

## High-precision Tungsten isotope analyses by multicollection N-TIMS

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The lithophile-siderophile  $^{182}\text{Hf}$ - $^{182}\text{W}$  decay pair is well-suited for constraining the timing of planetary core-formation events by coupling the isotopic growth of  $^{182}\text{W}$  resulting from the decay of  $^{182}\text{Hf}$  ( $T_{1/2} = 8.9$  Myr) and elemental Hf/W ratios. In addition, the presence of late-accretion extraterrestrial and/or core-derived materials can be detected through high-precision analyses down to the ppm-level [1]. High-precision Tungsten isotope analyses by MC-ICP-MS [2] are now challenged by N-TIMS analyses that yield a long-term reproducibility of 5 ppm on  $^{182}\text{W}/^{184}\text{W}$  [1]. However, a residual correlation has been observed between mass fractionation corrected  $^{182}\text{W}/^{184}\text{W}$  and  $^{183}\text{W}/^{184}\text{W}$  that has been attributed to mass dependent variability of O isotopes during analysis and from run to run, thus causing some inaccuracy in W and Re oxide correction.

The present study on a Thermo Scientific TRITON *Plus* is aimed at investigating the residual mass bias correlation further by monitoring  $^{18}\text{O}/^{16}\text{O}$  during W isotopic analysis so as to provide an in-run correction of W and Re oxides. Sample loads of 3  $\mu\text{g}$  were run for > 820 cycles of 8 s integration at average  $^{182}\text{W}^{16}\text{O}_3$  signals of 0.6-3.5V in single static mode, using  $10^{11}$   $\Omega$  amplifiers on W and Re oxide beams, with rotation of the amplifier-cup association (to average out amplifier gain biases).  $^{186}\text{W}^{18}\text{O}^{16}\text{O}_2$  beams of 4-10mV were measured with  $10^{13}$   $\Omega$  or  $10^{12}$   $\Omega$  amplifier for highly precise and accurate  $^{18}\text{O}/^{16}\text{O}$  ratios determination. Data are corrected for W and Re oxide interferences by using the measured  $^{18}\text{O}/^{16}\text{O}$  ratios, and for instrumental mass fractionation, by normalizing to  $^{186}\text{W}/^{184}\text{W}$  or  $^{186}\text{W}/^{183}\text{W}$ . The internal precision on  $^{182}\text{W}/^{184}\text{W}$  and  $^{183}\text{W}/^{184}\text{W}$  normalized to  $^{186}\text{W}/^{184}\text{W}$  is 3-8ppm (2RSE). External reproducibility is 17ppm and 12 ppm (2RSD). The internal precision on  $^{182}\text{W}/^{183}\text{W}$  and  $^{184}\text{W}/^{183}\text{W}$  normalized to  $^{186}\text{W}/^{183}\text{W}$  is 2-8ppm (2RSE). External reproducibility is 15ppm and 8 ppm (2RSD). The preliminary data do not seem affected by secondary residual mass bias and have not been doubly normalized [1]. Further analyses will allow assessment of how determination of  $^{18}\text{O}/^{16}\text{O}$  in samples will allow for improvement of long-term reproducibility.

[1] Touboul and Walker (2012) *Int.J.Mass Spectrom.* **309**, 109-117 [2] Holst *et al* (2013) *PNAS* **110**, 8819-8823