

Determination of Trace Anions in High-Purity Waters by Ion Chromatography with the IonPac[®] AS17 Using High-Volume Direct Injection with the EG40

INTRODUCTION

This application note describes an ion chromatographic method using the microbore IonPac[®] anion-exchange AS17 column to determine trace concentrations of common inorganic anions, low-molecular-weight organic acids, as well as acrylate, methacrylate, benzoate, and phthalate. These contaminants can come from some of the following sources: cleaning agents, adhesives, oils, mold release agents, and solder fluxes.^{1,2} Anionic contamination is known to cause corrosion in microelectronic circuitry.³ The analytes are detected by suppressed conductivity with a 2-mm ASRS[®] ULTRA operated in the “Gas-Assisted Recycle” mode. Two different gradient methods are described: a 27-min separation and 41-min separation. The EG40 generates high-purity and carbonate-free hydroxide eluents on-line to improve the method performance for determination of target analytes at trace levels.^{4,5} The high-volume, direct-injection technique is used to achieve sensitive detection at low- to sub- $\mu\text{g/L}$ levels without the use of a concentrator column or loading pump and valve.⁶⁻⁸ This application note expands on work presented in Dionex Technical Note 48, *Determination of Trace Anions in High-Purity Water by High-Volume Direct Injection with the EG40*.⁹

EQUIPMENT

Dionex DX-600 ion chromatography system consisting of:

- GS50 Gradient Pump
- CD25 Conductivity Detector
- LC30 Chromatography Enclosure equipped with Rheodyne Model 9126 injector PEEK, rear-loading (P/N 052291)

Columns:

- IonPac AS17 analytical, 2 × 250 mm (P/N 055683)
- IonPac AG17 guard, 2 × 50 mm (P/N 055685)

Trap Column:

- ATC-1 Trap, 9 × 24 mm (P/N 037151) placed after pump

Suppressor:

- ASRS ULTRA 2-mm (P/N 053947)

EG40 Eluent Generator with EluGen[®] EGC-KOH cartridge (P/N 053921)

Gas-Assisted Regeneration Kit (P/N 056886)

Pressurized sample vessel (P/N 37053) and low-pressure 4-way valve, 10-32 fittings (P/N 45010), optional

Chromeleon[®] Chromatography Workstation
300 cm of green 0.75-mm (0.030-in.) PEEK tubing to make a 1000- μL sample loop

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REAGENTS AND STANDARDS

Deionized water, Type I reagent-grade,
18 MΩ-cm resistance

ACS reagent-grade materials for preparing
anion standards (VWR or other)

Sodium hydroxide 50% w/w aqueous solution
(Fisher Scientific)

Potassium hydroxide 45% w/w aqueous solution
(Fisher Scientific) (optional in place of
sodium hydroxide)

Fluoride standard 1000 mg/L, 100 mL
(Dionex P/N 037158)

Chloride standard 1000 mg/L, 100 mL (Dionex
P/N 037159)

Sulfate standard 1000 mg/L, 100 mL (Dionex
P/N 037160)

Nitrate standard 1000 mg/L, 100 mL (Dionex
P/N 056497)

Phosphate standard 1000 mg/L, 100 mL (Ultra
Scientific, VWR P/N ULICC-005)

Bromide standard 1000 mg/L, 100 mL (Ultra
Scientific, VWR P/N ULICC-001)

CONDITIONS

Eluent: Potassium hydroxide
(EG40 as the source)

Temperature: 30 °C

EG40 Offset Volume: 0 µL

Eluent Flow Rate: 0.5 mL/min

Detection: Suppressed conductivity,
ASRS ULTRA Gas-Assisted
Recycle mode

ASRS Current Setting: 100 mA

Expected Background Conductivity:
1 µS (40 mM KOH)

Expected System Backpressure:
15.2–16.6 MPa
(2200–2400 psi)

Sample Volume: 1 mL

Pump Program Method 1

Time (min)	Flow (mL/ min)	A (%)	Valve	EG40 Conc. (mM)	Comments
Initial	0.50	100	Load	20.0	20 mM KOH
0.00	0.50	100	Load	1.0	Load sample loop, equilibrate 1.0 mM KOH for 5 min
5.00	0.50	100	Inject	1.0	Inject
15.00	0.50	100	Inject	1.0	1 mM KOH
19.00	0.50	100	Load	12.0	12 mM KOH
25.00	0.50	100	Load	20.0	20 mM KOH
27.00	0.50	100	Load	20.0	20 mM KOH

Pump Program Method 2

Time (min)	Flow (mL/ min)	A (%)	Valve	EG40 Conc. (mM)	Comments
Initial	0.50	100	Load	40.0	40 mM KOH
0.00	0.50	100	Load	0.3	Load sample loop, equilibrate 0.3 mM KOH for 5 min
5.00	0.50	100	Inject	0.3	Inject
11.00	0.50	100	Inject	0.3	0.3 mM KOH
13.00	0.50	100	Inject	1.0	1 mM KOH
24.00	0.50	100	Load	10.0	10 mM KOH
41.00	0.50	100	Load	40.0	40 mM KOH

PREPARATION OF SOLUTIONS AND REAGENTS

Standard Solutions

Stock Anion Standard Solution (1000 mg/L)

Several of the analytes of interest are available as 1000 mg/L anion standard solutions from Dionex or other commercial sources. In cases where commercial standards are not available, 1000 mg/L standards can be prepared by dissolving the appropriate amounts of the corresponding mass for the target analytes in 1000 mL of deionized water according to Table 1. We recommend making a 100-mL final volume of 1000 mg/L stock standards in 125-mL high-density polyethylene (HDPE) containers. Concentrated standards are stable for at least one month when stored at 4 °C.

Table 1. Amounts of Compounds Used to Prepare 1 L of 1000-mg/L Ion Standards

Anion	Compound	Mass (g)
Fluoride	Sodium fluoride (NaF)	2.210
Acetate	Sodium acetate (CH ₃ COONa • 3H ₂ O)	2.305
Formate	Sodium formate (HCOONa)	1.511
Acrylate	Sodium acrylate (H ₂ C=CHCO ₂ Na)	1.324
Methacrylate	Sodium methacrylate (H ₂ C=C(CH ₃)CO ₂ Na)	1.270
Chloride	Sodium chloride (NaCl)	1.648
Nitrite	Sodium nitrite (NaNO ₂)	1.499
Bromide	Sodium bromide (NaBr)	1.288
Nitrate	Sodium nitrate (NaNO ₃)	1.371
Benzoate	Sodium benzoate (C ₆ H ₅ CO ₂ Na)	1.190
Sulfate	Sodium sulfate (Na ₂ SO ₄)	1.479
Oxalate	Sodium oxalate (Na ₂ C ₂ O ₄)	1.522
Phthalate	Phthalic acid (C ₆ H ₄ -1,2-(CO ₂ H) ₂)	1.000
Phosphate	Potassium phosphate, monobasic (KH ₂ PO ₄)	1.433

Composite Standard Solution

Composite standards at lower analyte concentrations are prepared from the 1000-mg/L standards above. Select a range similar to the expected analyte concentrations in the samples. Take aliquots from this dilute standard to make working standards at the low- μ g/L (ppb) down to the high ng/L (ppt) range. Dilute stock standards at the low-mg/L (ppm) levels should be prepared fresh weekly. Working standards at the low- μ g/L (ppb) range should be made fresh daily.

ATC Regeneration Solution

2 M Sodium Hydroxide

Dilute 160 g of 50% (w/w) sodium hydroxide with degassed, deionized water (with a specific resistance of 18 M Ω -cm) to a final weight of 1080 g in an eluent bottle. Avoid the introduction of carbon dioxide from the air. Note that 2 M potassium hydroxide can be used instead of 2 M sodium hydroxide. Preparation is the same as above, except that 249 g of 45% potassium hydroxide is used for a final weight of 1090 g.

SYSTEM PREPARATION AND SETUP

This section describes the procedures for the initial installation and start-up of the ASRS ULTRA, ATC, and EGC-KOH EluGen cartridge. Prepare the ASRS according to the *Quickstart Instructions for the ASRS ULTRA* (Document No. 031368). Prepare the ATC for use by rinsing it with 200 mL of 2 M KOH or 2 M NaOH at 2.0 mL/min. This preparation can be done off-line without the GS50 by pressurizing an eluent bottle with helium at 34.5 kPa (5 psi). Rinse the ATC with deionized water at 2.0 mL/min for 20 min and then stop.

Install the EGC-OH EluGen cartridge according to the instructions in the *Operator's Manual for the EG40 Eluent Generator System* (Document Number 031373). Place the 4-mm ATC-1 column between the GS50 outlet and the EGC-KOH cartridge inlet, as shown in Figure 9, "EG40 Flow Path Diagram" in the EG40 Eluent Generator system manual.

Program the EG40 to generate the highest concentration of KOH that will be used by the method and equilibrate the separation column for 30 min at the method flow rate.

To operate the ASRS for the "Gas-Assisted Recycle" mode, use the Gas-Assisted Regeneration Kit (P/N 056886). Connect the "Regen-In" line (shorter tubing from the tee) to the SRS "Regen-In" port. Connect the line from the "Cell-Out" on the conductivity cell to the "From-Cell" line (longer tubing from the tee). Connect a gas line to the regulator. Compressed air can be used in place of helium or nitrogen if it is purified with suitable traps to remove moisture, hydrocarbons, carbon dioxide, and other particulate matter. For more information, see the *SRS Gas-Assisted Regeneration Kit Installation and Use Instructions* (Document No. 031665-02).

Table 2. Volume per Unit Length for Various Tubing Internal Diameters

Material	Color	Internal Diameter		Est. Vol.
		(inches)	(mm)	($\mu\text{L}/\text{cm}$)
PEEK	Red	0.005	0.125	0.126
PEEK	Black	0.010	0.250	0.506
PEEK	Orange	0.020	0.500	2.022
PEEK	Green	0.030	0.750	4.550

Make a 1000- μL sample loop by cutting a 220-cm portion of the green 0.030-in. (0.75-mm) i.d. PEEK tubing. If a loop or tubing with different internal diameter is desired, refer to Table 2 to calculate the length needed. The volume of a loop can be verified by measuring the weight difference between the sample loop filled with deionized water and the empty loop. The inside diameter of the PEEK tubing can vary by as much as 20% (for example, 0.010 ± 0.002 in.).

Connect the columns and suppressor in the IC system by using the red 0.005-in. (0.125-mm) or yellow 0.003-in. (0.075-mm) PEEK tubing. Keep the lengths of connecting tubing as short as possible to minimize the system void volume. Minimizing ensures efficient 2-mm column operation. Carefully use a Dionex plastic tubing cutter (P/N 45257) to ensure that the surfaces of the tubing cuts have straight, smooth surfaces. Irregularity on the surface of a tubing end can result in unwanted additional dead volume.

SYSTEM OPERATION

Turn on the gradient pump to begin the flow of eluent through the system. If the system backpressure is below 14 MPa (2000 psi), a length of yellow PEEK 0.003-in. (0.075-mm) tubing should be added between the outlet of the degas assembly in the EG40 and inlet of the injection valve. For optimal EG40 performance, maintain a system backpressure of 15.2–16.6 MPa (2200–2400 psi). Confirm that no leaks exist anywhere in the chromatographic pathway. For more information, see the *Operator's Manual for the EG40 Eluent Generator System* (Document Number 031373).

After stable eluent flow has been established, apply 10–15 psi gas pressure to the “Regen-In” port of the

ASRS. A mixture of gas and liquid will be observed exiting the “Regen-Out” port. Turn the power on to supply current to the ASRS; the gas flow rate across the regenerant chamber will be about 100 mL/min at a set gas pressure of 10–12 psi.

Turn on the EG40 to deliver the highest eluent concentration required by the method using Chromeleon's run program. Allow the LC30 oven to stabilize at 30 °C. Determine the quality of the blank by measuring the short-term noise. In a representative 1-min level portion of the chromatogram, a peak-to-peak measurement should be less than 10 nS. For trace analysis, it may take 12 h or more for the system to equilibrate to a stable low background conductivity. It is a good practice to run a system overnight to equilibrate for use the following day.

The sample is loaded with either a syringe or pressurized reservoir. When using a syringe, take care not to introduce contamination by contact of the sample with the syringe. The black rubber plunger in disposable plastic syringes can be a source of significant contamination. To avoid contamination, the syringe should be used to pull sample into the loop when placed at the waste port. Take care not to introduce bubbles into the loop by pulling too hard.

When loading sample with a pressurized reservoir, a low-pressure, double-stack valve at the waste port regulates when the sample is loaded into the loop. More details about sample loading strategies can be found in Dionex Technical Note 48, *Determination of Trace Anions in High-Purity Water by High-Volume Direct Injection with the EG40*. An AS40 or AS50 autosampler is not suitable for the determination of anions at concentrations below 10 $\mu\text{g}/\text{L}$ (ppb) because the pathway of an autosampler contributes unwanted anionic contamination.

DISCUSSION AND RESULTS

The EG40 Eluent Generator electrolytically produces high-purity KOH eluents using deionized water as the carrier stream.⁴ The EG40 can generate eluents that are free of carbonate contamination. For gradient separations, the EG40 provides negligible baseline shifts, greater retention time reproducibility, and better method precision. Background conductivity is lower, providing the best signal-to-noise ratio. These features enhance the ion chromatographic performance for the determination of anions with the EG40 at trace levels.⁵

Table 3. Anion Retention Times by High-Volume Direct-Injection Ion Chromatography with the EG40 and the IonPac AS15-5 μ m

Anion	Retention time, min
Fluoride	6.2
Glycolate	7.0
Acetate	7.7
Formate	8.1
Acrylate	10.1
Chloride	10.8
Nitrite	12.0
Carbonate	13.0
Benzoate	13.3
Methacrylate	13.4
Sulfate	14.2
Oxalate	14.5
Bromide	15.2
Nitrate	16.0
Phosphate	16.5
Phthalate	21.2

Conditions (see Application Update 142, *Improved Determination of Trace Anions in High Purity Waters by High-Volume Direct Injection with the EG40*).

The use of the IonPac AS15-5 μ m column was evaluated for the determination of anions at trace levels in a previous study.¹⁰ Table 3 summarizes the retention time results for the analytes of interest using a gradient from 7 to 60 mM KOH. Notice that several analytes are not well resolved: acrylate (10.1 min), chloride (10.8 min), carbonate (13.0 min), benzoate (13.3 min), and methacrylate (13.4 min). However, this column has very good resolution of fluoride and the weakly retained organic acids such as acetate, glycolate, and formate. For more information about this separation, see Application Update 142, *Improved Determination of Trace Anions in High-Purity Waters by High-Volume Direct Injection with the EG40*.¹⁰

The IonPac AS17 column is chosen for this method because it provides the best selectivity for the analytes of interest to the electronics industry: common inorganic anions, low-molecular-weight organic acids,

acrylate, methacrylate, benzoate, and phthalate. For trace analysis, the large-loop injection technique is used with a weak starting eluent concentration. The ASRS ULTRA delivers low background and noise for sensitivity at trace levels. The “Gas-Assisted Recycle” mode is chosen to supply the regenerant solution to the ASRS. This mode provides the benefit of low noise without the need for external water, significantly reducing water usage and waste. The “Gas-Assisted Recycle” mode reduces peak-to-peak noise from an ASRS by a factor of 4 to 10 while using the eluent as the source for regenerant. The use of the DS-3 conductivity cell minimizes the effects of cell drift and temperature fluctuations.

Two gradient methods were developed in this study: Method 1 has a separation time of 27 min and Method 2 takes 41 min. Method 1 is best suited for well-characterized samples where fast sample throughput is important. Method 2 provides better separation of analytes and is suited for samples containing target analytes at disparate concentrations.

Both methods begin with a 5-min equilibration at the starting eluent concentration. This dilute eluent is used to elute the weakly retained ions such as fluoride, acetate, and formate. A gradient to a higher KOH concentration is used to separate the more strongly retained ions, such as sulfate and phosphate. The chromatographic baseline shift during the gradient is typically less than 200 nS when using the EG40. The separations are performed at 30 °C to provide consistent retention times during trace analysis. Typical chromatograms for low-level standards for Method 1 and Method 2 are shown in Figures 1 and 2, respectively.

A representative deionized water blank for both methods is shown in Figures 3 and 4, respectively. Determining a blank establishes a starting point above which trace-level anion determinations can be made. No significant contaminants were detected except an unidentified peak and carbonate. The use of a high-quality deionized water purification system minimizes the amount of carbonate.

Method detection limits (MDLs) for the target analytes were established using Method 1 (Table 4). Method 2 yielded comparable results. The MDLs for the target analytes were determined by multiplying the standard deviation of seven replicate injections for the lowest-level standard with the Student’s *t* value for the 99% confidence level.¹¹ Calibration curves were ob-

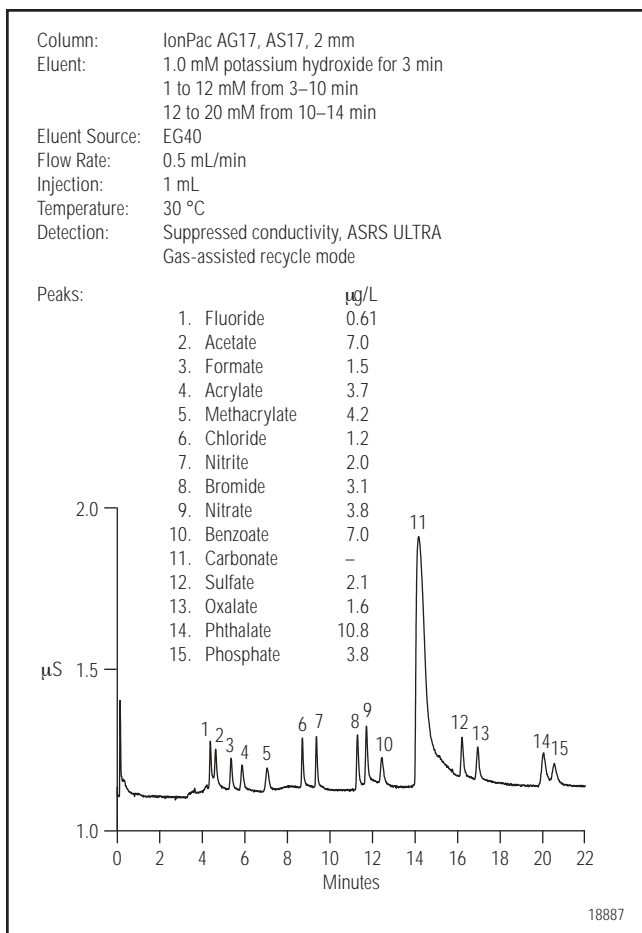


Figure 1. Trace anion determination using the IonPac AS17 with Method 1.

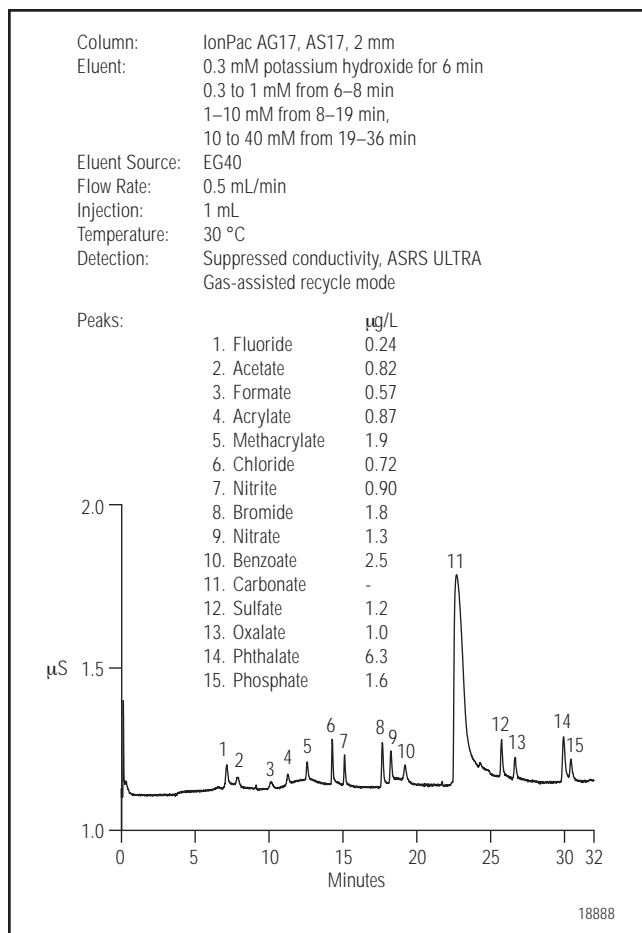


Figure 2. Trace anion determination using the IonPac AS17 with Method 2.

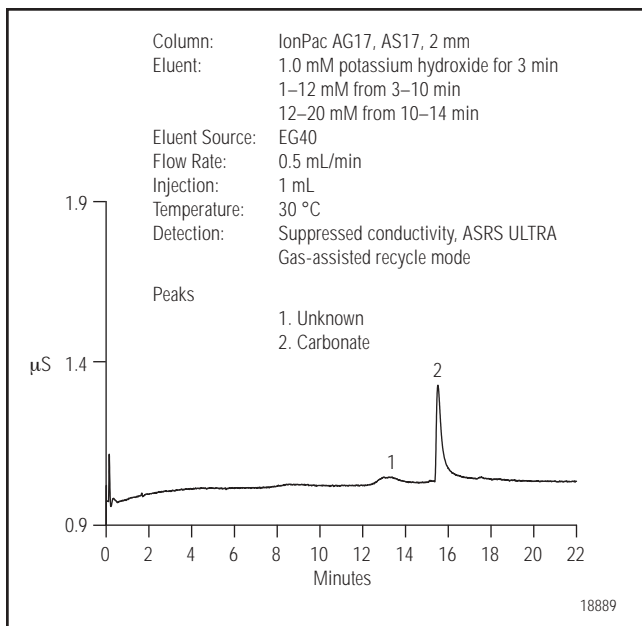


Figure 3. Representative blank for trace analysis with Method 1.

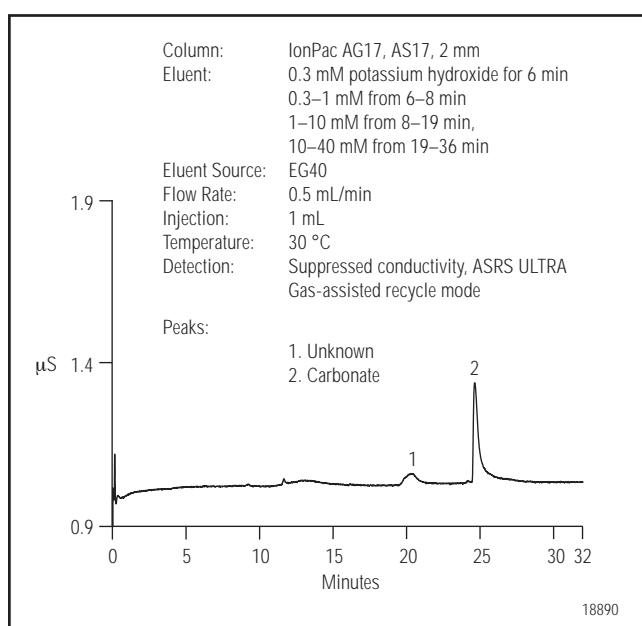


Figure 4. Representative blank for trace analysis with Method 2.

Table 4. Method Detection Limits^a for Anions by High-Volume Direct-Injection Ion Chromatography with the EG40 and the IonPac AS17 2 mm

Anion	MDL ^a µg/L (ppb)
Fluoride	0.040
Acetate	1.1
Formate	0.85
Acrylate	0.44
Methacrylate	0.35
Chloride	0.12
Nitrite	0.23
Bromide	0.28
Nitrate	0.12
Benzoate	0.76
Sulfate	0.23
Oxalate	0.21
Phthalate	0.84
Phosphate	0.88

^aMDL = (S.D.) x (t)_{99%}, where (t)_{99%} is for a 99% single-sided Student's *t* test distribution for n = 7

tained using standards prepared in deionized water. The analyte concentrations in the calibration standards are listed in Table 5. Three replicate injections were used at each concentration level. Results for the anions of interest yielded a linear response with coefficients of determination (*r*²) greater than 0.99. To accurately determine the area of a peak at trace levels, it may be necessary to manually draw the baselines using tools in the Chromeleon software. Remember that the lowest quantifiable analyte concentration is generally 3 to 5 times greater than the lowest detectable concentration.¹²

PRECAUTIONS

Special care must be taken to minimize contamination when performing trace analysis. The quality of the blank will be an important determining factor to the concentration of ions that can be reliably determined in a sample. It is very important to use only the highest quality deionized water. When conducting analysis at trace levels, the sources of contamination are numerous. To minimize contamination, wear disposable, powder-free PVC gloves. After putting them on, rinse with

Table 5. Calibration Curve Concentrations (µg/L) for Anions by High-Volume Direct-Injection Ion Chromatography with the EG40 and the IonPac AS17 2 mm

Anion	Levels		
	1	2	3
Fluoride	0.1	0.3	1.0
Acetate	1.0	3.0	10.0
Formate	1.0	3.0	10.0
Acrylate	0.3	1.0	3.0
Methacrylate	1.0	3.0	10.0
Chloride	0.1	0.3	1.0
Nitrite	0.1	0.3	1.0
Bromide	0.3	1.0	3.0
Nitrate	0.3	1.0	3.0
Benzoate	1.0	3.0	10.0
Sulfate	0.3	1.0	3.0
Oxalate	0.3	1.0	3.0
Phthalate	1.0	3.0	10.0
Phosphate	1.0	3.0	10.0

deionized water and air dry. Do not dry with paper towels. All containers should be dedicated for this analysis and copiously rinsed with 18 MΩ-cm deionized water before use. Exercise caution when handling anything that could have contact with the blank, unknown, or standards. The flow path of the chromatographic instrumentation (eluent containers, injector, pump, valves, tubing, columns, suppressor, and conductivity cell) are all potential sources of contamination. For this reason, the flow path should be minimized wherever possible. Take care when switching from a system setup that had previously seen significant concentrations of anions. Rinse with high-purity water to reduce residual contamination.

The ATC should be periodically regenerated with the procedure described in the “System Preparation and Setup” section. Monitoring the blank for any significant increase in anionic contamination will indicate when regeneration is necessary. Monitoring the baseline shift during the EG40 hydroxide gradient will indicate when regeneration is necessary. A significant increase beyond

200 nS indicates that the ATC has exceeded its capacity to trap ionic contaminants and should be regenerated. The frequency of regeneration depends on the quality of the deionized water and usage rate of the instrument. Since the completion of this work, a new ATC—the ATC-HC (P/N 059604)—was introduced that extends the time between ATC regeneration.

If unexpected broad peaks appear in the chromatogram, it may be necessary to modify the gradient. These peaks can result from retained species of previous injections. Confirm this possibility by using an eluent concentration higher than the ending concentration of the gradient. It is best to run with this higher concentration for a period of time after all the peaks of interest have eluted. To confirm that the column has been cleaned of retained species, rerun a sample to observe the quality of the separation.

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