig. 1), including any associated with Fe-Ni metal. There has been a reduction of over 50% of the nitrogen which was released up to 500 °C and presumed present in a carbonaceous component, without significant change in $\delta^{15}\text{N}$ value or C/N ratio. The most visible difference between results from the whole-rock and HF/HCl-resistant residue is that the combustion temperature of N_c has decreased to 550–700 °C, with a concomitant drop in $\delta^{15}\text{N}$ from +1584‰ to +1274‰. It is unlikely that a minor (even heavier) sub-fraction of the ^{15}N -rich material has been removed; now that N_c combusts at a temperature closer to the more abundant “organic” nitrogen, it is probable that mixing lowers the observed $\delta^{15}\text{N}$. N_c itself does not seem to

