

On the tracks of man-made radionuclides in the Indonesian Throughflow sediments

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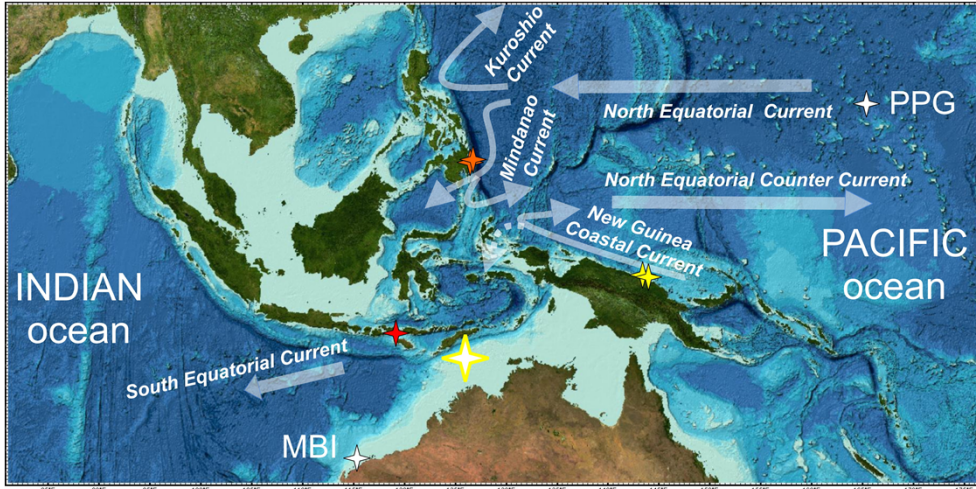


Fig. 1: Map of the study area. PPG = Pacific Proving Grounds, MBI = Monte Bello Islands.

★ Sediment site GeoB 10065

- Non-destructive gamma spectrometry was used for analyzing natural and anthropogenic radionuclides in sediment cores
- Summing spectra of 4-5 following individual samples allowed improving of counting statistics (Fig. 2)
- In individual spectra ¹³⁷Cs was below decision threshold, it could be quantified in summed up intervals (Fig. 2, 3)
- Unusually high concentrations of ²⁴¹Am were observed (Fig. 3)

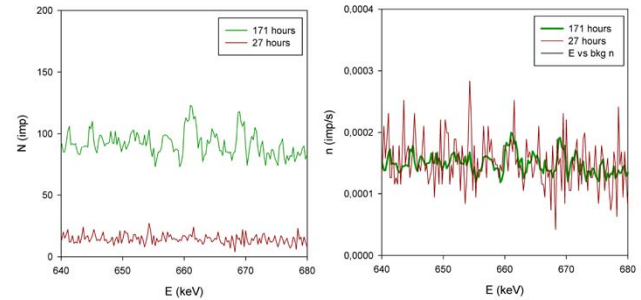


Fig. 2: Close-up of gamma spectra in the area of 662 keV (¹³⁷Cs) before (red) and after (green) summing

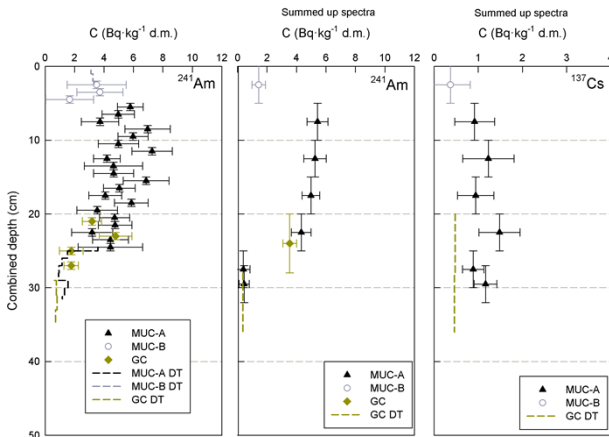


Fig. 3: Left: Depth profile of ²⁴¹Am. Center and right: depth profiles of ²⁴¹Am and ¹³⁷Cs based on summed up intervals.

Outlook

- Contributions of individual sources preserved in the sediment archives will be studied at several ITF locations:
 - ★ the North Pacific inflow portal (the effect of the PPG is expected to be the strongest)
 - ★ the South Pacific inflow portal (the global fallout should dominate)
 - ★ Timor Sea shelf and slope
- Gamma spectrometry: ²⁴¹Am, ¹³⁷Cs
- Alpha spectrometry: ²³⁸Pu, ^{239,240}Pu
- Mass spectrometric methods (AMS, ICP-MS): ²⁴⁰Pu/²³⁹Pu, ²⁴¹Pu

Sources of artificial radionuclides

- The ²⁴¹Am/¹³⁷Cs ratios do not correspond to the global fallout signature (expected value: 0.01)
- High ²⁴¹Am/¹³⁷Cs ratios in sediments were found within 800 km range of the Pacific proving grounds (PPG), however these decreased significantly below 1 in more distant locations in the NW Pacific^[2]
- Possible regional radionuclide contributions:
 - U.S. tests performed at the Pacific Proving Grounds (1946-62)
 - British bomb tests at Australian Monte Bello Islands (1952-56)
 - the SNAP-9A satellite carrying plutonium batteries burnt in the atmosphere (1964)
 - others?
- Each of these sources has an unique isotopic fingerprint

²⁴¹ Am	141 ± 14 Bq·m ⁻²
¹³⁷ Cs	55 ± 10 Bq·m ⁻²
²⁴¹ Am / ¹³⁷ Cs	2.6 ± 0.6

- **Global nuclear weapon fallout:** The testing of nuclear weapons in the atmosphere between 1945 and 1980 (with maximum in 1963) was the most significant source of artificial radionuclides in the environment.
- **¹³⁷Cs:** A major fission product and one of the long lived nuclear fallout radioisotopes (half-life of 30 years) is easily detectable by non-destructive gamma spectrometry.
- **²⁴¹Am:** A less common nuclear fallout radioisotope (half-life of 432 years), present usually as a decay product of ²⁴¹Pu, which is formed during a nuclear explosion by neutron activation.