Method Guide: 40729

Method Guide - As in Potable waters

Key Words

- Arsenic
- Atomic Absorption
- Hydride generation
- HGAAS
- Potable water

Principle:

The sample is pre-reduced using a mixture of potassium iodide and L-ascorbic acid. Arsenic is determined by Hydride Generation Atomic Absorption Spectrometry (HGAAS) using the VP100 Continuous Flow Vapour System.

Reagents:

Hydrochloric acid (AnalaR grade, 50 %v/v)

Sodium borohydride solution (AnalaR grade, 0.5 % m/v in 0.5 % m/v sodium hydroxide)

Pre-reducing solution (AnalaR grade, 10 % m/v potassium iodide + 10 % m/v L-ascorbic acid)

Arsenic master standard (1000 mg/L)

Arsenic sub-stock standard solution (200 µg/L)

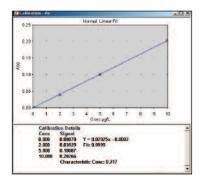
Transfer 0.2 mL of arsenic master standard solution to a 1 L volumetric flask and add 10 mL of pre-reducing solution, then dilute to volume with deionised water.

Working standards

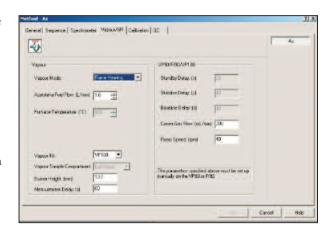
Transfer 0, 1.0, 2.5 and 5.0 mL of the arsenic sub-stock standard solution into a series of 100 mL volumetric flasks. Add 10mL of deionised water and 10 mL of hydrochloric acid to each flask and dilute to volume with deionised water. The working standards will contain 0, 2, 5 and 10 µg/L of arsenic.

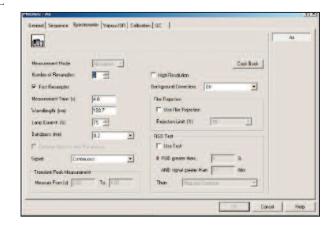
Sample Preparation:

Into a test tube, add 16.0 mL of drinking water sample, 2.0 mL of concentrated hydrochloric acid and 2.0 mL of the pre-reducing solution and mix thoroughly. Allow the mixture to stand for 1-2 hours at room temperature. The sample will then be ready for analysis. Tap water samples were spiked with 0.25, 0.5, 1.0 and 2.0 μ g/L of As and the spike recovery was calculated.



Instrument Parameters:







Results:

The calibration is perfectly linear and the characteristic concentration is 0.217 µg/L. The unspiked tap water sample = $0.61 \pm 0.02 \,\mu\text{g/L}$ and the average spike recovery (see table 1.) for As was $103.2 \pm 2.6 \%$

SAMPLE	SPIKE RECOVERED
Tap Water + 0.25 μg/L As	0.27 μg/L
Tap Water + 0.50 μ g/L As	0.48 µg/L
Tap Water + 1.00 μg/L As	1.03 µg/L
Tap Water + 2.00 μg/L As	2.13 μg/L

The method of sample treatment described in this publication should be performed only by a competent chemist or technician trained in the use of safe techniques in analytical chemistry. Users should acquaint themselves with particular hazards which may be incurred when toxic materials are being analysed and handled in the instruments, and the instrument must be used in accordance with the operating and safety instructions given in the Operators manual.

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