

## Lead

## Function: Differential Pulse Stripping Voltammetry (DPS/a)

| Start Potential     | (mV)   | -800     |
|---------------------|--------|----------|
| End Potential       | (mV)   | -200     |
| Current range       |        | 1,024 μΑ |
| Scan Speed          | (mV/s) | 30       |
| Deposition time     | (s)    | 120      |
| Deposition Pot.     | (mV)   | -800     |
| Number of cycles    |        | 3        |
| Delay before swee   | p (s)  | 5        |
| Purge and stir time | e (s)  | 20       |
| Stirring speed      | (rpm)  | 300      |
| Drop Size           | (a.u.) | 60       |

#### Lead concentrated standard solution (1 g/l)

Dissolve1.5986 g of Pb(NO<sub>3</sub>)<sub>2</sub> (pure and dried), in 1 l of 1 % HCl, in a volumetric flask.  $(MM_{Pb(NO_3)_2} = 331.21; MM_{Pb} = 207.2)$ .

## **Supporting Electrolyte**

A- 37% HCl

## B- 1 M H<sub>2</sub>C<sub>2</sub>O<sub>4</sub> and 2 M HCl solution

Dissolve 90 g of H<sub>2</sub>C<sub>2</sub>O<sub>4</sub> (or 126 g of H<sub>2</sub>C<sub>2</sub>O<sub>4</sub> · H<sub>2</sub>O) and 167 ml of 37% HCl in 1 l of distilled water. Store in a polythene bottle.

## **Procedure**

Add 20 µl of 37 % HCl to 10 ml of neutralised sample.

Alternatively: add 1 ml of B solution (especially if copper has to be analysed in the same solution).

Analyse sea water, high salt content sample and acidic solution (at pH between 1 and 3) avoiding the addition of the supporting electrolyte.

Samples at pH above 7 are to be neutralised before the addition of the supporting electrolyte.

## Working standard solution (1 mg/l)

Dilute 100  $\mu$ l of 1 g/l Pb standard solution in 100 ml of distilled water, in a volumetric flask. Add also 20  $\mu$ l of Cd concentrated standard solution if cadmium has to be analysed in the same scanning.

## **Alternative supporting electrolytes**

HCl or KCl or NaCl solution from 0.1 up to 1 M

- 0.1 M Acetate buffer pH 4.5 or 0.1 M citrate buffer a pH 3
- 0.1 M Tartrate buffer H 9 (when zinc has to be analysed in the same solution)

Avoid any addition of HNO<sub>3</sub> to the sample, because this acid could raise up the analytical peak. Use HCl instead of HNO<sub>3</sub>. If HNO<sub>3</sub> has to be used, dry the sample solution first and after add the supporting electrolyte.



# **Analytical report**

Analysis: tap water

Sample Concentration =  $4.40 \mu g/l$ 

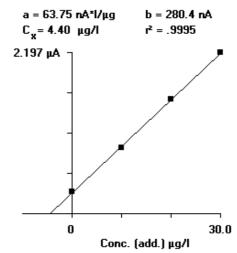
Method: 3 additions

## Volumes Table

Solvent Volume 0 (ml)
Supporting Sol. 0.01 (ml)
Sample Volume 10 (ml)
Standard Conc. 1000 (µg/l)

## Height Table

|   | 11018111111111 |          |
|---|----------------|----------|
| # | Peak Pot.      | Height   |
| 0 | -318.4         | 295.0 nA |
| 1 | -322.8         | 882.4 nA |
| 2 | -322.9         | 1.529 μΑ |
| 3 | -323.8         | 2.130 uA |



**AMEL 433** 

## Regression Data

| # | Add.Conc.   | Height x dilution |                               |
|---|-------------|-------------------|-------------------------------|
| 0 | $0 \mu g/l$ | 295.3 nA          | y = ax + b                    |
| 1 | 10.0 "      | 892.1 nA          | $a = 63.75 \text{ nA*l/\mug}$ |
| 2 | 20.0 "      | 1.562 μΑ          | b = 280.4  nA                 |
| 3 | 30.0 "      | 2.197 μA          | $r^2 = .9995$                 |
|   |             |                   |                               |

