





prep*FAST* MC™



Automating Sample Purification



prep <i>FAST</i> MC™	
Brief and system schematics2	
Uranium Isotopes3	
Strontium Isotopes6	
Boron Isotopes9	

Brief

The prepFAST MC[™] is a fully automated, low pressure chromatography system that isolates elements of interest from the sample matrix and collects multiple discrete eluent fractions for precise isotopic analysis. The syringe-driven system allows sample loading, multiple acid washes, column conditioning and elution cycles all at user-defined intervals (time, volume and flow rate).

Features:

- · Fully automated
- Syringe control
 - Load exact volume
 - Dispense exact volume
 - Precise flow rates
- · All fluoropolymer flow path
- · Multiple destination locations
- · Flexible chemistry

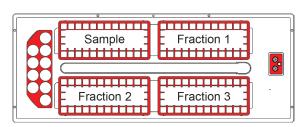
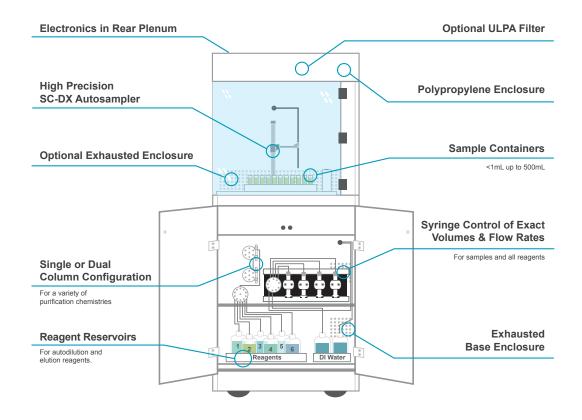
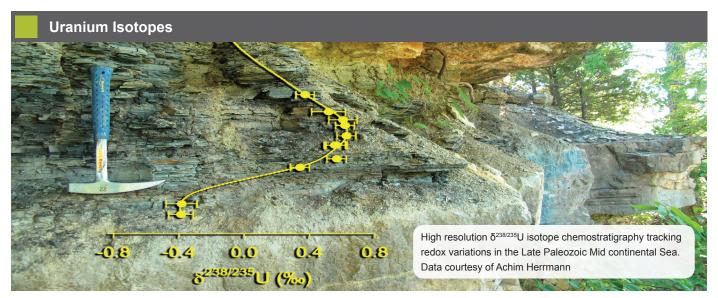


Figure 1. SC-4 DX top view with locations for Sample, Fraction 1, Fraction 2, and Fraction 3. Vial sizes, styles and rack configurations are flexible.





Author: Paul Field (field@icpms.com)

High-resolution isotope ratio stratigraphy requires the routine analysis of large numbers of samples. To reduce interferences and minimize matrix effects, extensive purification procedures are used to isolate elements from their natural matrix. Current purification protocols require manually feeding gravity-driven separation columns, a process that is both costly and time consuming. This laboratory bottleneck is eliminated with the prep*FAST* MC[™], an automated, low-pressure ion exchange chromatography system that can process from 1 to 60 samples in unattended operation. The syringe-driven system automatically isolates elements of interest and collects up to 3 discrete fractions at user-defined intervals (time, volume and flow rate). The combination of maximizing sample throughput and minimizing costs associated with personnel and consumables provides an opportunity to greatly expand research horizons in fields where large isotopic data sets are required, including archeology, geochemistry, climate/environmental science, biomedical sciences and food authentication.

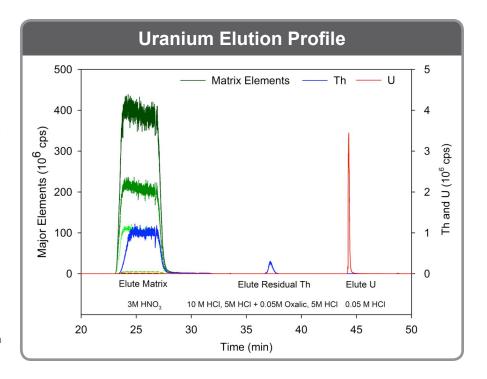
Figure 2. Fully-automated chromatographic separation of uranium (red) from all other matrix elements in BCR-2 (Columbia River Basalt) using the prepFAST MC™ system. Elution profile was collected online in real-time using a Thermo XSERIES ICP-MS. Data courtesy of Stephen Romaniello and Gwyneth Gordon (Arizona State University).



In collaboration with:



Stephen Romaniello, Gwyneth W. Gordon, Achim Herrmann and Ariel D. Anbar



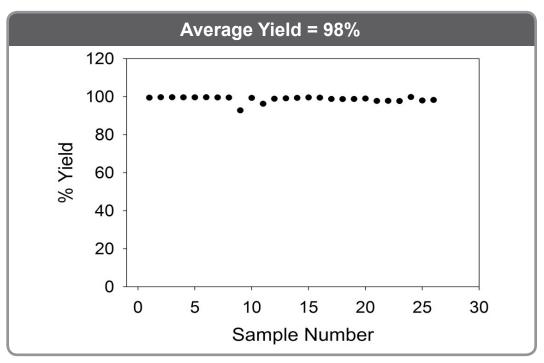


Figure 3. Uranium recovery for 26 consecutive samples with fully-automated cleaning and reuse of the separation column. The average yield was 98% and the minimum yield was 92%. Data courtesy of Stephen Romaniello and Gwyneth Gordon (Arizona State University).

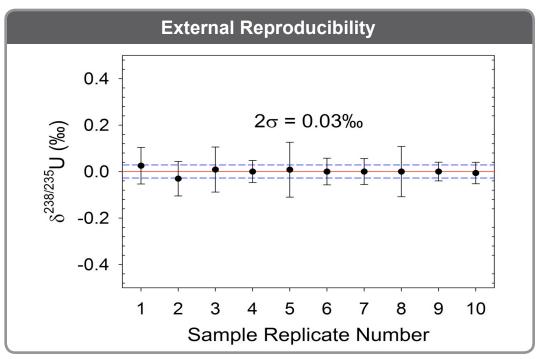


Figure 4. External reproducibility of $\delta^{238/235}$ U for 10 aliquots CRM145 independently processed through chemistry over a 1 week period interspersed with natural samples using the prep*FAST* MC[™]. Errors bars indicate the 2 σ precision of replicate measurements on a single sample aliquot (n≥3). Blue dashed lines indicate the 2 σ precision for the means of all sample aliquots. Data collected using a Thermo Neptune MC-ICPMS and courtesy of Stephen Romaniello and Gwyneth Gordon (Arizona State University).

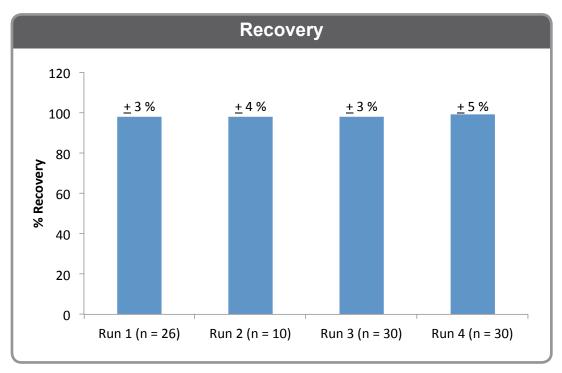


Figure 5. Data collected over four runs of thirty automated extractions illustrates constant, complete recovery for within run, and run to run extractions. The column was replaced for each run. Within run replicates are indicated by "n" and recovery determinations are within analytical uncertainty.

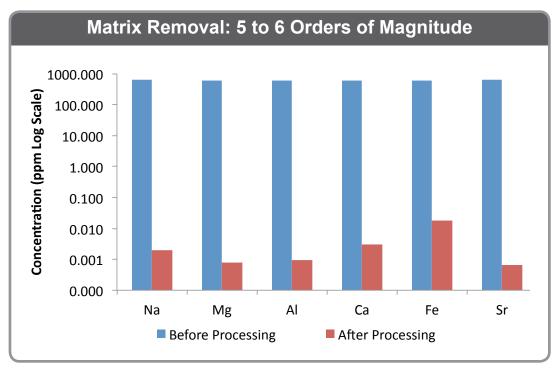


Figure 6. The concentration of major matrix elements determined in final extractions indicates greater than 99.999% matrix removal. Initial concentration of 500 ppm matrix elements is reduced to less than 20 ppb or lower.



Authors: Paul Field (field@icpms.com) and Patrick Sullivan

High-precision isotope analyses in forensic anthropology, nuclear forensics and food authentication applications, requires the generation of large data sets. A new, fully automated Sr purification method for the prepFAST MC $^{\text{TM}}$ eliminates the sample preparation bottle neck. Reproducibility, reliability, recovery, blank and carry over are at required levels for the determination of Sr isotopes in a variety of natural samples. Potential applications of prepFAST MC $^{\text{TM}}$ are illustrated by accurate and precise determination of Sr isotopes in rock, bone, and seawater reference materials. Unattended, the prepFAST MC $^{\text{TM}}$ can process up to 48 samples in 24 hours and, (depending on chemistry), collect several purified extractions from each sample.

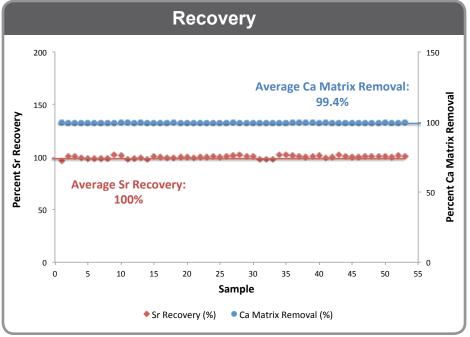
Figure 7. Strontium recovery for 55 consecutive, fully automated extractions is plotted. 100% recovery and near complete removal of Ca provides pure fractions for accurate and precise determination of isotopic ratios.



In collaboration with:



Stephen Romaniello, Gwyneth W. Gordon and Ariel D. Anbar



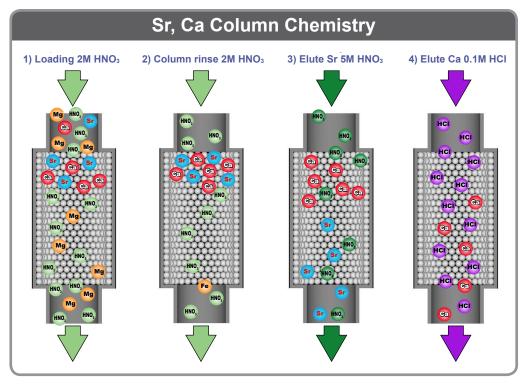


Figure 8. Illustration of fully-automated chromatographic separation of Sr from Ca and other matrix components.

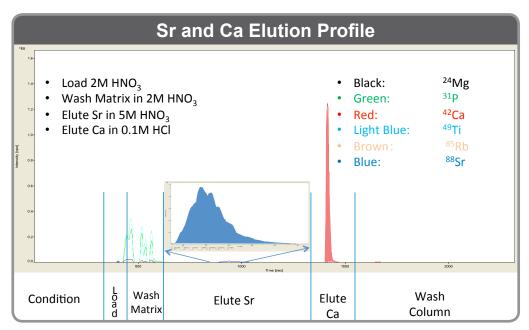


Figure 9. Online, real-time elution profile collected on the Element 2 (Thermo) illustrates the fully-automated chromatographic separation of Sr and Ca from matrix components. In carbonate samples it is possible to also collect the Mg fraction during sample load and wash cycles.

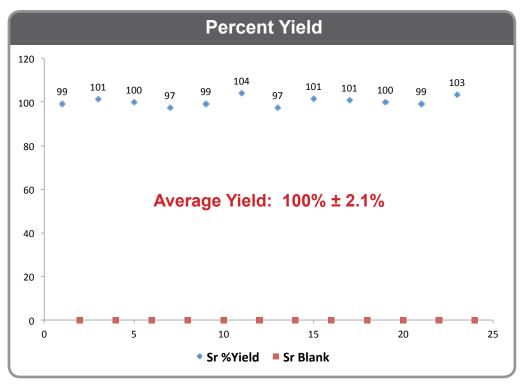


Figure 10. Strontium recovery for 24 consecutive, fully automated extractions with cleaning and reuse of the separation column. Constant high yields and no carryover (below detection) illustrate the ability to reuse the resin.

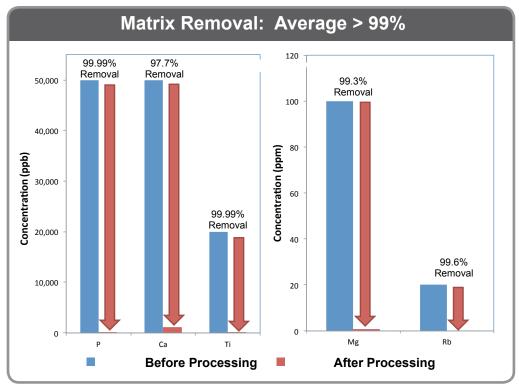
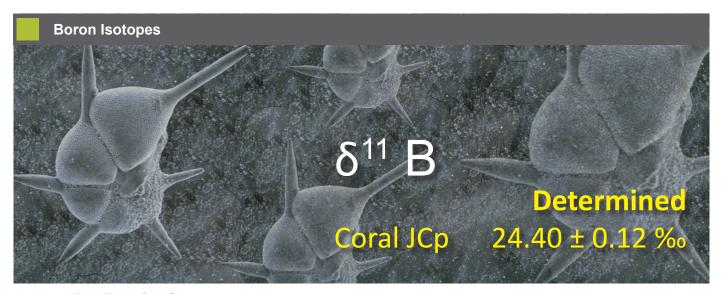


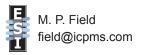
Figure 11. The concentration of major matrix elements determined in final extractions indicates > 99% matrix removal (P, Ti, Mg, Rb). Ca removal is sufficient for accurate determination of Sr isotope ratios and can be further reduced with additional wash cycles. The Ca fraction can also be collected for isotope ratio measurement, if desired.



Authors: Paul Field (field@icpms.com)

The merits of a new prepFAST MC TM method for the purification of boron from carbonate samples are demonstrated. The fully automated syringe-driven system cleans the column (0.5M HNO $_3$), conditions the column (DI H $_2$ O), loads the sample (0.5M HNO $_3$) buffered with Na acetate/acetic acid), elutes the matrix (DI H $_2$ O) and elutes the sample (0.5M HNO $_3$) from the column. User-defined reagent volumes and flow rates result in near complete removal of sample matrix (>99% of Ca) and the quantitative recovery of boron in ~ 600 μ L (~ 100% \pm 2.7%, n=10). Full automation, precise/accurate syringe control, identical flow path, and one column/resin bed all contribute to the reproducibility of extraction. The concept of resin reuse is validated by > 30 replicate extractions with high yield and minimal carryover (< 0.01%; below our detection). The prepFAST MC TM processes a sample in approximately 30 mins (48 samples per day) automating timely extraction techniques that will aid the advance of boron isotopes as a mainstream research tool.

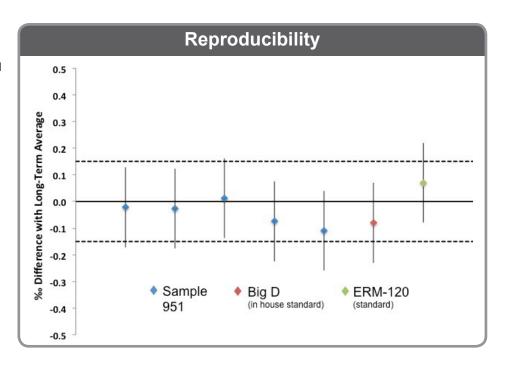
Figure 12.Comparison of prep*FAST* MC[™] purified samples to National Oceanography Centre long term average.



In collaboration with:



Miguel A. Martinez-Boti, Eleni Anagnostou and Gavin Foster National Oceanography Centre Southampton University, UK



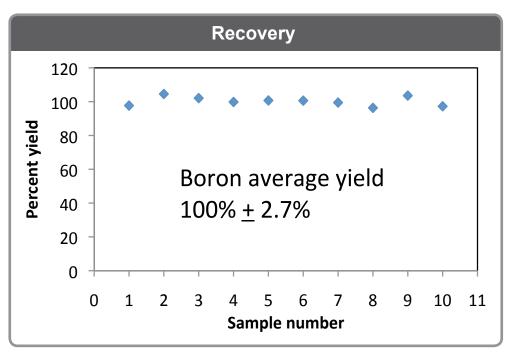


Figure 13. Data collected from twenty automated extractions, alternating between sample and blank using the prep*FAST* MC[™] boron method, are plotted. The data illustrate the ability to maintain both consistent and complete recovery of boron from multiple injections on one column.

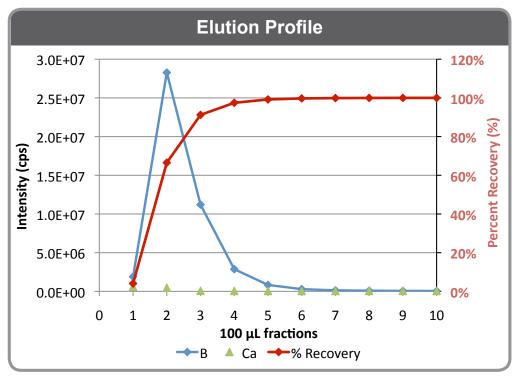


Figure 14. Ten 100 μ L fractions collected from the prep*FAST* MCTM are used to generate an elution profile. Baseline calcium and high recovery (> 99.9% in 600 μ L) indicate sufficient matrix removal and quantitative boron elution. Quantitative recovery in a small final volume maintains boron concentrations at sufficient levels for isotopic analysis.

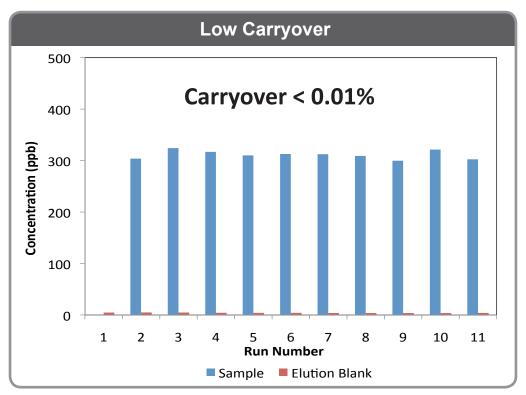


Figure 15. Analysis of alternating blank - sample fractions from the prep*FAST* MC[™] indicate boron values return to baseline concentrations for each blank. When baseline boron concentration is subtracted from each blank the data indicate greater than 4 orders of magnitude washout. Less than 0.01% carryover immediately after processing a 300 μg/L sample demonstrates no cross contamination from column reuse.

Benefits:

- · Fully automated
- User defined
 - sample size
 - elution volume
 - wash/load/elute rates
- Multiple fractions
- High throughput (30-50 samples/day)
- Stand alone system runs 24 hrs unattended





© Elemental Scientific | 7277 World Communications Drive | Omaha, NE 68122 Tel: 402-991-7800 | sales@icpms.com | www.icpms.com