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Distribution of heavy p-process isotopes in extraterrestrial materials: Reconciling uniform 174 Hf with 180 W heterogeneities

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Iron meteorites carry excesses of p-process ¹⁸⁰W with respect to terrestrial materials, which correlate with the accretion ages of the parent bodies (Schulz et al., 2010, 2011). The favoured explanation for these excesses is nucleosynthetic heterogeneity in the early solar system, in which case also other heavy p-process isotopes are expected to show anomalies in extraterrestrial materials. We therefore investigated the relative abundance of ¹⁷⁴Hf, the first lighter stable p-process nuclide next to ¹⁸⁰W, in silicate materials of different ages and from different regions within the inner solar system. All samples were measured with high-precision MC-ICPMS in setup with standard sample and X-skimmer cone, typically consuming ~ 60 ng Hf at precisions of ± 70 ppm (2σ). In order to successfully correct for interference by isobaric ¹⁷⁴Yb, we optimised the existing hafnium purification protocol by Münker et al. (2001). Mass bias corrections were made based on different hafnium isotope ratios in order to monitor unlikely s- and r-process heterogeneities and deviations caused by secondary neutron capture reactions. It is found that relative abundances of ¹⁷⁴Hf, as well as of other non-radiogenic hafnium isotopes, for enstatite chondrites, H and L ordinary chondrites, one CV chondrite, eucrites, and one lodranite sample are indistinguishable from terrestrial materials. A silicate inclusion of the El Taco IAB iron meteorite and one EL6 chondrite (Pillistfer) exhibit higher 174 Hf/ 177 Hf ratios, namely 190 \pm 72 ppm and 210 \pm 75 ppm respectively. These ratios correlate with higher ¹⁷⁸Hf/¹⁷⁷Hf ratios (18±5 and 21±5 ppm, respectively) and are therefore likely to reflect secondary neutron capture on ¹⁷⁷Hf that is related to the high iron content of these particular materials (Sprung et al., 2010). Together, the presented sample suite indicates a uniform distribution of ¹⁷⁴Hf in the solar system. Two possible scenarios can reconcile this conclusion with a nucleosynthetic origin of ¹⁸⁰W excesses, namely (1) the original carrier phases of ¹⁷⁴Hf and ¹⁸⁰W were different; or (2) existing ¹⁷⁴Hf heterogeneities were homogenised by the time of parent body formation of the analysed silicates. An alternative explanation would be that ¹⁸⁰W excesses are not of nucleosynthetic origin and result from a different process that did not affect ¹⁷⁴Hf abundances. A possible process is secondary neutron capture, but we did not identify plausible pathways for the production of ¹⁸⁰W. We therefore prefer the two scenarios of nucleosynthetic origin of the anomalies, and aim distinguishing between them by determination of ¹⁷⁴Hf in older sample materials than those analysed so far.