FRANKLIN CRUISES FR 8/90, 5/92 AND 8/93 DATA DOCUMENTATION JGOFS WESTERN EQUATORIAL PACIFIC PROCESS STUDY

[1] General:

Parameter: Profiles of ²²⁶Ra, ²¹⁰Pb, and ²¹⁰Po

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E-Mail Address: jdsmith@primus.com.au List of Parameters: 226 Ra, 210 Pb, and 210 Po List of Units: Specific activity (Bq m⁻³)

[2] Sampling:

Gear (e.g. CTD, pump, etc.): CTD; 10 litre Niskin bottles

Standard Depths: Hydrochemistry depths: see Hydrochemistry data Chemicals used: Spiked with ¹³³Ba (for ²²⁶Ra), stable Pb, and ²⁰⁸Po. Special Procedures: See method. Samples from each depth were

pressure-filtered through cellulose nitrate

membrane filters (0.45µM pore diameter) and the

filtered water collected

Comments and Notes: Sampled at 10°N, 5°N, Equator, 5°S and 10°S

along 155°E.

[3] Analysis:

Instrument: Alpha spectrometry (²¹⁰Po); atomic absorption

spectrometry (Pb); and liquid scintillation counting (²²⁶Ra)

Method: See method below.

Precision: Given in Table based on one standard deviation from

counting statistics.

Comments: FR 9205: data from only one station

FR 9308: data for dissolved ²¹⁰Po not available due to

very low recovery rates. Particulate data fine.

[4] Results:

Quality of Data: FR 9308: good.

FR 9205: from station at 3°S, 155° E: good

Known Problems: Low recovery rates for dissolved ²¹⁰Po on FR 9308 meant that it

was not possible to calculate residence times in each depth bin.

[5] Brief description of analytical methods

Water samples were collected in 10-L Niskin bottles, using 4 bottles per depth. The water was pressure filtered through 0.45 μm cellulose nitrate membrane filters

and collected in separate plastic bins. The filters, containing particulate material, were folded and stored in plastic bags for processing ashore. Dissolved ²²⁶Ra, ²¹⁰Pb, and ²¹⁰Po were preconcentrated from the filtered seawater samples (Towler et al., 1996) after being spiked with yield tracers (¹³³Ba, stable Pb, ²⁰⁸Po) by adsorption onto MnO₂-coated magnetite. The loaded magnetic adsorbent was collected by siphoning the water containing the loaded magnetic adsorbent through two bottles in series, each held between two magnets. The collected, loaded adsorbent was returned to the laboratory ashore for analysis. ²¹⁰Po was spontaneously plated onto an Ag disk that was presented for alpha-spectrometry. The activity of ²¹⁰Pb in the seawater sample was determined from a second measurement after ingrowth of the ²¹⁰Po daughter in solution. Pb recovery was measured by atomic adsorption spectrometry and the appropriate correction made to the ²¹⁰Pb activity. ²²⁶Ra in the residual solution was measured by liquid scintillation counting after successive precipitations with PbSO₄ and BaSO₄ followed by re-solution with EDTA. ²²⁶Ra concentrations were corrected according to the recovery of the ¹³³Ba spike measured by gamma-spectrometry.

The filters containing particulate ²¹⁰Pb and ²¹⁰Po were spiked with stable Pb and ²⁰⁸Po and digested in boiling 2M HCl. The samples were centrifuged and the ²¹⁰Po plated onto an Ag disk from the supernatant solution. ²¹⁰Pb was determined in this solution as described above.

[6] References:

- Towler, P.H. (1995). Naturally occurring radionuclides in the marine environment. PhD thesis, University of Melbourne, Melbourne, Victoria, Australia, 165pp.
- Towler, P.H. and J.D. Smith (1997). Distribution of 226Ra and 210Pb in the mixed layer of the western equatorial Pacific Ocean. *Marine and Freshwater Research* **48**, 371-375.
- Towler, P.H., Smith, J.D., and D.R. Dixon (1996). Magnetic recovery of radium, lead and polonium from seawater samples after preconcentration of manganese dioxide coated magnetite. *Analytica Chimica Acta* **328**, 53-59.

[7] Comments:

A new method (Towler et al., 1996) was used to preconcentrate ²²⁶Ra, ²¹⁰Pb, and ²¹⁰Po from filtered seawater samples. Recovery of dissolved ²¹⁰Po was low, leading to unreliable results. Without information on this shorter-lived radionuclide it was not possible to calculate particulate residence times in finer layers corresponding to the depths sampled.