

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

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**ABSTRACT:** The introduction of rapid-response laser ablation cells and sample transport technologies to laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS) has enabled signal pulse durations for a single laser ablation shot of less than 10 ms. These developments have resulted in marked improvements in analytical throughput, resolution and sensitivity vital for the generation of large, highly spatially-resolved elemental maps. The focus on mapping, particularly bioimaging, has obscured the possibility of applying the sensitivity advantage of rapid-response technologies to other LA-ICP-MS applications, such as high precision isotope ratio analysis on multicollector (MC-)ICP-MS. In this work a commercially available rapid-response sample transport system and a conventional configuration were compared for LA-MC-ICP-MS analysis. Ablation of known reference materials demonstrated 'sensitivity' or sample ion yield of 7-9% using the rapid-response sample transport system, more than double that for the conventional set-up. This increase in efficiency was demonstrated to improve precision for the Pb isotope ratio analysis of the MPI-DING reference glasses and improve the spatial resolution of Hf isotope ratio analysis of reference zircons.

The achievable precision during mass spectrometry is a function of the amount of material sampled and the detection efficiency of the combined sample introduction system and mass spectrometer. Without limitation from the amount of sample, high precision can be easily achieved, even with a system featuring low detection efficiency, simply by measuring the analyte for a prolonged period of time. For some challenging applications, and sample introduction techniques such as laser ablation, however, the amount of sample material which is available is fundamentally limited. Here using a high detection efficiency method enables smaller amounts of sample to be used whilst still maximizing precision. In geoscience using laser ablation as the sample introduction technique for an inductively coupled plasma mass spectrometer (LA-ICP-MS), many commonly analyzed isotopes can be rapidly measured in situ within containing minerals. To confidently interpret changes in zonation requires constraining the area or volume ablated. Therefore to maintain the analytical precision required under these spatial constraints requires the highest efficiency.

Recent years have seen the development<sup>1-3</sup>, and subsequent commercialization (Teledyne CETAC™ ARIS™ Aerosol Rapid Introduction System, ESI NWR™ Bloodhound™), of rapid response laser ablation cells and sample transport technologies. Their designs are aimed towards minimizing aerosol dispersion in the cell and substantially reducing the washout time<sup>4</sup>. Upon leaving the cell the low dispersion aerosol must be transported to the ICP-MS. As aerosol dispersion also occurs within the transport tubing and injector of the ICP torch, rapid response laser ablation cell designs have also incorporated changes to these elements of the LA-ICP-MS configuration<sup>1-4</sup>. Reductions

in transport tubing diameter and elimination of injector dead volumes have been seen to minimize aerosol dispersion<sup>3,4</sup>. However reducing the diameter of the transport tubing decreases the mass flux outflow from the cell. If the ablated particle density exceeds what the transport tubing can efficiently extract then the excess material can either be lost (*e.g.*, gravitational sedimentation or inertial deposition), reducing efficiency, or is only released after a significant delay, reducing speed and throughput efficiency or productivity<sup>5</sup>.

The key driver in these developments has been to improve the utility of LA-ICP-MS for sample mapping, primarily for determining the distribution and concentration of drugs or tagged biomarkers in tissue sections or single cells. For ablation cell designs with washout  $\approx 500$  ms, high resolution sample mapping requires using a slow laser pulse repetition rate in order to restrict signal overlap between adjacent image pixels: resulting in long analysis times. In practice high repetition rates have also been used, sacrificing pixel resolution in favor of time. The prototype designs on which the current commercially available rapid response ablation cells were based both reported sub-5 ms washout<sup>2,3</sup>. As such completely resolved pixels could be collected at rates  $>150$  Hz, greatly reducing the analysis time (commercial designs 40 Hz<sup>6</sup>). The corresponding improvements in signal-to-noise ratio, lowering the limit of detection, allows the use of smaller spot sizes further improving resolution. 5  $\mu\text{m}$  resolution mapping of the binding of platinum-based anti-cancer compounds to both cochlea<sup>7</sup> and multi-cellular tumor spheroids<sup>8</sup> have recently been reported. Van Malderen et al. (2017) produced a fully resolved trace elemental distribution map of a crustacean at 2  $\mu\text{m}$  resolution<sup>9</sup>. Although rapid response cells

were designed for bio-imaging applications, they also have applicability to the more established LA-ICP-MS applications of geochemistry and isotope ratio analysis. Petrus et al. (2017) produced large, 5.5 x 5.0 mm, trace elemental distribution maps of serpentinized peridotite and augite in less than 2 hours<sup>10</sup>.

To be best utilized, rapid response laser ablation cells and transport technologies demand an ICP-MS with a short duty cycle. For trace elemental analysis this condition can be met by a fast scanning ICP-Q-MS or ICP-TOF-MS, but for isotope ratio analysis there is another option. Multi-Collector (MC-)ICP-MS has become a well-established technique for high precision isotope ratio measurements, especially in the field of geochemistry. Here the issues of spectral skew observed for rapid response ablation cells on scanning mass spectrometers<sup>6</sup> should be eliminated as every isotope is collected simultaneously. It should be noted this is only true if the same type of detector is used for each isotope as a form of spectral skew has been reported for MC-ICP-MS with a mixed SEM/Faraday cup detector array<sup>11</sup>. LA-MC-ICP-MS is routinely used to measure a variety of isotopic systems, including Hf<sup>12,13</sup> and U/Pb<sup>14-16</sup> in zircon geochronology, at a variety of precision levels. Developments such as the Jet Interface and 10<sup>13</sup> Ω amplifier technology<sup>17,18</sup> for the Thermo Scientific™ NEPTUNE Plus™ and enhanced sensitivity HR-ES<sup>19</sup> for the Nu Instruments™ Nu Plasma™ MC-ICP-MS have already delivered significant improvements in signal-to-noise ratio, and hence LA-MC-ICP-MS spatial resolution. Rapid response laser ablation cells have already been reported to deliver significant improvements in signal-to-noise for LA-ICP-MS and here we report on the application of one of these designs to LA-MC-ICP-MS.

**Sample Ion Yield.** ICP-MS instrument sensitivity is typically reported as the number of ions detected against the concentration of the measured isotope in the sample (cps/ppm for Q-ICP-MS or V/ppm for MC-ICP-MS). However, reporting sensitivity in this manner is not necessarily the best unit of comparison as it ignores the extent to which the signal intensity is dependent on the rate at which the sample is introduced<sup>20</sup>. Nominally, an analyte in a solution introduced at a flow rate of 200 µL/min would yield a peak signal and apparent sensitivity (in cps/ppm) twice that using a solution introduced at 100 µL/min, making comparison between different ICP-MS configurations difficult if flow rate isn't taken into account. For LA-ICP-MS, sample introduction rate is dependent on a complex series of factors (including fluence, spot size, and repetition rate and sample matrix). Another reporting method is therefore clearly required, ideally one appropriate for both solution and laser ablation sample introduction.

The sample ion yield (also referred to elsewhere as 'useful yield' and 'detection efficiency'), the ratio of the number of ions detected against the number of atoms sampled, is a method for reporting sensitivity commonly used in TIMS and SIMS mass spectrometry. A significant advantage of reporting the sample ion yield is that it is independent of the rate at which the sample is introduced; nominally the ratio of ions detected to atoms sampled should remain the same. The number of atoms can be calculated for a measured volume of material provided the elemental concentration and isotopic composition of the material are known. The mass response curve of the mass spectrometer (as well as other factors) however, determines that the sample ion yield is dependent on the isotope that is measured. The highest sample ion yields are usually reported for the highest mass elements, such as Pb and U (in MC-ICP-MS and SIMS).

LA-MC-ICP-MS is already amongst the more sensitive mass spectrometry techniques. Schaltegger et al. (2015) reported sample ion yields of 2 % and 2.8 % for Pb and U respectively<sup>14</sup>. A complementary technique, SIMS, when optimized for transmission, can achieve similar sample ion yields for these elements<sup>21</sup>. The sample ion yield reported for TIMS is more dependent on the element ionization potential (and the filament loading technique) than ICP-MS. U & Pb sample ion yields can be significantly higher than for MC-ICP-MS (≈10 %) <sup>22</sup>, but are usually broadly equivalent<sup>23</sup>.

The only previous application of a rapid response ablation cell to a MC-ICP-MS, focused on a single application and with an experimental prototype cell, suggested LA-MC-ICP-MS may be capable of significantly greater sample ion yields<sup>11</sup>. Here we report the first coupling of a MC-ICP-MS (NEPTUNE Plus) to a commercially available (ARIS) rapid response sample transport technology. The impact on instrument sensitivity (as sample ion yield) was quantified for a suite of elements (Li, Sr, Nd, Hf, Pb and U) across the total mass range and compared to that provided by a more conventional laser ablation cell. Subsequently the configuration was tested for Hf and Pb isotope ratio analysis for a series of well characterized reference materials.

## EXPERIMENTAL SECTION

All MC-ICP-MS measurements were carried out on the NEPTUNE Plus at Thermo Fisher Scientific (Bremen) GmbH. In order to maximize instrument ion yield across the mass range the Jet Interface option was used with the X skimmer and Jet sampler cone. N<sub>2</sub> addition was also used. The laser ablation system used was a Teledyne Photon Machines™ Analyte G2™.

In the high efficiency, rapid response ablation cell configuration, the inner cell of the HelEx™ ablation cell was coupled directly to the MC-ICP-MS torch injector via the ARIS. The transfer line consisted of 1.5 m of PEEK tubing, 1 mm internal diameter. Ar and N<sub>2</sub> were added via the ARIS adapter.

In the conventional configuration the output of the inner cell was not connected directly to the ICP torch injector, but via a fast washout switch (Teledyne Photon Machines™ CleanShot™) by 2 m of 4 mm outer diameter, 2 mm internal diameter, Teflon® tubing. The single valve within the CleanShot was thought to have minimal impact on the transport efficiency and allowed more efficient extraction of atmospheric gases from the transport system. The outlet of the fast washout switch was connected to the ICP injector by 1.5 m of ¼" outer diameter Teflon coated Tygon™ tubing. Ar and N<sub>2</sub> make-up gasses were added via two nylon y-pieces.

The methodology applied to tuning a LA-MC-ICP-MS is dependent on the isotope ratios being measured. For experiments which focus on intra-element isotope ratios, such as those for Pb or U, issues of laser-induced elemental fractionation or oxide formation rates are not necessarily principle factors during instrument tuning. In these instances the system can be tuned for maximum 'sensitivity' and these therefore represent the best achievable sample ion yield. However for Hf isotopic analysis, the required corrections for Lu and Yb interferences are negatively impacted at high oxide conditions due to the different oxide formation rates of each element. Furthermore potential Gd and Dy oxide interferences are more likely to occur in high oxide conditions<sup>24</sup>. Nd isotopic analysis is similarly affected by the required <sup>144</sup>Sm correction on both the determined <sup>143</sup>Nd/<sup>144</sup>Nd and <sup>146</sup>Nd/<sup>144</sup>Nd used for mass bias correction<sup>25</sup>. For

these isotope systems the LA-MC-ICP-MS needs to be optimized to minimize laser-induced elemental fractionation and rates of oxide formation at the expense of maximum sample ion yield. The sample ion yield was therefore determined for the two possible states, optimized and maximum U ion yield, with

both the conventional and high efficiency configurations (Table 1).

		Conventional Optimized	Conventional Maximum Ion yield	High Efficiency Optimized	High Efficiency Maximum Ion yield
Thermo Scientific NEPTUNE Plus MC-ICP-MS					
Cool Gas	L min <sup>-1</sup>	16	16	16	16
Aux Gas	L min <sup>-1</sup>	0.75	0.90	0.75	0.95
Sample Gas	L min <sup>-1</sup>	0.72	0.90	0.95	1.085
Z	mm	-1.95	-1.45	-1.95	-1.95
Power	W	1200	1300	1300	1300
Integration Time	ms	131	131	131	131
Photon Machines Analyte G2 193 nm excimer laser ablation system					
He – Outer Cell	L min <sup>-1</sup>	0.50	0.60	0.425	0.20
He – Cup Flow	L min <sup>-1</sup>	0.30	0.35	0.15	0.15
N <sub>2</sub> Addition	mL min <sup>-1</sup>	9.0	9.0	8.5	9.5
UO/U	%	0.107	0.930	0.092	
<sup>238</sup> U/ <sup>232</sup> Th		1.131	1.794	1.088	

**Table 1.** LA-MC-ICP-MS parameters on the NEPTUNE Plus MC-ICP-MS and Analyte G2 laser ablation system. Tuning was performed for a line raster of NIST<sup>TM</sup> SRM@610, 4 J cm<sup>-2</sup> fluence, 7 Hz repetition rate, 35 μm spot size. Three isotopes <sup>232</sup>Th, <sup>238</sup>U and <sup>238</sup>UO were monitored simultaneously. The optimized conditions (Conventional and High Efficiency) were tuned to minimize fractionation between <sup>232</sup>Th and <sup>238</sup>U and reduce <sup>238</sup>UO/<sup>238</sup>U as far as possible. The maximum ion yield conditions were tuned to maximize the response for <sup>238</sup>U.

The sample ion yield was measured for six isotopic systems (Li, Sr, Nd, Hf, Pb and U) using the cup configurations provided in Table 2. Four replicates of fifty shots were fired at a rate of 1 Hz, fluence 4 J cm<sup>-2</sup>, into a mounted piece of SRM@610 glass from NIST. Five different spot sizes ranging from 10 μm to 110 μm were measured. As the glass density and concentration of the six elements measured is known for SRM610, it was possible to calculate the number of atoms introduced for each element from the volume of the crater ablated. The volume of each crater was measured using a 3D optical microscope (InfiniteFocus<sup>TM</sup>, Aliconia, Sevenoaks, UK).

#### **Pb Isotopic Analysis of MPI-DING Reference Materials.**

The determination of the isotope ratios of lead (<sup>208</sup>Pb/<sup>206</sup>Pb, <sup>207</sup>Pb/<sup>206</sup>Pb) was used to evaluate the performance of the high efficiency rapid response cell when tuned for maximum sample ion yield. Ten spots were each ablated (3 J cm<sup>-2</sup> fluence, 40 μm circular spot size, 16 Hz repetition rate) in three of the MPI-DING series of reference glasses (ATHO-G, StHs6/80-G, GOR132-G). For each analysis a further ten spots on the NIST SRM@612 glass were used as the primary reference analyses.

A high repetition rate is required with rapid response ablation cells in order to ensure a stable signal profile<sup>7,10</sup>. Measuring on the central SEM of the MC-ICP-MS the washout time of the high efficiency configuration was estimated to be less than

100 ms (50 ms, full peak width at 10 % maximum), requiring 20 Hz or greater repetition rate to achieve a signal profile which did not resolve the individual pulse tops. However, analogue Faraday detectors are principally used in MC-ICP-MS which introduce a significant degree of smoothing (dependent on the resistance of the associated amplifier) to the signal profile when compared to an SEM. A detector array equipped only with Faraday cup detectors can achieve a stable signal profile for repetition rates less than 10 Hz, due to the signal smoothing from the associated amplifier.

<sup>206</sup>Pb and <sup>207</sup>Pb were measured on Faraday cup detectors equipped with 10<sup>13</sup> Ω amplifiers, all other isotopes were measured on 10<sup>11</sup> Ω amplifiers. Tau correction of the 10<sup>13</sup> Ω amplifiers was performed within the Neptune/Triton v3.3.0 software. All data were processed within Iolite<sup>TM</sup> v3.63 using an in-house data reduction scheme.

#### **Hf isotope ratio and U/Pb geochronology of reference zircons.**

The determination of <sup>176</sup>Hf/<sup>177</sup>Hf in two reference zircons, 91500 and Plešovice, was used to evaluate the utility of the high efficiency rapid response cell for analyses which need to be optimized for low oxide formation. The laser ablation parameters (Table 3) followed the small-volume method reported in Bauer & Horstwood (2017)<sup>13</sup>. For each analysis twenty spots on Plešovice and twenty spots on the reference material 91500 zircon were ablated.

Cup Configuration	L4	L3	L2	L1	C	H1	H2	H3	H4
Li					<sup>7</sup> Li				
Sr		<sup>82</sup> Kr	<sup>83</sup> Kr	<sup>84</sup> Sr	<sup>85</sup> Rb	<sup>86</sup> Sr	<sup>87</sup> Sr	<sup>88</sup> Sr	
Nd		<sup>142</sup> Nd	<sup>143</sup> Nd	<sup>144</sup> Nd	<sup>145</sup> Nd	<sup>146</sup> Nd	<sup>147</sup> Sm	<sup>148</sup> Nd	
Hf		<sup>171</sup> Yb	<sup>173</sup> Yb	<sup>175</sup> Lu	<sup>176</sup> Hf	<sup>177</sup> Hf	<sup>178</sup> Hf	<sup>179</sup> Hf	
Pb		<sup>202</sup> Hg	<sup>203</sup> Tl	<sup>204</sup> Pb	<sup>205</sup> Tl	<sup>206</sup> Pb	<sup>207</sup> Pb	<sup>208</sup> Pb	
U			<sup>234</sup> U	<sup>235</sup> U	<sup>236</sup> U		<sup>238</sup> U		
U/Pb	<sup>206</sup> Pb	<sup>207</sup> Pb	<sup>208</sup> Pb				<sup>232</sup> Th	<sup>235</sup> U	<sup>238</sup> U

**Table 2.** Table of MC-ICP-MS cup configurations used. <sup>206</sup>Pb and <sup>207</sup>Pb were measured on 10<sup>13</sup> Ω amplifiers in both the Pb and U/Pb configurations. All other isotopes were measured with 10<sup>11</sup> Ω amplifiers.

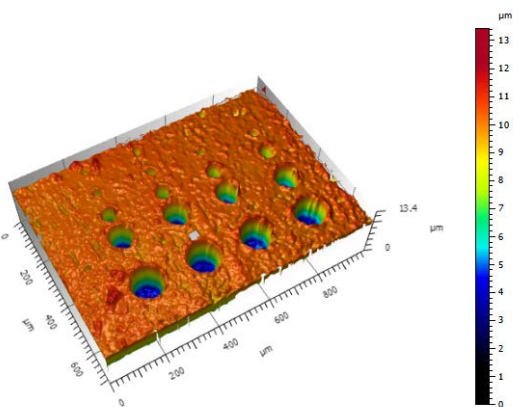
Further spots were used to collect U/Pb for the same two zircons, but with a lower laser energy and duration than for Hf. All data were processed within Iolite v3.63 using an in-house Hf data reduction scheme and the supplied U-Pb\_Geochronology\_4 data reduction scheme. 1 s was cropped from the beginning and end of each analysis.

**Table 3.** Laser ablation parameters for Hf and U/Pb isotope analysis of 91500 and Plešovice zircons.

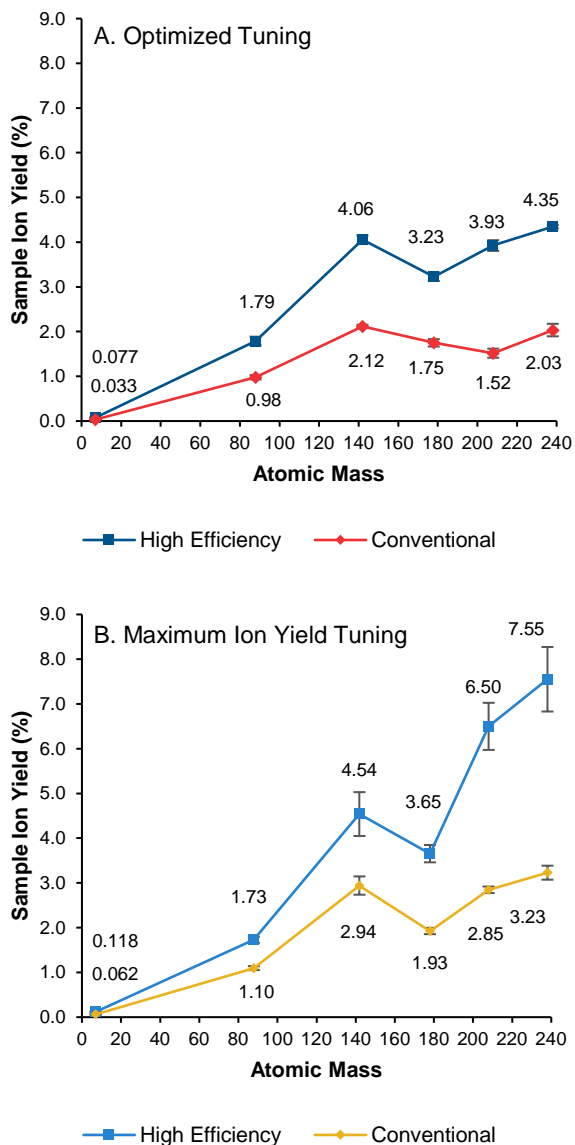
Parameter	Hf		U/Pb
Fluence	6.5 J cm <sup>-2</sup>	6.5 J cm <sup>-2</sup>	3 J cm <sup>-2</sup>
Spot Size	25 μm circle	20 μm circle	20 μm circle
Repetition Rate	10 Hz	10 Hz	10 Hz
Duration	10 s	10 s	5 s

## RESULTS & DISCUSSION

**Sample Ion Yield.** The sample ion yield of the high efficiency configuration was found to be dependent on the spot size ablated (Table 4). The sample ion yield of the conventional configuration was the same regardless of spot size (Table 5). This effect has been observed before with high efficiency laser ablation designs: at the largest spot sizes the transient ablated particle density is greater than the mass flux outflow of the cell into the transport tubing of the high efficiency design and thus not all of the material can be efficiently extracted<sup>5,11</sup>.



**Figure 2.** False color 3D image, created with an InfiniteFocus 3D microscope, of spot craters in SRM610 glass.



**Figure 1.** Calculated sample ion yield (%) on SRM610 v. atomic mass for the high efficiency and the conventional LA-MC-ICP-MS configurations. 50 μm circle, 4 J cm<sup>-2</sup> fluence, 1 Hz, 50 shots. A. LA-MC-ICP-MS optimally tuned for low oxide formation. B. LA-MC-ICP-MS tuned for maximum ion yield. Error bars are 1SD, n = 4.

**Table 4. Average sample ion yield for six elements based on 50 shots of SRM610, fluence 4 J cm<sup>-2</sup>, 1 Hz repetition rate, tuned for optimized conditions. Uncertainties reported are 1SD, n = 4.**

Spot Size		U	Pb	Hf	Nd	Sr	Li
110 μm	High Efficiency	4.00 ± 0.17	3.65 ± 0.08	1.71 ± 0.07	2.59 ± 0.04	1.08 ± 0.02	0.054 ± 0.001
	Conventional	2.03 ± 0.07	1.48 ± 0.02	1.57 ± 0.10	2.10 ± 0.03	0.92 ± 0.05	0.034 ± 0.000
	% Improvement	97	146	9	23	16	61
85 μm	High Efficiency	4.14 ± 0.18	3.65 ± 0.09	1.97 ± 0.08	2.90 ± 0.13	1.26 ± 0.07	0.057 ± 0.001
	Conventional	2.27 ± 0.04	1.59 ± 0.09	1.82 ± 0.13	2.33 ± 0.08	1.03 ± 0.01	0.037 ± 0.001
	% Improvement	83	129	8	24	23	55
50 μm	High Efficiency	4.35 ± 0.04	3.93 ± 0.12	3.23 ± 0.10	4.06 ± 0.09	1.79 ± 0.10	0.077 ± 0.007
	Conventional	2.03 ± 0.14	1.52 ± 0.10	1.75 ± 0.08	2.12 ± 0.04	0.98 ± 0.05	0.033 ± 0.001
	% Improvement	114	159	85	92	83	131
20 μm	High Efficiency	6.37 ± 1.28	6.82 ± 1.37	3.94 ± 0.27	4.14 ± 0.30	1.78 ± 0.18	0.073 ± 0.010
	Conventional	2.05 ± 0.15	1.46 ± 0.21	1.42 ± 0.10	1.72 ± 0.29	1.08 ± 0.35	0.029 ± 0.002
	% Improvement	210	366	177	140	65	150
10 μm	High Efficiency	5.43 ± 2.58	7.24 ± 2.45	3.86 ± 0.23	5.78 ± 3.50	2.47 ± 0.80	0.064 ± 0.018
	Conventional	1.82 ± 0.42	1.00 ± 0.07	1.07 ± 0.19	1.08 ± 0.48	0.80 ± 0.24	0.022 ± 0.003
	% Improvement	263	626	262	436	208	194

**Table 5. Average sample ion yield for six elements based on 50 shots of SRM610, fluence 4 J cm<sup>-2</sup>, 1 Hz repetition rate, tuned for maximum sample ion yield. Uncertainties reported are 1SD, n = 4.**

Spot Size		U	Pb	Hf	Nd	Sr	Li
110 μm	High Efficiency	5.11 ± 0.22	4.38 ± 0.11	1.21 ± 0.03	2.46 ± 0.11	0.80 ± 0.03	0.084 ± 0.003
	Conventional	3.52 ± 0.11	2.88 ± 0.06	1.45 ± 0.05	2.27 ± 0.05	1.05 ± 0.04	0.063 ± 0.003
	% Improvement	45	52	-17	8	-24	33
85 μm	High Efficiency	5.91 ± 0.30	5.11 ± 0.16	1.73 ± 0.04	2.95 ± 0.15	0.99 ± 0.03	0.089 ± 0.005
	Conventional	3.39 ± 0.13	2.99 ± 0.15	1.68 ± 0.05	2.51 ± 0.13	1.16 ± 0.07	0.065 ± 0.003
	% Improvement	74	71	3	17	-14	37
50 μm	High Efficiency	7.55 ± 0.72	6.50 ± 0.53	3.65 ± 0.19	4.54 ± 0.49	1.73 ± 0.07	0.118 ± 0.007
	Conventional	3.23 ± 0.16	2.85 ± 0.07	1.93 ± 0.07	2.94 ± 0.20	1.10 ± 0.04	0.062 ± 0.002
	% Improvement	134	104	90	54	58	91
20 μm	High Efficiency	10.69 ± 1.86	9.94 ± 0.92	3.15 ± 0.23	4.12 ± 0.40	2.39 ± 0.71	0.109 ± 0.008
	Conventional	5.43 ± 2.37	3.47 ± 0.38	2.26 ± 0.48	3.23 ± 0.22	1.15 ± 0.14	0.068 ± 0.008
	% Improvement	97	187	40	28	107	61
10 μm	High Efficiency	10.00 ± 1.23	17.22 ± 8.46	2.75 ± 0.99	4.60 ± 1.74	2.13 ± 0.80	0.095 ± 0.035
	Conventional	4.80 ± 3.14	4.25 ± 2.39	1.94 ± 0.16	5.89 ± 4.39	1.00 ± 0.37	0.068 ± 0.030
	% Improvement	108	306	41	-22	112	39

The reduction in sample ion yield at larger spot sizes was also associated with an increase in washout time. The spot sizes 50 μm and below consequently showed the highest sample ion yields. At a spot size of 50 μm the sample ion yield for U averaged 7.55 % when tuned for maximum signal and 4.35 % in optimized (for low laser-induced elemental fractionation and oxide formation) tune conditions. The equivalent for the conventional configuration was 3.23 % and 2.03 % respectively. Using the high efficiency configuration effectively doubled the sample ion yield which could be achieved by the LA-MC-ICP-MS (Figure 1). Increases in sample ion yield were not

limited to U, but observed across the entire mass range at spot sizes smaller than 50 μm.

The sample ion yield of Hf was less than would be expected from its atomic mass, interpolating between Nd and Pb in Figure 1, but still approximately two times larger (3.65 % against 1.93 %) than for the conventional configuration. Due to its refractory nature and high ionization potential, Hf requires a significant amount of energy to fully ionize. In theory the residence time of the ablated material inside the ICP may have been insufficient to ionize all of the Hf atoms introduced (compared proportionately to Nd and Pb).

**Table 6. 20 Hf spot ablations on the Plešovice zircon, 25  $\mu\text{m}$  spot size, 6.5 J  $\text{cm}^{-2}$  fluence, 10 Hz repetition rate, 8 s duration.**

Spot	$^{173}\text{Yb}$ (V)	Total Hf (V)	$^{178}\text{Hf}/^{177}\text{Hf}$	2SE	$^{179}\text{Hf}/^{177}\text{Hf}$	2SE	$^{176}\text{Yb}/^{177}\text{Hf}$	2SE	$^{176}\text{Lu}/^{177}\text{Hf}$	2SE	$^{176}\text{Hf}/^{177}\text{Hf}_{\text{Yb}}$	2SE	$^{176}\text{Hf}/^{177}\text{Hf}_{\text{CF}}$	2SE	$\beta_{\text{Hf}}$	$\beta_{\text{Yb}}$
Plešovice_1	0.064	44	1.467231	0.000046	0.740047	0.000039	0.006314	0.000046	0.0001562	0.0000009	0.282440	0.000021	0.282440	0.000024	-0.911	-0.710
Plešovice_2	0.064	42	1.467197	0.000052	0.739948	0.000059	0.006640	0.000160	0.0001562	0.0000008	0.282462	0.000020	0.282466	0.000021	-0.899	-0.653
Plešovice_3	0.062	41	1.467244	0.000045	0.739905	0.000041	0.006510	0.000130	0.0001565	0.0000004	0.282435	0.000020	0.282435	0.000022	-0.894	-0.690
Plešovice_4	0.057	41	1.467265	0.000045	0.739905	0.000054	0.006020	0.000140	0.0001442	0.0000005	0.282456	0.000025	0.282456	0.000027	-0.894	-0.686
Plešovice_5	0.060	41	1.467206	0.000046	0.739937	0.000048	0.006290	0.000140	0.0001533	0.0000004	0.282448	0.000015	0.282445	0.000016	-0.897	-0.718
Plešovice_6	0.064	42	1.467217	0.000048	0.739950	0.000050	0.006570	0.000100	0.0001584	0.0000009	0.282435	0.000025	0.282436	0.000024	-0.899	-0.683
Plešovice_7	0.062	42	1.467218	0.000046	0.739965	0.000042	0.006394	0.000075	0.0001590	0.0000008	0.282446	0.000021	0.282446	0.000022	-0.901	-0.691
Plešovice_8	0.059	41	1.467204	0.000047	0.739930	0.000032	0.006290	0.000150	0.0001495	0.0000005	0.282470	0.000025	0.282475	0.000027	-0.897	-0.648
Plešovice_9	0.056	41	1.467186	0.000039	0.739865	0.000051	0.005900	0.000170	0.0001370	0.0000005	0.282440	0.000020	0.282442	0.000023	-0.889	-0.670
Plešovice_10	0.061	40	1.467216	0.000051	0.739893	0.000051	0.006670	0.000140	0.0001578	0.0000003	0.282457	0.000018	0.282456	0.000021	-0.892	-0.702
Plešovice_11	0.061	43	1.467225	0.000050	0.740049	0.000044	0.006187	0.000048	0.0001538	0.0000008	0.282456	0.000020	0.282457	0.000022	-0.911	-0.677
Plešovice_12	0.060	43	1.467257	0.000049	0.740013	0.000045	0.006076	0.000051	0.0001503	0.0000010	0.282465	0.000020	0.282471	0.000022	-0.907	-0.639
Plešovice_13	0.060	43	1.467187	0.000042	0.739999	0.000044	0.006029	0.000068	0.0001479	0.0000006	0.282463	0.000021	0.282466	0.000024	-0.905	-0.666
Plešovice_14	0.059	43	1.467256	0.000043	0.739988	0.000041	0.006016	0.000059	0.0001485	0.0000006	0.282463	0.000019	0.282461	0.000024	-0.904	-0.710
Plešovice_15	0.060	43	1.467254	0.000054	0.740047	0.000038	0.006118	0.000029	0.0001525	0.0000011	0.282452	0.000022	0.282460	0.000024	-0.911	-0.613
Plešovice_16	0.059	43	1.467238	0.000037	0.740045	0.000048	0.005978	0.000048	0.0001484	0.0000005	0.282450	0.000021	0.282450	0.000019	-0.910	-0.690
Plešovice_17	0.059	43	1.467291	0.000054	0.740015	0.000035	0.005962	0.000044	0.0001480	0.0000006	0.282457	0.000020	0.282457	0.000020	-0.907	-0.690
Plešovice_18	0.059	43	1.467224	0.000046	0.740049	0.000042	0.006016	0.000050	0.0001491	0.0000005	0.282443	0.000019	0.282452	0.000022	-0.911	-0.605
Plešovice_19	0.059	43	1.467237	0.000049	0.740008	0.000041	0.005968	0.000055	0.0001478	0.0000010	0.282458	0.000019	0.282461	0.000021	-0.906	-0.670
Plešovice_20	0.059	43	1.467219	0.000047	0.740007	0.000037	0.005992	0.000049	0.0001482	0.0000010	0.282452	0.000021	0.282452	0.000024	-0.906	-0.690
	$^{173}\text{Yb}$ (V)	Total Hf (V)	$^{178}\text{Hf}/^{177}\text{Hf}$	2SE	$^{179}\text{Hf}/^{177}\text{Hf}$	2SE	$^{176}\text{Yb}/^{177}\text{Hf}$	2SE	$^{176}\text{Lu}/^{177}\text{Hf}$	2SE	$^{176}\text{Hf}/^{177}\text{Hf}_{\text{Yb}}$	2SE	$^{176}\text{Hf}/^{177}\text{Hf}_{\text{CF}}$	2SE	$\beta_{\text{Hf}}$	$\beta_{\text{Yb}}$
Mean	0.060	42	1.467229	0.000047	0.739978	0.000044	0.006197	0.000088	0.0001511	0.0000007	0.282452	0.000021	0.282454	0.000022	-0.902	-0.675
2SD			0.000056		0.000046						0.000020		0.000022			
2RSD (ppm)			38		63						72		79			

All reported sample ion yields were based on the average of four 50 shot ablations. However, when ablating a crater using a static spot, the amount of signal generated decreases over time - ablation efficiency decreases as the hole depth increases below the depth of field of the laser focus. For the smallest (10  $\mu\text{m}$ ) spot size determined, a 48 % decrease in measured signal was observed between the first 5 shots and last 5 shots ablated. There was a corresponding 22 % signal decrease for the 20  $\mu\text{m}$  spot size and a 7 % signal decrease at 50  $\mu\text{m}$ . Compared to the overall efficiencies reported in Table 6, assuming consistent ablation, we can calculate the first five shots were 36%, 11% and 1.5% more efficient for 10  $\mu\text{m}$ , 20  $\mu\text{m}$  and 50  $\mu\text{m}$  spot sizes respectively. The higher peak sample ion yields are important considerations given single pulse laser ablation<sup>26,27</sup> and sample surface rastering, e.g. for mapping, where these peak sample ion yields would be maintained. More generally, higher peak sample ion yields for near surface ablation advocates increasing utilization of shallower, better spatially constrained, laser ablation methodologies which can still achieve required precisions.

A sample ion yield >7 % for U represents a significant increase on values which have been previously reported for MC-ICP-MS, either by laser ablation or any other sample introduction technique. The maximum sample ion yield of U ( $\leq 3.5$  %) previously reported using a desolvating nebulizer system<sup>28</sup> are similar to the value measured here for laser ablation using the conventional configuration. We can hypothesize that a further optimization of sample introduction using desolvating nebulizers, based on the principles developed for high efficiency laser ablation, may yield similar improvements in sample ion yields for solution analysis.

#### **Pb Isotopic Analysis of MPI-DING Reference Materials.**

A normal LA-MC-ICP-MS analysis would require a faster repetition rate than the 1 Hz this study used to determine the sample ion yield in order to achieve a smooth signal profile. The fast washout time of the ARIS on a single collector ICP-MS would require a very high repetition rate (>40 Hz) to achieve a smooth signal profile. Ultimately the utility of rapid response ablation cells to routine LA trace elemental analysis of the complete mass range with a scanning, single collector instrument, is limited by spectral skew until the introduction of a signal smoothing device. The slower tau response of the Faraday cup detectors in the MC-ICP-MS already smooth the signal profile such that rates >8 Hz are sufficient. By eliminating the requirement for high repetition rates, which lead to rapid ablation through a sample, Faraday detectors allow rapid response laser ablation cells to be used in routine LA-MC-ICP-MS analysis, taking advantage of the higher sample ion yields prevalent at lower ablation rates. Note if a mixed detector array of Faraday and SEM detectors was used a higher repetition rate would be needed to avoid spectral skew on the SEM portion of the measurement.

Similarly to when calculating the sample ion yield, the Pb isotope ion yield was between 85 – 105 % higher for the SRM612 standard when using the high efficiency configuration (Table 7.) whilst maintaining equivalent  $^{208}\text{Pb}/^{206}\text{Pb}$  and  $^{207}/^{206}\text{Pb}$  ac-

curacy compared to the conventional laser configuration, allowing for the determined uncertainty. For ATHO-G and GOR132-G the  $^{208}\text{Pb}/^{206}\text{Pb}$  repeatability was nearly identical for both configurations; for StHs6/80-G the 2SD was lower for the conventional configuration (0.13 v 0.17 %). The repeatability of  $^{207}\text{Pb}/^{206}\text{Pb}$  was much improved by the increased signal of the high efficiency configuration for all three reference glasses.

#### **Hf isotope ratio analysis and U/Pb geochronology of reference zircons.**

The demanding precision requirements of  $^{176}\text{Hf}/^{177}\text{Hf}$  analysis requires sufficient ablated material to satisfy counting statistics. Typically this is achieved by using relatively large spot sizes of 35 – 50  $\mu\text{m}$ . A potential consequence of using a larger spot size in complex zircons is that the measured  $^{176}\text{Hf}/^{177}\text{Hf}$  may be decoupled from the age measured at the same or an adjacent location. Efforts have therefore been ongoing to reduce the amount of material required for  $^{176}\text{Hf}/^{177}\text{Hf}$  analysis in zircons.

Bauer & Horstwood (2017) reported achieving the required precision on  $^{176}\text{Hf}/^{177}\text{Hf}$  with a LA-MC-ICP-MS spot ablation, 100 shots at 10 Hz, 25  $\mu\text{m}$  circle, 6.5  $\text{Jcm}^{-2}$ . For the common reference zircon Plešovice this equated to an average of 25 V/s total Hf signal for 40 ng of total zircon ablated. Using the same laser conditions in conjunction with the high efficiency rapid response laser ablation cell yielded 42 V/s of total Hf signal (Table 6) maintaining  $^{176}\text{Hf}/^{177}\text{Hf}$  uncertainties <1  $\epsilon$  (2s). Reducing the spot size further to 20  $\mu\text{m}$  the total Hf measured for Plešovice was still 29 V/s (supplementary data). Assuming the same ablation rate for these fluences between the two systems, this is equivalent to just 26 ng of total ablated zircon.

Using an identical 20  $\mu\text{m}$  spot size Pb/Pb ages were determined for the same reference zircons. The signal intensity on the 91500 zircon was 123 % higher for the high efficiency laser ablation configuration compared to the conventional configuration. This corresponded to a reduction in uncertainty on the  $^{207}\text{Pb}/^{206}\text{Pb}$  age from 2.26% to 0.80 % (2SD, n=10).

## **CONCLUSION**

Coupling LA-MC-ICP-MS with high efficiency laser ablation designs has been demonstrated to double the sample ion yield. We demonstrate these designs can be used successfully to measure routine LA-MC-ICP-MS applications with better isotope ratio precision and spatial resolution.

A sample ion yield of greater than 7 % for U (and potentially up to 8-9% for U and Pb) represents a new maximum in the field of MC-ICP-MS and confirms that limitations in ion yield for this technique are principally related to the sample introduction system. Incorporating some of the technological developments from high efficiency laser ablation designs into other sample introduction systems within ICP-MS may lead to similar improvements in ion yield. This would be of great benefit to analyses, in varied fields such as cosmochemistry, environmental monitoring and nuclear safeguards where the amount of sample material available is limited.



		<sup>206</sup> Pb (mV)	<sup>207</sup> Pb (mV)	<sup>208</sup> Pb (mV)	<sup>208</sup> Pb/ <sup>206</sup> Pb	2SD	2RSD (%)	<sup>207</sup> Pb/ <sup>206</sup> Pb	2SD	2RSD (%)
<b>ATHO-G</b> (5.5 µg/g)	High Efficiency	7.6	6.5	15.8	2.0746	0.0056	0.27	0.8427	0.0012	0.14
	Conventional	3.3	2.8	6.6	2.0723	0.0058	0.28	0.8419	0.0025	0.30
					(Ref: 2.0735)			(Ref: 0.8421)		
<b>GOR132-G</b> (20.25 µg/g)	High Efficiency	32.2	26.4	64.4	2.0081	0.0023	0.11	0.8160	0.00044	0.054
	Conventional	16.6	13.6	32.5	2.0087	0.0023	0.11	0.8166	0.00054	0.066
					(Ref: 2.0095)			(Ref: 0.8163)		
<b>StHs6/80-G</b> (10.25 µg/g)	High Efficiency	19.72	16.40	38.70	2.0368	0.0034	0.17	0.8259	0.00038	0.046
	Conventional	7.21	5.99	14.33	2.0364	0.0026	0.13	0.8262	0.00092	0.111
					(Ref: 2.0380)			(Ref: 0.8263)		

**Table 7.** <sup>208</sup>Pb/<sup>206</sup>Pb and <sup>207</sup>Pb/<sup>206</sup>Pb isotope ratio results for three MPI-DING glasses. 40 µm spot size, 3 J cm<sup>-2</sup> fluence, 16 Hz repetition rate, 22 s duration.

## ASSOCIATED CONTENT

### Supporting Information

Sample ion yield signal intensity, Hf and U/Pb, enhanced single crater image, data reduction schemes for Pb and Hf (.docx)  
Crater volume measurement (PDF)

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### Notes

Dr. Grant Craig and Dr. Nicholas Lloyd are employed by Thermo Fisher Scientific (Bremen) GmbH. Dr. Ciprian Stremtan is employed by Teledyne CETAC Technologies.

## REFERENCES

- Wang, H. A. O.; Grolimund, D.; Giesen, C.; Borca, C. N.; Shaw-Stewart, J. R. H.; Bodenmiller, B.; Günther, D. *Anal. Chem.* 2013, **85**, 10107–10116.
- Van Malderen, S. J. M.; van Elteren, J. T.; Vanhaecke, F. *J. Anal. At. Spectrom.* 2015, **30**, 119–125.
- Douglas, D. N.; Managh, A. J.; Reid, H. J.; Sharp, B. L. *Anal. Chem.* 2015, **87**, 11285–11294.
- Van Malderen, S. J. M.; Managh, A. J.; Sharp, B. L.; Vanhaecke, F. *J. Anal. At. Spectrom.* 2016, **31**, 423–439.
- Managh, A. J.; Horstwood, M. S. a.; Reid, H. J.; Sharp, B. L. In *13th European Workshop on Laser Ablation*; Ljubljana, 2016.
- Van Malderen, S. J. M.; van Elteren, J. T.; Šelih, V. S.; Vanhaecke, F. *Spectrochim. Acta - Part B At. Spectrosc.* 2018, **140**, 29–34.
- Breglio, A. M.; Rusheen, A. E.; Shide, E. D.; Fernandez, K. A.; Spielbauer, K. K.; McLachlin, K. M.; Hall, M. D.; Amable, L.; Cunningham, L. L. *Nat. Commun.* 2017, **8**, 1654–1662.
- Theiner, S.; Van Malderen, S. J. M.; Van Acker, T.; Legin, A.; Keppler, B. K.; Vanhaecke, F.; Koellensperger, G. *Anal. Chem.* 2017, **89**, 12641–12645.
- Van Malderen, S. J. M.; Laforce, B.; Van Acker, T.; Nys, C.; De Rijcke, M.; De Rycke, R.; De Bruyne, M.; Boone, M. N.; De Schampelaere, K.; Borovinskaya, O.; De Samber, B.; Vincze, L.; Vanhaecke, F. *Anal. Chem.* 2017, **89**, 4161–4168.
- Petrus, J. A.; Chew, D. M.; Leybourne, M. I.; Kamber, B. S. *Chem. Geol.* 2017, **463**, 76–93.

(11) Craig, G. PhD. Dissertation, Loughborough University, 2015.

(12) Fisher, C. M.; Vervoort, J. D.; Dufrane, S. A. *Geochemistry, Geophys. Geosystems* 2014, **15**, 121–139.

(13) Bauer, A. M.; Horstwood, M. S. A. *Chem. Geol.* 2017, **476**, 85–99.

(14) Schaltegger, U.; Schmitt, A. K.; Horstwood, M. S. A. *Chem. Geol.* 2015, **402**, 89–110.

(15) Johnston, S.; Gehrels, G.; Valencia, V.; Ruiz, J. *Chem. Geol.* 2009, **259**, 218–229.

(16) Horstwood, M. S. A.; Košler, J.; Gehrels, G.; Jackson, S. E.; McLean, N. M.; Paton, C.; Pearson, N. J.; Sircombe, K.; Sylvester, P.; Vermeesch, P.; Bowring, J. F.; Condon, D. J.; Schoene, B. *Geostand. Geoanalytical Res.* 2016, **40**, 311–332.

(17) Kimura, J.-I.; Chang, Q.; Kanazawa, N.; Sasaki, S.; Vaglarov, B. S. *J. Anal. At. Spectrom.* 2016, **31**, 790–800.

(18) Lloyd, N. S.; Sadekov, A. Y.; Misra, S. *Rapid Commun. Mass Spectrom.* 2018, **32**, 9–18.

(19) Cottle, J. M.; Burrows, A. J.; Kylander-Clark, A. R.; Freedman, P. A.; Cohen, R. S. *J. Anal. At. Spectrom.* 2013, **28**, 1700–1706.

(20) Hieftje, G. M.; Mermet, J.-M.; Hill, S. J.; Fisher, A.; Foulkes, M.; Sharp, B. L.; O'Connor, C.; O'Connor, G.; Evans, E. H.; Vanhaecke, F.; Balcaen, L.; Taylor, P.; Hansen, H. R.; Pergantis, S. A. *Inductively Coupled Plasma Spectroscopy and its Applications*, 2nd ed.; Hill, S. J., Ed.; Blackwell Publishing: Oxford, 2007.

(21) Ranebo, Y.; Hedberg, P. M. L.; Whitehouse, M. J.; Ingeneri, K.; Littmann, S. *J. Anal. At. Spectrom.* 2009, **24**, 277–287.

(22) Gerstenberger, H.; Haase, G. *Chem. Geol.* 1997, **136**, 309–312.

(23) Bürger, S.; Riciputi, L. R.; Bostick, D. A.; Turgeon, S.; McBay, E. H.; Lavelle, M. *Int. J. Mass Spectrom.* 2009, **286**, 70–82.

(24) Payne, J. L.; Pearson, N. J.; Grant, K. J.; Halverson, G. P. *J. Anal. At. Spectrom.* 2013, **28**, 1068–1079.

(25) Fisher, C. M.; McFarlane, C. R. M.; Hanchar, J. M.; Schmitz, M. D.; Sylvester, P. J.; Lam, R.; Longrich, H. P.; *Chem. Geol.*, 2011, **284**, 1–20.

(26) Cottle, J. M.; Horstwood, M. S. a.; Parrish, R. R. *J. Anal. At. Spectrom.* 2009, **24**, 1355–1363.

(27) Cottle, J. M.; Kylander-Clark, A. R.; Vrijmoed, J. C. *Chem. Geol.* 2012, **332–333**, 136–147.

(28) *Neptune Plus Multicollector ICPMS, Another step ahead in MC-ICP-MS*, Thermo Scientific BR30197-E01/12G, Thermo Fisher Scientific, Bremen, 2011.

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