

RiO5 METHOD (38)

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Po-210 in otolith: analytical method

The following is a historical method provided by J. Smith at BIO, Dartmouth, NS and dates to pre-1985. Additional information regarding this method may be found in the following publications:

Campana, S.E., K.C.T. Zwanenburg and J.N. Smith, 1990. Pb-210/Ra-226 determination of longevity in redfish. Canadian Journal of Fisheries and Aquatic Sciences, 46: 163-165.

Bennett, J.T., G.W. Boehlert, and K.K. Turekian, 1982. Confirmation of longevity in (*Sebastes diploproa*) (*Pisces:Scorpaenidae*) from $^{210}\text{Pb}/^{226}\text{Ra}$ measurements in otoliths. Mar. Bio. 71:209-215

Disclaimer

It is the responsibility of the analyst to follow established safety and health practices. Although each laboratory identified as the source has tested the methods, each user should perform an individual validation procedure.

Po-210 in Otolith:- analytical method

Laboratory Equipment

- 30 ml. beaker and watch glass
- Small stirring bar
- Glass rod
- Cap and screw neck piece from 30 ml. Nalgene LDPE plastic vial.
- 1 plastic disc to fit into the cap.
- Same size silver disk.(1.7 cm.)
- Stirrer hot plate
- Millipore filtratoin mechanism used in SR method.

Reagents

- 30 % H₂O₂
- 0.1 N HCL
- 1/2 Conc. HCL+HNO₃
- Conc. HCL
- 0.5 n. HCL
- Po-208 tracer
(act. =.592469
dpm/ml @30.12.86)

* Equipment and glasswear are boiled and cleaned in mild acid to reduce contamination. double-distilled acids are also essential. (Seastar quality)

Procedure

1) Clean otoliths

- Take approximately 1 to 2 grams of otoliths into 30 ml. beaker and add 2 ml. of 30 % H₂O₂. Soak for 30 min.
- Add 10 ml. 0.1 N HCL and rinse for 10 to 15 seconds.
- Pour off liquid and add 10 ml. of H₂O to rinse out excess residue. repeat * 2. Try to remove all possible film left on the otoliths.
- Put in the oven @ 40 deg. C. to dry over night.

2) Weigh otoliths and record weight.

3) Add 0.1 ml. Po-~~208~~ tracer.

4) Add 10 ml. 1/2 conc HCL+HNO₃, very slowly (1 ml. at a time) to avoid overflow from frothing.

5) When foaming subsides, completely evaporate the solution on the hot plate. (temperature set at approximately 150 - 200 deg C.)

6) Add 10 ml. conc. HCL

7) Evaporate the solution down again. Repeat, and cool.

8) Add 1 ml. H₂O₂ and leave over night.

9) Add 30 ml. 0.5 N. HCL.

Filter solution through the Millipore filtration system. Use filter type HA 0.45 um.

- 10) Plate solution at 85-90 deg. C. for approximately 4-6 hours onto silver disc encased in Nalgene 30 ml. LDPE plastic bottle cap. Make sure that it is totally immersed and is free of air bubbles. (Use a small stirring bar and a glass rod.)
- 11) Save solution for the Radium-226 analyses.
- 12) Count disc in L-spectrometer for approximately two to three weeks.
- 14) Calculate activity with APPLE-2 or P.C. computer using GWBASIC with program named PBOH, or PBOHP
(for printing out results.)

Blanks

- .1 ml tracer → 30 ml beaker
- + 10 ml 1/2 conc HCl + HNO₃ → dry
- 10 ml conc HCl → dry
- 10 ml conc HCl → dry
- + 1 ml H₂O₂ leave overnight
- + 30 ml .5M HCl → plating