ASSIMILATION AND RETENTION OF HEAVY METALS AND RADIONUCLIDES IN SEASTARS

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From both ecological and toxicological viewpoints, it is important to understand the transfer and cycling of trace elements and contaminant heavy metals and radionuclides through marine food chains. Studying these aspects in the natural system or using elevated concentrations of the target elements under simulated conditions usually entails greatly perturbing the elemental composition of the water, subsampling from a large population of organisms, and carrying out lengthy chemical separations for eventual elemental and/or radionuclide analyses. The use of high specific activity or carrier-free gamma-emitting radiotracers of these elements circumvents these problems and allows rapid radioanalyses of live organisms or tissues which have been exposed to the contaminants at concentrations more likely to be present in the surrounding waters.

tissues which have been exposed to the contaminants at concentrations more likely to be present in the surrounding waters. For our studies, we have developed a multi-isotope analytical technique which allows measuring simultaneously seven gamma-emitting radiotracers in the same experimental organisms. Use of this multi-tracer technique reduces inter-experimental variation which occurs between separate treatments labelled with single radioisotopes. This report summarizes results from laboratory radiotracer experiments aimed at quantifying the assimilation and retention of some key heavy metals and radionuclides in carnivorous seastars following contaminant transfer via a twical three step food chain (phytoplawthen a binalwe seastar)

typical three-step food chain (phytoplankton - bivalve - seastar). Carrier-free or high specific activity solutions of the gamma emitters ¹⁰⁹Cd, ⁶⁵Zn, ¹¹⁰mAg, ⁶⁰Co (inorganic) and ⁵⁷Co (cobalamine) were employed in all experiments. In addition, two radionuclides of current interest, ²⁴¹Am and ¹³⁴Cs, were also used in In addition, two radionuclides of current interest, ²⁴¹Am and ¹⁵⁴Cs, were also used in the multi-isotope mixture. Mussels (*Mytilus edulis*) were labelled for 96 hours in sea water containing a suspension of phytoplankton cells (*Isochrysis galbana*, 5x10³ cells/ml) and radiotracers of the selected elements. During this period, the labelling medium was changed every 24 hours. Following the contamination period, the mussels were rinsed and their soft parts removed and counted for radionuclide content. The mussel soft parts were then fed to asteroids (*Marthasterias glacialis*) which promptly ingested the food ration. After radio-labelled feeding, the seastars were periodically whole body counted live over the next several weeks in order to assess excretion rates for the different elements. General experimental protocols, radio-labelling techniques and whole body gamma spectrometric analyses (GeLi

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Element/	Ag	Cd	Co(in.)	Co(org.)	Zn	^{134,137} C	²⁴¹ Am
RN						S	
Tb½ (days)	57	47	40	53	101	78	44

Table 1. Biological half-lives (Tb1/2) of selected elements and radionuclides in th Marthasterias glacialis following a single ingestion of radiolabelled food. in the seastar

Other noteworthy features were the enhanced retention of organic Co over the inorganic form (Fig. 1) and the stronger retention of the monovalent radiocaesium compared to trivalent ²⁴¹Am. This latter observation merits further investigation particularly in view of the many studies which report longer biological half-lives for ²⁴¹Am than radiocaesium in marine organisms.

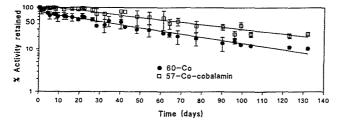


Fig. 1.Long-term excretion of inorganic ⁶⁰Co and ⁵⁷Co²cobalamine in seastars following a single ingestion of radiolabelled food. T = 15±1°C; S = 37‰; Bars =±1σ.

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