

Characterization of cesium isotopic composition in offshore seawater in May 2011 after Fukushima nuclear accident

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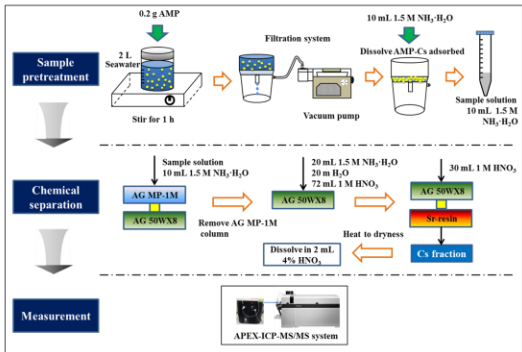
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Introduction

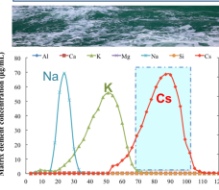
- Since the Fukushima Daiichi nuclear power plant (FDNPP) accident in 2011, the activity ratio of ¹³⁴Cs/¹³⁷Cs has been widely used as a tracer for contamination source identification, for tracing the migration of released ¹³⁷Cs in the North Pacific and study on the mode water evolution. However, due to the short half-life of ¹³⁴Cs (2.06 y), this tracer will become unavailable in the future.
- The ¹³⁵Cs/¹³⁷Cs isotopic ratio can be considered as a new powerful tracer for long-term source identification and environmental behavior studies (Zheng et al., 2014).
- To use Fukushima accident released ¹³⁵Cs as new tracer for tracing the transport of released radionuclides in the Pacific Ocean and other potential oceanography studies, we recently analyzed activities of ¹³⁷Cs and ¹³⁵Cs, and Cs isotopic ratios (¹³⁵Cs/¹³⁷Cs, ¹³⁵Cs/¹³³Cs) in Fukushima offshore seawaters collected in May 2011 using ICP-MS/MS analytical method with 2 L seawater.

Experimental

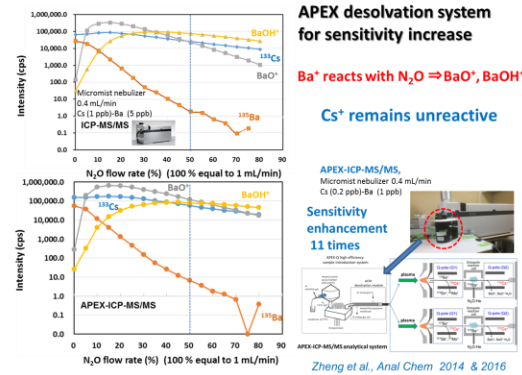
Analytical method for Cs isotope analysis in small volume seawater



成分	化学式	質量%	溶質%
ナトリウムイオン	Na ⁺	1.0556	30.61
マグネシウムイオン	Mg ²⁺	0.1272	3.69
カルシウムイオン	Ca ²⁺	0.0400	1.16
カリウムイオン	K ⁺	0.0380	1.10
ストロンチウムイオン	Sr ²⁺	0.0008	0.03
塩化イオン	Cl ⁻	1.8980	55.05
硫酸イオン	SO ₄ ²⁻	0.2649	7.68
炭酸水素イオン	HCO ₃ ⁻	0.0140	0.41
フッ化物イオン	F ⁻	0.0001	0.003
水素量	H ₂ O	0.0026	0.07



Chemical resolution of Cs from interfering Ba in ICP-MS/MS



With 2 L seawater, the developed APEX-Q/ICP-MS/MS analytical system can measure accurately ¹³⁷Cs activity at > 0.1 Bq/L level, and ¹³⁵Cs concentration at > 0.05 pg/L. The developed method was validated by the analysis of IAEA-443 Seawater reference materials.

Results and discussion

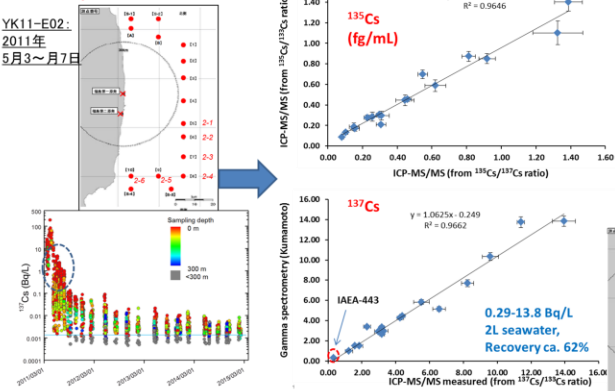
YK11-E02 2 L seawater, ¹³⁵Cs, ¹³⁷Cs analysis, in process

30-60 km off the FDNPP

ICP-MS/MS analysis of ¹³⁵Cs and ¹³⁷Cs

Preliminary results

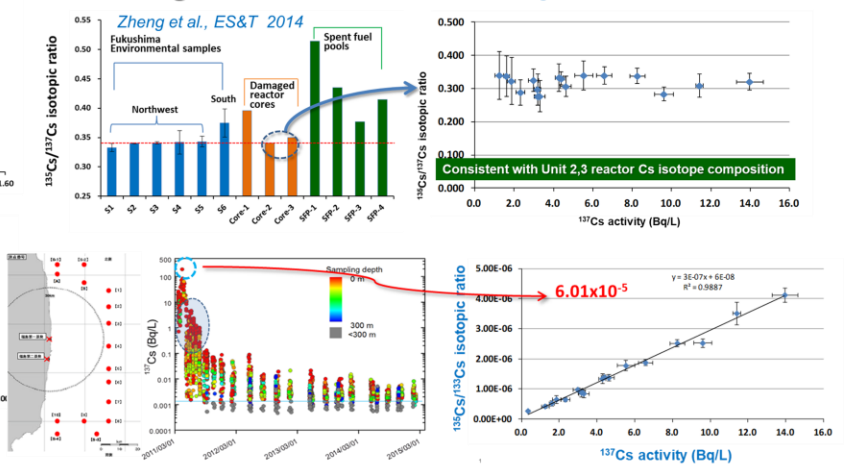
YK11-E02: 2011年 5月3~7日



Quantification of ¹³⁷Cs and ¹³⁵Cs concentration activities can be obtained using ¹³³Cs as yield tracer

Cs isotope fingerprint in seawater (2011 May): long-term tracer

1.64 – 2.30 kg ¹³⁵Cs released into ocean



Conclusion: The analysis of radioCs isotopes in the source term seawater after the accident will provide an opportunity to re-construct the distributions of released radioCs isotopes in the coastal waters off Fukushima. The obtained data will be of benefit to verify the developed models for better understanding the dispersion/migration of accident-released radionuclides in the NW Pacific Ocean. It can be expected that ¹³⁵Cs, as a new geochemical tracer, can be widely applied in oceanography study in near future.