Concentration Coefficients of Radioiodine in Different Chemical Forms from Seawater to Fishery Products

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Abstract

Radioiodine takes various chemical forms in the environment. Ion forms of both I⁻ and IO₃⁻ were found in the ocean for ¹²⁹I discharged from the first Japanese commercial nuclear fuel reprocessing plant located in Rokkasho. Since the concentration factor of iodine from seawater to marine products strongly depends on the chemical form of iodine, it is necessary to use the concentration factor of each chemical form of iodine for realistic assessment of radiation dose from the discharged radioiodine via marine products. This study aims to establish the concentration factor of radioiodine in I⁻ and IO₃⁻ for marine products (seaweed, shellfish and benthos). In FY 2013, the concentration coefficients of I⁻ and IO₃⁻ for brown algae (*Sargassum horneri*) were measured using an iodine radiotracer and the chemical form of stable iodine in *S. horneri* was analyzed by using X-ray absorption fine structure (XAFS) analysis.

The concentration factor of I^{-} or IO_{3}^{-} for *S. horneri* was measured with ¹²⁵I radiotracer. The seaweed samples were incubated for 7 d in the seawater with added ¹²⁵I⁻ or ¹²⁵IO₃⁻. The biological activity of the seaweed samples was checked by their ¹³C absorption. The total ¹²⁵I and ¹²⁵I⁻ concentrations in seawater were monitored during the incubation. The difference between the total ¹²⁵I and ¹²⁵I⁻ concentrations was assumed as ¹²⁵IO₃⁻ concentration. Iodine-125 concentration in *S. horneri* was almost saturated after 3 d exposure. The rate constant of ¹²⁵I⁻ or ¹²⁵IO₃⁻ transfer from seawater into *S. horneri* and the transfer constant of ¹²⁵I from *S. horneri* into seawater were obtained by fitting experimental data to a compartment model with three compartments: ¹²⁵I in *S. horneri*, ¹²⁵I⁻ and ¹²⁵IO₃⁻ in seawater. Organic ¹²⁵I, which was possibly synthesized by the seaweed, was included in the ¹²⁵IO₃⁻ fraction in seawater but neglected here. However, the fitting result showed that organic ¹²⁵I should be considered to obtain an accurate rate constant.

The analysis of chemical forms of stable iodine in *S. horneri* samples by XAFS showed that most of the iodine in *S. horneri* was in organic iodine. A sample of *S. horneri* incubated in seawater with added Γ (iodine concentration of 2.5 mg L⁻¹), was also analyzed and no significantly different chemical form from the natural one was found.



Fig. 1 Measured ¹²⁵I concentration in *S. horneri* and seawater after adding ¹²⁵I⁻ to the cultivation system. Solid circles show ¹²⁵I concentration in *S. horneri*, and solid squares and triangles show concentrations of ¹²⁵I⁻ and ¹²⁵IO₃⁻ in seawater, respectively.



Fig. 2 Measured iodine concentrations in *S. horneri* and seawater after adding ¹²⁵IO₃⁻ to the cultivation system. Solid circles show radioiodine concentration in *S. horneri*, and solid squares and triangles show concentrations of ¹²⁵I⁻ and ¹²⁵IO₃⁻ in seawater, respectively.



Fig. 3 Measured XAFS spectra of NaIO₃, NaI, indophenol, *S. horneri* incubated in seawater with elevated I⁻ concentration and natural *S. horneri*.