137Cs and 210Po in Pacific Walrus and Bearded Seal from St. Lawrence Island, Alaska

T. F. Hamilton, D. J. Seagars, T. Jokela, D. Layton

April 8, 2005

Marine Pollution Bulletin
DISCLAIMER

This document was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor the University of California nor any of their employees, makes any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or the University of California, and shall not be used for advertising or product endorsement purposes.
137Cs and 210Po in Pacific Walrus and Bearded Seal from St. Lawrence Island, Alaska

Terry Hamilton1,∗, Dana Seagars2, Terry Jokela1, and David Layton1

1Environmental Science Division, Lawrence Livermore National Laboratory, P.O. Box 808, Livermore, CA 94550-0808, USA
2U.S. Fish and Wildlife Service, Marine Mammals Management Field Office, 1011 East Tudor Road, Anchorage, AK 99503, USA

Abstract

The activity concentration of Cesium-137 (137Cs) and naturally-occurring Polonium-210 (210Po) were measured in the muscle tissue, kidney and liver of Pacific walrus (Odobenus rosmarus divergens) and bearded seal (Erignathus barbatus) collected by native hunters from the Bering Sea. The mean 137Cs concentrations in muscle, liver and kidney of Pacific walrus were 0.07, 0.09 and 0.07 Bq kg\(^{-1}\) (N= 5, wet weight), respectively, and 0.17, 0.10, and 0.17 Bq kg\(^{-1}\) (N=2, wet weight), respectively, in bearded seal. In general, 137Cs tissue concentrations are significantly lower than those previously reported for mammals from other regions. By comparison, 210Po activity concentrations appear to be higher than those reported elsewhere but a larger variation. The mean 210Po concentration in the muscle tissue, liver and kidney of Pacific walrus (N=5, wet weight) were 28.7, 189, and 174 Bq kg\(^{-1}\), respectively. This compares with 210Po concentration values (N=2, wet weight) of 27, 207, and 68 Bq kg\(^{-1}\) measured in the muscle tissue, liver and kidney, of bearded seal, respectively. Estimated bioaccumulation factors—as defined by the radionuclide concentration ratio between the target tissue to that in sea water—were two to three orders of magnitude higher for 210Po that those of 137Cs. We conclude from radiological dose estimates that ingestion of 137Cs in foods derived from walrus and seal will pose no threat to human health. This work has important implications for assessing health risks to Alaskan coastal communities concerned about the dumping of nuclear waste in the Russia Arctic.

Keywords: 137Cs, 210Po, Bering Sea, marine mammals

∗ Corresponding author, e-mail: hamilton18@llnl.gov
Introduction

Cesium-137 (\(^{137}\text{Cs}\)) is one of the most abundant anthropogenic radionuclides produced during atmospheric nuclear weapons testing programs (1945-81) and deposited into the world’s ocean (Hamilton, 2004). \(^{137}\text{Cs}\) also forms an important constituent of nuclear waste inventories dumped in the Kara Sea and the North-Western Pacific Ocean (Layton et al., 1997a) and at contaminated sites in the North Siberian Basin (Bradley and Jenquin, 1995). Moreover, radioactive contaminants derived from Russian Arctic regions may be transported along the Siberian continental shelf and slope, and eventually impact on North American waters. The focus of this study was to provide high quality baseline data on radionuclide concentrations in marine mammals as a basis for conducting a risk assessment on Alaskan coastal communities exposed to radionuclides in subsistence diets. Potential dietary exposures to \(^{137}\text{Cs}\) have also been compared with those derived from polonium-210 (\(^{210}\text{Po}\))—a naturally occurring radionuclide and an important contributor to the dose to man through consumption of marine foods (Pentreath and Alington, 1988). The concentration of any contaminant in a marine organism reflects the sum of all bioaccumulation, metabolic, and elimination processes affecting the transfer of the contaminant to a whole organism and/or target tissue. The main source of \(^{137}\text{Cs}\) in the Pacific Ocean can be attributed to worldwide fallout deposition from atmospheric nuclear weapons testing (Hamilton, 2004). A large fraction of the \(^{137}\text{Cs}\) oceanic deposit has been retrained in surface waters with latitudinal and/or localized inputs becoming more obscure over time (Hamilton et al., 1996). Consequently, marine mammals living within a defined region such as the Bering Sea have been exposed to near steady-state \(^{137}\text{Cs}\) conditions (food + water) from which field-derived radionuclide concentration factors (CF) may be calculated. Similarly, field-derived CF values can be estimated for \(^{210}\text{Po}\) and/or other radionuclides, and the results used to determine the criticality of different radionuclides and/or exposure pathways in contributing to the risk to populations from potential transfers or releases of radionuclides into the marine environment. Over the past decade there has been a large international effort under the Arctic Monitoring and Assessment Programme (AMAP) to assess the state of the Arctic environment with respect to pollution issues (AMAP, 1998a) including those pertaining to dumping of nuclear waste and other sources of marine radioactivity in the Arctic Ocean. Samples of fish, seals and whales collected in Greenland and Icelandic waters have been analyzed for \(^{137}\text{Cs}\) since the early 1960s (Rissanen et al., 1997; AMAP, 1998b) but precise measurements of radionuclide concentrations in tissues of marine mammals from the Arctic regions of North America are relatively few. Moreover, recommended CF values used in radiological assessments are often based on extrapolated average values taken over broadly characterized groups (e.g., mammals, fish, mollusks, etc.) and/or geographical regions, and may not be directly applicable to coastal communities of Alaska.

The animals used in this study—five individuals of Pacific Walrus (\(\text{Odobenus rosmarus divergens}\)) and two individuals of Bearded Seal (\(\text{Erignathus barbatus}\))—were caught by Alaskan hunters for food. Pacific walrus primarily consume bivalve mollusks, although they are known to feed on a wide variety of benthic and epibenthic invertebrates such as gastropods, polychaetes, and molting crabs; also demersal fishes and other pinnipeds may be consumed (Fay et al., 1977; Fay, 1982; Fay and Stoker, 1982a,b; Lowry et al., 1980; Fay et al., 1984). Bearded seal eat a diversity of epifaunal and infaunal invertebrates such as crabs, shrimp, clams, and snails, as well as demersal and pelagic fish such as capelin (\(\text{Mallotus villosus}\)), Arctic cod (\(\text{Boreogadus saida}\)), and scuplins (Burns, 1967; Lowry et al., 1979; Finely and Evans, 1983; Antonelis et al., 1994). Therefore, dietary habits of bearded seal and Pacific walrus alone...
may lead to real differences in the body burden of radionuclides between the two species, and from those of more pelagic and piscivorous feeding marine mammals. Furthermore, food consumption of seal and walrus (and thus, exposure to a particular contaminant) in the Bering and Chukchi Seas varies with their seasonal distribution (Burns and Frost, 1979; Lowry et al., 1980; Fay et al., 1984). Both species move throughout the relatively shallow (<100 m) waters of the Beringian platform as the pack ice advances and recedes each season (Fay, 1982; Fedoseev, 1990) but utilize different forms and types of ice (Fay 1974; Braham et al., 1984). With the break-up and retreat of the pack ice in spring, the majority of the Pacific walrus population migrates from the Bering Sea into the Chukchi Sea, returning in the fall to over-winter. About 25-50% of the sub-adult and adult male walrus forego the northward migration, remaining on summer haul-out sites along the Russian Bering Sea coast and in Bristol Bay, Alaska (Gilbert et al., 1992). Bearded seals are commonly found along the southern extent of the pack ice in the Bering Sea during winter, and through spring into summer follow the southern margin of the retreating pack ice in the northern Chukchi Sea. The authors recognize that the value of this study is limited by the small population size. However, the high quality of the data coupled with our knowledge of the feeding habits and seasonal distributions of mammals in the Bering and Chukchi Seas do provide a meaningful basis for evaluating trends in levels of radionuclide contaminations between species, and in providing reliable predictions about potential impacts of radionuclide releases on coastal communities concerned about exposures to radioactive contaminants associated with the dumping of nuclear waste in the Arctic Seas.

Sections of muscle, liver and kidney from each animal were collected under the careful supervision of personnel from the U.S. Fish and Wildlife Service—Marine Mammals Management Field Office and the samples transported to the Lawrence Livermore National Laboratory (LLNL) for analysis. Here we present the results of $^{137}$Cs and $^{210}$Po analyses on muscle tissue, liver and kidney. Our data are used to estimate field derived Concentration Factor (CF) values—as defined by the radionuclide concentration ratio between the whole organism or target tissue to that in seawater—and the results compared with those CF values used in dose assessment models. Consequently, our baseline CF values could be used with predicted concentrations of radionuclides in North American waters to provide a better insight into the potential health and ecological impacts of radionuclide releases in the Russian Arctic on Alaskan coastal communities.

Materials and Methods

Sample Collection & Storage

Tissue sections were collected from Pacific walrus and bearded seal during May 1996. Alaskan hunters caught the animals by off St. Lawrence Island in the Bering Sea (Fig. 1) as part of their normal subsistence hunting practices. Meristics, sex, morphological as well as other data relevant to the health and maturity of walrus and seals were collected within minutes of post-mortem. Tissues were sampled within 30 minutes of death by working cooperatively with native hunters as they butchered animals for meat. Up to 10 kg of muscle tissue, and 1-4 kg chunks of kidney, liver, and "coak" (skin with some underlining muscle/fat) were collected for radionuclide measurements, and each sample immediately placed in labeled plastic bags. Sample collection was standardized as follows: muscle—trepesius and latissmus groups, kidney—whole (bearded seal, most walrus) or a nearly whole large chunk, liver—large section taken from the central lobe, and coak—sternal region between anterior and posterior margins of the fore flippers. Samples were held in a cooler at near freezing temperature during transport (ca. 2-4 hours) from sea ice to field quarters. Frozen samples
were subsequently shipped to the Lawrence Livermore National Laboratory (LLNL) for immediate analysis. Tissue samples (~1 cm³) for subsequent histological examination were also collected within 30 minutes post-mortem, and placed in 10% buffered formalin; analysis was conducted by the Armed Forces Institute of Pathology (Department of Veterinary Pathology, Washington, D.C.). For walrus, two canine teeth were collected for subsequent laboratory age determination following the procedures described in Seagars et al., (1995) and Fay et al., (1986). For bearded seals, standard lengths were taken and the lower jaw (for teeth) and claws were collected where possible for subsequent age determination following procedures of Bengston and Siniff (1981), Burns and Frost (1979), Burns (1967), Scheffer (1967), Laws (1962), and McLaren (1958).

At LLNL, the tissue samples were partially thawed and either coarsely ground in a meat mincer or diced into small pieces. The samples were then packed in plastic cartons, the sample cartons placed in a large capacity freeze drying unit, and the material dried to a constant weight over a period of 5 to 6 weeks. Each sample was then reconstituted into a single bulk sample and homogenized in a laboratory blender.

**Determination of ⁴¹⁷ Cs**

Up to 2 kg of dry, homogenized material was used for each analysis. Each sample was dry ashed at 450°C for 4-5 days, and then leached with concentrated HNO₃ at 70°C for 3-4 hours. Any remaining insoluble material was removed by filtration. Ten mg of stable Cs was then added to the sample leachate (as a chemical yield tracer) and ⁴¹⁷ Cs (Cs) separated from the bulk solution by batch extraction onto 1-2 g of the microcrystalline cation exchanger, ammonium molybdophosphate (AMP), at pH 2-3. The precipitate was then recovered by centrifugation and dissolved in a minimum quantity of 10 M NaOH to which an additional 10 ml of 2% EDTA-0.75 M NaOH solution was added. This solution was then passed through a cation exchange column containing 20 ml of BioRex-40 cation exchanger (20-40 mesh) preconditioned with 100 ml of 3 M HCl, 150 ml of 5% NaCl solution, and 50 ml of distilled water. The ion-exchange column was allowed to drain, washed with 60-70 ml of distilled water followed by 160 ml of 0.75 M HCl to remove any potassium (K) and/or rubidium (Rb). Cs was eluted from the column using 125 ml of 3 M HCl and the eluant evaporated to dryness on a hotplate. The residue was dissolved in 2-4 ml of water containing 1-2 drops of 8 M HNO₃, and the solution carefully transferred to a clean 50 ml C-tube using an additional 4-6 ml of distilled water. A 1 ml aliquot of 10 M NaOH was then added to the C-tube, the solution chilled in an ice-bath and Cs precipitated by drop wise addition of 2 ml of 0.12 M chloroplatinic acid (H₂PtCl₆). The precipitate (Cs₂PtCl₆) was collected on a pre-weighed GF/A filter (Whatman, 25 mm), rinsed with 2-3 ml of water, followed by 2-3 ml of acetone, and dried under an I.R. heat lamp to constant weight (±0.01 mg). The sample filters along with associated blanks and standards were all mounted on a standard ring/disc assembly, covered with a mylar film, and counted on a low background gas proportional beta counter (G5000 alpha/beta counter, Gamma Products, Inc.). The Minimum Detectable Activity (MDA) for this technique, as determined on a series of 12 reagent blank analyses, was approximately 0.01 Bq. Calibrations were performed using a standard ⁴¹⁷ Cs solution that was indirectly traceable to the United States National Institute of Standards and Technology (NIST). The total uncertainty of the measurements was propagated from the sample counting error (taken from the standard deviation between five consecutive sample counts), the background counting error, and estimates of uncertainties associated with instrument calibration and gravimetric determination of the chemical recovery. Overall chemical recoveries varied between 48 and 87%.
**Determination of $^{210}$Po**

A 1-10 g aliquot of dried sample material was spiked with $^{209}$Po tracer and 10 mg of stable Pb carrier, and dissolved in a mixture of HNO$_3$-HCIO$_4$ under reflux. The sample residue was converted to a chloride form by repeated evaporation with 6 M HCl, and finally dissolved in 35-40 ml of 0.5 M HCl using gentle heat. The solution was heated to 75-85°C, a few hundred milligrams of ascorbic acid added to reduce any iron (Fe) present, and Po spontaneously deposited onto the surface of a spinning silver (Ag) disc (Hamilton & Smith, 1986). $^{210}$Po and $^{209}$Po were measured by alpha-spectrometry using surface barrier detectors. The total uncertainty of the measurements was propagated from the sample and background counting errors, and uncertainty in calibration and addition of the $^{209}$Po tracer spike.

**Results**

**Morphological Data**

Tissue samples were obtained from 3 male and 2 female adult Pacific walrus, and 1 male and 1 female bearded seal (Table 1). A recently born calf accompanied both female walrus. Available reports of walrus histological samples revealed no clinically significant lesions (animals G96-0005, G96-0006, and G96-0009, T. Lipscomb, per comm., 1997). Based on physical examination and age-length relationships (Burns, 1967) both the seals were young adults. More exact determination of seal age based on layers in the teeth and claw awaits further analysis. A calf accompanied the female seal. While a visual examination indicated no obvious adverse health conditions, the seals were at the lower bound of average blubber thickness for the late spring period (after Burns and Frost, 1979). Histological material from the female found commonly occurring nematode *Parafilaroides sp.* present in the lungs, mild subacute hepatitis, and hepatic capsular fibrosis; none of which were considered to have seriously comprised the health of the animal (T. Lipscomb, per comm., 1997).

$^{137}$Cs and $^{210}$Po Activity Concentrations in Tissues of Walrus and Seal

The activity concentration of $^{137}$Cs and $^{210}$Po in muscle, liver and kidney of Pacific walrus and bearded seal are shown in Table 2a&b. Decay corrections for $^{210}$Po were based on a supported $^{210}$Pb activity concentration of 1 Bq kg$^{-1}$ (dry wt.). The majority of samples were processed within about 120 days of collection. The dry tissue weights (after lyophilization) to wet tissue weight ratios have been used to calculate radionuclide concentrations and results reported on a wet weight basis (Bq kg$^{-1}$, wet wt.). The mean activity concentrations of $^{137}$Cs in tissues of Pacific walrus were 0.07 Bq kg$^{-1}$ in muscle, 0.09 Bq kg$^{-1}$ in liver, and 0.07 Bq kg$^{-1}$ in kidney (Table 2a). Similarly, mean $^{137}$Cs activity concentrations in tissues of bearded seal were 0.17 Bq kg$^{-1}$ in muscle, 0.10 Bq kg$^{-1}$ in liver, and 0.18 Bq kg$^{-1}$ in kidney. The activity concentrations of $^{210}$Po in tissues of Pacific walrus and bearded seal are considerably higher and span a much broader range compared with levels of $^{137}$Cs (Table 2b). The mean (and range) of $^{210}$Po concentrations in the muscle tissue, liver and kidney of Pacific walrus were 28.7 Bq kg$^{-1}$ (12 to 58 Bq kg$^{-1}$), 189 Bq kg$^{-1}$ (70 to 428 Bq kg$^{-1}$), and 174 Bq kg$^{-1}$ (101 to 276 Bq kg$^{-1}$), respectively. The mean (and range) of $^{210}$Po concentration in muscle tissue, liver and kidney of bearded seal were 27 Bq kg$^{-1}$ (25.5-28.7 Bq kg$^{-1}$), 207 Bq kg$^{-1}$ (185-228 Bq kg$^{-1}$) and 68 Bq kg$^{-1}$ (23-113 Bq kg$^{-1}$), respectively.

**Discussion**

There has been widespread concern over the possible health and environmental impacts of radioactive waste dumped in the Kara and Barents Seas and/or held in wastewater storage ponds and reservoirs in the West Siberian Basin (Yablokov et al., 1993;
Strand & Cooke, 1995; Layton et al., 1997a). The $^{137}$Cs concentration in tissues of Pacific walrus and bearded seal from the Bering Sea are generally lower than previously reported concentrations in the muscle tissue of mammals from other regions (Roos et al., 1992; Dahlgaard, per. comm., 1996). Roos et al. (1992) published some comparable baseline data for different species of seal collected from Antarctic waters but these data are typically reported as ‘not detected’ or with large uncertainties. The measured $^{137}$Cs concentration in muscle tissue from a single sample of crab eater seal was $2.5 \pm 0.8$ Bq kg$^{-1}$, dry weight. This compares with a mean average dry weight $^{137}$Cs concentration in Bearded Seal collected from the Bering Sea of $0.63 \pm 0.03$ Bq kg$^{-1}$ (recalculated from Table 2A). The Risø National Laboratory in Denmark has been measuring $^{137}$Cs concentrations in mammals (largely whale and seal) from Greenland since the early 1960s (Dahlgaard, per. comm., 1996). During the period between 1982 and 1991, the $^{137}$Cs concentration in muscle tissue averaged 0.4 Bq kg$^{-1}$, wet weight (N=19). The combined average $^{137}$Cs concentration in muscle tissue of mammals collected from the Bering Sea during 1996 was 0.1 Bq kg$^{-1}$ (wet wt., N=7) or a factor of 2-6 times lower concentration. Similarly, grey seal ($Halichoerus grypus$) collected from the North Sea and north-east Atlantic Ocean during 1987 appear to have contained much higher levels of $^{137}$Cs with mean concentrations in muscle and liver ranging between 14.3 and 27.5 Bq kg$^{-1}$, and 6.4 and 14.6 Bq kg$^{-1}$, respectively (Anderson et al., 1990). These findings were not unexpected because of localized inputs of radioesium into the marine environment from transport of radioactive waste from European reprocessing facilities, riverine discharges into the Arctic Seas from large Siberian rivers, dumping of radioactive waste and inputs from the Chernobyl accident (Hamilton, 2004).

The concentration of $^{137}$Cs in muscle tissue, kidney, and to a lesser extent, in liver of bearded seal appear to be lower than those concentrations measured in comparison tissues of Pacific walrus. The mean $^{137}$Cs concentration in the muscle tissue of bearded seal $(0.17 \pm 0.03$ Bq kg$^{-1}$, wet wt.) was more than double that observed in Pacific walrus $(0.07 \pm 0.02$ Bq kg$^{-1}$, wet wt.). The observed differences in radionuclide concentrations may be related to the small sample size, the age or sex of the animals, different rates of deposition and elimination from tissues, food habits, and/or different seasonal patterns of distribution. We assume that walrus and bearded seal acquire the subject elements through ingestion of prey. While bearded seal and Pacific walrus feed predominately on benthic invertebrates and there is a strong dietary overlap (in particular for the clam, $Serripes greonlandicus$) bearded seals are euryphagous and opportunistic feeders while walrus are stenophagous and specialized for infaunal feeding. Therefore, dietary habits coupled with differences in seasonal distributions of walrus and seal in the Bering and Chukchi Seas may well lead to real differences in $^{137}$Cs body burdens between the two species.

In general, the activity concentrations of naturally occurring $^{210}$Po in walrus and seal are between 2 and 4 orders of magnitude higher than those of $^{137}$Cs. The range of $^{210}$Po concentrations observed between species was also much greater than observed for $^{137}$Cs. The mean dry weight $^{210}$Po concentration in the muscle tissue of bearded seal is about $96 \pm 6$ Bq kg$^{-1}$ (N=2) (recalculated from Table 2b). This compares with $^{210}$Po concentrations reported by Roos et al. (1992) of $5 \pm 0.3$, $4.1 \pm 0.6$ and $16 \pm 8$ Bq kg$^{-1}$ dry weight in the muscle tissue of leopard seal, crab eater seal and Weddel seal, respectively, from the Antarctica. Based on this small population size, bearded seal from the Bering Sea appear to contain significantly higher levels of $^{210}$Po compared with other species of seal from the Antarctic.

The enrichment of $^{210}$Po in marine species over that of other radionuclides including the precursor $^{210}$Pb has been well documented (Carvalho, 1988; Cherry and Shannon, 1974). Noshkin et al. (1994), Pentreath et al. (1979) and others have also noted similar large variations in $^{210}$Po concentrations among different species and across the same species.
These differences are probably related to the feeding habits of marine species and the variable 210Po content of any food consumed but, in general, the mechanisms are poorly understood. 210Po has a relatively short half-life (138 days) and any seasonal variations in 210Po uptake by mammals (e.g., through ingestion of different prey or prey containing different concentrations of 210Po) should show in the level of 210Po in tissue samples. Of those pinnipeds living in the Bering and Chukchi Sea, only the bearded seal and Pacific walrus feed predominately on benthic invertebrates (Lowly et al., 1980). It is not unreasonable to assume that 210Po body burdens may be very different to those of piscivorous-feeding and/or mixed-feeding mammals.

An interesting observation is that the highest 210Po concentrations in our samples appear to be associated with females of both species (Table 1 & 2B; specimens G96-0009, G96-0748, and G96-0002). The female walrus we sampled were younger than the males, and, while the total number of samples analyzed is small, these results appear to differ from those of other workers reporting bioaccumulation of trace elements with age in walrus and other Arctic pinnipeds (Warburton & Seagars, 1993; Wagemann & Stewart, 1994; Seagars et al., 1994; Mackey et al., 1996;) and transfer of various contaminants from female pinnipeds to their pups during lactation (Wagemann et al., 1988; Addison & Brodie, 1987; Anderson et al., 1990) or trans-placental transmission to the fetus (Eisler, 1988). In the absence of other data, it is possible that the higher accumulation of 210Po in female walrus results from undetermined sex related or species-specific factors. Diet, age and sex are known to influence the concentration of organochlorines in marine mammals (Addison and Brodie, 1987) and may be equally important for radioactive contaminants. However, a more plausible explanation might well be related to the summer-fall distributions of these groups into the northern regions of the Chukchi Sea and regional differences in the distribution of 210Po.

Field-Derived Concentration Factors

The bioconcentration factors for 137Cs and 210Po in Pacific walrus and bearded seal from the Bering Sea are shown in Table 3. Field-derived CF values have been expressed in m³ kg⁻¹, and as a unitless measure to help facilitate comparisons with data from the literature. CF values for 137Cs and 210Po were estimated using associated seawater concentrations of 2 and 1 Bq m⁻³, respectively (Povinec et al., 1996; Ellis et al., 1995).

The full range of CF values (unitless) for 137Cs are as follows: 24-47 for muscle tissue, 41-49 for liver, and 28-44 for kidney in Pacific walrus, and 71-100 for muscle tissue, 47-57 for liver, and 85-90 for kidney in bearded seal. The mean CF values for muscle tissue in walrus and seal are 36 and 86, respectively. Our data for the muscle tissue of marine mammals from the Bering Sea are in general agreement with a world-wide consensus value of 100 reported by Povinec et al. (1996) and compares with CF values of 100 for fish flesh, and 30 for molluscs and crustaceae, as previously recommended by IAEA (IAEA, 1985).

For 210Po, CF values for each tissue type are very similar across the two species (Table 3). The mean 210Po CF values (unitless) for muscle, liver and kidney tissues were 29,000, 190,000 and 174,000, in Pacific Walrus, and 27,000, 206,000 and 208,000, in bearded seal, respectively. Recommended CF values for different groups of marine organisms in the Arctic Seas were recently reported by Templeton et al., (1997) for use in risk assessments. Based on limited data from Roos et al. (1992), the recommended CF values for mammals are given as 1,700 for muscle tissue and 22,000 for liver. Measured 210Po CF values for Pacific walrus and bearded seal from the Bering Sea are up to an order of magnitude higher for each of these two tissue types.
Exposure to Radionuclides in Subsistence Diets

In evaluating doses to individuals exposed to radionuclides in subsistence diets, it is important to have detailed information on dietary patterns of subject populations as well as accurate radiometric data. Detailed information related to the subsistence diets of Alaskan Natives and/or Alaskan coastal communities were not available for this assessment. However our studies clearly show that the concentration of $^{137}$Cs in tissues of Pacific walrus and bearded seal are several orders of magnitude lower than those of naturally occurring $^{210}$Po. Existing background concentrations in waters of the Chukchi Sea are in order of 2 Bq m$^{-3}$ (Ellis et al., 1995). The highest predicted $^{137}$Cs concentration in seawater resulting from an instantaneous release from nuclear wastes in the Russian Arctic is estimated to be around 0.03 Bq m$^{-3}$ (Layton et al., 1997b) or well below existing background levels. As such, the potential risks associated with ingestion of foods derived from walrus and seal containing $^{137}$Cs derived from Russian nuclear wastes will be extremely low, and pose no threat to human health. Based on the activity concentrations of $^{137}$Cs and $^{210}$Po measured in seal and walrus from the Bering Sea, and using dose conversion factors adopted from the International Commission of Radiological Protection (ICRP) for ingestion of radionuclides, i.e., $1.2 \times 10^{-8}$ Sv Bq$^{-1}$ for $^{137}$Cs (ICRP 30, 1978) and $1.2 \times 10^{-6}$ Sv Bq$^{-1}$ for $^{210}$Po (ICRP 67, 1994), the committed effective dose for adults from intakes of $^{137}$Cs will be insignificant or about 3-4 orders of magnitude lower compared with that from naturally-occurring $^{210}$Po. In evaluating these data, it is important to recognize that unsupported $^{210}$Po has a relatively short half-life ($T_{1/2} = 138$ days), and a decay factor between catch and consumption should be used. More extensive studies will also be needed to evaluate the relatively high $^{210}$Po CF values in mammals from the Bering Sea compared with values reported from other regions. Furthermore, polar bear are known to track directly across the Bering Sea from the Asian continent during winter providing an important source of meat for coastal communities. It would be useful to supplement the data presented here with a more detailed study of the levels and transfer pathways of radionuclides in Alaskan arctic regions, e.g., through consumption of polar bear, whale, and caribou meat. These data could be used to develop a more comprehensive assessment of background radiation doses for critical population groups, and yield more accurate dose estimates for any potential releases and/or transfers of radionuclides into the region.

Conclusion

The focus of this study was to provide high quality baseline data on radionuclide concentrations in marine mammals as a basis for conducting an integrated risk assessment on Alaskan coastal communities exposed to radionuclides in subsistence diets. The activity concentration of $^{137}$Cs and naturally occurring $^{210}$Po were measured in the muscle tissue, kidney and liver of Pacific walrus and bearded seal collected from the Bering Sea. The mean $^{137}$Cs concentrations (wet weight) in tissues of Pacific walrus and bearded seal were 0.07 and 0.17 Bq kg$^{-1}$ in muscle, 0.09 and 0.10 Bq kg$^{-1}$ in liver, and 0.07 and 0.175 Bq kg$^{-1}$ in kidney, respectively. Bioconcentration factors for muscle and kidney tissues in Pacific walrus appear to be significantly lower than for bearded seal, and values for both species are generally lower than those reported for other pinniped species. Observed differences in the concentration of $^{137}$Cs in muscle and kidney tissue between the two species can be explained by differences in prey utilized by these two benthic feeding mammals, and in turn, may be coupled to differences in their seasonal distribution within the Bering and Chukchi Seas. The activity concentrations and associated bioconcentration factors for naturally-occurring $^{210}$Po are much higher than those of $^{137}$Cs. Consequently, the committed effective dose from ingestion of $^{137}$Cs in foods derived from walrus and seal will be insignificant or about 3-4 orders of magnitude lower...
compared with that from $^{210}$Po. Using a worst-case release scenario, we also conclude that any potential risks associated with consumption of marine foods containing $^{137}$Cs derived from the dumping of nuclear wastes will be extremely low and pose no treat to human health. There is, however, a general lack of measurement data and information on the levels and transfer behaviors of radionuclides through Arctic food chains leading to man. A key to more accurate dose estimates will be the availability of high quality radiometric data covering all potential exposure pathways, and detailed dietary information for critical population groups.

**Acknowledgement**

This study formed part of a contribution to the Arctic Nuclear Waste Assessment Program (ANWAP) and was made possible through the cooperation of the U.S. Fish and Wildlife Service (Marine Mammals Management Field Office-AK). We gratefully acknowledge the assistance of boat captains Mr. Tason Nowpowkahok, Mr. Dennis James, Sr., H.V. Slwooko, Jr., Mr. David Angie, and L. Apangoqlook for sample collection. We also thank Marshall Stuart for sample preparation; Jennifer Luna, Patricia Lopez, and Rayla Bradsher for secretarial support, and Lynn Wilder for preparing graphics. Work performed under the auspices of the U.S. Department of Energy by the University of California at the Lawrence Livermore National Laboratory under contract W-7405-Eng-48.

**References**


Arctic Monitoring and Assessment Programme (AMAP) (1998b), AMAP Assessment Report: Arctic Pollution Issues, Chapter 8: Radioactivity, P. Strand et al., (editors), Oslo, Norway, xii + 525-619.


Finely, K.J., & Evans, C.R. 1983. Summer diet of the bearded seal (*Ergiathus barbatus*) in the Canadian high Arctic. *Arctic* 36, 82-89.


Wagemann R., & Stewart R.E.A. 1994. Concentrations of heavy metals and selenium in tissues and some foods of walrus (Odobenus rosmarus rosmarus) from the eastern Canadian Arctic and sub-arctic, and associations between metals, age and gender. Canadian Journal of Fisheries and Aquatic Science 51, 426-436.


Table 1. Sample collection data for Pacific Walrus and Bearded Seal from the Bering Sea

<table>
<thead>
<tr>
<th>USFWS field log #</th>
<th>Collection date</th>
<th>Species type</th>
<th>Animal physiology</th>
</tr>
</thead>
<tbody>
<tr>
<td>G96-0003</td>
<td>10-May-96</td>
<td>Pacific Walrus</td>
<td>Adult male</td>
</tr>
<tr>
<td>G96-0005</td>
<td>11-May-96</td>
<td>Pacific Walrus</td>
<td>Adult male, 322 cm in length</td>
</tr>
<tr>
<td>G96-0006</td>
<td>14-May-96</td>
<td>Pacific Walrus</td>
<td>Adult male, 305 cm in length</td>
</tr>
<tr>
<td>G96-0009</td>
<td>19-May-96</td>
<td>Pacific Walrus</td>
<td>Young adult female, 290 cm in length</td>
</tr>
<tr>
<td>G96-0748</td>
<td>20-May-96</td>
<td>Pacific Walrus</td>
<td>Young adult female, 270 cm in length</td>
</tr>
<tr>
<td>G96-9001</td>
<td>12-May-96</td>
<td>Bearded Seal</td>
<td>Adult male, 215 cm in length</td>
</tr>
<tr>
<td>G96-9002</td>
<td>17-May-96</td>
<td>Bearded Seal</td>
<td>Adult female, 202 cm in length</td>
</tr>
</tbody>
</table>
Table 2. Concentration of $^{137}$Cs and $^{210}$Po in walrus and bearded seal from the Bering Sea

<table>
<thead>
<tr>
<th>USFWS field log #</th>
<th>Muscle Tissue</th>
<th>Liver</th>
<th>Kidney</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Activity Concentration (Bq kg$^{-1}$, wet wt.)</td>
<td>Activity Concentration (Bq kg$^{-1}$, wet wt.)</td>
<td>Activity Concentration (Bq kg$^{-1}$, wet wt.)</td>
</tr>
<tr>
<td></td>
<td>$^{137}$Cs (%)</td>
<td>$^{210}$Po (%)</td>
<td>$^{137}$Cs (%)</td>
</tr>
<tr>
<td>Pacific Walrus (<em>Odobenus rosmarus divergens</em>)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>G96-0003</td>
<td>27.3</td>
<td>0.071±0.004</td>
<td>0.094±0.006</td>
</tr>
<tr>
<td>G96-0005</td>
<td>29</td>
<td>0.047±0.003</td>
<td>0.084±0.003</td>
</tr>
<tr>
<td>G96-0006</td>
<td>26.7</td>
<td>0.066±0.004</td>
<td>0.098±0.004</td>
</tr>
<tr>
<td>G96-0009</td>
<td>29.9</td>
<td>0.080±0.003</td>
<td>0.089±0.003</td>
</tr>
<tr>
<td>G96-0748</td>
<td>29.5</td>
<td>0.095±0.004</td>
<td>0.082±0.003</td>
</tr>
<tr>
<td>Bearded Seal (<em>Erignathus barbatus</em>)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>G96-9001</td>
<td>29.1</td>
<td>0.14±0.01</td>
<td>0.093±0.004</td>
</tr>
<tr>
<td>G96-9002</td>
<td>31.6</td>
<td>0.20±0.01</td>
<td>0.113±0.004</td>
</tr>
</tbody>
</table>

Notes: $^{210}$Po was decay corrected to the date of sampling assuming a supported $^{210}$Pb concentration of 1 Bq kg$^{-1}$ (dry wt.).
### Table 3. Bioconcentration factors for $^{137}$Cs and $^{210}$Po in Pacific Walrus and Bearded Seal from the Bering Sea

<table>
<thead>
<tr>
<th>Radionuclide</th>
<th>Species</th>
<th>Tissue type</th>
<th>Tissue concentration, Bq kg$^{-1}$ (dry wt.)</th>
<th>Bioconcentration factor m$^3$ kg$^{-1}$</th>
<th>Unitless</th>
</tr>
</thead>
<tbody>
<tr>
<td>$^{137}$Cs</td>
<td>Pacific Walrus (<em>Odobenus rosmarus divergens</em>) (N=5)</td>
<td>muscle</td>
<td>0.05-0.1</td>
<td>0.024-0.047</td>
<td>0.036</td>
</tr>
<tr>
<td></td>
<td></td>
<td>liver</td>
<td>0.08-0.1</td>
<td>0.041-0.049</td>
<td>0.045</td>
</tr>
<tr>
<td></td>
<td></td>
<td>kidney</td>
<td>0.06-0.09</td>
<td>0.028-0.044</td>
<td>0.034</td>
</tr>
<tr>
<td></td>
<td>Bearded Seal (<em>Erignathus barbatus</em>) (N=2)</td>
<td>muscle</td>
<td>0.14-0.20</td>
<td>0.071-0.10</td>
<td>0.086</td>
</tr>
<tr>
<td></td>
<td></td>
<td>liver</td>
<td>0.09-0.11</td>
<td>0.047-0.057</td>
<td>0.052</td>
</tr>
<tr>
<td></td>
<td></td>
<td>kidney</td>
<td>0.17-0.18</td>
<td>0.085-0.090</td>
<td>0.087</td>
</tr>
<tr>
<td>$^{210}$Po</td>
<td>Pacific Walrus (<em>Odobenus rosmarus divergens</em>) (N=5)</td>
<td>muscle</td>
<td>13-58</td>
<td>13-58</td>
<td>29</td>
</tr>
<tr>
<td></td>
<td></td>
<td>liver</td>
<td>70-428</td>
<td>70-428</td>
<td>190</td>
</tr>
<tr>
<td></td>
<td></td>
<td>kidney</td>
<td>101-276</td>
<td>101-276</td>
<td>174</td>
</tr>
<tr>
<td></td>
<td>Bearded Seal (<em>Erignathus barbatus</em>) (N=2)</td>
<td>muscle</td>
<td>25-29</td>
<td>25-29</td>
<td>27</td>
</tr>
<tr>
<td></td>
<td></td>
<td>liver</td>
<td>185-228</td>
<td>185-228</td>
<td>206</td>
</tr>
<tr>
<td></td>
<td></td>
<td>kidney</td>
<td>113-142</td>
<td>113-142</td>
<td>208</td>
</tr>
</tbody>
</table>

Notes: Bioconcentration factors for $^{137}$Cs and $^{210}$Po have been calculated using associated seawater concentrations of 2 and 1 Bq m$^{-3}$, respectively.
Fig. 1. Map showing sample locations off St. Lawrence Island in the Bering Sea.