sapphire
Collision Cell Multi-Collector
ICP-MS

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Sapphire: Technologies Collide for Unparalleled Performance

Sapphire: Collision Cell Multi Collector Inductively Coupled Plasma Mass Spectrometer

Introduction

The application of non-traditional stable isotope systems such as Mg, Si, S, K, Ca, Ti, V, Cr, Fe, Ni and Se in geochemistry, cosmochemistry and life sciences has been spurred by the advancement of the multi-collector ICP-MS (MC-ICP-MS) over the last two decades. This versatile technique, although widely considered as the standard method for the high precision isotopic analysis of such systems, is limited by the molecular species generated by the argon ICP ion source which cause significant interferences on the mass spectrum of these elements. These molecular species can interfere directly with the atomic ions of the same nominal mass, leading to inaccurate isotope ratio determination. A typical solution is to use the high-resolution capabilities of MC-ICP-MS to resolve or partially resolve the isotopes of interest from their respective interferences, but at the cost of significantly reduced ion transmission. Furthermore, analysis of certain isotopes such as $^{40}$Ca is not possible on the traditional MC-ICP-MS instruments, as the $^{40}$Ar interference cannot be resolved.

Sapphire is a collision/reaction cell MC-ICP-MS incorporating both a “high energy” ion path (traditional MC-ICP-MS) and a separate “low energy” ion path (collision/reaction cell). The instrument offers an ideal solution for the precise and accurate isotopic analysis of both the traditional isotope systems, and the non-traditional isotope systems mentioned above.

Sapphire has combined the Plasma 3 mass analyser with the dual path transfer optics (patent pending) offering the benefits of collision/reaction cell without compromising performance for classical geological applications.

The ‘dual path’ design allows the cell to be used to remove isobaric interferences for specific applications and can be fully bypassed when analyzing classical isotope systems that do not require removal of interferences or for systems where the high resolution features are preferred.

Application Areas

- planetary
- life science
- geochemistry

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Key features:

- Double focusing high precision isotopic measurement mass spectrometer
- Enhanced Sensitivity Interface as standard
- Dual Path transfer optics, incorporating:
  - Low energy path with collision / reaction cell for interference removal
  - High energy path for classical multi-collector performance
- Up to 4 independently controlled reaction gas inlets
- Fully dry pumped configuration available for lower hydrocarbon backgrounds
- Sixteen Faraday detectors, with optional software switchable resistor amplifiers and up to six ion-counting detectors (with SEM / Daly options) giving maximum user flexibility and dynamic range
- High abundance deceleration filters available for improved abundance sensitivity
- High resolution and pseudo high resolution capabilities
- Variable Zoom Optics allow for instantaneous switching between isotopic systems during analysis
Dual Path Transfer Optics

The dual path transfer optics retain the high sensitivity of MC-ICP-MS by extracting the ions at a high energy (~6keV). The ion beam can then be directed along two distinct paths.

The ‘High Energy Path’ maintains the high ion energy throughout the transfer optics before focusing into the defining slit of the double focusing mass spectrometer, retaining all the key performance of traditional MC-ICP-MS.

The ‘Low Energy Path’ diverts the ion beam along a second route before the ions are decelerated into the collision / reaction cell. Gas is introduced into the RF cell to produce reactions with the low energy ions. On exiting the cell, the ions are accelerated back to the high energy and focused into the defining slit of the double focusing mass spectrometer.

The ‘dual path’ design provides an effective solution to mass discrimination problems potentially caused by RF multi-pole devices associated with other reaction cell instruments.

Interference Removal

The collision / reaction cell enables removal of isobaric interferences such as Ar⁺ or polyatomic interferences such as ArH⁺, ArN⁺ or ArO⁺ which can interfere directly with the atomic ions of the same nominal mass for example K, Ca, Fe.

Such interferences can be removed via charge exchange reactions with Hydrogen such as:

\[
\text{Ar}^+ + \text{H}_2 \rightarrow \text{Ar} + \text{H}_2^+ \\
\text{ArH}^+ + \text{H}_2 \rightarrow \text{Ar} + \text{H}_3^+ \\
\text{ArO}^+ + \text{H}_2 \rightarrow \text{Ar} + \text{H}_2\text{O}^+
\]

These reactions reduce the Argon ion signal by more than 9 orders of magnitude while other analytes are unaffected.

The removal of these interferences allows the measurement of non-traditional stable isotopes such as potassium and calcium at low resolution, without reducing transmis-
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