

## Development of high-precision La isotope analysis by MC-ICP-MS

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The long-lived radionuclide  $^{138}\text{La}$  is of exclusive supernova origin and therefore can be used as a tracer of supernova-derived material in the early Solar System. However, the very low isotope abundance of  $^{138}\text{La}$  (0.09%) makes its precise measurement difficult. Shen and Lee (2003) [1] successfully detected La isotope variation among CAIs using TIMS. Yet, the analytical precision in the pioneering work ( $2\sigma = 8\text{--}16\epsilon$ ) is not high enough for investigating potential isotope variations among whole-rock meteorites. In this study, we have developed a method for precise La isotope analysis using MC-ICP-MS. This has been achieved by (1) using  $10^{13}$  and  $10^{10}$   $\Omega$  amplifiers on 138 and 139 amu, (2) correcting for instrumental mass bias by the Nd external calibration technique, and (3) developing a novel chemical separation scheme for effective removal of isobaric elements, Ba and Ce.

The precision of La isotope measurements was tested by analyzing a La standard solution NIST 3127a. The use of  $10^{13}$  and  $10^{10}$   $\Omega$  amplifiers improved the precision by a factor of two compared with using  $10^{11}$   $\Omega$  amplifier. The external calibration technique improved the precision by an order of magnitude compared with the standard bracketing method. As a result of these improvements, an internal precision of  $\pm 1.4\epsilon$  (2SE) was achieved. We also found that  $^{138}\text{Ba}$  and  $^{138}\text{Ce}$  isobaric interferences can be accurately corrected when Ba/La and Ce/La are as low as  $2 \times 10^{-3}$  and  $2 \times 10^{-4}$ , respectively.

Our La chemical separation technique consists of 3-step column chromatographic procedures. In the first step, matrix elements and Ba were separated from rare earth elements (REEs) by using AG 50W-X8 (100–200 mesh) cation exchange resin with elution using HCl and  $\text{HNO}_3$  successively. In the second step, REEs were further purified from Ba with TODGA resin (50–100  $\mu\text{m}$ ) by direct loading from the former column. In the last step, La was separated from other REEs using Ln resin (25–50  $\mu\text{m}$ ) by eluting with 0.15 M HCl. For 100 mg JB-1b basaltic reference material, the total recovery of La was >97% while Ba/La, Ce/La, Nd/La were  $< 3 \times 10^{-4}$ ,  $< 5 \times 10^{-5}$  and  $< 2 \times 10^{-4}$ , respectively.

[1] Shen and Lee (2003), *ApJL* **596**, L109–L112.