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Distribution and sedimentation of radionuclides in the Barents Sea

by

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Abstract

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Analysis of sediment samples for cesium-137 from a wide range of positions have demonstrated an uneven distribution. The causes, pathways and the possible origin of the radionuclide are discussed. The importance of the sedimentation processes for the distribution pattern is demonstrated. The mean values of the upper centimetre of the bottom sediment are between non detectable and eight becquerel per kilo dry weight.

Introduction

The Chernobyl accident in April 1986 and the following speculations about possible contamination of the marine environment similar to the pronounced contamination observed in mountain lakes and on land in some areas of the Nordic countries, initiated a monitoring programme at the Institute of marine research, IMR, aimed at establishing the degree of contamination in areas important for norwegian fisheries.

The sinking of the nuclear submarine «Komsomolets» in April 1989 and the information about the dumping of radioactive material in the Barents and Kara Seas by the former USSR (Anon, 1993a) increased the public and the political attention to the possible problems of radioactive contamination in the marine environment.

In 1991 we started sampling of sediments, water and biota from the Barents Sea. The programme was later included in IMR's part of the AMAP (Arctic Monitoring and Assessment Programme). We decided, in the first hand, to concentrate our measurements to sediments. The ability of sedimentary particles to remove most of the radionuclides from the water phase is referred to by among others Duursma and Gross (1971) and Meili (1994). As a first approach to determine a possible radioactive contamination of the Barents Sea, analysis of sediments were therefor chosen as most likely to give the best estimate of the actual situation.

Materials and methods

Sediment samples were collected from a set of 102 stations during cruises in the years of 1991, 92 and 93 (fig. 1). Sampling procedure followed the guidelines established for the monitoring in the North Sea (ICES, 1992). At each station three boxcorer shot were taken. The boxcorer have an inner dimension of 30 cm x 30 cm. From each boxcorer approximately 100 cm² of the upper 1 cm was collected. The samples from each station were mixed and stored at - 20 °C until further preparation.



Figure 1. Sediment sampling stations in the Barents Sea and the hydrographic section A, B, C.

In the laboratory the samples were freeze dried, homogenized and filled in 215 ml polyethylene counting boxes before gamma-counting. The weight of the samples varied between 100 g and 200 g. The gamma-counting were performed on a Canberra HpGe-detector with 30 % efficiency, an electric cryostat cooling system and a 10 cm lead shielding. Counting time was approximately 22 hours for each sample.

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To establish the particle composition of the sediments, analysis were done to give the weight percentage of each sample in the following fractions: <63 μ m, 63 - 125 μ m, 125 - 250 μ m, 250 - 500 μ m, 500 - 1000 μ m, 1000 - 2000 μ m and >2000 μ m.

Results and discussion

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Fig. 2 presents the results of our determination of ¹³⁷Cs in sediments as becquerel per kg dry weight (Bq·kg⁻¹ d.w.). The overall mean value of the 102 stations in the Barents Sea and at the west coast of Spitsbergen is 3,2 Bq·kg⁻¹ d.w., ¹³⁴Cs was not detected. The ¹³⁷Cs values varied between <1,0 and 8,6 Bq·kg⁻¹ d.w. with the highest values found close to Spitsbergen.



Figure 2. Cesium-137 values in becquerel per kilo dry weight (Bq·kg⁻¹ d.w.) of the first cm of sediment samples from the Barents Sea.

For comparison values in sediments of the North Sea, as reported in the North Sea Quality Status Report 1993 (Anon, 1993b), were found, in 1987, to be at the highest along the North Sea coast of U.K. with values of 65 Bq·kg⁻¹ d.w. and 6,0 Bq·kg⁻¹ d.w. for ¹³⁷Cs and ¹³⁴Cs respectively. Sediment surface values for ¹³⁷Cs in the German Bight (in 1990) and the Skagerrak were measured up

to 20 and 30 $Bq\cdot kg^{-1}$ d.w. respectively (Anon, 1993b). The Quality Status Report (Anon, 1993b) states that there was a decrease in the sediment values of radiocesium between 1986 and 1991 in the North Sea as a major source of contamination was the Chernobyl fall-out in 1986.

In the Barents Sea the main sources of anthropogenic radioactive contaminants are;

- atmospheric fall-out (including Chernobyl fall-out),
- discharges from nuclear industry in Europe,
- assumed contribution from land based sources in northern Russia,
- dumped radioactive material of the former USSR,
- contribution from the Chernobyl accident via the Baltic.

As can be seen from fig. 2 there is an uneven distribution of ¹³⁷Cs in the sediments of the Barents Sea. Most pronounced are the relatively high values close to Spitsbergen and the low values in the southeastern part of the Barents Sea. From our observations it may be interpreted that of the many sources contributing to the radioactive contamination of the Barents Sea sediments some have to be more important than others.

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Global fall-out originate from the nuclear bomb tests beginning in the 1940s with a pronounced peak period in 1961 and 1962. The major test took place in the northern hemisphere and especially in the Barents Sea area where there was observed an immediately response in the radioactive contamination of fish (Føyn, 1991, Føyn and Sværen, 1995). It may be assumed that the fall-out contribution from the nuclear bomb tests is evenly distributed throughout the Barents Sea. Kautsky (1980) reported from measurement in 1969 only small variations of ¹³⁷Cs in sea water from the whole Barents Sea. However, fall-out on the ice-covered islands i.e. the Spitsbergen islands, Franz Josefs land and Novaya Zemlya will be slowly transported towards the sea by glaciers and entered into the water through the melting of the ice. High ¹³⁷Cs values (>60 Bq·kg⁻¹) in sediments in some Spitsbergen fjords observed last year by Ivanov (pers. com. 1995) may indicate transport via glaciers to the fjords.

The transport of radiocesium, discharged from Sellafield into the Irish Sea, to the Barents Sea is well documented (Kautsky, 1980 and Kautsky, 1988). The transport routes of radiocesium observed by Kautsky (1988) follows the main current systems of the Atlantic current and the Norwegian coastal current, i.e. at the entrance of the Barents Sea one branch is towards east and one moving north towards Spitsbergen, fig. 3. Kautsky (1988) found the highest values (40 - 50 Bq·m⁻³ of $^{134}Cs + ^{137}Cs$) in the coastal current close to Norway as far east as $^{30^{\circ}}$ E in his observations i 1979 and values between 10 and 20 Bq·m⁻³ to the west of Spitsbergen.

From this distribution it can be assumed that the main part of the radiocesium was transported eastward into the Barents Sea. The distribution pattern observed by Kautsky (1988) is in good agreement with the common transport pattern of fish larvae. Why do not our observations of ¹³⁷Cs in the sediments reflect the same pattern ?

The content of radiocesium in the sediments reflects the grain size distribution in the actual sediment samples, and as pointed out by Meili (1994) cesium is often found in the clay fraction of sediments. Fig. 4 shows a plot of the ¹³⁷Cs values against percentage of particles <63 μ m in the individual samples. As can be seen there is an overall increase in ¹³⁷Cs content in the sediments with



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Figure 3. The main pattern of the surface water movements in the Barents Sea.

increasing percentage of particles <63 μ m in the samples. However, some outlayers disturb this picture of which one station with the lowest value of contamination and with >80% of the particles <63 μ m represent the vicinity of the sunken nuclear submarine «Komsomolets» at 1670 m depth. The samples with relatively high contamination and a varying fairly low percentage of particles <63 μ m represent the area around Spitsbergen.

To further elaborate the relation between the ¹³⁷Cs content in the samples and

the particle composition a cluster analysis (STATISTICA, Clusteranalysis, Kmeans clustering) of the grain size data was performed. The program forms



Figure 4. The observed 137 Cs values plotted against the percentage of particles in the sediment samples <63 μ m.



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Figure 5. A cluster plot of the means for each of the three clusters according to the percentage of the samples in the defined particle size fractions.

three clusters with the goal of (1) minimize variability within clusters and (2) maximize variability between clusters to get the most significant ANOVA results. The plot of means for each of the three clusters are given in fig. 5. From the plot it can be seen that cluster no. 1 contains sample with particles mainly <63 μ m, but also a relatively large part of particles >2000 μ m. Cluster no. 2 contains samples with most of the particles in the fraction 125-250 μ m.

Cluster no. 3 contains samples with particles mainly $<63 \mu m$.

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Fig. 6 shows the ¹³⁷Cs values plotted according to how the particle composition of the corresponding samples are related to the three clusters.

- The sediment samples having most of their particles in the fraction <63 μ m (cluster no. 3) have an average ¹³⁷Cs content of 4,0 Bq·kg⁻¹ d.w. The major part of the samples belong to this type.
- The samples with the major part of the particles in the grain size 125-250 μ m and the smallest amount of particles <63 μ m (cluster no. 2) have an average ¹³⁷Cs content of 1,2 Bq·kg⁻¹ d.w. These samples are mostly from the southeastern part of the Barents Sea but also with someone from the area close to Novaya Zemlya.
- The samples with the major part of the particles in the fraction >2000 μ m (cluster no. 1) have an average ¹³⁷Cs content of 3,6 Bq·kg⁻¹ d.w. These samples represent for the most stations around Spitsbergen.



Figure 6. The ¹³⁷Cs values of the sediment samples plotted in relation to the three clusters.

It seems that the samples collected from the area around Spitsbergen do not fit into the general view as expressed by Meili (1994) that small sized particles, i.e. clay, will contain the most of the cesium. As we find the highest values in this area where the sediments are supposed to contain less due to the grain size and also the fact that this area should be less affected according to the normal water transport pattern, other explanations have to be found.

Dahlgaard (1994) summarizes concentration factors (CF) between phytoplankton and seawater for some elements giving a CF for cesium of 20. In comparison Dahlgaard (1994) for other major anthropogenic radionuclides refer to plutonium and americium having CF of 1×10^5 and 2×10^5 respectively while strontium has a CF of 3. This indicate that phytoplankton may play an important role in the contamination process of the sediments.

The positions of the sediment stations around Spitsbergen are located in a very geographically stabile polar front area and according to observations by Rey (pers.com. 1995) in this special regime the phytoplankton may actively act as scavenging elements. An example of his (Rey, pers.com. 1995) observations, from 1984, are presented in fig. 7. The actual hydrographic section, A, B, C, is shown in fig. 1. As can be seen there is an extensive



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Figure 7. Temperature and chlorophyll <u>a</u> distribution in the hydrographic section A, B, C (fig. 1) August 1984 (Rey, pers. com. 1995).

primary production in the area, and high chlorophyll concentrations are found all the way to the bottom. The corresponding temperature profiles show clearly a non stratified water mass which indicate an active transport of phytoplankton down to the bottom. The phytoplankton consists mainly of diatoms and the process is according to Rey (pers.com. 1995) not a normal sedimentation process where only the ungrazed phytoplankton sinks out in a slow sedimentary process which is the normal for other parts of the Barents Sea.

If we assume that this active scavenging process is able to remove the cesium present in the water column throughout the production period and actively deposit the cesium at the bottom, this can explain the higher degree of ¹³⁷Cs contamination in this area compared to the rest of the Barents Sea. The main sources of this contamination are most likely to be the water masses transported from south along the norwegian coast, i.e. discharges from european nuclear industry and from the Chernobyl contribution to the Baltic.

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The branch of the Atlantic and coastal water moving eastward in the Barents Sea is spread and mixed with Arctic water as is indicated on fig. 2. The Barents Sea is partly covered by ice during the winter. As the ice starts to melt in spring and the ice-edge is slowly retreating northwards the melt water flows on top of the more saline water creating favourable conditions for primary production (Rey and Loeng, 1985). This spring bloom is distributed in a fan like manner along the ice edge (polar front) as the ice melts and recedes northwards.

The rich phytoplankton production is heavily grazed by zooplankton (Hassel et *al.*, 1991) and mainly ungrazed phytoplankton together with faecal pellets of zooplankton sinks to the bottom. As this production zone is moving northwards during the melting season the sedimentation and thereby the transfer of radiocesium to the sediments will be more widely distributed. This distribution is in accordance with our findings of ¹³⁷Cs in the sediment of the north-eastern part of the Barents Sea.

The potential of the dumped radioactive waste, mostly in the Kara Sea, and the possible contribution from land based sources in Russia to contaminate the Barents Sea is due to a transport westwards from the Kara Sea to the Barents Sea. The main potential of a pronounced contamination is from the nuclear industrial complexes upstream of the rivers Ob and Yenesei in the northeastern part of the Kara Sea. In addition considerable potential sources of radioactive contamination of the Barents Sea are also in found in the many nuclear submarines berthed in fjords of the Kola peninsula waiting for decommissioning.

Føyn and Nikitin (1994) reported preliminary results from the third norwegianrussian expedition to the Kara Sea and fjords at the east coast of Novaya Zemlya, that contamination from the dumped solid radioactive material was only demonstrated in close vicinity to the dumped material.

Vinje and Kvambekk (1991) estimate that there is an average annual ice transport of 500 km³ from the Kara Sea to the Barents Sea through the strait

between Frans Josefs Land and Novaya Zemlya, in comparison they refer to the Transpolar Ice Drift Stream poring 4000 to 5000 km³ of ice per year into the Greenland Sea. If some of the ice that enters the Barents Sea is frozen in the estuaries of the rivers Ob and Yenesei, there is a potential possibility that this ice may transport sediments frozen into the ice in the estuaries and thereby contribute to the radioactive contamination of the Barents Sea given that the sediments of the two estuaries are contaminated.

A possible transport of contaminated sediments by the ice should be reflected in the ice melting zone where possible sediments contained in the ice will be released. Our data do not allow for any conclusion regarding a possible transport of radionuclides from the actual estuaries and the Kara Sea by ice to the Barents Sea. But it may be assumed that if such sediment transports occur some signals of particular contamination of the sediments in the ice melting zone would be seen.

Conclusion

The sediments of the Barents Sea are far less contaminated by cesium-137 than the North Sea. The observed contamination is unevenly distributed which is in accordance with the particle composition of the sediments, i.e. in sediments with a high percentage of fine particles (<63 μ m) relatively high values of 137Cs are found. However, in the area close to Spitsbergen the highest values of 137Cs contamination are found. It is assumed that the particular primary production taking place in this area, due to the behaviour of the polar front, actively transport phytoplankton to the sediments and thereby scavenge the water for radiocesium throughout the production period.

To interpret a variable distribution of radionuclides in the sediments of a wide area as the Barents Sea, good knowledge of the hydrographic situation and the primary production processes as well are needed.

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