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## **Doubling sensitivity in MC-ICP-MS using high-efficiency, rapid-response laser ablation technology.**

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Laser ablation multicollector inductively coupled mass spectrometry (LA-MC-ICP-MS) is one of the most sensitive mass spectrometry techniques. Detection efficiencies (number of ions detected/number of atoms sampled) of 2 % for Pb and 2.8 % for U [1] have been reported for the Thermo Scientific™ NEPTUNE *Plus* MC-ICP-MS and are comparable to those typically reported for solution introduction using a desolvating nebulizer system.

Recent developments have seen the introduction of rapid-response laser ablation cells and sample transport technologies, enabling signal pulse durations for a single laser ablation shot of less than 10 ms [2]. This has resulted in marked improvements in analytical throughput, resolution and sensitivity [3,4] vital for the generation of large, highly spatially-resolved elemental maps. The focus on mapping, particularly bioimaging, in their development has until now [5] mostly obscured the possibility of using the sensitivity advantage of rapid-response technologies for other LA-ICP-MS applications.

Here we report the coupling of the NEPTUNE *Plus* to a Teledyne CETAC™ Analyte G2 equipped with just such a rapid-response sample transport system, the Aerosol Rapid Introduction System (ARIS™), to achieve higher detection efficiencies than previously reported for LA-MC-ICP-MS. Compared to a conventional two-volume configuration we achieved a twofold increase in sensitivity across the entire mass range (Li – U). The increase in sensitivity was utilised to improve a number of example applications – Pb isotope ratio analysis, small spot (20 µm) Hf analysis of zircons and Rb/Sr in carbonates.

[1] Schaltegger (2015) *Chem. Geo.*, **402**, 89-110 [2] van Malderen (2016) *J. Anal. At. Spectrom.*, **31**, 423-439. [3] Douglas (2015) *Anal. Chem.*, **87**, 11285–11294. [4] van Malderen (2018) *Spectrochim. Acta - Part B At. Spectrosc.*, **140**, 29-34. [5] Petrus (2017) *Chem. Geo.*, **463**, 76-93.