

Fast determination of low mass, inorganic cations in a ground water sample using IC-MS

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Introduction

Reliable analytical methods for food and beverage samples are required in order to report ingredients for food labeling requirements, to maintain product quality, and to ensure the absence of food contamination and spoilage. Food and beverage samples are almost invariably one of the most complex sample matrices and, therefore, are among the most challenging samples to analyze.

Ion chromatography (IC), using eluent generation and suppressed conductivity detection, provides chromatographic selectivity, low chemical noise, and high compatibility with a mass spectrometer (MS).^{1,2} Additionally, the analytes leave the IC system as ions, further increasing MS compatibility.

Mass spectrometry coupled to IC provides higher selectivity and often, better detection limits. In addition, MS is one of the most universal detectors, providing powerful screening, structural, and confirmatory information.² The heated electrospray ionization (HESI), with applied high temperature aiding voltage, transforms the aqueous IC stream into a fine spray that can enter the MS detector.¹

In this application brief, separations of four low-mass, inorganic cations in a diluted ground water sample were achieved within six minutes. The analytes were detected serially, by suppressed conductivity and single quadrupole mass spectrometry in full scan (FS) and SIM (select ion monitoring) modes without the aid of any desolvation agent during separation or ionization.

This application brief is an update to Application Note 269.³ Here, the application is performed using a Thermo Scientific™ Dionex™ Integrion™ HPIC™ system coupled to a Thermo Scientific™ ISQ™ EC single quadrupole mass spectrometer.

Experimental Ion Chromatography

- Dionex Integrion HPIC system, RFIC™ model with a second six-port high-pressure divert valve and conductivity detector
- Thermo Scientific™ Dionex™ AS-AP autosampler
- Thermo Scientific™ Dionex™ AXP-MS auxiliary pump to supply water for the suppressor

Mass Spectrometry

- ISQ EC single quadrupole mass spectrometer
- Thermo Scientific syringe pump for method optimization
- HESI II probe

Software

Thermo Scientific™ Chromeleon™ Chromatography Data System (CDS) software, 7.2 SR 6

Methods

Columns:	Thermo Scientific™ Dionex™ IonPac™ CG12A-5µm, CS12A-5µm, 3 mm i.d.	
Eluent:	33 mM Methanesulfonic acid (MSA)	
Eluent Source:	Thermo Scientific™ Dionex™ EGC 500 MSA cartridge, Thermo Scientific™ Dionex™ CR-CTC 600 trap column, high pressure degasser module	
Flow Rate:	0.5 mL/min	
Injection Volume:	100 µL	
Detection 1:	Suppressed conductivity, Dionex CERS-500e suppressor, external water mode at 0.7 mL/min by the AXP-MS auxiliary pump	
Typical Conductance Background:	< 1 µS-min	
MS Detection:*	+ESI, +3000 V, Full Scan, 18-250 <i>m/z</i> and SIM, HESI II	
Temperatures:	Vaporizer: 250 °C; Ion Transfer: 300 °C	
Flow (N ₂):	Sheath: 60 psi, Aux: 26 psi; Sweep: 0.5 psi	
Desolvation agent:	None	
SIM mode:	Ion <i>m/z</i>	CID (V)
Sodium as Na•2H ₂ O	59	10
Ammonium as NH ₄ •H ₂ O	36	2
Potassium	39	45
Magnesium as Mg ₂ •H ₂ O	66	5
Calcium	40	45
Dimethylamine	46	10
Ethylamine	46	45
Monoethanolamine	62	15
Diethylamine	74	15
Triethanolamine	150	25

*Note: The optimum HESI ion source settings and responses may vary between instruments.

Results

Figure 1 shows the IC-MS results of inorganic cations in a diluted ground water sample. Small m/z analytes are challenging because of the low mass. The ions were detected in SIM mode from m/z 39 to 66 as bare ions or water-adducts. The IC and SIM peaks are symmetrical and have strong responses, E2 to E5 ion counts (in SIM mode). Bare, unsolvated calcium (peak 5) m/z 40 had the lowest response at 5e2 (500) counts. This application brief demonstrates the low mass detection capabilities of the ISQ EC mass spectrometer. More information can be found in the Thermo Scientific AppLabs Library of Analytical Applications.⁴

References

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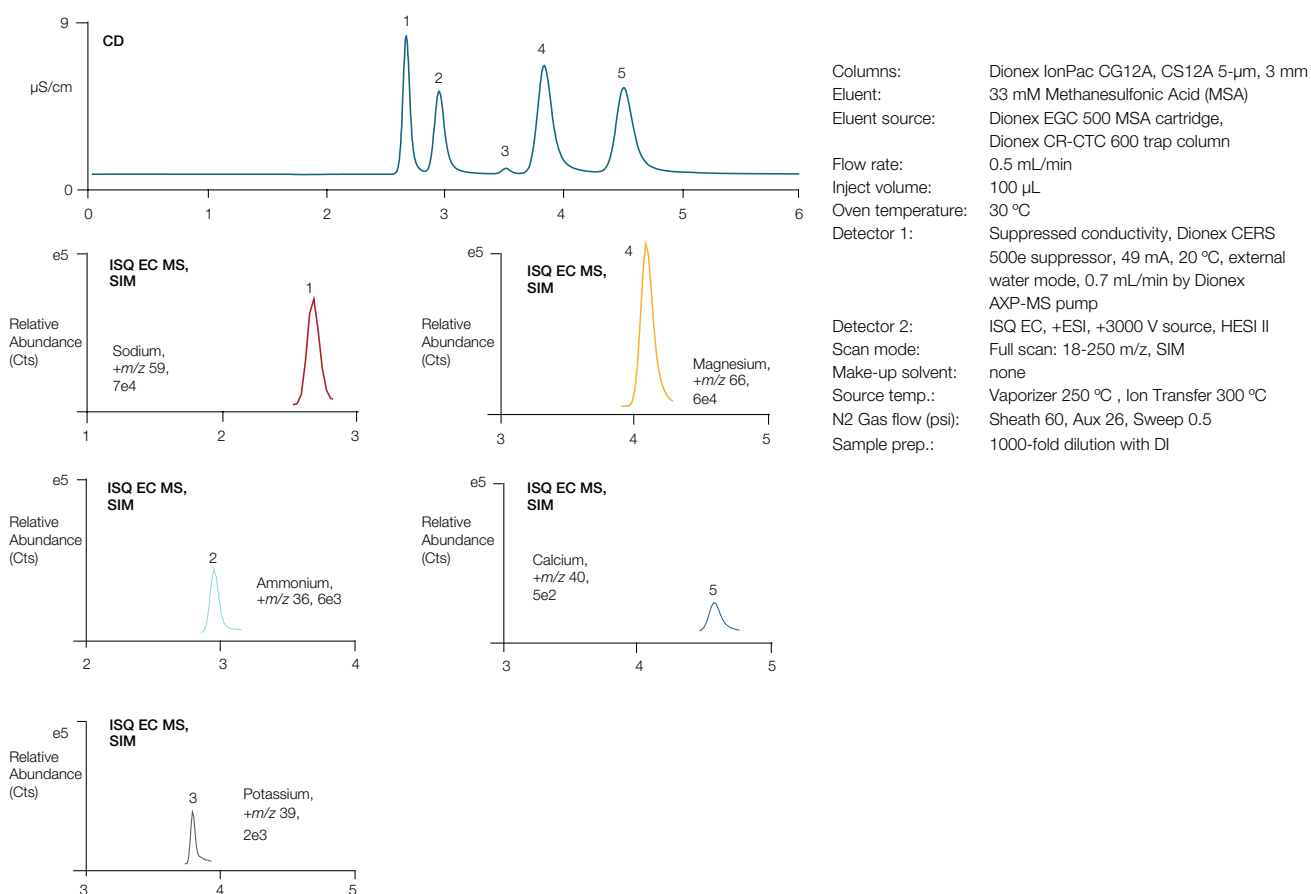


Figure 1. Determination of low-mass cations in a ground water sample.

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