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# AQUATIC ENVIRONMENT MONITORING REPORT



## NUMBER 11

## RADIOACTIVITY IN SURFACE AND COASTAL WATERS OF THE BRITISH ISLES, 1982

G.J. HUNT

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Figure 1. UK nuclear establishments giving rise to principal discharges of liquid radioactive waste.

### 1. Introduction

This report presents the results of the environmental monitoring programme carried out during 1982 by staff of the Directorate of Fisheries Research, Lowestoft. The monitoring programme is part of this Ministry's responsibilities under the Radioactive Substances Act, 1960 (Great Britain - Parliament, 1960). The programme is set up to verify the satisfactory control of liquid radioactive waste discharges to the aquatic environment, and to ensure that the resulting public radiation exposure is within nationallyaccepted limits. The monitoring is independent of similar programmes carried out by nuclear site operators as a condition of their authorisations to discharge radioactive wastes. This report also includes results of monitoring carried out on behalf of departments of the Scottish Office, the Welsh Office, the Department of the Environment for Northern Ireland, the Channel Islands States and the Republic of Ireland. Where appropriate, the information presented is supplemented by results from our extensive programme of research into the behaviour of radioactivity in the aquatic environment.

To set the monitoring results in proper context, liquid radioactive discharges from UK nuclear establishments to the aquatic environment in 1982 are first summarised. Before exposition of the monitoring results, an explanatory section gives details of how the results are presented and interpreted in terms of public radiation exposures.

#### 2. Discharges of radioactive waste

Following the Government's response (Great Britain – Parliament, 1977) to the 6th Report of the Royal Commission on Environmental Pollution (1976), data on radioactive discharges are published annually by the Department of the Environment (DOE). Data for 1982 have been published (DOE, 1983) but to enable the results of environmental monitoring presented in this report to be considered readily in the context of relevant discharges, a summary is included here.

#### 2.1 Liquid radioactive waste

Table 1 lists the principal discharges of liquid radioactive waste from UK nuclear establishments during 1982. The locations of these establishments are shown in Figure 1. Discharge data are derived from the operators' returns. Table 1 also lists the discharge limits which are authorised or, in the case of Crown establishments, administratively agreed. Discharges are given both in terabecquerels (see Section 3.1) and curies. The limits are given in the units specified in the relevant authorisation. In some cases, the authorisations specify limits in greater detail than can be summarised in a single table: in particular, where periods shorter than one year are specified the annual equivalent has been used. The limits are lower (often very much lower) than the activities which could be released without exceeding the dose limits recommended by the International Commission on Radiological Protection (ICRP), embodied in national policy (Great Britain – Parliament, 1982). For each discharge the percentage of the authorised (or agreed) limit taken up in 1982 is also stated in Table 1.

For completeness, data are included here on the very small discharges into Holy Loch from the US Navy Submarine Base. Radiological safety for the Holy Loch base is the responsibility of the US Navy in association with the Ministry of Defence who have supplied the following information. For the year 1982 the radioactivity released into the waters of Holy Loch was less than 0.04 GBq (1 mCi) of long-lived gamma radioactivity, primarily cobalt-60; less than 0.04 GBq (1 mCi) of fission product radionuclides; and less than 0.4 GBq (10 mCi) of tritium.

#### 2.2 Solid radioactive waste

In addition to receiving most of the above liquid discharges the marine environment also receives low specific activity packaged solid waste. This is disposed of not in coastal waters but in an area of the deep Atlantic Ocean. Solid radioactive waste from some other West European countries is also disposed of in the same area. The current disposals conform to the requirements of the Convention on the Prevention of Marine Pollution by Dumping of Wastes and Other Matter (The London Convention) and are organised within the Multilateral Consultation and Surveillance Mechanism operated by the Nuclear Energy Agency (NEA) of the Organisation for Economic Cooperation and Development (OECD). This Mechanism makes provision for consultation between member states before a disposal operation takes place, and the operation itself is subject to surveillance by a representative of the OECD(NEA). In 1982 the waste was disposed of within an area defined by the two lines of longitude 16°W and 17°30W and two lines of latitude 10 nautical miles north and 10 nautical miles south respectively of latitude 46°N. The continued suitability of this site was reviewed in 1979 (OECD(NEA). 1980) and is due for review again in 1984. Following previous practice, the 1982 UK disposal operation was carried out by the Atomic Energy Research Establishment (AERE) Harwell according to the conditions laid down by this Ministry and the DOE. These conditions embody internationally-agreed safeguards. The operation was observed by a representative of the OECD(NEA). The waste was from several establishments and totalled 2833 packages of 2697 tonnes gross weight containing 47 TBq (1264 Ci) of alpha activity and 3756 TBq (101512 Ci) of beta/gamma activity, including 2345 TBq (63383 Ci) tritium. Routine environmental monitoring does not provide an effective means of assessing public radiation exposure from these disposals. Research data (Pentreath, 1983) show no caesium-137 attributable to sea dumping present in the deep sea fish Coryphaenoides (Nematonurus) armatus collected from near the sea bed in the disposal

Table 1 Principal discharges of liquid radioactive waste from UK nuclear establishments, 1982

Establishment	Radioactivity	Discharge	Discharges du	ring 1982		
		(annual equivalent), Ci*	TBq	Ci	% of limit utilised	
BRITISH NUCLEAR FUELS plc	· · · · · · · · · · · · · · · · · · ·					
Sellafield Sea pipeline	Total beta Ruthenium-106 Strontium-90 Total alpha	300 000 60 000 30 000 6 000	3 528 419 319 28	95 352 11 316 8 628 769	32 19 29 13	
Seaburn sewer	Total activity	4	0.0089	0.24	6.0	
Springfields	Total alpha	360	0.68	18.3	5.1	
	Total beta	12 000	174	4 709	39	
Chapelcross	Total activity <sup>l</sup>	700	4.1	112	16	
	Tritium	150	0.74	20	13	
Capenhurst Rivacre Brook Meols outfall	Total activity <sup>2</sup> Technetium-99	0.04 4	0.00052 0.0020	0.014 0.055	35 1.4	
UNITED KINGDOM ATOMIC ENERGY AUTHORITY						
Winfrith	Total activity	30 000	111	3 005	10	
	Ruthenium-106	9 000	0.18	4.9	<1	
	Strontium-90	1 200	0.11	3.1	<1	
	Total alpha	1 200	0.027	0.74	<1	
Harwell	Total activity <sup>l,3</sup> Tritium	240 240	0.75	20 37	8.4 16	
Dounreay	Total activity	24 000	38	1 019	4.2	
	Strontium-90	2 400	5.7	154	6.4	
	Total alpha	240	0.48	13	5.4	
CENTRAL ELECTRICITY GENERATING BOARD						
Berkeley	Total activity <sup>l</sup>	200	0.67	18	9.0	
	Tritium	1 500	1.7	47	3.1	
Bradwell	Total activity <sup>l</sup>	200	1.0	27	13	
	Zinc-65	5	0.0011	0.03	<1	
	Tritium	1 500	2.0	53	3.5	
Dungeness	Total activity <sup>l</sup>	200	1.2	32	16	
"A" Station	Tritium	2 000	0.74	20	1.0	
"B" Station <sup>4</sup>	Total activity <sup>l,5</sup>	4 TBq	<0.00029	<0.0078	<1	
	Sulphur-35	25 TBq	<0.000024	<0.00065	<1	
	Tritium	650 TBq	<0.00025	<0.0067	<1	
Hinkley Point <sup>6</sup>	Total activity <sup>l</sup>	200	2.6	72	36	
"A" Station	Tritium	2 000	0.67	18	<1	
"B" Station	Total activity <sup>l,5</sup>	100	0.037	1	1.0	
	Sulphur-35	700	0.85	23	3.3	
	Tritium	18 000	232	6 260	35	
Oldbury	Total activity <sup>l</sup>	100	2.0	54	54	
	Tritium	2 000	0.96	26	1.3	
Sizewell	Total activity <sup>l</sup>	200	1.0	28	14	
	Tritium	3 000	1.3	35	1.2	
Trawsfynydd	Total activity <sup>l</sup>	40	0.48	13	33	
	Caesium-137	7	0.026	0.7	10	
	Tritium	2 000	3.7	100	5.0	
Wylfa	Total activity <sup>l</sup> Tritium	65 4 000	0.11 18.6	3 502	4.6 13	
SOUTH OF SCOTLAND ELECTRICITY BOARD						
Hunterston	Total activity <sup>l</sup>	432	8.7	235	54	
"A" Station <sup>7</sup>	Tritium	1 200	1.5	40	3.3	
"B" Station	Total activity <sup>1,5</sup>	100	0.22	6.0	6.0	
	Sulphur-35	700	3.0	82	12	
	Tritium	40 000	256	6 930	17	

#### Table 1 (continued)

Establishment	Radioactivity	Discharge	Discharges during 1982				
		limit (annual equivalent), Ci*	ТВq	Ci	% of limit utilised		
MINISTRY OF DEFENCE (PROCUREMENT EXECUTIVE)							
Aldermaston	Total activity <sup>1,3</sup> Tritium	156 156	0.14 0.83	3.9 22.4	2.5 14		
MINISTRY OF DEFENCE (NAVY DEPARTMENT)							
Chatham	· Total activity <sup>l</sup> Cobalt-60 Tritium	20 10 20	0.0023 0.0023 0.004	0.062 0.062 0.11	<1 <1 <1		
Devonport	Total activity <sup>1</sup> Cobalt-60 Tritium	4 1 10	0.00057 0.00039 0.016	0.015 0.011 0.42	<1 1.1 4.2		
Faslane	Total activity <sup>1</sup>	1	0.000055	0.0015	<1		
Rosyth	Total activity <sup>1</sup>	30	0.0046	0.124	<1		
AMERSHAM INTERNATIONAL plc							
Amersham	Total activity <sup>1,3</sup> Tritium	72 400	0.35 0.093	9.5 2.5	13 <1		
Cardiff	Beta/gamma activity <sup>8</sup> Carbon-14 Tritium	96 GBq 2 TBq 1 400 TBq	0.018 1.1 64	0.49 30 1 700	19 56 4.5		

LExcluding tritium.

<sup>2</sup>Excluding uranium and its decay products.

<sup>3</sup>Authorisation or agreement specifies a control formula in which the total activity is calculated in equivalent curies, intended to allow for the relative radiotoxicities of different nuclides. The sums of the actual discharges in curies were lower than the values indicated. Column 4 gives equivalent terabecquerels. <sup>4</sup>Discharges commenced in December 1982.

<sup>5</sup>Excluding sulphur-35.

 $^{6}$ A single site authorisation applies at Hinkley Point. The table format represents the way in which it has been agreed that the authorisation should be apportioned in practice.

<sup>7</sup>The total activity limit was changed in June 1980 and during the period of this report both this limit and the one for tritium have referred to the 12 month period commencing on 16 June each year.

<sup>8</sup>Excluding tritium, carbon-14 and radioisotopes of calcium and strontium.

\*Unless otherwise stated.

area. These fish are not exploited commercially and their content of radioactivity therefore has no direct significance in terms of public radiation exposure. Concentrations of caesium-137 in these fish are not significantly different from those in the same species at locations remote from the disposal area. The activity is consistent with caesium-137 found through worldwide deposition of fallout and the concentrations are similar to those due to caesium-137 of fallout origin found in commercial fish species from Icelandic waters (section 4). The environmental impact of these disposals, as indicated by calculations using appropriate models, is negligible (OECD (NEA),1980).

#### Methods of analysis and of presentation and inter-3. pretation of results

#### 3.1 SI units

In this report data are presented using the SI (Systeme Internationale) radiological units recommended for use in the UK by the British Committee on Radiation Units and Measurements (BCRU, 1978). Table 2 summarises the radiological units used in this report, and provides relevant conversion factors to relate SI units to the old radiological units.

#### 3.2 Summary of analytical methods

Although some of the analytical methods which we have used are detailed elsewhere (Dutton, 1968, 1969), a very brief summary is given here in support of the measurements and the method of their presentation. The tables of results mostly include measurements of total beta radioactivity and of specific gamma-emitting nuclides. Pure beta emitters and alpha emitters (including transuranics) are also measured in appropriate cases.

Total beta radioactivity is measured using thin sources with a potassium-40 standard (Dutton, 1968). The efficiency of the method is nearly constant over a wide range of beta energies and the result gives a measure of the total radioactivity of the beta emitters present. However, agreement with the total as derived from isotopic analysis is not expected to be exact. The main advantage of total beta measurements is that they can be carried out quickly to give an early warning of any change in radioactivity concentrations which might require further investigation.

Gamma-emitting nuclides are analysed by gamma spectrometry. This is carried out using both NaI(T1) and Ge(Li)

Table 2 Radiological units used in this report

Quantity	New SI unit and symbol	Definition	Old unit and symbol	Definition	Conversion data
Radioactivity	becquerel (Bq)	disintegration per second	curie (Ci)	3.7 10 <sup>10</sup> disintegra- tions per second	1 Ci = $3.7 \ 10^{10}$ Bq 1 Bq $\approx 2.7 \ 10^{-11}$ Ci = $27$ pCi
Notes: l The	terabecquerel (I	Bq) is used in this re	port for radio	active discharges:	$1 \text{ TBq} = 10^{12} \text{ Bq} \approx 27 \text{ Ci}$
2 Rad	loactivity concen	trations are given in	becquerels per	kilogram (Bq kg <sup>−1</sup> );	1 Bq kg <sup>-1</sup> = 1 mBq g <sup>-1</sup> $\approx$ 27 pCi kg <sup>-1</sup> 1 pCi g <sup>-1</sup> = 37 Bq kg <sup>-1</sup>
Absorbed dose	gray (Gy)	J kg <sup>-l</sup> (joule per kilogram)	rad (rad)	10 <sup>-2</sup> J kg <sup>-1</sup>	$1 \text{ rad} = 10^{-2} \text{ Gy}$ $1 \text{ Gy} = 10^2 \text{ rad}$
Dose equivalent	sievert (Sv)	J kg <sup>-l</sup> x (modify- ing factors)	rem (rem)	10 <sup>-2</sup> J kg <sup>-1</sup> x (modify- ing factors)	$1 \text{ rem} \approx 10^{-2} \text{ Sv} = 10 \text{ mSv}$ $1 \text{ Sv} = 10^2 \text{ rem}$

detectors, calibrated using suitable reference sources. The spectra are reduced by computer-aided techniques to give radioactivity concentrations of detected nuclides. For samples of biota and sediments, searches are routinely made for, amongst others, the following artificial gamma emitters: manganese-54, cobalt-60, zinc-65, zirconium-95 plus niobium-95, ruthenium-106, silver-110m, antimony-124 and -125, caesium-134 and -137, and cerium-144. In the tables of results for these materials the absence of a column for any of these nuclides indicates non-detectability in each sample in that table.

Pure beta emitters, such as strontium-90 and technetium-99, are chemically separated from samples before beta counting.

Transuranic nuclides are chemically separated and analysed by alpha spectrometry using silicon surface-barrier detectors. Radiochemical procedures are generally labourintensive and are carried out on samples in which these nuclides are of particular relevance, often on an annual bulk (section 3.3).

#### 3.3 Methods of presentation of measurements

The tables of monitoring results generally contain summarised values of observations obtained during the year under review. Observations of a given quantity may vary throughout the year; in general any variations are larger than the analytical errors inherent in the observations. The variations may, for example, be due to changes in rates of discharge or to different dispersion conditions in the receiving environment. The presentation of the summarised results reflects the purpose of this monitoring which is interpretation in terms of public radiation exposures. The method of interpretation is described more fully in section 3.4. The appropriate integration period for comparison with recommended limits is one year; standard practice is to combine annual rates of consumption or occupancy of members of the public more highly exposed (the critical group) with the arithmetic means of observed radioactivity concentrations or dose rates respectively during the year. The use of, say, the highest observed (but unsustained) radioactivity concentration with an annual consumption rate would not provide a realistic comparison with the recommended limits which already embody a number of maximising assumptions. Therefore, the tables present the arithmetic means of observations made during the year.

The frequency of sampling reflects the resolution (implying the accuracy) judged to be necessary in the assessment or, as is largely self-evident, its radiological importance. So the number of sampling observations during the year is also given. Observations on biota consist of the results of analysing suitably large samples of material: for fish and shellfish a sufficient number of individual animals is sampled and analysed for each observation so as to allow for statistical variations. The number of individuals sampled also reflects the radiological importance. Thus, as in previous years, the number of individuals sampled within an observation varied - up to several hundred for fish and molluscs from near Sellafield. For gamma dose rates, which are measured using portable instruments, each observation consists of the mean of a number of individual readings at a given location. This number again depends upon the radiological importance of the observation; the locations chosen are generally those where there is likely to be occupancy by persons as determined by habits surveys (see section 3.4).

Analyses requiring radiochemical separation may be carried out on individual samples directly or on bulks made up of a number of individual samples collected over an extended period; in tables combining the results of gamma spectrometry and radiochemical analysis the extended period is one year unless otherwise stated.

Measurements on biota are given in terms of concentrations in wet material as collected. For fish and shellfish, because the purpose is assessment of internal exposure of the consumer, the concentrations apply to the edible fractions. For sediments, whose water content is more variable, dry concentrations are given.

The results for certain measurements, particularly total beta radioactivity concentrations and gamma dose rates, include a contribution due to natural radioactivity. Further analysis of samples (usually by gamma spectrometry) indicates the component of total beta radioactivity which is due to artificial sources and the component due to natural radionuclides (mainly potassium-40 and the decay products of uranium and thorium). In the case of gamma dose rates, an indication of the natural background component can be gained from measurements at similar locations remote from nuclear activities or from experience before these activities

environmental materials and natural background dose rates around the British Isles											
Material	Total beta radioactivity concentration (wet)*										
	Bq kg <sup>-1</sup>	Comments									
Fish	40 to 100	D Mostly <sup>40</sup> K									
Shellfish	40 to 100	) "									
Seaweed	200 to 600	) "									
Sand	200 to 400	) $40$ K and decay products of U and Th									
Mud	700 to 1000	) "									
Gamma dose rates	in air ove	r intertidal sediments: $\mu$ Gy h <sup>-1</sup>									
		Sand, shingle 0.03 to 0.05									
		Mud 0.05 to 0.1									

Table 3 Natural radioactivity concentrations of various

\*Except sediments for which dry concentrations apply.

began. For both types of measurement, however, experience is also useful. Table 3 lists representative values to be expected from natural sources.

#### 3.4 Methods of interpretation

The monitoring results in this report are interpreted in terms of radiation exposures of the public. The standards against which these exposures are judged are the recommendations of the International Commission on Radiological Protection (ICRP). For many years these recommendations have been endorsed for use in the UK by appropriate advisory bodies. UK practice relevant to the general public is now mainly based on the recent recommendations of ICRP as set out in ICRP Publication 26 (ICRP, 1977). The dose limitation system therein embodied has been accepted as national policy (Great Britain -Parliament, 1982). UK legislation will comply with the Euratom Directive on basic radiation safety standards, the current version of which (Commission of the European Communities, 1980) is based on the recommendations of ICRP Publication 26. In this report, results have been interpreted also on the basis of these recommendations.

The effect of these recommendations on the interpretation of the results will be briefly described. Emphasis is given to the principle that "all exposures shall be kept As Low as Reasonable Achievable ......" (ALARA). Thus the recommendations of ICRP Publication 26 underline the importance of consideration of collective doses in radiological control procedures. As in previous reports in this series, collective doses from liquid radioactive waste discharges continue to be kept under review. ICRP Publication 26 does not recommend a dose limit for populations;

such a limit might be regarded as suggesting the acceptability of a higher population exposure than is either necessary or probable. The ICRP concludes that its system of dose limitation is likely to ensure that the annual dose equivalent averaged over the population from all sources, excluding natural and medical irradiation, will not exceed 0.5mSv. The NRPB considers (NRPB, 1978) that maintenance of the annual dose equivalent below this value when averaged over the whole UK population is a reasonable objective; further, that the contribution from all UK waste management practices is unlikely to exceed one tenth of this, that is 0.05 mSv year<sup>-1</sup>. In this report an annual average dose equivalent of 0.05 mSv has been used for reference purposes regarding collective doses from radioactive waste discharges. By comparison, the annual average dose equivalent in the UK from natural radiation is approximately 2 mSv (NRPB, 1981).

ICRP Publication 26 recommends that doses should meet the ALARA objective, subject to compliance with appropriate individual dose limits. Control of individual exposures is intended to prevent non-stochastic (threshold) effects and to limit stochastic effects (i.e. those whose probability depends on the dose) to an acceptable level. To prevent non-stochastic effects, a dose equivalent limit for the public of 50 mSv year<sup>-1</sup> to any one organ or tissue is prescribed. For stochastic effects, it is recommended that the risk should be equal whether the whole body is irradiated uniformly or non-uniformly; weighting factors proportional to the risk are defined for different organs. The weighted sum is called the effective dose equivalent, and the ICRP-recommended limit for members of the public is 5 mSv year<sup>-1</sup>. It is these limits with which ICRP recommends that exposures of critical groups should be compared. This is the procedure which is followed in this report. ICRP also suggests that, in any rare cases where a few individuals were actually found to be receiving high rates of exposure over prolonged periods, it would be prudent to take measures to restrict their lifetime dose so that it corresponds to no more than 1 mSv year<sup>-1</sup> of lifelong whole body exposure. Consideration of this secondary objective has also been given in this report. The NRPB (NRPB, 1978) notes that the use of a limit of 5 mSv year<sup>-1</sup> combined with the technique of optimisation (the ALARA principle) will in most cases result in an average dose equivalent to a critical group of less than 1 mSv year<sup>-1</sup> of whole body exposure over a lifetime.

The ICRP also recommends secondary limits for internal and external irradiation. For internal irradiation, the limits are expressed as Annual Limits of Intake (ALIs). Values for radiation workers for a number of elements have been published in ICRP Publication 30 (ICRP, 1979a, 1980, 1981a). In this report environmental monitoring results are interpreted in terms of doses to members of the public. Thus the data on committed doses per unit intake, published in supplements to ICRP Publication 30 (ICRP 1979b, 1981b, 1982a, 1982b), have been used. The following points should be noted. First, metabolic differences may exist between certain age groups of the public and radiation workers. In this report appropriate allowances have been made where children are known to be members of critical groups. Secondly, in advance of a review by ICRP, the NRPB has recently published (NRPB, 1984) advice on gut uptake factors for actinides. This advice is that, for adult members of the public ingesting low concentrations of plutonium in food, an appropriate value of absorption factor by the gut is a factor of 5 higher than that currently used in ICRP Publication 30 for relevant forms of plutonium, except when a lower value can be justified. The effect is to enhance doses from plutonium essentially by this factor, and these higher doses are given in this report; alongside are given, in important cases, the doses derived using the unenhanced gut uptake factors used in ICRP Publication 30. Thirdly, especially for nuclides with long body retention times, such as the transuranics, committed doses are not completely received in the year of intake but over a longer period; a limit to integration of 50 years, typical of a working lifetime, is used by ICRP. Following ICRP recommendations, it is the committed dose equivalents from radionuclide intakes which are compared in this report with the recommended dose equivalent limit. Fourthly, in addition to the estimation of the committed effective dose equivalent for comparison with the ICRPrecommended dose limit based on stochastic effects, non-stochastic effects also require consideration, and this is given in this report. However, provided that the secondary objective relating to lifetime exposures is met, ICRP considers (ICRP, 1983) that non-stochastic effects will be avoided.

In the case of external exposure to penetrating radiation, uniform whole body exposure has been assumed. The measured quantity is absorbed dose rate in air. When interpreting this in terms of radiological effect, an absorbed dose rate in air of 1  $\mu$ Gy h<sup>-1</sup> has been taken as producing an effective dose equivalent rate of 0.87  $\mu$ Sv h<sup>-1</sup> (Spiers *et al.*, 1981).

In order to interpret monitoring results in terms of the recommendations of the ICRP, the remaining data required are, as appropriate, rates of food consumption or occupancy of areas relevant to external exposure. These are obtained by habits surveys specific to and generally near each nuclear establishment of interest. The results are kept under review and the surveys are repeated at intervals. The main purpose of the surveys is to identify a group (the critical group) of persons most highly exposed through a particular pathway or pathways. The critical pathway approach has been in use for many years, and is still embodied in the recommendations of the ICRP. In this report, critical group habits data relevant to a given establishment are combined with the results of environmental monitoring and appropriate dosimetric data as above to estimate the annual dose to the critical group. This is then expressed as a percentage of the appropriate ICRP-recommended dose limit for members of the public.

### 4. British Nuclear Fuels plc (BNFL)

BNFL is concerned mainly with the design and production of fuel for nuclear reactors and its reprocessing after irradiation. The company also operates nuclear power plant supplying electricity to the national grid. We regularly monitor the environmental consequences of discharges of liquid radioactive waste from four BNFL sites, namely Sellafield (until 1981, known as Windscale and Calder), Springfields, Capenhurst and, on behalf of Scottish Departments, Chapelcross.

#### 4.1 Sellafield, Cumbria

Operations and facilities at this establishment include fuel element storage and decanning, the Windscale nuclear fuel reprocessing plant and the Calder Hall magnox-type nuclear power station. The most significant liquid radioactive waste discharges are from the fuel element storage ponds and the reprocessing plant, through which passes all the irradiated fuel from the UK nuclear power programme. Most of the nuclear waste separated from the fuel is presently stored on site; relatively small quantities of radioactivity are discharged to the north-east Irish Sea, through pipelines which terminate 2.1 km beyond low-water mark. Discharges during 1982 are summarised in Table 1, and were within the limits set by the authorising Departments. Discharges of total beta activity, at 32% of the authorised limit, were less than in 1981 (35%). Total beta discharges are substantially dependent upon releases of radiocaesium which mainly originate from the fuel element storage ponds. In 1982 caesium-137 pipeline discharges totalled 2000 TBq, a lower total than in 1981 (2357 TBq). This reduction was brought about by the continued use of

Sampling area/landing point	Sample	No. of sampling	Mean radioa centration	(wet),	con- Bq kg-1
		observa- tions <sup>3</sup>	Total beta	134Cs	<sup>137</sup> Cs
Sellafield shoreline area <sup>1,3</sup>	Cod	4	1200	57	1100
	Whiting	1	680	29	620
	Mackerel	1	600	28	590
Sellafield offshore area <sup>1,3</sup>	Plaice	12	510	23	4 <b>9</b> 0
	Dab	4	500	22	500
	Flounder	1	420	18	430
	Whiting	1	/10	38	820 370
	cou	1	400	10	570
Ravenglass <sup>2</sup>	Cod	7	680	35	660
	Plaice	5	480	25	490
	Flounder	I	1100	40	1100
Morecambe Bay <sup>1</sup>	Flounder	4	410	17	380
Whitehaven <sup>2</sup>	Cod	4	360	12	260
	Plaice	4	240	8.2	200
	Herring	2	280	7.5	160
Flootwood <sup>2</sup>	Cod	4	320	10	220
LIEELWOOD -	Plaice	4	160	5.9	140
		•	250		240
Cumbrian rivers <sup>4</sup>	Sea trout	3	350	13	340
Isle of Man <sup>2</sup>	Cod	4	170	3.7	74
	Plaice	1	120	1.6	3/
	Herring	3	92	J.0 05	0/ 19
	Lemon sole	1	150	2.6	60
	Dover sole	ī	NA	2.3	57
1		,			50
Irish Seat (Research vegeel erwise)	Plaice	4		2.2	28 86
(Research vesser cluise)	Whiting	5	••	6.6	160
	Herring	1	••	1.7	60
	Ray	4		1.0	33
	Hake	1		0.7	19
	Monkfish	2		4.3	64
	Mackerel	1		3.5	76
Tanan Osland	6-1	,	130	ND	4 1
Inner Solway	Flounder	3	380	11	300
_					
North Anglesey <sup>1</sup>	Plaice	1	140	ND 13	32
	Dab	1	150	1.5	
Northern Ireland <sup>2</sup>	Whiting	4	230	4.2	120
Minchl	Plaice	3	120	0.9	23
Minen-	Cod	3	160	1.0	23
	Herring	1	140	ND	8.6
Northern North Sea <sup>1</sup>	Plaice	4	170	ND	3.8
	Cod	9	140	0.09	6.1
	Haddock	5	NA	ND	3.0
	Herring	1			3.3
	Saithe Norway pout	2	110	•	1.2
Mid-North Sea <sup>1</sup>	Plaice	10	100	0.05	4.6
	Haddock	4	NA NA	0.1	6.5
	Herring	1	100	0.4	10
	Whiting	2	NA	0.2	15
Southern North Seal	Plaico	4	94	ND	2.8
Journern morrn Jea*	Cod	4	140	0.1	5.6
	Whiting	2	NA	ND	4.4
Numeration Oct	0-1	,	110	ND	
Norwegian Sea <sup>1</sup>	Uod Haddock	1	130	ND "	2.2
		-			2.0
Barents Sea <sup>l</sup>	Cođ	1	NA	••	1.6
Iceland area <sup>1</sup>	Cod	2	100	••	0.4
	Haddock	1	120	**	0.2

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Table 4	Beta/gamma	radioactivity	in	fish	from	the	Irísh	Sea	vicinity	and	further
	afield, 198	32									

ND = not detected; NA = not analysed; <sup>1</sup>Sampling area; <sup>2</sup>Landing point; <sup>3</sup>See section 3.3 for definition; <sup>4</sup>Samples collected from a number of rivers by the North West Water Authority.

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zeolite skips in the ponds to absorb caesium, (Hunt, in press) in order to ensure that doses were ALARA (section 3.4) as required by the authorising Departments. Strontium-90 discharges in 1982 were slightly more than in 1981. Discharges of ruthenium-106, which derives mainly from operations other than in the ponds, were slightly less in 1982 than in 1981. Discharges of alpha-emitting plutonium isotopes in 1982 totalled 20.8 TBq, very slightly more than in 1981 (20.3 TBq) but were still at a lower level than in previous years. Americium-241 discharges in 1982 were less than in 1981, and this reduction was reflected in an overall decrease in total alpha activity discharged.

We maintained a substantial monitoring effort during 1982. The two critical radiation exposure pathways continued to be from consumption of fish and shellfish and from external exposure. Following established practice, the largest monitoring effort has been expended on these pathways. In 1982, as in previous recent years, there was no harvesting of Porphyra in the immediate Sellafield vicinity for manufacture of laverbread, but monitoring was continued because the pathway remains potentially important. An extensive research programme was also continued. The aims of this programme are to improve our knowledge of the distribution and behaviour of radionuclides in the marine environment, especially in relation to the critical exposure pathways, and also to provide a means of assessing other pathways, of lower current importance, thereby assisting in keeping all exposure pathways under review. Some of the research was supported by contract with the Commission of the European Communities. Results from our research programme are included where relevant.

#### 4.1.1 The fish and shellfish consumption pathway

Public radiation exposure from Sellafield discharges by consumption of fish is predominantly due to radiocaesium. Concentrations of total beta activity and caesium-134 and -137 in fish from the vicinity of the Irish Sea and from further afield are given in Table 4. Data are listed by location of sampling or landing point, in approximate order of increasing distance from Sellafield. So as to be representative of consumption by the public, samples are generally obtained from commercial sources. However, to minimise the risk of underestimating exposures, and as certain materials may not be available commercially, we also carry out specific surveys sampling fish and shellfish from the Sellafield vicinity. The location "Sellafield Shoreline Area" is close inshore in this vicinity. "Sellafield Offshore Area" represents a rectangle, one nautical mile wide and two nautical miles long, situated south of the pipeline with the long side parallel to the shoreline; this Area averages about 5 km from the pipeline outlet.

The results reflect the progressive dilution of radiocaesium with increasing distance from Sellafield. They also reflect

the age of the radioactivity, such that the ratio of caesium-137 to caesium-134 (half-lives 30 years and 2 years respectively) increases with distance. At large distances, and remote from the smaller discharges from elsewhere, concentrations of artificial radioactivity tend towards those from weapons-test fallout. For caesium-137 in fish, measurements remote from land run-off indicate a value of about 0.2-0.4 Bq kg<sup>-1</sup> from this source. Variations between species for a given area, while not large, are mainly to be explained in terms of residence time in the area as well as feeding habits. These variations are likely to be most apparent in the results close to Sellafield because of the relatively steep concentration gradient of radiocaesium in sea water. Because the purpose of the result is dose estimation, results are based on observations which include large numbers of individual fish (section 3.3).

Concentrations of radiocaesium in 1982 were generally less than in 1981 for fish from both the Irish Sea and further afield. This is attributed to reduced concentrations in sea water, following the decreasing trend in radiocaesium discharges from Sellafield since 1975; also, for the Irish Sea, improved dispersion occurred in 1982 as compared with 1981, as described later in this section.

Radiation exposure from consumption of shellfish is due in part to radiocaesium, but other nuclides also make significant contributions owing to higher concentration factors in these foods than in fish. Table 5 lists concentrations of total beta activity and beta/gamma-emitting nuclides in shellfish from the Irish Sea and further afield. Winkles are of particular radiological importance to the critical group of shellfish consumers near to Sellafield, as described later in this section. In 1982 additional samples of winkles were obtained from consumers who collected supplies at Coulderton and Nethertown, locations typical of the supplies of this critical group.

Concentrations of radionuclides in shellfish, as with fish and as in previous years, diminished with increasing distance from Sellafield; the rate of reduction was least for nuclides which are conservative to sea water, such as isotopes of caesium. There were also, as previously, substantial variations between species: in general, molluscs tend to concentrate the less conservative nuclides to a greater extent than do crustaceans, whilst in contrast molluscs concentrate radiocaesium somewhat less than do crustaceans, which in turn tend to concentrate it less than fish.

Concentrations of radiocaesium in shellfish in 1982, as for fish, showed general reductions as compared with 1981. There were also general reductions in concentrations of ruthenium-106, reflecting decreased discharges of this nuclide. Concentrations of other beta/gamma-emitting radionuclides were generally similar to those in 1981. Table 5 Beta/gamma radioactivity in shellfish from the Irish Sea vicinity and further afield, 1982

Sampling area/landing point	Sample	No. of sampling	Mean radioactivity concentration (wet), Bq kg <sup>-1</sup>											
		tions <sup>3</sup>	Total beta	<sup>60</sup> Co	<sup>95</sup> Zr + <sup>95</sup> Nb	103 Ru	106 Ru	<sup>110</sup> mAg	125 Sb	<sup>134</sup> Cs	<sup>137</sup> Cs	<sup>144</sup> Ce	<sup>154</sup> Eu	155 Eu
Sellafield shoreline area <sup>1,3</sup>	Crabs Lobsters	7 3	540 940	3.2 0.6	ND "	ND	350 71	20 7.1	0.8 ND	8.4 15	180 320	3.4 ND	ND	0.3 ND
Sellafield offshore area <sup>1</sup>	Whelks	2	770	ND			520	9.3		4.1	95			
Coulderton <sup>1</sup>	Limpets Winkles Winkles <sup>4</sup>	1 10 5	2100 4000 5400	5.6 14 27	430 550 1100	 28 8.1	1800 2500 4300	20 55 70	 6.8 14	15 22 25	350 380 480	63 66 110	 5.2 3.6	12 5.9 5.4
Nethertown <sup>1</sup>	Winkles <sup>5</sup>	4	4200	17	1900	12	3200	57	9.1	20	400	130	10	6.5
Ravenglass <sup>1</sup>	Cockles Mussels	4 4	1600 1000	13.0 4.3	350 180	13 4.7	1100 980	ND ND	ND 2.3	8.7 8.1	170 170	74 10	7.9 1.0	14 3.8
St Bees <sup>1</sup>	Limpets Mussels Winkles	4 4 4	2700 3900 3300	4.3 10 11	190 800 300	5.2 56 7.0	990 3500 2000	19 2.4 44	11 6.1 ND	10 8.0 11	220 180 200	32 36 28	3.4 1.8 ND	2.4 4.2 ND
Whitehaven <sup>2</sup>	Nephrops	2	320	ND	ND	ND	ND	ND	•	9.6	130	ND	•	
Morecambe Bay <sup>1</sup>	Shrimps Cockles	4 4	220 230	 1.1		 	 54		" 0.5	7.9 3.8	160 85	" 2.2	 	 0.3
Isle of Man <sup>2</sup>	Scallops Queens	4 1	120 110	ND "			0.5 ND	1.0 2.4	ND 	0.3 0.4	14 9.0	ND "	 	ND "
Inner Solway <sup>1</sup>	Shrimps	3	200	•		•	3.8	ND		6.3	140		•	
Kirkcudbright <sup>2</sup>	Scallops Queens Winkles	3 4 4	65 67 270	  1.2	••		ND 	" 0.3 2.0	 	0.1 0.2 2.7	6.3 9.9 60	 	  	  
North Solway coast <sup>1</sup>	Winkles	5	260	0.6		•	110	3.2	0.5	2.3	64	-		•
Wirral <sup>1</sup>	Shrimps Cockles	2 3	210 110	ND 	•	•• ••	ND 4.3	ND 	ND "	3.6 0.9	87 36	•• ••	 	 0.2
Conwy <sup>2</sup>	Mussels	2	76	•			3.0	•		0.6	18			ND
North Anglesey <sup>1</sup>	Crabs	1	100		-		ND	•		2.1	16			
Northern Ireland <sup>2</sup>	Nephrops	4	140	•		•				1.3	27			
Irish Sea <sup>l</sup> (Research vessel cruise)	<i>Nephrops</i> Queens Squid Whelks	4 4 2	NA  	" " 0.3	•• •• ••	  	 94 ND 61	 12 ND 5.5		1.4 0.7 1.5 1.1	51 18 29 25	  	  	  
Northern North Seal	Nephrops	3	100	ND			ND	ND	•	0.1	6.7			
Mid North Sea <sup>l</sup>	Nephrops Mussels	1 1	110 45	•	 	•• ••	 	 	 	ND 	5.4 1.1	 	 	 
Southern North Sea <sup>1</sup>	Cockles Mussels	4 3	26 32			 	 	 	 	 	0.7	 	 	

NA = not analysed; ND = not detected.

<sup>1</sup>Sampling area; <sup>2</sup>Landing point; <sup>3</sup>See section 3.3 for definition; <sup>4</sup>Samples collected by Consumer A; <sup>5</sup>Samples collected by Consumer B.

Public radiation exposure from transuranic nuclides in fish is lower than from radiocaesium. Analyses for transuranics are also labour-intensive. Therefore, only a selection of samples of fish and shellfish chosen mainly on the basis of potential radiological significance are analysed for transuranic nuclides. Analyses are often carried out on samples bulked annually (section 3.3). The data for 1982 are presented in Table 6. Transuranics are less conservative to sea water than is radiocaesium; this is reflected in higher concentrations of transuranics in shellfish as compared with fish, and a rapid reduction with distance in concentrations of transuranics, particularly in shellfish.

Concentrations of transuranics in fish and shellfish from the Irish Sea were generally lower in 1982 than in 1981. Despite the slightly increased discharges of alpha-emitting plutonium nuclides in 1982, concentrations diminished slightly; this was a reflection of the improved dispersion described later, and was also probably because the nonconservative nature of plutonium caused a delayed effect of the overall reductions in discharges since 1979. The reductions in concentrations of americium-241 reflected the lower discharges in 1982 and a smaller contribution due to grow-in from the parent plutonium-241, discharges of which have also reduced since 1979.

The radiation dose to consumers of fish and shellfish depends upon the product of the mass of foodstuff consumed and its radioactivity concentration. Because of variations in these two quantities between individual consumers, a wide range of annual doses is to be expected. The critical group approach, which is well established in the UK and recommended by ICRP for control purposes, is based on identifying groups of individuals in exposed populations subject to the highest radiation dose rates. Of the two main variables, radioactivity concentrations in fish and shellfish are highest in the coastal area in the vicinity of the pipeline. Hence, eaters of fish and shellfish within the local fishing community represent one exposed population whose consumption rates we have studied and kept under review. As regards the other main variable, consumption rates, surveys have shown that, in addition to the Cumbrian coastal community, the larger population in Cumbria and north Lancashire of those associated with

Table 6 I	fransuranic	radioactivity in	fish and	shellfish	from th	ne Irish	Sea	vicinity	and	further	afield,	1982
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Sampling area/landing point	Sample	No. of sampling	Mean radioactivity concentration (wet), Bq $kg^{-1}$							
		observa- tions <sup>3</sup>	238Pu	$239_{Pu} + 240_{Pu}$	241 Pu	241 Am	242 Cm	<sup>243</sup> Cm + <sup>244</sup> Cm		
Sellafield shoreline area <sup>1,3</sup>	Cod	2	0.011	0.048	NA	0.043	0.00084	0.00032		
	Crabs	2	0.61	2.7		7.3	ND	0.038		
	Lobsters	2	0.25	1.0	•	4.8	0.030	0.016		
Sellafield offshore area <sup>1,3</sup>	Plaice Whelks	3 2	0.032 NA	0.13 NA	 	0.13 16	0.0016 NA	0.00072 NA		
Coulderton <sup>1</sup>	Winkles	1	19	72		67	1.1	0.40		
ooditerton	Winkles <sup>4</sup>	î	30	113		106	2.3	0.78		
	Limpets	1	23	88		95	1.5	0.78		
St Bees <sup>1</sup>	Winkles	1	9.0	32	<b>99</b> 0	32	0.35	0.25		
	Mussels	1	12	50	NA	37	0.35	0.27		
	Limpets	4	NA	NA	••	30	NA	NA		
Ravenglass <sup>1</sup>	Cockles	1	16	65		93	1.3	0.65		
	Mussels	1	9.0	35		53	0.24	0.22		
Ravenglass <sup>2</sup>	Plaice	1	0.016	0.066	••	0.064	0.0023	0.00047		
	Cod	1	0.0031	0.012		0.017	0.00048	0.00012		
Whitehaven <sup>2</sup>	Plaice	1	0.0031	0.015		0.014	ND	0.00008		
	Cod	1	0.0018	0,0068		0.0074		0.00003		
	Nephrops	1	0.28	1.2		2.6	0.014	0.0096		
Morecambe Bay <sup>1</sup>	Shrimps	1	0.018	0.077		0.079	ND	0.00052		
	Cockles	ī	0.92	4.0		6.0		0.037		
Isle of Man <sup>2</sup>	Mixed									
	flatfish	1	0.00065	0.0031		0.0036		ND		
	Cod	1	0.00034	0.0012		0.0015	••			
	Herring	1	0.00070	0.0026		0.0029		"		
	Scallops	1	0.018	0.094		0.037				
Irish Sea <sup>l</sup> (research vessel cruise)	Whelks	2	NA	NA	••	1.2	NA	NA		
Inner Solway <sup>1</sup>	Salmon	1	0.00014	0.00056	н	0.00053	ND	ND		
Kirkcudbright <sup>2</sup>	Scallops	1	0.037	0.18	••	0.070		0.00014		
-	Queens	1	0.020	0.097	••	0.070		0.00049		
	Winkles	1	0.63	2.7	"	3.4	0.027	0.016		
North Solway coast $^{1}$	Winkles	1	0.95	4.2	•	5.0	0.019	0.024		
Wirral <sup>1</sup>	Cockles	3	NA	NA		1.8	NA	NA		
Conwy <sup>2</sup>	Mussels	1	0.029	0.16	•	0.18	ND	0.00068		
Northern Ireland <sup>2</sup>	Nephrops	1	0.0025	0.012	"	0.026	0.00089	0.00026		
Minch <sup>1</sup>	Cod	1	0.00020	0.00092	NA	0.00087	ND	ND		
Northern North Sea <sup>1</sup>	Cod	1	0.00012	0.00048		0.00057				
	Norway pout	1	0.00049	0.0038	"	0.0026	"			
	Nephrops	1	0.0031	0.016		0.0082	0.00027	0.00011		
Mid North Sea <sup>1</sup>	Nephrops	1	0.0023	0.0099		0.0077	ND	ND		
Southern North Sea <sup>1</sup>	Mussels	1	0.0013	0.0083	•	0.0023	0.00027	0.00007		
	Cockles	I	0.0023	0.011		0.0052	ND	0.00013		
Iceland area <sup>1</sup>	Cod	1	0.00015	0.00065		0.00048	••	ND		

ND = not detected. NA = not analysed.

<sup>1</sup>Sampling area.

<sup>2</sup>Landing point.

<sup>3</sup>See section 3.3 for definition. <sup>4</sup>Samples collected by consumer A.

commercial fisheries based primarily at Whitehaven, Fleetwood and in the Morecambe Bay area contains consumers of large quantities of fish and shellfish. This therefore represents a second exposed population which is kept under review, even though, in general, the relevant fishing grounds are further afield than the Cumbrian

coastal area and concentrations of radioactivity in fish landed are lower.

The consumption rates of the Cumbrian coastal community described above were reassessed in 1982 (Hunt, 1983). Techniques used in the collection of data have continued

Exposed population	Consumption assessment (	rate used see text)	in	Nuclide	Committed effective dose equivalent (as % of ICRP- recommended dose limit of 5 mSv year <sup>-1</sup> for members of the public)			
		kg y <sup>-1</sup>	(g d <sup>-1</sup> )		On basis of current ICRP recommenda- tions	Effect of Pu enhanced by a factor of 5 (see text)		
Consumers in local fishing community	fish: crustaceans: molluscs:	36.5 6.6 16.4	(100) (18) (45)	90 Sr 106 Ru 134 Cs 137 Cs 238 Pu 239 Pu + 240 Pu 241 Pu 241 Am	0.5 5.9 0.5 7.1 0.7 2.8 1.7 13.9	0.5 5.9 0.5 7.1 3.3 13.9 8.7 13.9		
				Total	34	54		
Consumers associated with commercial fisheries (Whitehaven, Fleetwood, Morecambe Bay)	fish: crustaceans: molluscs:	131 26 18	(360) (70) (50)	90 Sr 106 Ru 134 Cs 137 Cs 238 Pu 239 Pu + 240 Pu 241 Pu 241 Pu 241 Am Total	0.3 0.1 0.6 8.9 0.04 0.2 0.1 1.4	0.3 0.1 0.6 8.9 0.2 0.9 0.6 1.4 13		
Typical member of the fish-eating public con- suming fish landed at Whitehaven/Fleetwood	fish:	15	(40)	l34Cs l37 <sub>Cs</sub> Total	0.05 0.8 0.9	0.05 0.8 0.9		

Table 7 Individual radiation exposures due to consumption of Irish Sea fish and shellfish, 1982

to include the use of consumption logging sheets particularly by members of critical groups (Leonard *et al.*, 1982; Leonard, in press). Consumption rate data have been interpreted using techniques based upon ICRP recommendations (Hunt *et al.*, 1982) to select appropriate critical groups of higher-rate consumers.

Radioactivity concentrations in fish and shellfish eaten by the two exposed populations will vary with the species involved, so in estimation of doses it is not sufficient to determine only the total consumption rates of fish and shellfish together. Our experience (illustrated by Tables 4--6) has shown, however, that for a given area, within each of the classes, fish, crustaceans and molluscs, the concentrations of given nuclides in representative samples are relatively constant. For each of the two exposed populations, therefore, critical sub-groups were identified for each class of foodstuff and the mean consumption rates of the sub-groups were determined. For the Cumbrian coastal community the 1982 reassessment indicated that these consumption rates were 36.5 kg year<sup>-1</sup> (100 g d<sup>-1</sup>) fish, 6.6 kg year<sup>-1</sup> (18 g d<sup>-1</sup>) crustaceans and 16.4 kg year<sup>-1</sup> (45 g d<sup>-1</sup>) molluscs (Hunt, 1983). In this report, consumption rates are expressed additionally in kilograms per year to emphasise the importance of assessment over a year, in line with ICRP recommendations.

The data obtained show that above-average consumers in each of the component sub-groups are not generally members of another component sub-group. However, members of more than one sub-group do exist, so to avoid underestimating the exposure of the overall critical group, this exposure is derived by combining the exposures of each sub-group additively. Comparison based on individual critical group members' exposures shows that this procedure is not excessively conservative. Plaice and cod are overwhelmingly the fish most eaten by the high-rate consumers, and the assessment of exposure of the critical group is based upon an equal mix of these species taken from the Sellafield Offshore Area and from landings at Ravenglass. Consumption data indicate that it is unreasonable to base the assessment on fish from the Shoreline Area. The exposure due to consumption of crustaceans is based upon an equal mix of crabs and lobsters from the Shoreline Area, whilst the exposure from consumption of molluscs is based upon averaged radionuclide concentrations in winkles from Coulderton and St Bees (and Nethertown insofar as data are available), including the data from samples collected by local consumers.

Table 7 summarises exposures in 1982. Provided the ICRP lifetime dose criterion is met, as addressed later in this section, the occurrence of non-stochastic effects will be avoided; thus the primary need is to consider stochastic

effects on the basis of committed effective dose equivalent (section 3.4). For each exposed population this quantity is given together with the contributions of individual nuclides. For simplicity, only the more important nuclides are listed; hence it is not to be expected that the sums of the listed contributions will necessarily equal the totals presented. The contributions due to strontium-90 were estimated from the discharges of this nuclide. The contributions due to plutonium-241 were derived on the basis of the measured value in winkles from St Bees; this gave a ratio of plutonium-241/plutonium-239+240 which was in good agreement with the ratio in the discharges, confirming the method of estimation in previous years (Hunt, 1983).

Comments in section 3.4 on the dose estimates for transuranics are relevant here; in particular the effect of applying the enhanced gut uptake factor for plutonium, following NRPB's advice, is shown in the last column of Table 7. Using this advice on gut uptake factors which are still under review by the ICRP, the committed effective dose equivalent to the critical group of local consumers in 1982 would have been at most 54% of the ICRP-recommended dose limit of 5 mSv year<sup>-1</sup> for members of the public. On the basis of dosimetric factors currently recommended by ICRP, but now subject to review, the committed effective dose equivalent would have been 34% of this limit. These results represent decreases from 69% and 46% respectively reported for 1981 (Hunt, 1983). The decreases are due to lower concentrations, particularly of radiocaesium in fish, and of ruthenium-106 and transuranic nuclides in winkles, as described earlier in this section.

The dose to the critical group has also been considered in the context of the ICRP's advice on lifetime exposure (section 3.4). Effective dose equivalent rates to the critical group in excess of 1 mSv year<sup>-1</sup> (20% of the ICRP-recommended dose limit for members of the public) have only been reported for the last few years. This is not long enough for lifetime exposures to have exceeded, on average, 1 mSv year<sup>-1</sup>. As a result of measures already implemented by BNFL, discharges of radiocaesium and actinides are declining, and further measures to reduce these discharges are planned. Nevertheless, the dose to the critical group in the context of lifetime exposure is being kept under review.

Habits surveys carried out in relation to the consumers associated with commercial fisheries based mainly on Whitehaven, Fleetwood and the Morecambe Bay area indicate critical sub-group consumption rates for fish, crustaceans and molluscs to be 131 kg year<sup>-1</sup> (360 g d<sup>-1</sup>), 26 kg year<sup>-1</sup> (70 g d<sup>-1</sup>) and 18 kg year<sup>-1</sup> (50 g d<sup>-1</sup>) respectively. As for the Cumbrian coastal community, the overall critical group has been defined by the maximising procedure of summing exposures due to these component consumption rates. The dose rate due to intake of fish has been assessed using activity concentrations of an equal mix of plaice and cod landed at Whitehaven and Fleetwood. Consumption of crustaceans has been based on shrimps from Morecambe Bay, and consumption of molluscs has been based on Morecambe Bay cockles. The effective dose equivalent to members of this critical group in 1982 is given in Table 7. The total of 13% of the ICRP-recommended dose limit for members of the public, on the basis of the enhanced gut uptake factor for plutonium, represents a decrease from 19% of this limit reported for 1981 (Hunt, 1983). The decrease was due to the lower concentrations in the Irish Sea of radiocaesium in fish and of transuranic nuclides in shellfish.

The effective dose appropriate to a consumption rate of 15 kg year<sup>-1</sup> (40 g d<sup>-1</sup>) of fish from landings at Whitehaven and Fleetwood is also given in Table 7. This consumption rate represents an average for typical fish-eating members of the public. The effective dose in 1982 was 0.9% of the ICRP-recommended dose limit for members of the public, which represents a decrease from 1.2% reported for 1981 (Hunt, 1983), due to the reduced concentration of radiocaesium in Irish Sea fish.

Collective doses from fish and shellfish have been estimated for 1982 for the UK and other European countries. In general, the method used has been to combine data on fish and shellfish landings from relevant sea areas with average radioactivity concentrations in fish and shellfish caught in these areas. Sea areas considered included the Irish Sea, Scottish waters, the North Sea, Baltic Sea, Norwegian Sea, Spitzbergen/Bear Island area and Barents Sea. Corrections were made for the fraction of fish or shellfish consumed. The contribution of weapons-test fallout to the radioactivity concentrations was subtracted. Consideration has been given to the pathway due to fish offal and industrial fisheries, the product of both of which is fish meal which is fed to pigs and poultry. Consumption of food products from these animals give rise to a small contribution to the collective dose, and this has been included. The results are presented in Table 8; the result for other European countries is provisional because final landings statistics for fish and shellfish in 1982 have not yet been published by the International Council for the Exploration of the Sea (ICES). The provisional result will be reviewed in a future report; it is unlikely to change markedly so can be used as a basis for comparison with previous years.

Table 8	Collective doses	from	fish	and	shellfish,
	1982 (provisional	.)			

Population	Size of population	Committed effective dose equivalent, man-Sv
UK	5.6 x $10^7$	90
Other European countries	$7.0 \times 10^8$	100

Liquid radioactive discharges from Sellafield are the main source of collective dose reported here; by comparison the effect of liquid discharges from other establishments is very small. Most of the collective dose is due to radiocaesium in edible fish; the contribution due to shellfish is minor. Also relatively small is the contribution, again mainly from radiocaesium, due to fish offal and industrial fisheries (Hunt and Jefferies, 1981). Other radionuclides which contribute to the collective dose, but in even smaller proportions, are strontium-90, through both fish and shellfish, and ruthenium-106 and the transuranics, mainly through shellfish. It should be noted that for transuranics the doses per unit intake allow for the long body halftimes, so that the small contributions estimated for the transuranics are committed in the future rather than already received (section 3.4). The contribution of pathways other than fish and shellfish consumption, e.g. external exposure, to the collective dose from Sellafield liquid discharges is relatively small (Hunt and Jefferies, 1981).



Figure 2. Concentration (Bq kg<sup>-1</sup>) of caesium-137 in filtered water from the Irish Sea, November 1982.



Figure 3. Concentration (Bq kg<sup>-1</sup>) of caesium-137 in filtered water from the North Sea, February-March 1982.

The result of 90 man-Sv for the UK in 1982 represents a decrease in collective dose as compared with 130 man-Sv in 1981 (Hunt, 1983). This decrease was due to the generally lower radiocaesium concentrations, noted above, in fish and shellfish from the Irish Sea and further afield. There were small increases in fish landings from the Irish Sea and Scottish waters in 1982 but these were not sufficient to offset the effect of decreased radiocaesium concentrations. The provisional value of the collective dose to inhabitants of other countries in 1982 was also less than in 1981, for which year 150 man-Sv was reported. This reduction is also due to the generally decreased radiocaesium concentrations in fish and shellfish in 1982.

The collective dose for the UK given in Table 8 may be compared with the annual dose equivalent averaged over the population of 0.05 mSv considered unlikely to be exceeded (NRPB, 1978) (see section 3.4) as a result of all waste management practices. In 1982 the UK collective doses through the fish and shellfish pathways as a result of liquid radioactive waste disposal operations amounted to less than 5% of this value.

It is clear from the statements above which compare the 1981 and 1982 results for both critical group and collective dose rates that an important factor determining exposures is the distribution of radioactivity in the marine environment. We maintain a continuing programme of research on marine behaviour and distributions (including budget assessments) of significant radionuclides. Data on the distribution of caesium-137 in sea water are regularly collected by research vessel cruises; the distribution observed in the Irish Sea in November 1982 is shown in Figure 2. Comparison with the data for March 1981 (Hunt, 1983) shows that concentrations of caesium-137 in sea water of the Irish Sea generally decreased in 1982. This is partly explained by the reductions in radiocaesium discharges from Sellafield in 1982. Because of the extent of the decrease it is also likely that there was a greater flow of sea water through the Irish Sea, following the reduced flow postulated for 1981 (Hunt, 1983). The distribution of caesium-137 in sea water of the North Sea during February and March 1982 is shown in Figure 3. This was generally similar to the distribution observed in August and September 1981 (Hunt, 1983); the intrusion off north-east Scotland of water slightly richer in caesium-137 is of minor radiological significance and is believed to be an effect of the flushing of the Irish Sea noted above. The lower concentrations of radiocaesium in fish from the North Sea in 1982 as compared with 1981 are probably due to the period required for uptake of radiocaesium by fish, such that concentrations in 1981 reflected the higher levels of radiocaesium in water observed in 1980 (Hunt, 1982). The reducing trend in radiocaesium concentrations in sea water and fish from the Irish Sea and further afield reflects the general decrease in radiocaesium discharges from Sellafield since 1975 achieved particularly by the use of zeolite skips in the magnox fuel storage ponds in pursuance of the ALARA principle (section 3.4) as required by the authorising Departments (Hunt, in press).

#### 4.1.2 External exposure

A further important pathway leading to radiation exposure as a result of Sellafield discharges arises from uptake of gamma-emitting radionuclides by intertidal sediments in areas frequented by the public. In general, it is the finegrained muds and silts prevalent in estuaries and harbours, rather than the coarser-grained sands to be found on open beaches, which adsorb the radioactivity more readily. Gamma dose rates currently observed are mainly due to radiocaesium, ruthenium-106 and zirconium-95 plus niobium-95.

We regularly monitor a range of coastal locations both in the Sellafield vicinity and further afield using portable gamma-radiation dosemeters. Locations are chosen on account of both dose rates themselves and levels of occupancy by members of the public. Table 9 lists the locations monitored together with the dose rates in air at 1m above ground level. Monitoring in Scotland is carried out on behalf of departments of the Scottish Office. Dose rates on Irish Sea shorelines near other nuclear establishments which reflect Sellafield discharges are given later in this report (see sections 4.2, 4.3, 4.4, 6.5, 6.10). Variations in sediment type account for the quite marked fluctuations in dose rate, superimposed on a general decrease with increasing distance from Sellafield. Dose rates over intertidal areas in 1982 were generally similar to those in 1981 (Hunt, 1983).

We also regularly monitor radioactivity concentrations in sediments. This is both because of relevance to dose rates and in order to keep under review distributions of adsorbed radioactivity. Concentrations of beta/gamma radioactivity and transuranics, in most cases at the same locations as the dose rate measurements, are given in Table 10. Variations similar in cause to those of the dose rates are observed, and comparison with results for 1981 (Hunt, 1983) shows generally similar concentrations, in line with the behaviour of dose rates. It is to be noted that these levels of radionuclide concentrations give rise to negligible exposure following inhalation of resuspended sediment (Pattenden *et al.*, 1981).

To identify those members of the public subject to the highest external exposures, occupancies of different locations need to be considered. We keep under review the amounts of time spent by members of the public on intertidal areas of coastline bordering the north-east Irish Sea, and, together with the consumption rates as described in section 4.1.1, these habits were re-surveyed in 1982. It is considered that, combining dose rates and occupancy times, the critical group for external exposure is represented by persons who live on board their boats in Whitehaven

Location	Ground type	No. of sampling observa- tions†	Mean gamma dose rate in air at l m, μGy h <sup>-]</sup>
Maryport harbour	Silt	4	0.34
Workington harbour		4	0.38
Whitehaven outer harbour		12	0.41
	Sand	12	0.28
Whitehaven yacht basin	Silt	12	0.79
Coulderton winkle beds	Rock	4	0.18
Braystones south	Sand	4	0.18
Sellafield		11	0.20
Drigg		4	0.18
Ravenglass - Salmon Garth		11	0.21
	Silt	12	0.53
	Mussel beds	12	0.55
Ravenglass - boats area	Sand	12	0.18
	Silt	12	0.36
Ravenglass - ford area		12	0.67
Newbiggin		13	0.86
Newbiggin - west of bridge	Sand/silt	12	0.45
"east""	Salt marsh	12	0.99
Haverigg	Sand	4	0.22
	Silt	4	0.33
Millom	Sand	4	0.11
	Silt	4	0.33
Walney Channel	Sand	4	0.22
	Silt	4	0.27
" west shore	Sand	4	0.098
Flookburgh	Sand	4	0.13
Fleetwood	"	4	0.097
Blackpool	**	4	0.088
Southport		4	0.089
New Brighton		4	0.088
Mersey (Rock Ferry)	Silt	4	0.19
Llandudno	Shingle	4	0.080
Prestatyn	Sand	4	0.066
Kippford - slipway	Silt	4	0.22
" - jetty	**	4	0.19
		4	0.28

Gamma radiation dose rates over intertidal areas of the Cumbrian coast and further afield, 1982 Table 9

†See section 3.3 for definition.

Table 10 Radioactivity in sediment from the Cumbrian coast and further afield, 1982

Sampling point and sediment typ	e		No. of sampling	Mea	an radioa	ctivit	y concentrati	on (dr	'y), Bq	kg-	1				
			tionst	Tot	al beta	<sup>60</sup> Co	952r + 95Nb	<sup>99</sup> Tc	103 Ru	100	<sup>6</sup> Ru	125Sb	<sup>134</sup> Cs	1	<sup>37</sup> Cs
Maryport	(sil	t)	4	13	000	33	2 300	NA	ND		700	89	210	5	600
Whitehaven	("	<u>)</u>	4	17	000	31	4 100	8.0	17	6	600	53	310	7	500
Newbiggin	ć"	)	4	20	000	66	6 000	NA	ND	11	000	200	260	6	300
Walney Island	("	)	4	5	100	17	1 100		••	2	600	37	74	1	900
Heysham	("	)	4	2	600	2.8	250				880	4.1	71	1	800
Sunderland Pt	("	)	4	1	800	0.5	24		••		310	ND	49	1	300
Fleetwood	(san	d)	4		320	ND	ND			ND			4.7		110
Blackpool	("	)	4		350			•		•• .			5.4		130
New Brighton	("	)	4		260		**			••			1.7		77
Rock Ferry	(sil	t)	4	3	200	3.2	••				230	••	69	2	300
Garlieston	("	)	4	1	700	4.0	66				420	0.9	34		<b>9</b> 10
Kippford slipway	( "	)	4	3	600	8.8	280		9.0	1	100	12	88	2	100
" merse	("	)	3	4	600	9.2	580		16	1	400	21	83	2	200
" jetty	("	)	4	3	000	6.2	90		ND		660	5.9	58	1	600

Sampling point and sediment type No. of sampling Mean radioactivity concentration (dry), Bq  $kg^{-1}$ 

				observa-								
				tionst	<sup>144</sup> Ce	154Eu	155 <sub>Eu</sub>	238 Pu	239 <sub>Pu</sub> + 240 <sub>Pu</sub>	241 Am	242 <sub>Cm</sub>	243 Cm + 244 Cm
Maryport	(s	il	t)	4	470	92	86	280	1 100	1 200	ND	5.0
Whitehaven	(	••	)	4	660	99	95	330	1 400	1 200	5.0	7.3
Newbiggin	(	••	)	4	1 300	230	180	680	2 700	2 500	17	13
Walney Island	(	••	)	4	270	51	37	NA	NA	570	NA	NA
Heysham	(		)	4	88	11	7.9	44	200	200	ND	0.89
Sunderland Pt	(	••	)	4	3.5	4.3	1.8	NA	NA	86	NA	NA
Fleetwood	(s.	and	1)	4	ND	ND	ND	0.9	4.9	4.1	ND	0.009
Blackpool	(	**	)	4				NA	NA	2.5	••	NA
New Brighton	(	••	)	4					•	2.2	NA	
Rock Ferry	(s:	il	t)	4		5.0	••	43	190	240	ND	0.78
Garlieston	(	••	)	4	27	5.3	4.4	21	97	130		0.41
Kippford slipway	7 ( '	••	)	4	63	21	19	NA	NA '	350	NA	NA
" merse	(	••	)	3	75	24	20	••		330	••	
" jetty	(		)	4	38	9.8	11	45	210	240	ND	0.95

NA = not analysed. ND = not detected. †See section 3.3 for definition.

harbour. Taking account of the time the boats are shielded from the mud by tidal effects and the shielding afforded by the boats themselves, their exposure is equivalent to that from spending 650 h year<sup>-1</sup> over unshielded mud. From Table 9, making an allowance for natural background, their external exposure in 1982 was 8% of the ICRP-recommended limit for members of the public. This result makes use (section 3.4) of the factor of 0.87 Sv Gy<sup>-1</sup> to convert absorbed dose rate 'free-in-air' to effective dose equivalent rate (Spiers et al., 1981). These persons also consume fish and shellfish, and an addition is necessary to derive total exposure related to Sellafield liquid discharges; other exposure pathways, such as handling of fishing gear, are negligible by comparison. This addition is estimated to be 4% of the ICRP-recommended dose limit for members of the public, on the basis of both values of gut uptake factor for plutonium described in section 3.4. Total exposure of the externally exposed critical group is thus estimated to be 12% of the appropriate ICRP-recommended limit. This exposure is less than that of the critical group of fish and shellfish consumers given earlier.

The converse situation, of the critical group of fish and shellfish consumers also receiving exposure from external pathways, also needs to be considered. Habits survey data indicate, however, that the external component is too small to make a significant difference to the result for their exposure already given in section 4.1.1; additions of this small order are considered to be adequately taken into account by the maximising process of summing exposures from the consumption of fish, crustaceans and molluscs.

### 4.1.3 Porphyra/laverbread pathway

No harvesting of *Porphyra* in the Sellafield vicinity for ultimate consumption was reported in 1982; this pathway has therefore remained essentially dormant. However, in view of its potential importance and the value of *Porphyra* as an indicator, monitoring has continued. Samples of *Porphyra* are regularly collected from selected locations along the Cumbrian coast. Results of analyses for 1982 are presented in Table 11. Samples of laverbread from the product of the major manufacturers are regularly collected from markets in South Wales and analysed for ruthenium-106. Results for 1982 are presented in Table 12. Because of the low concentrations, results are given as limits of detection. The exposure of critical individuals was less than 0.1% of the ICRP-recommended dose limit, confirming the virtual abeyance of this pathway.

#### 4.1.4 Other surveys

In addition to the monitoring described above which is related to the more (or potentially more) significant radiation exposure pathways as a consequence of Sellafield

Table 11 Radioactivity in Porphyra from the Cumbrian coast, 1982

Sampling point	No. of sampling	Mean radioactivity concentration (wet), Bq kg $^{-1}$										
	observa- tions†	Total	beta	<sup>60</sup> Co	<sup>95</sup> Zr + <sup>95</sup> Nb	103 <sub>Ru</sub>	106 R	125	Sb <sup>134</sup> Cs	<sup>137</sup> Cs		
Braystones South Seascale	12 29*	5 100 NA		1.8	320 100	64 19	4 60 2 70	0 6.7 0 4.7	9.8 6.5	160 120		
Sampling point	Mean r	Mean radioactivity concentration (wet), Bq $kg^{-1}$										
	observa- tions†	<sup>144</sup> Ce	154Eu	238Pu	$2^{39}Pu + 24$	<sup>0</sup> Pu 2	4 <sup>1</sup> Am	242 Cm	243 Cm + 2	<sup>44</sup> Cm		
Braystones South Seascale	12 29*	30 9.2	2.0	7.1 NA	29 NA	2	6 5	0.67 NA	0.18 NA			

NA = not analysed.

<sup>†</sup>See section 3.3 for definition.

\*These samples are counted wet to provide a rapid indication.

Manufacturer	No. of sampling	<sup>106</sup> Ru concentra- tion (wet)			
	tions†	Bq kg <sup>-1</sup>			
A	3	< 4			
В	4	< 9			
С	4	< 9			

Table 12 Radioactivity in laverbread from South Wales, 1982

<sup>†</sup>See section 3.3 for definition.

discharges, we undertake a number of further investigations. Some of these are of a research nature; however, they also enable pathways of lower current importance to be kept under review.

Seaweeds are useful indicator materials; they may concentrate certain radionuclides so that they greatly facilitate measurement and assist in the tracing of these radionuclides in the environment. Table 13 presents the results of measurements in 1982 on seaweeds from UK shorelines of the Irish Sea. Radioactivity concentrations in *Porphyra* 

Type of seaweed and sampling point	No. of sampling	M	ean radioa	ctivit	y conc	entration (we	et), Bq	kg <sup>-1</sup>	
	tions†	T	otal beta	54Mn	<sup>60</sup> Co	$95_{Zr} + 95_{Nb}$	<sup>99</sup> Tc	<sup>103</sup> Ru	106 <sub>Ru</sub>
Porphyra							<u> </u>		
Larbrax	4		230	ND	ND	ND	NA	ND	45
Port William	4		230	••				**	52
Garlieston	4		240		••	"	••	"	72
Fucus vesiculosus									
Sellafield	12	4	200	0.2	16	570	3 500	2.2	540
Heysham	4		860	ND	0.3	ND	NA	ND	12
Port William	4		750	••	0.4	••			5.2
Garlieston	4		730		0.8	22			36
Auchencairn	4	1	100	"	1.0	24		••	17
Portrush	4		340	••	ND	ND	**		ND
Rhodymenia palmata									
Millisle	2	1	100	••	••	••			19
Strangford Lough	1		670	••	••		••	"	ND
Ascophyllum nodosum Ardglass	2		550	••					
Fucus spiralis Ardglass	2		330		••			••	

Table is Radioaccivity in beawcoab from on bioterines of the fitbh bear i	Table 13	Radioactivity	in	seaweeds	from	UK	shorelines	of	the	Irish	Sea,	1
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Type of seaweed and sampling point	No. of sampling	Mean ra	Idioacti	vity co	ncentrati	on (wet)	, Bq kg	5-1	
	observa- tions†	110m <sub>Ag</sub>	125Sb	<sup>134</sup> Cs	137 <sub>Cs</sub>	<sup>144</sup> Ce	<sup>154</sup> Eu	155 <sub>Eu</sub>	241 Am
Porphyra									
Larbrax	4	ND	ND	0.3	15	ND	ND	ND	0.4
Port William	4		••	0.9	21				0.5
Garlieston	4		2.8	1.0	31	••		**	1.7
Fucus vesiculosus									
Sellafield	12	7.1	0.5	47	<b>89</b> 0	11	0.3	0.8	19
Heysham	4	ND	ND	12	280	ND	ND	ND	1.2
Port William	4			3.9	83	••	••		0.4
Garlieston	4		0.7	6.3	150			н	4.8
Auchencairn	4		0.4	9.6	220	0.5	••	••	4.9
Portrush	4		ND	0.2	11	ND		n	ND
Rhodymenia palmata									
Millisle	2			2.0	54	••		н	
Strangford Lough	1	"	••	1.9	42		••		
Ascophyllum nodosum Ardglass	2			0.8	22		••		
Fucus spiralis									
Ardglass	2			1.2	32				

NA = not analysed. ND = not detected. †See section 3.3 for definition.

are reported in this section for areas relatively remote from Sellafield because of the value of this seaweed as an indicator, particularly for ruthenium-106. Although small quantities of *Porphyra* and *Rhodymenia palmata* from these locations may be eaten, radioactivity concentrations are of negligible radiological significance. *Fucus* seaweeds are also useful indicators particularly of fission product radioactivity other than from ruthenium-106; samples of *Fucus vesiculosus* are collected both in the Sellafield vicinity and further afield, and the results are presented here. Monitoring in Scotland is carried out on behalf of departments of the Scottish Office. Analyses of samples collected in Northern Ireland are carried out on behalf of DOE(NI).

#### 4.2 Springfields, Lancashire

This establishment is mainly concerned with manufacture of fuel elements for nuclear reactors and production of uranium hexafluoride. Radioactive waste arisings are small and consist mainly of uranium and its daughter products; liquid discharges are made by pipeline to the Ribble Estuary. Public radiation exposure in this vicinity as a result of these discharges is very low; there is, however, a greater contribution due to Sellafield discharges. The critical pathway is external exposure, due to adsorption of radioactivity on the muddy areas of river banks. The amounts of time for which members of the public are subject to such exposure is kept under review. In 1982 the critical group consisted of people who live on houseboats moored in muddy creeks of the Ribble Estuary. We regularly monitor dose rates in relevant areas; Freckleton is typical of an area of muddy creeks where houseboats are moored. Dose rates are also monitored close to the Springfields outfall, and these measurements are supported by analyses of mud.

Results for 1982 are shown in Table 14. The only detectable radionuclide due to Springfields discharges is protactinium-234m; other radionuclides present are mainly from Sellafield. Exposure of the critical group of houseboat dwellers in 1982, including the Sellafield component, was about 7% of the ICRP-recommended dose limit, similar to the result for 1981 (Hunt, 1983). The exposure is mainly due to Sellafield discharges; the contribution due to Springfields would have been a small fraction of the total.

Table 14(a) Radioactivity in sediment near the Springfields pipeline, 1982

Location	No. of sampling observa- tions†	Mean radioactivity concentration (dry), Bq kg <sup>-1</sup>										
		Total beta	60Co	$95_{Zr} + 95_{Nb}$	106 <sub>Ru</sub>	125 SĐ	<sup>134</sup> Cs	<sup>137</sup> Cs	<sup>144</sup> Ce			
Pipeline outlet	4	42 000	14	180	940	45	190	4 200	87			

Location	No. of sampling observa- tions†	Mean r	Mean radioactivity concentration (dry), Bq kg <sup>-1</sup>									
		<sup>154</sup> Eu	234mPa	238 <sub>Pu</sub>	$239_{Pu} + 240_{Pu}$	241 Am	242Cm	243 Cm + $244$ Cm				
Pipeline outlet	4	17	130 000	93	380	430	2.0	1.2				

†See section 3.3 for definition.

Table 14(b)	Gamma dose rates in air at
	l m over intertidal areas
	near the Springfields pipe-
	line, 1982

No. of sampling observa- tions†	µGy h <sup>−1</sup>
4	0.34
4	0.31
4	0.25
	No. of sampling observa- tions† 4 4 4

<sup>†</sup>See section 3.3 for definition.

#### 4.3 Capenhurst, Cheshire

The main function of the Capenhurst Works is enrichment of uranium. Radioactive waste arisings, mainly of uranium and its daughter products, are very small; the Works have an authorisation to dispose of liquid wastes to the Rivacre Brook. Recently, uranium recovered from irradiated fuel has been recycled; this may contain small quantities of fission products, of which technetium-99 is the only component of potential significance. Waste arisings in this second category are again very low; their disposal to Liverpool Bay from the North Wirral outfall at Meols is regulated by authorisation. It is not expected that the environmental consequences of these small disposals would be detectable above background levels due both to

ſable 15	Radioactivity i	n environmental	. materials	in	the	vicinity	of	the	Wirral,	198	32
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laterial	Sampling point	No. of sampling	Mean radioa	ctivity	concen	tration	(wet)*	, Bq kg	-1	
		tions†	Total beta	99Tc	106 Ru	125 Sb	<sup>134</sup> Cs	137Cs	155 <sub>Eu</sub>	241 Am
Shrimps	Hoylake	2	210	8.4	ND	ND	3.6	87	ND	ND
Cockles	Hoylake Dee Estuary	2 1	140 44	2.7 1.2	3.0 6.7	 	1.0 0.7	41 25	0.3 ND	1.8 2.0
Fucus spiralis "	Hoylake Little Orme	3 2	3 <b>9</b> 0 430	92 220	ND	0.5 ND	4.5 2.4	100 52	••	0.5 ND
Sand	Hoylake	3	260	5.2	••		1.3	80		••

ND = not detected.

\*Except for sand where dry concentrations apply.

†See section 3.3 for definition.

natural sources of radioactivity and to Sellafield discharges. However, we have established an environmental monitoring programme which reflects the potentially critical pathway due to consumption of locally-caught shellfish. Fucus-type seaweed is also sampled, being a good indicator for technetium-99. It is to be noted that the programme is much more extensive than is technically justified by the potential radiological hazard from Capenhurst discharges.

Results for 1982 are presented in Table 15. The concentrations of artificial radioactivity are mainly due to Sellafield discharges and are consistent with values to be expected at

this distance from Sellafield. Technetium-99 concentrations were lower than in 1981 (Hunt, 1983), reflecting the much reduced discharges of technetium-99 from Sellafield since the latter part of 1980, when decay-stored liquors were released. Discharges of technetium-99 from Capenhurst were reduced in 1982 as compared with 1981. Exposure of critical shellfish consumers in the vicinity of the Wirral in 1982 amounted to less than 2% of the ICRPrecommended dose limit; this was mainly due to radiocaesium and transuranic nuclides from Sellafield. Only a tiny fraction of this exposure was due to technetium-99, which was almost entirely from Sellafield discharges.

Radioactivity in environmental materials in the vicinity of Chapelcross, 1982 Table 16(a)

faterial	Sampling point	No. of sampling	Mean radioa	g <sup>-1</sup>							
		observa- tions†	Total beta	<sup>60</sup> Co	$95_{Zr} + 95_{Nb}$	103 Ru	106 Ru	125 SB	<sup>134</sup> Cs	137 Cs	<sup>144</sup> Ce
Flounder	Seafield	3	380	ND	ND	ND	ND	ND	11	300	ND
Salmon	••	1	130	•				"	ND	4.1	
Shrimps	"	3	200	•			3.8		6.3	140	"
Fucus vesiculosus	Waterfoot	4	750	••			8.6		7.9	210	
	Seafield	4	800	1.0		"	20	"	9.6	240	0.9
Sediment	"	4	2800	8.5	480	14	1100	2.9	8.2	2200	55

laterial	Sampling point	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg <sup>-1</sup>								
		observa- tions†	<sup>154</sup> Eu	155 <sub>Eu</sub>	238 <sub>Pu</sub>	239 <sub>Pu</sub> + <sup>240</sup> Pu	241 Am	242 <sub>Cm</sub>	243 Cm + 244 Cm		
Flounder	Seafield	3	ND	ND	NA	NA	ND	NA	NA		
Salmon		1	•	"	0.00014	0.00056	0.00053	ND	ND		
Shrimps		3	•		NA	NA	ND	NA	NA		
Fucus vesiculosus	Waterfoot	4					2.5				
	Seafield	4		0.3			5.9		••		
Sediment	••	4	13	11		•	200				

ND = not detected. NA = not analysed.

\*Except for sediment where dry concentrations apply.

†See section 3.3 for definition.

	the vicinity of 1982	Chapelcross,
Location	No. of sampling observa- tions†	µGy h <sup>−1</sup>
Seafield	4	0.23
Browhouses	4	0.15
Waterfoot	4	0.13
Torduff Point	4	0.12
Battle Hill	4	0.13

Gamma dose rates in air at

<sup>†</sup>See section 3.3 for definition.

#### 4.4 Chapelcross, Dumfriesshire

Table 16(b)

At this establishment BNFL operates a magnox-type nuclear power station. Liquid waste arisings are discharged to the Solway Firth under authorisation of the Scottish Development Department. Discharges increased in 1982 as compared with 1981, due to emptying of a fuel element storage pond, but were still well within authorised limits. Pathways leading to public radiation exposures were reinvestigated during 1982. Two pathways were confirmed as being of potential importance. These were internal irradiation from consumption of locally-caught fish and shellfish and external exposure from use of intertidal areas by fishermen and turf cutters; fishermen continue to constitute a critical group in view of their regular occupancy of intertidal areas and consumption of local catches. Our monitoring, which is carried out on behalf of departments of the Scottish Office, continued to reflect these pathways. Samples of Fucus vesiculosus, as a useful indicator, are also analysed. The results of monitoring in 1982 are presented in Table 16.

Concentrations of artificial radionuclides in the Chapelcross vicinity are mostly due to Sellafield discharges, and the general levels given in Table 16(a) are consistent with values to be expected at this distance from Sellafield. Radio-caesium concentrations in 1982 were generally similar to those in 1981. Exposure of the critical group in 1982, making the maximising assumption of additivity of the two pathways, amounted to less than 5% of the ICRP-recommended dose limit. The magnitude of the Chapelcross discharges indicate that the local contribution would have been a tiny fraction of this exposure; most is due to Sellafield discharges.

#### 5. United Kingdom Atomic Energy Authority

We regularly monitor the environmental impact of liquid radioactive discharges from two UKAEA sites. These are the Atomic Energy Establishment, Winfrith and the Dounreay Nuclear Power Development Establishment. Liquid radioactive wastes also arise at the Atomic Energy Research Establishment, Harwell. In common with such wastes from other nuclear establishments in the Thames Valley area, these are discharged into the River Thames, and the critical exposure pathway is from drinking water. Monitoring in respect of these discharges is therefore carried out by the Department of the Environment rather than this Ministry.

#### 5.1 Atomic Energy Establishment, Winfrith, Dorset

The principal installation at which liquid radioactive wastes arise at this establishment is the Steam Generating Heavy Water Reactor. Most of the activity is due to tritium from the moderator and coolant, but small amounts of activation products, including manganese-54, cobalt-60 and zinc-65, are removed during decontamination of the reactor pressure circuit. These wastes are disposed of under authorisation to deep water in Weymouth Bay. It is the activation products rather than tritium which are of greater, but still small, environmental significance. Reconcentration of activation products by shellfish, followed by local consumption, constitutes the critical exposure pathway; this is reflected in our monitoring programme. Monitoring of the indicator material, Fucus serratus, provides additional information on the distribution of activation products. Data are presented in Table 17.

The impact of Winfrith discharges was, as in previous years, mainly observed in the activation product concentrations. Radiocaesium concentrations were similar to those to be expected from fallout; local discharges were likely to give rise to a negligible contribution. In 1982 the total radiation dose to critical consumers near this establishment was low, at most about 1% of the ICRP-recommended dose limit.

5.2 Dounreay Nuclear Power Development Establishment, Caithness

Liquid radioactive waste discharges from this establishment are made to the Pentland Firth under authorisation of the Scottish Development Department. Discharges include a minor contribution from the adjoining reactor site (Vulcan Naval Nuclear Propulsion Test Establishment) operated by the Ministry of Defence (Procurement Executive). Reprocessing of Prototype Fast Reactor (PFR) fuel has taken place since 1980. Our monitoring near Dounreay is carried out on behalf of departments of the Scottish Office.

There are two critical exposure pathways, both involving external radiation. The first pathway is due to radioactivity adsorbed mainly on fine particulate matter becoming

Table 17	Radioactivity	in environmental	materials	from the	vicinity of	Winfrith,	1982
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Material	Sampling point	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$									
		tions†	Total beta	54Mn	<sup>58</sup> Co	<sup>60</sup> Co	65 Zn	137 <sub>Cs</sub>	<sup>238</sup> Pu			
Whiting	Lulworth	1	100	ND	ND	1.9	18	1.0	NA			
Crabs		2	78		"	57	170	ND	0.0020			
Oysters	Poole	2	61		••	4.8	130	•	0.0016			
Fucus	Kimmeridge	2	380	34	17	460	46		NA			
serratus	Osmington Mills	2	190	12	4.9	230	25					
	Weymouth	2	330	13	4.5	250	24		••			
	Portland	2	300	7.4	2.4	100	6.4	ND				
	Swanage	2	380	8.5	5.2	310	26					
	Hengistbury Head	2	2 <b>9</b> 0	3.2	2.0	240	13					
	Bognor Regis	1	230	ND	ND	46	10	н				
	Sandgate	1	280	ND		16	ND	1.1				
Fucus vesiculosus	Bognor Regis	1	190	0.4		22		0.3				
Sediment	Poole Harbour	2	500	12	"	120	"	13				

Material	Sampling point	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg <sup>-l</sup>							
		tions†	239Pu + 240Pu	241 Am	242 <sub>Cm</sub>	<sup>243</sup> Cm + <sup>244</sup> Cm				
Whiting	Lulworth	1	NA	ND	NA	NA				
Crabs		2	0.0070	0.014	0.00046	0.00020				
Oysters	Poole	2	0.0061	0.019	0.0013	0.00052				
Fucus	Kimmeridge	2	NA	ND	NA	NA				
serratus	Osmington Mills	2			••					
	Weymouth	2		••	••					
	Portland	2								
	Suapago	2								
	Hengistbury Head	2	••							
	Bognor Pogie	1				н				
	Sandgate	1		••	"					
Fucus vesiculosus	Bognor Regis	1		•						
Sediment	Poole Harbour	2	"			••				

Mean gamma dose rate in air at 1 m over intertidal sediments in Poole Harbour (4 sampling observations†): 0.075  $_\mu Gy~h^{-1}$ 

ND = not detected. NA = not analysed. \*Except for sediment where dry concentrations apply. †See section 3.3 for definition.

Table 18 Radioactivity in environmental materials from the vicinity of Dounreay, 1982

Sampling point and material	No. of sampling	Mean ra	Mean radioactivity concentration (wet), Bq kg $^{-1}$										
	observa- tions†	Total	beta	54Mn	60Co	95Z	r + <sup>95</sup> Nb	106 Ru	110mAg	g 125 Sb	<sup>134</sup> Cs	<sup>137</sup> Cs	
Sandside Bay													
Winkles	3	210		0.2	3.1	ND		68	59	2.8	0.6	6.4	
Limpets	11	2 <b>9</b> 0		0.04	1.9			150	21	6.2	ND	6.4	
Fucus vesiculosus	11	410		3.7	8.2	70		14	7.4	0.4	0.4	15	
Sampling point and material	No. of sampling	Mean r	adioac	tivit	/ cond	centr	ation (we	et), 3q	kg <sup>-1</sup>				
	observa- tions†	<sup>144</sup> Ce	154Eu	155	2 2	<sup>38</sup> Pu	239Pu +	240 Pu	241 Am	242 <sub>Cm</sub>	<sup>243</sup> Cm +	244 Cm	
Sandside Bay													
Winkles	3	34	ND	ND	0	.64	1.40		2.3	0.16	0.011		
Limpets	11	40	0.2	0.7	1	.3	3.2		7.8	0.69	0.026		
Fucus vesiculosus	11	21	0.2	0.4	N	A	NA		4.6	NA	NA		

ND = not detected. NA = not analysed. †See section 3.3 for definition.

entrained on fishing gear which is regularly handled. This results in skin dose, mainly from beta particles, to the hands and forearms of fishermen. The critical group is a small number of people who operate a salmon fishery from Sandside Bay, close to Dounreay. Our regular measurements prior to 1981 have shown that at current rates of discharge the average dose rates on nets will be low, such that monitoring by the UKAEA is sufficient. Its surveys (Flew, 1983) have confirmed that the exposure of these fishermen was low, at less than 1% of the ICRPrecommended dose limit.

The second critical pathway arises also from the uptake of radioactivity by particulate material which accumulates in rocky areas of the foreshore and presents a potential source of exposure mainly to gamma radiation of those who visit these areas. Monitoring of foreshore dose rates is also carried out by the UKAEA (Flew, 1983). Public radiation exposure via this pathway was low, at less than 1% of the ICRP-recommended dose limit.

We sample winkles from Sandside Bay to enable the subcritical pathway of shellfish consumption to be kept under direct review. Additionally, as in previous years, limpets and *Fucus vesiculosus* were sampled as indicator materials. Results are presented in Table 18. Radiocaesium concentrations are mostly due to discharges from Sellafield. Other radionuclides detected, including transuranics, mainly reflect Dounreay discharges. Concentrations were generally similar to those in 1981, except for increases in ruthenium-106 and silver-110m, due to small increases in discharges of these nuclides within the total activity discharged under authorisation, and a decrease in americium-241 in *Fucus vesiculosus*, reflecting reduced discharges of this nuclide; consequent reductions of americium-241 in shellfish are expected later. The radiological significance of shellfish consumption continued to be low; for high-rate winkle consumers the radiation dose was less than 0.4% of the ICRP-recommended dose limit. This pathway therefore remained of sub-critical importance.

## 6. Nuclear power stations operated by the electricity boards

All but one of these power stations are in England and Wales and are operated by the Central Electricity Generating Board. The Scottish power station at Hunterston is operated by the South of Scotland Electricity Board. Results are also presented for two power stations not operational in 1982, namely Hartlepool and Heysham, where our monitoring had already commenced.

#### 6.1 Berkeley, Gloucestershire and Oldbury, Avon

Liquid radioactive wastes from both of these stations are generally similar in composition and are discharged to the same stretch of the Severn Estuary. The stations are therefore considered together for the purpose of our environmental monitoring. The two critical pathways for public radiation exposure are internal irradiation following consumption of locally-caught fish and shellfish, and external exposure from occupancy of muddy intertidal areas. We therefore analyse samples of fish and shellfish and monitor beach gamma dose rates. In addition, measurements of external exposure are supported by analyses of intertidal mud, and *Fucus vesiculosus* is collected as an indicator material.

Data for 1982 are presented in Table 19. The only artificial radioactivity detected in fish and shellfish was due to radiocaesium. Concentrations of radiocaesium represent the combined effect of discharges from the stations and fallout,

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$									
	observa- tions†	Total beta	54 <sub>Mn</sub>	<sup>60</sup> Co	<sup>134</sup> Cs	137 <sub>Cs</sub>	<sup>144</sup> Ce	155Eu			
Flounders	2	67	ND	ND	ND	2.4	ND	ND			
Mullet	1	160	••			2.4		**			
Eels	2	80	"	**		0.3	"	••			
Shrimps	2	100		"		2.7					
Fucus vesiculosus	2	210		1.1	0.6	14					
Mud: area of outfalls area upstream	3 3	700 860	0.4 ND	ND "	1.8 2.5	76 70	" 3.0	1.0 1.2			

Table 19Radioactivity in environmental materials and gamma dose rates near Berkeley and<br/>Oldbury nuclear power stations, 1982

Mean gamma dose rate in air at 1 m over intertidal mud (9 sampling observations†): 0.11  $\mu Gy~h^{-1}$ 

ND = not detected.

\*Except for mud where dry concentrations apply.

<sup>†</sup>See section 3.3 for definition.

and possibly include a small Sellafield-derived component, but apportionment is difficult at the low levels detected. Public radiation exposure, however, was very low, at less than 0.1% of the ICRP-recommended limit to the critical group of fish and shellfish consumers. Very small concentrations of other artificial radionuclides, in addition to radiocaesium, were detected in mud and seaweed but taken together were of negligible radiological significance. Directly-measured gamma dose rates over intertidal mud continued to be indistinguishable from the natural background.

#### 6.2 Bradwell, Essex

Radioactive liquid effluent from this power station is discharged to the estuary of the River Blackwater. There are two critical pathways, via consumption of locallycaught fish and shellfish, and external exposure of people who live in houseboats moored in muddy areas of the estuary. Our environmental monitoring reflects these pathways. Gamma dose rate measurements are supported by analyses of intertidal mud, and *Fucus vesiculosus* is analysed as an indicator material.

Measurements for 1982 are summarised in Table 20. In fish, the only artificial radioactivity detected was due to radiocaesium, for which concentrations represent the combined effects of discharges from the station, Sellafield discharges and fallout. Apportionment is difficult because of the low levels detected. The dose to members of the critical group of fish and shellfish consumers, however, was low, totalling less than 0.3% of the ICRP-recommended dose limit for members of the public. The concentrations of zinc-65 and transuranic nuclides in oysters were low, such that the contributions to dose from these nuclides remained small. Concentrations of artificial radionuclides detected in mud and seaweed were also low and of negligible radiological significance. Gamma dose rates, as directly measured, were indistinguishable from the natural background.

#### 6.3 Dungeness, Kent

There are two, essentially separate, "A" and "B" nuclear power stations at this establishment: the "A" station is powered by magnox-type reactors and the "B" station by AGRs. Liquid radioactive waste discharges from the "B"

Table 20Radioactivity in environmental materials and gamma dose rates nearBradwell nuclear power station, 1982

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg <sup>-1</sup>										
	observa- tions†	Total beta	<sup>60</sup> Co	<sup>65</sup> Zn	106 <sub>Ru</sub>	125Sb	<sup>134</sup> Cs	137Cs				
Mixed flatfish	2	120	ND	ND	ND	ND	0.2	4.8				
Oysters	2	80		4.7			ND	2.5				
Whelks	1	110	0.5	ND		1.9	**	1.4				
Fucus vesiculosus	2	320	1.5		••	ND	0.6	12				
Sediment	4	640	5.0		9.4	••	5.0	52				

Material	No. of sampling	Mean ra	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$								
	observa- tions†	238 <sub>Pu</sub>	$239_{Pu} + 240_{Pu}$	241 <sub>Am</sub>	242Cm	243Cm + <sup>244</sup> Cm					
Mixed flatfish	2	NA	NA	ND	NA	NA					
Ovsters	2	0.0056	0.018	0.042	0.0016	0.0016					
Whelks	1	NA	NA	ND	NA	NA					
Fucus vesiculosus	2	**	••								
Sediment	4	••				**					

Mean gamma dose rate in air at 1 m over intertidal sediments (8 sampling observations†): 0.084  $\mu Gy\ h^{-1}$ 

ND = not detected.

NA = not analysed.

<sup>\*</sup>Except for sediment where dry concentrations apply.

<sup>†</sup>See section 3.3 for definition.

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$										
	tions†	Total beta	54Mn	<sup>60</sup> Co	65 <sub>Zn</sub>	106 <sub>Ru</sub>	125 <sub>Sb</sub>	<sup>134</sup> Cs	137 Cs	155 <sub>Eu</sub>		
Plaice Whelks	3	160 120	ND 	ND 4.4	ND 6.2	ND 7.2	ND 2.4	ND 	2.5 ND	ND 		
Fucus serratus Sand	2 2	280 290	0.5	16 6.5	ND 	ND 4.9	ND "	0.3	2.3	0.5		

Table 21 Radioactivity in environmental materials and gamma dose rates near Dungeness nuclear power station, 1982

Mean gamma dose rate in air at 1 m over intertidal sediment (10 sampling observations†): 0.057  $\mu Gy~h^{-1}$ 

ND = not detected.

\*Except for sand where dry concentrations apply.

<sup>†</sup>See section 3.3 for definition.

station, which began raising power at the end of 1982, were very small indeed (see Table 1). Discharges from both "A" and "B" stations are made via the same outfall and for the purposes of our environmental monitoring are considered together. There are two critical radiation exposure pathways as a result of liquid radioactive waste discharges: internal irradiation due to consumption of locally-caught fish, and external exposure from occupancy of the foreshore. Our monitoring programme therefore includes analyses of fish and shellfish and gamma dose rate surveys of the generally sandy beach. Samples of sand are also collected and analysed. Local whelks and seaweed have been analysed mainly for their value as indicator materials. The results for 1982 are given in Table 21.

Concentrations of caesium-137 in plaice are attributable to discharges from the station and from Sellafield, with a small contribution due to fallout. Apportionment is difficult at these low levels. The radiation dose to members of the critical group of fish consumers was very low, at less than 0.1% of the ICRP recommended dose limit. Gamma dose rates over sand were indistinguishable from natural background. Whelks, seaweed and sand all showed trace levels of cobalt-60, and whelks showed trace levels of zinc-65. The indicator sampling programme described in section 5.1 shows that AEE Winfrith rather than Dungeness may be the source of these nuclides. Trace amounts of ruthenium-106 were also detected in whelks. Our monitoring programme in the Channel Islands (section 9) shows that the French reprocessing plant at Cap de la Hague may be the source of this nuclide. The concentrations of radionuclides in whelks were, however, of negligible radiological significance.

### 6.4 Hartlepool, Cleveland

This station, with its two AGRs, was not yet operational in 1982. However, our monitoring had already begun in order to investigate background levels and to establish reliable sources of environmental materials. Potential critical pathways of radiation exposure for the public near this station, likely to be associated with liquid discharges, are internal irradiation following consumption of local fish and shellfish and external exposure from occupancy of intertidal areas. Collectors of small coal, which is washed ashore along this stretch of coast, account for the highest beach occupancies, but the highest external exposures are likely to be of fishermen who operate in muddy areas near the mouth of the Tees.

Results of our monitoring programme carried out in 1982 are shown in Table 22. Concentrations of radiocaesium and transuranics were due to discharges from Sellafield and to fallout; the radiation exposure of the potentially critical group of local fish and shellfish consumers from these existing sources was low, at less than 0.4% of the ICRP-recommended dose limit for members of the public.

#### 6.5 Heysham, Lancashire

This establishment, which will comprise two, essentially separate, nuclear power stations both powered by AGRs, was under construction in 1982. Our monitoring had begun for similar reasons as for the station at Hartlepool; in addition, information on radiation exposures and on the distribution of a range of radionuclides as a result of Sellafield discharges is to be gained. The potential critical radiation exposure pathways from liquid radioactive discharges from Heysham are likely to be internal irradiation following consumption of locally-caught fish and shellfish (mainly shrimps and cockles) and external exposure from occupancy of intertidal areas. Our monitoring programme includes analyses of fish and shellfish and measurements of beach gamma dose rates. Samples of sediment are also analysed, and Fucus vesiculosus is monitored as an indicator material.

The results for 1982 are given in Table 23. These mainly reflect discharges from Sellafield; it is unlikely that the effect of future discharges from Heysham will be detectable

Table 22	Radioactivity in environmental materials and gamma dose rate	tes
	near Hartlepool nuclear power station, 1982	

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg <sup>-1</sup>								
	tions†	Total beta	<sup>134</sup> Cs	137 <sub>Cs</sub>	155 <sub>Eu</sub>	238 <sub>Pu</sub>				
Cod	4	140	0.3	11	ND	NA				
Crabs	2	58	ND	1.8	••	0.00059				
Fucus vesiculosus	4	260	••	5.5		NA				
Sand	4	150	••	3.6	**	••				
Mud	4	720	1.9	66	2.0	**				

Material	No. of
	sampling
	observa-

Mean radioactivity concentration (wet)\*, Bq kg<sup>-1</sup>

	tions $239_{Pu} + 240_{I}$		241 <sub>Am</sub>	242 <sub>Cm</sub>	<sup>243</sup> Cm + <sup>244</sup> Cm
Cod	4	NA	ND	NA	 NA
Crabs	2	0.0029	0.0015	ND	ND
Fucus vesiculosus	4	NA	ND	NA	NA
Sand	4				
Mud	4	**	••	••	

Mean gamma dose rate in air at 1 m over intertidal sediment (12 sampling observations†): 0.072  $\mu$ Gy h<sup>-1</sup>

NA = not analysed.

ND = not detected.

\*Except for sediments where dry concentrations apply.

†See section 3.3 for definition.

Radioactivity in environmental materials and gamma dose rates near Heysham nuclear power station, 1982 Table 23

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$									
	observa- tions†	Total beta	<sup>60</sup> Co	<sup>95</sup> Zr + <sup>95</sup> Nb	106 Ru	125 SB	<sup>134</sup> Cs	<sup>137</sup> Cs	<sup>144</sup> Ce		
Flounders	4	410	ND	ND	ND	ND	17	380	ND		
Shrimps	4	220					7.9	160			
Cockles	4	230	1.1		54	0.5	3.8	85	2.2		
Fucus vesiculosus	4	860	0.3		12	ND	12	280	ND		
Sediment:											
Sunderland Point	4	1800	0.5	24	310		49	1300	3.5		
Half Moon Bay	4	2600	2.8	250	880	4.1	71	1800	88		

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq $kg^{-1}$								
	tions†	<sup>154</sup> Eu	155 <sub>Eu</sub>	238 <sub>Pu</sub>	$239_{Pu} + 240_{Pu}$	241 <sub>Am</sub>	242 Cm	<sup>243</sup> Cm + <sup>244</sup> Cm		
Flounders	4	ND	ND	NA	NA	ND	NA	NA		
Shrimps	4		••	0.018	0.077	0.079	ND	0.00052		
Cockles	4	••	0.3	0.92	4.0	6.0	••	0.037		
Fucus vesiculosus	4	•	ND	NA	NA	1.2	NA	NA		
Sediment:										
Sunderland Point	4	4.3	1.8			86	••	••		
Half Moon Bay	4	11	7.9	••	11	220		••		

Mean gamma dose rate in air at 1 m over intertidal sediment: Heysham vicinity (12 sampling observations†): 0.15  $\mu$ Gy h<sup>-1</sup> Sunderland Point ( 4 sampling observations†): 0.12  $\mu$ Gy h<sup>-1</sup>

NA = not analysed. ND = not detected.

\*Except for sediments for which dry concentrations apply.

†See section 3.3 for definition.

above the Sellafield-derived background. Estimates of the radiation exposure in 1982 of members of the critical group of fish and shellfish consumers associated with commercial fisheries (which include the Morecambe Bay area) are given in section 4.1.1. External exposure of members of the public was less than 1% of the ICRP-recommended dose limit.

#### 6.6 Hinkley Point, Somerset

At this establishment there are two essentially separate "A" and "B" nuclear power stations; the "A" station is powered by magnox-type reactors and "B" station by AGRs. Liquid radioactive discharges are made via the same outfall and for the purposes of our environmental monitoring are considered together. There are two critical radiation exposure pathways associated with liquid radioactive waste discharges: consumption of locally-caught fish and shrimps gives rise to internal irradiation, while external exposure results from occupancy of the foreshore. Our monitoring programme includes analyses of locally-caught fish and shrimps. External exposure is monitored by means of gamma dose rate measurements, supported by analyses of sediment. In addition, *Fucus vesiculosus* is monitored as an indicator.

The results for 1982, presented in Table 24, indicate concentrations of radiocaesium representing the combined effect of discharges from the station and from

3

2

2

Sellafield, in addition to fallout. Apportionment is difficult in view of the low levels detected. The total radiation exposure of members of the critical group through the fish and shellfish pathway was low, at less than 0.2% of the ICRP-recommended dose limit. The concentrations in shrimps of transuranic nuclides from the station and from Sellafield were of negligible radiological significance. Gamma radiation dose rates over intertidal sediment close to the station were indistinguishable from the natural background.

#### 6.7 Hunterston, Ayrshire

This establishment also comprises "A" and "B" stations, of which the latter is powered by AGRs. Liquid radioactive waste discharges are made to the Firth of Clyde under authorisation of the Scottish Development Department. For various reasons, storage of irradiated magnox fuel resulted in higher than usual concentrations of radionuclides (mainly radiocaesium) in pond water. The authorisation for the "A" station was changed in 1980 to allow increased discharges of these radionuclides, and this authorisation was renewed in 1981 and again in 1982, each time for a further year (see Table 1). There are two critical radiation exposure pathways: fish and shellfish consumption leading to internal irradiation, and occupancy of intertidal areas leading to external exposure. We regularly monitor, on behalf of departments of the

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0.4

ND

0.1

0.6

4.4

2.6

4.8

48

0.3

10

1.1

NA

ກບ	clear power station	, 1982								
Material	No. of sampling	Mean radioa	ctivit	y conc	entrati	on (we	et)*, Bq	kg <sup>-1</sup>		
	tionst	Total beta	<sup>54</sup> Mn	<sup>60</sup> Co	106 <sub>Ru</sub>	<sup>90</sup> Sr	110mAg	<sup>134</sup> Cs	<sup>137</sup> Cs	<sup>144</sup> C
Flounders	3	115	ND	ND	ND	NA	ND	0.3	3.0	ND
Eels	1	91						0.2	7.6	

..

0.9

0.4

...

1.1

4.2

...

0.2

ND

110

300

630

Table 24 Radioactivity in environmental materials and gamma dose rates near Hinkley Point nuclear power station, 1982

Material	No. of sampling		Mean radioactivity concentration (wet)*, Bq kg <sup>-1</sup>								
	tions†	155 <sub>Eu</sub>	238Pu	$239_{Pu} + 240_{Pu}$	241 Am	242Cm	243 Cm + $244$ Cm				
Flounders	3	ND	NA	NA	ND	NA	NA				
Eels	1				••						
Shrimps	3		0.0066	0.020	0.018	0.014	0.0019				
Fucus vesiculosus	2		NA	NA	ND	NA	NA				
Sediment	2	0.8	••			**	"				

Mean gamma dose rate in air at 1 m over intertidal sediment (8 sampling observations†): 0.10  $\mu Gy~h^{-1}$ 

NA = not analysed.

Shrimps

Sediment

Fucus vesiculosus

ND = not detected.

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\*Except for sediment where dry concentrations apply.

<sup>†</sup>See section 3.3 for definition.

Table 25 Radioactivity in environmental materials and gamma dose rates near Hunterston nuclear power station, 1982

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$										
	observa- tions†	Total beta	<sup>54</sup> Mn	<sup>60</sup> Co	<sup>65</sup> Zn	<sup>95</sup> Zr + <sup>95</sup> Nb	106 Ru	110 <b>mAg</b>	125 Sb	<sup>134</sup> Cs	137 Cs	
Cod	4	190	ND	ND	ND	ND	ND	ND	ND	3.2	79	
Turbot (fish farm A)	6	130								3.4	37	
Eel (fish farm B)	2	78		0.3	"				"	1.8	16	
Cockles	3	240	1.0	8.1	3.3		99	2.5	1.1	9.6	42	
Winkles	4	280	0.6	15	3.9		150	13	1.8	12	52	
Fucus vesiculosus	1	750	7.7	25	10	19	76	2.4	ND	33	150	
Fucus spiralis	3	610	5.0	24	11	ND	43	1.4	••	20	96	
Sand	4	340	0.6	3.6	ND		26	ND	"	12	86	

No. of Mean radioactivity concentration (wet)\*, Bq kg<sup>-1</sup>

	observa								
	tionst	<sup>144</sup> Ce	154Eu	155 <sub>Eu</sub>	238 <sub>Pu</sub>	239 <sub>Pu</sub> + 240 <sub>Pu</sub>	241 Am	242Cm	243Cm + 244Cm
Cod	4	ND	ND	ND	NA	NA	ND	NA	NA
Turbot (fish farm)	6		н			•	••	**	"
Eel (fish farm)	2								0
Cockles	3	49			0.31	0.52	0.41	1.9	0.14
Winkles	4	48		0.4	NA	NA	ND	NA	NA
Fucus vesiculosus	1	110	2.9	1.9			0.8		
Fucus spiralis	3	24	ND	ND			ND		
Sand	4	53	0.9	0.9			"		

Mean gamma dose rate in air at 1 m over intertidal sediment (12 sampling observations†): 0.11  $\mu$ Gy h^{-1}

NA = not analysed.

Material

ND = not detected.

\*Except for sand where dry concentrations apply.

See section 3.3 for definition.

Scottish Office, samples of fish and shellfish and carry out gamma dose rate measurements on the foreshore. Samples of sand are analysed together with *Fucus spiralis* as indicators. The results of monitoring in 1982 are shown in Table 25.

The concentrations of artificial radioactivity in this area are predominantly due to Sellafield discharges, the general values being consistent with those to be expected at this distance from Sellafield. However, the resulting public radiation exposure in 1982 was low, at less than 2% of the ICRP-recommended dose limit to members of the critical group of fish and shellfish consumers. Radiocaesium concentrations detected in fish from farms which are supplied by station cooling water were lower than in fish caught in the open sea; this is because the farmed fish are fed on manufactured food brought from further afield. No effects were detectable from the increased level of radiocaesium discharges. The concentrations of activation products observed in molluscs, seaweed and sand were due to discharges from the "B" station. However, they gave rise to but a small fraction of the above exposure and their radiological significance was negligible.

#### 6.8 Sizewell, Suffolk

Our monitoring near this station reflects the two critical radiation exposure pathways of fish and shellfish consumption leading to internal irradiation, and occupancy of intertidal areas giving rise to external exposure (Leonard and Smith, 1982). The results of this monitoring in 1982 are shown in Table 26.

Table 26	gamma dose ra station, 1982	tn environme ites near Size	well n	uclear	s and power				
Material	No. of sampling	Mean radioactivity concen- tration (wet), Bq kg <sup>-1</sup>							
	observa- tions†	Total beta	<sup>60</sup> Co	<sup>134</sup> Cs	137 <sub>Cs</sub>				
Mixed fish	2	98	ND	0.1	4.9				
Shrimps	2	100	0.2	ND	5.3				
Crabs	2	70	0.4		1.5				
Lobsters	1	90	ND		2.2				
Mussels	2	78	••		1.6				

Mean gamma dose rate in air at 1 m over intertidal sand/shingle (20 sampling observations†): 0.047  $\mu Gy~h^{-1}$ 

ND = not detected.

†See section 3.3 for definition.

The radiocaesium concentrations in fish and shellfish represent the combined effect of discharges from the station and from Sellafield, as well as of fallout. Apportionment is difficult at the low levels measured. Trace levels of cobalt-60 detected in some shellfish are likely to have been due to discharges from the station, but their radiological significance was negligible. The total radiation exposure of local fish and shellfish consumers was low, at less than 0.2% of the ICRP-recommended dose limit. Gamma dose rates, as in previous years, were indistinguishable from the natural background.

#### 6.9 Trawsfynydd, Gwynedd

Discharges from this station are made to the freshwater Lake Trawsfynydd under authorisation of the Welsh Office. Because of the limited volume flow for dispersion they are of greater radiological significance than those from the other UK nuclear power stations which discharge to estuarine or coastal waters. The critical radiation exposure pathway is due to consumption of fish caught in the lake, leading to internal irradiation; the important radionuclides are those of caesium and, to a lesser extent, strontium-90. Species of fish consumed are brown trout, rainbow trout and, in very small amounts, perch. Perch and most brown trout are indigenous to the lake but rainbow trout are regularly introduced from a hatchery. Because of the limited period they spend in the lake, rainbow trout generally exhibit lower radiocaesium concentrations than indigenous fish.

Our monitoring programme reflects the exposure pathways. Samples of rainbow trout, brown trout and perch are regularly analysed. As part of our research programme, mud and peat from the lake bed are also analysed; these materials contribute to the fishes' diet. Additional information is gained from analyses of the moss *Fontinalis* which is a sensitive indicator for a number of radionuclides, and from analyses of lake water. The results of these measurements for 1982 are shown in Table 27.

Table 27 Radioactivity in environmental materials near Trawsfynydd nuclear power station, 1982

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$									
	tions†	Total beta	<sup>54</sup> Mn	<sup>60</sup> Co	90Sr	<sup>106</sup> Ru	125 <sub>Sb</sub>	<sup>134</sup> Cs	<sup>137</sup> Cs		
Rainbow trout	3	190	ND	ND	17	ND	ND	3.1	28		
Brown trout	3	440			18	"		37	360		
Perch	1	1500			NA			<b>9</b> 0	750		
Mud	2	2300		1.9		**	150	27	1500		
Peat	2	3600	••	21		45	470	54	1700		
Fontinalis											
Afon Prysor	2	150	1.6	ND	**	ND	ND	ND	7.2		
Gwylan Stream	2	710	2.4	38		20	44	8.2	140		
Water											
Hot Lagoon	3	NA	NA	NA	0.37	NA	NA	0.018	0.18		
Cold Lagoon	3				0.38	••	••	0.031	0.25		

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$								
	observa- tions†	<sup>144</sup> Ce	155Eu	238 <sub>Pu</sub>	<sup>239</sup> Pu+ <sup>240</sup> Pu	241 Am	242 Cm	<sup>243</sup> Cm+ <sup>244</sup> Cm		
Rainbow trout	3	ND	ND	0.00010	0.00040	0.00057	ND	ND		
Brown trout	3			0.00020	0.00074	0.00068	"			
Perch	1	••		0.00066	0.0026	0.0030	0.00018	0.000050		
Mud	2			NA	NA	6.9	NA	NA		
Peat	2	29			"	29		"		
Fontinalis										
Afon Prysor	2	6.8	1.5	••	••	ND	**			
Gwylan Stream	2	7.1	2.1			**	"			
Water										
Hot Lagoon	3	NA	NA	••		NA	•1			
Cold Lagoon	3									

NA = not analysed.

ND = not detected.

\*Except for mud and peat where dry concentrations apply.

†See section 3.3 for definition.

Radiocaesium concentrations in fish in 1982 were lower than in 1981 (Hunt, 1983), reflecting reduced discharges of radiocaesium. However, concentrations of strontium-90 in fish were slightly higher in 1982 owing to increased discharges of this nuclide. Changes between 1981 and 1982 in concentrations of radiocaesium and strontium-90 in fish showed a similar pattern to concentrations in lake water. As in previous years, low concentrations of transuranic nuclides from station operations were observed in fish; these continued to be of negligible radiological significance.

It is estimated that in 1982 members of the critical group of fish consumers received at most about 7% of the ICRPrecommended dose limit. This reduced exposure as compared with 1981 (Hunt, 1983) reflects the lower radiocaesium concentrations in fish noted above.

### to the Irish Sea under authorisation of the Welsh Office. The two critical pathways are due to consumption of local fish and shellfish and to occupancy of intertidal areas. Monitoring is carried out in respect of these pathways. Samples of mud are analysed in support of the gamma dose rate measurements, and the indicator seaweed *Fucus vesiculosus* is also sampled. The results of monitoring in 1982 are presented in Table 28.

The effects of discharges from this station are masked by Sellafield-derived radioactivity. Concentrations of artificial radionuclides in environmental materials were consistent with those to be expected at this distance from Sellafield.

The total radiation exposure of members of the critical group in 1982 was less than 3% of the ICRP-recommended dose limit. The magnitude of discharges from the station indicate that the local contribution would have been a small fraction of this exposure. Gamma dose rates continued to be indistinguishable from the natural background.

#### 6.10 Wylfa, Gwynedd

Liquid radioactive wastes from this station are discharged

Table 28 Radioactivity in environmental materials and gamma dose rates near Wylfa nuclear power station, 1982

Material	No. of sampling	Mean radioa Bq kg <sup>-l</sup>	Mean radioactivity concentration (wet)*, Bq kg <sup>-1</sup>								
	tions†	Total beta	106 <sub>Ru</sub>	<sup>134</sup> Cs	137 <sub>Cs</sub>	<sup>144</sup> Ce	155 <sub>Eu</sub>				
Flatfish	2	140	ND	0.67	38	ND	ND				
Crabs	1	100	"	2.1	16	"					
Mussels	2	76	3.0	0.6	18	"					
Fucus vesiculosus	4	400	ND	0 <b>.9</b> 4	29						
Mud	2	1700	67	29	1000	11	4.5				

Material	No. of sampling	Mean r	Mean radioactivity concentration (wet)*, Bq kg <sup>-1</sup>							
	observa- tions†	238 <sub>Pu</sub>	$239_{Pu} + 240_{Pu}$	241 <sub>Am</sub>	242 <sub>Cm</sub>	<sup>243</sup> Cm + <sup>244</sup> Cm				
Flatfish	2	NA	NA	ND	NA	NA				
Crabs	1			••	••					
Mussels	2	0.029	0.16	0.18	ND	0.00068				
Fucus vesiculosus	4	NA	NA	ND	NA	NA				
Mud	2			54						

Mean gamma dose rate in air at 1 m over intertidal sediment (8 sampling observations†): 0.078  $\mu Gy \ h^{-1}$ 

NA = not analysed.

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ND = not detected.
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*Except for mud where dry concentrations apply.
†See section 3.3 for definition.
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Table 29 Radioactivity in environmental materials and gamma dose rates near naval establishments, 1982

Establishment	Material	No. of sampling	Mean radioactivity concentration, (wet)*, Bq $\rm kg^{-1}$							Mean gamma dose rate in air at l m	
		observa- tions†	Total beta	54Mn	60 Co	106 Ru	<sup>134</sup> Cs	137 <sub>Cs</sub>	155 <sub>Eu</sub>	No. of sampling observa- tions†	µGy h <sup>−1</sup>
Chatham	Sediment	6	780	ND	7.9	ND	1.7	45	0.8	19	0.063
Devonport.	Mussels Limpets Fucus vesiculosus	1 1 2	44 97 180	 	ND 	  	ND "	ND 0.6 0.2	ND "	NP  	NP 
	Sediment	6	800	0.2	0.3			6.5	0.6	24	0.083
Faslane	Sediment	4	670	ND	18		8.7	270	•	20	0.077
Rosyth	Sediment	2	420	0.3	2.5	••	1.4	49	1.0	5	0.074
Holy Loch	Fucus spiralis Fucus vesiculosus	2 2	250 270	ND 	ND 0.1	"	1.4	22 37	ND "	NP "	NP "
	Sediment	2	250		4.8	8.0	2.0	58	1.0	15	0.083

ND = not detected. NP = not applicable.

\*Except for sediment where dry concentrations apply.

†See section 3.3 for definition.

#### 7. Naval establishments

Liquid wastes containing relatively small quantities of radioactivity are discharged from the following establishments: Chatham, Devonport, Faslane and Rosyth, all of which are operated by the Ministry of Defence (Navy Department). The US naval base at Holy Loch also discharges small quantities of radioactive waste. We monitor the effects of all these discharges, in the case of Faslane and Rosyth on behalf of departments of the Scottish Office.

The critical pathway for public radiation exposure due to these discharges is via external exposure from occupancy of intertidal areas, the nuclide of main importance being cobalt-60. We therefore regularly carry out measurements of gamma dose rates: these are supported by analyses of sediments. Indicator shellfish and seaweed are also analysed.

Results of monitoring in 1982 are presented in Table 29. The small concentrations of activation product nuclides including cobalt-60 mainly reflect discharges from the establishments; levels of other artificial nuclides are largely due to fallout and to discharges from Sellafield. Gamma dose rates over intertidal sediments remained indistinguishable from the natural background, such that public radiation exposure was very low, at less than 0.1% of the ICRP-recommended dose limit.

### 8. Amersham International plc

Amersham International plc is engaged in the manufacture of radioactive materials for use in industry, research and medicine. The company's parent establishment is located in Amersham, Buckinghamshire, from which radioactive discharges are made into the catchment of the River Thames. As explained in section 5, environmental monitoring in respect of these discharges is carried out by the DOE. A further laboratory, situated near Cardiff, is engaged in the production of labelled compounds used in research and of diagnostic kits used in medicine for the *in vitro* testing of clinical samples. An authorisation issued by the Welsh Office regulates disposals of liquid radioactive wastes from this establishment to a sewer discharging into the Severn Estuary.

Our monitoring programme, carried out on behalf of the Welsh Office, reflects the two potentially critical pathways due to consumption of fish and shellfish and to external exposure over muddy intertidal areas. Measurements of external exposure are supported by analyses of intertidal sediment, and *Fucus* seaweed is collected as an indicator material. The environmental consequences of discharges from this establishment are very small and difficult to detect above the background levels due to fallout, other nuclear facilities which discharge small amounts of radioactive wastes to the Severn Estuary and the Bristol Channel, and possibly Sellafield.

The results of monitoring in 1982 are presented in Table 30. Artificial radioactivity detected was due to radiocaesium and other nuclides. However, none of these nuclides was processed or discharged by this establishment in 1982: the results were therefore due to the combined background effects noted above. Small amounts of iodine-131 detected in seaweed are likely to have been due to discharges from a local hospital. The exposure of the critical group of fish and shellfish consumers due to these effects in 1982 was very low, at less than 0.1% of the ICRP-recommended dose limit for members of the public. Gamma dose rates over sediment were indistinguishable from those to be expected from natural background.

Table 30	Radioactivity in environmental materials and gamma dose rates	
	near the outfall of the sewer serving Amersham International plc	,
	Cardiff, 1982	

Material	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg $^{-1}$								
	observa- tions†	Total beta	131 <sub>I</sub>	<sup>134</sup> Cs	137 <sub>Cs</sub>	<sup>144</sup> Ce	<sup>155</sup> Eu			
Flounders	2	180	ND	ND	1.3	ND	ND			
Mussels	2	100			0.6					
Fucus spiralis	3	NA	16	0.09	0.7					
Fucus vesiculosus	1		6.0	ND	1.4		"			
Sediment	4	<b>9</b> 80	ND	1.2	57	3.8	0.6			

Mean gamma dose rate in air at 1 m over intertidal sediment (8 sampling observations<sup>†</sup>): 0.090  $\mu$ Gy h<sup>-1</sup>

ND = not detected.

NA = not analysed.

\*Except for sediment where dry concentrations apply. †See section 3.3 for definition.

#### 9. Channel Islands monitoring

We have continued to analyse marine environmental samples provided by the Channel Islands States in surveillance of the effects of radioactive liquid discharges from the French reprocessing plant at Cap de la Hague. Fish and shellfish are monitored in relation to the internal irradiation pathway; sediment is analysed with relevance to external exposures. Seaweeds are sampled as indicator materials.

The results for 1982 are given in Table 31. Concentrations of caesium-137 in fish and shellfish were not significantly in excess of those to be expected from other sources, including fallout. However, the presence of transuranics in fish and shellfish may be attributed to discharges from the plant at Cap de la Hague. The presence of ruthenium-106 in environmental materials may also be attributed to this plant. However, the concentrations of artificial radionuclides in each of these materials were of negligible radiological significance.

### 10. Summary and Conclusions

A summary of estimated public radiation exposures in 1982 resulting from liquid radioactive waste discharges from nuclear establishments which we monitor is presented in Table 32. The exposures are expressed in terms of the committed effective dose equivalent to members of the critical group as percentages of the ICRP-recommended dose equivalent limit for members of the public, and incorporate the higher gut uptake factor for plutonium (section 3.4), except where otherwise indicated.

All exposures remained within the ICRP-recommended limit for members of the public. Discharges from Sellafield have, as in previous years, given rise to the highest exposures. The most important contribution to these exposures was due to transuranic radionuclides from the reprocessing operations; a further important contribution was from radiocaesium which is discharged mainly from the fuel element storage ponds. Exposures near Sellafield decreased in 1982 as compared with 1981, following the generally reducing trend in discharges and greater dispersion from the Irish Sea during 1982. Radioactivity from Sellafield also contributed to exposures near many other nuclear establishments. Since apportionment of exposure to radioactivity of local origin is often difficult, the exposures from all sources (including the small contribution due to fallout) are quoted in Table 32, with appropriate footnotes.

As in previous years, collective doses from UK liquid radioactive discharges have also been considered. The most significant discharges giving rise to collective dose, compared with which all other discharges may be disregarded, were those from Sellafield, radiocaesium being the most significant component. Details are given in section 4.1.1. The committed collective effective dose equivalent to the UK population in 1982 was 90 man-Sv, a decrease from 130 man-Sv reported for 1981. For the population of other European countries the provisional value of the committed collective effective dose equivalent was 100 man-Sv in 1982, representing a decrease from 150 man-Sv for 1981. These reductions were due to generally lower radiocaesium concentrations in fish and shellfish from the Irish Sea and further afield, following the reducing trend in discharges of radiocaesium from Sellafield. This trend is mainly a result of the optimised use, as required by the authorising Departments, of zeolite skips in the magnox fuel element storage ponds.

Table 31	Radioactivity in marine	environmental	materials	from the	e Channel	Islands,	1982

Material	Sampling area	No. of sampling	Mean radioactivity concentration (wet)*, Bq kg <sup>-1</sup>								
		observa- tions†	Total bet	a <sup>60</sup> Co	<sup>65</sup> Zn	<sup>90</sup> Sr	106 Ru	<sup>110 m</sup> Ag	125 Sb	<sup>137</sup> Cs	<sup>144</sup> Ce
Ray	Guernsey	1	73	ND	ND	NA	ND	ND	ND	5.3	ND
Crabs	Guernsey	1	98	2.4	6.6				••	ND	
	Jersey	1	100	0.8	ND		11	3.8		0.6	
Oysters	Jersey	1	94	ND	3.6		47	5.1		ND	
Limpets	Jersey	1	74		ND		27	ND			••
	Guernsey	1	84				ND	••			
	Alderney	1	130				25	1.0			
Porphyra	Jersey										
	Greve de Lecq	4	240	0.2		••	33	ND			••
	La Rozel	4	230	ND	••		35			0.1	
	Guernsev										
	Fort Doyle	3	220	••	••		9.0	••		ND	•
	Fermain Bay	4	230		"		22			0.2	
	Alderney										
	Telegraph Bay	4	320				37			0.2	
	Quenard Point	4	380	••			100	••	"	0.4	3.9
Burgur	Tamaan										
rucus serratus	Jersey La Rozel	4	300	1.4		0.38	9.7	•		0.6	ND
	Guernsey Fermain Bay	4	290	1.1		0.52	10	•		0.1	•
	Alderney Quenard Point	4	410	5.0		0.46	38	••	0.4	1.0	3.0
Sediment	Jersey St Helier Harbour	1	740	1.2	•	NA	200	•	7.0	11	61
	Guernsey Bordeaux Harbour	1	380	ND	•		4.5		ND	2.4	3.5
	Braye Harbour	1	620	•	"		15		2.4	4.7	4.2
Material	Sampling area	No. of sampling	Mean radi	.oactivi	ty cond	centra	tion (w	et)*, Bo	kg <sup>-1</sup>		
		tions†	238 <sub>Pu</sub>	239 <sub>Pu</sub> +	240 Pu	241 Ar	n 24:	<sup>2</sup> Cm <sup>24</sup>	<sup>3</sup> Cm + 2 <sup>4</sup>	+4 Cm	
Ray	Guernsey	1	0.00003	0.00016		0.000	017 ND	NE			
Crabs	Guernsev	1	0.0011	0.0029		0.00	o6 "	0	0005		
	Jersey	ì	0.0026	0.0056		0.01	i 0.	0016 0.	.0032		
Oysters	Jersey	1	0.023	0.055		0.030	0.0	0033 0.	0056		
Limpets	Jersey	1	0.0130	0.029		0.016	5 0.0	0018 0.	0025		
-	Guernsey	1	0.0049	0.014		0.009	93 0.	0015 0.	0018		
	Alderney	1	0.018	0.037		0.047	7 0.0	0053 0.	019		

	Alderney	1	0.018	0.037	0.047	0.0053	0.019
Porphyra	Jersey						
	Greve de Lecq	4	NA	NA	ND	NA	NA
	La Rozel	4	••		••		
	Guernsey						
	Fort Doyle	3	••	••			
	Fermain Bay	4	•				
	Alderney						
	Telegraph Bay	4		•			
	Quenard Point	4	-				••
Fucus	Jersey						
serratus	La Rozel	4	0.050	0.12	0.034	0.0039	0.0055
	Guernsey						
	Fermain Bay	4	0.042	0.11	0.035	0.0036	0.0061
	Alderney						
	Quenard Point	4	0.11	0.25	0.12	0.019	0.035
Sediment	Jersey						
	St Helier Harbour	1	1.7	4.4	3.2	0.24	0.47
	Guernsey						
	Bordeaux Harbour	ı	0.056	0.26	0.12	0.012	0.014
	Alderney						
	Braye Harbour	1	0.15	0.64	0.38	0.025	0.036
NA = not a	malvsed.						
ND = not d	etected.						
*Except fo	or sediment where dry co	ncentrati	ons apply.				
†See secti	on 3.3 for definition.						

Establishment	Radiation exposure pathway	Critical group	Committed effective dose equivalent (as % of ICRP recommended dose limit of 5 mSv year <sup>-1</sup> for members of the public)
BRITISH NUCLEAR FUELS I	.IMITED		
Sellafield	Fish and shellfish consumption	Local fishing community Commercial fishing community	54(34)* 13(12)*
	External	Whitehaven boat dwellers	12
	Porphyra/laverbread consumption	Consumers in South Wales	<0.1
Springfields	External	Houseboat dwellers	7 <sup>a</sup>
Capenhurst (Meols outfall)	Shellfish consumption	Local fishing community	<2 <sup>a</sup>
Chapelcross	External Fish and shellfish consumption	Local fishermen	<5 <sup>a</sup>
UNITED KINGDOM ATOMIC H	ENERGY AUTHORITY		
Winfrith	Fish and shellfish consumption	Local fishing community	<1
Dounreay	External to hands: fishing gear External Shellfish consumption	Local fishermen Local community Local fishing community	<1 <sup>b</sup> <1 <sup>b</sup> <0.4 <sup>b</sup>
	•		
NUCLEAR POWER STATIONS	OPERATED BY THE ELECTRICITY BOARD	5	
Berkeley and Oldbury	Fish and shellfish consumption External	Local fishing community	<0.1 <sup>b</sup>
Bradwell	Fish and shellfish consumption External	Local fishing community Houseboat dwellers	<0.3 <sup>b</sup>
Dungeness	Fish consumption External	Local fishing community	<0.1
Hartlepool <sup>C</sup>	Fish and shellfish consumption External	Local fishing community Coal collectors	<0.4 <sup>a</sup>
Heysham <sup>C</sup>	Fish and shellfish consumption External	Local fishing community	13 <sup>a</sup> <1 <sup>a</sup>
Hinkley Point	Fish and shellfish consumption External	Local fishing community	<0.2 <sup>b</sup>
Hunterston	Fish and shellfish consumption External	Local fishing community	<2 <sup>a</sup>
Sizewell	Fish and shellfish consumption External	Local fishing community	<0.2 <sup>b</sup>
Trawsfynydd	Fish consumption	Local fishing community	7
Wylfa	Fish and shellfish consumption External	Local community	<3 <sup>a</sup>
NAVAL ESTABLISHMENTS			
Chatham	External	Houseboat dwellers	<0.1
Devonport	External	Bait diggers	<0.1
Faslane	External	Boatyard workers	<0.1 <sup>b</sup>
Rosyth	External	Dredgermen	<0.1 <sup>b</sup>
Holy Loch	External	Local community	<0.1 <sup>b</sup>
AMERSHAM INTERNATIONAL	plc		
Cardiff	Fish and shellfish consumption External	Local fishing community	<0.1 <sup>a</sup>

#### Summarised estimates of public radiation exposure from discharges of liquid radioactive waste in the UK, Table 32 1982

\*See section 4.1.1. The first value is based on the enhanced gut uptake factor for plutonium; the value using the ICRP-recommended factor follows in parentheses. <sup>a</sup>Mainly due to discharges from Sellafield. <sup>b</sup>Partly due to discharges from Sellafield. <sup>c</sup>No radioactive discharges made in 1981. Potential critical pathways given; exposures are due to other sources of artificial radioactivity.

artificial radioactivity.

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