



Regional Fukushima fallout in Germany – data and models

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Introduction

Following the Fukushima NPP accident radionuclide air releases, samples of regional environmental media – rain water, river sediment, soil, grass and cow milk were taken in NW Germany and analyzed by gamma spectrometry [1]. The sampling in part contributed to the German state routine environmental radioactivity monitoring network IMIS [2], from which nationwide data could be retrieved later.

Sampling

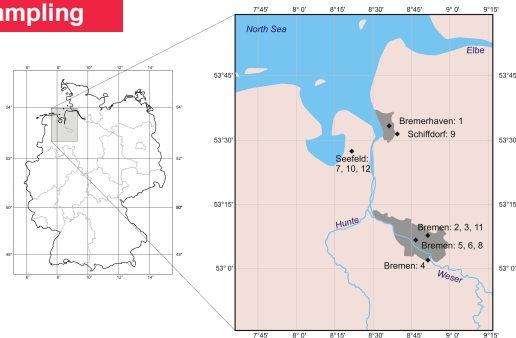


Figure 1: Location of sampling points and corresponding sample numbers (Table 1).

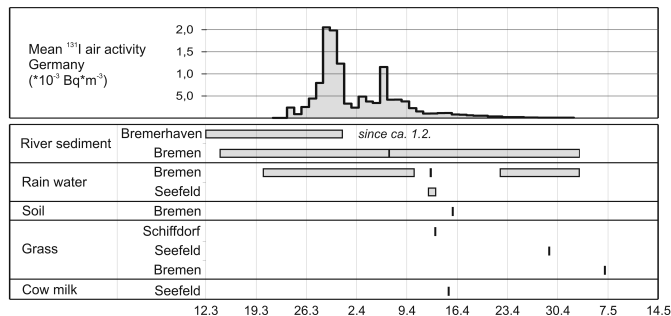


Figure 2: Timeline of the sampling campaign. In the upper panel, averaged ^{131}I air concentrations reported by 4 German air activity monitoring stations [3-5] are plotted.

Measured values

Table 1: Measured values in $\text{Bq}\cdot\text{kg}^{-1}$ wet mass. Data are decay corrected to the end of the sampling period, unless noted otherwise.

No.	Type of sample	Sampling period	^{131}I	^{137}Cs	^{134}Cs
1	River sediment (Bremerhaven)	ca. 1.2. – 31.3.	0.113 ± 0.046	2.67 ± 0.07	< 0.084
2 ¹	River sediment (Bremen)	14.3. – 6.4.	0.45 ± 0.06	1.76 ± 0.04	< 0.06
3	River sediment (Bremen)	6.4. – 3.5.	< 0.12	1.73 ± 0.05	< 0.05
4	Soil (Bremen)	15.4.	0.068 ± 0.025	3.00 ± 0.07	< 0.069
5 ¹	Rain water (Bremen)	20.3. – 10.4.	0.43 ± 0.03	< 0.04	< 0.04
6	Rain water (Bremen)	12.4.	0.10 ± 0.02	< 0.04	< 0.04
7	Rain water (Seefeld)	11.4. – 13.4.	0.14 ± 0.02	< 0.03	< 0.03
8 ²	Rain water (Bremen)	22.4. – 3.5.	0.031 ± 0.008	< 0.02	< 0.02
9	Grass (Schiffdorf)	13.4.	3.58 ± 0.13	1.59 ± 0.07	0.32 ± 0.03
10	Grass (Seefeld)	28.4.	0.31 ± 0.04	0.26 ± 0.04	0.062 ± 0.014
11	Grass (Bremen)	6.5.	0.12 ± 0.03	0.18 ± 0.03	< 0.08
12	Milk (Seefeld)	14.4.	0.08 ± 0.02	< 0.02	< 0.03

¹Decay corrected to the date of the maximum rainfall (3.4.2011). ²Mixed with old rain water.

Isotope ratios

$^{131}\text{I}/^{137}\text{Cs}$:

In NW Germany, a residual deposition of about $2 \text{ kBq}\cdot\text{m}^{-2}$ ^{137}Cs remains from bomb test fallout and Chernobyl. Therefore the ratio $^{131}\text{I}/^{137}\text{Cs}$ can differ from the atmospheric data depending on the type of sample. At the time of maximal concentration (end of March), this ratio was about 10. For the sediment samples the maximum value is 0.256, indicating a strong contribution from "old" ^{137}Cs . In fact, ^{137}Cs had been detected at these locations earlier and in similar concentrations.

$^{134}\text{Cs}/^{137}\text{Cs}$:

In two grass samples ^{134}Cs could be detected together with ^{137}Cs and ^{131}I . This offers the possibility to discriminate between "recent" and "old" Cs isotopes, using the $^{134}\text{Cs}/^{137}\text{Cs}$ ratio from atmospheric measurements. This value was close to 1 in most published air concentration data [3-5]. Assuming this value, it could be concluded that the main ^{137}Cs contribution in the grass samples was "old".

Modeling

Deposition with rain

Wet deposition by rain is known to be the most effective transfer path for airborne radioisotopes to ground and water bodies. It can be estimated [6] as:

$$D = C_{\text{air}} \Lambda_0 t_{\text{rain}} h \quad (1)$$

D areal deposition
 C_{air} isotope concentration in air
 Λ_0 standard washout coefficient ($=7 \cdot 10^{-5} \text{ s}^{-1}$)
 T_{rain} rainfall duration
 h contaminated air column height ($=1000 \text{ m}$, assuming constant concentration within the boundary layer)

$$C_{\text{rain}} = \frac{D}{P} \quad (2)$$

C_{rain} activity concentration in rain water
 P precipitation ($\text{l}\cdot\text{m}^{-2}$)

Transfer of radioiodine from grass to milk

can be predicted by applying emergency models [7].

$$C_{\text{milk}} = C_{\text{grass}} M_{\text{grass}} T_{\text{milk}} \quad (3)$$

C_{milk} activity concentration in milk
 C_{grass} activity concentration in grass
 M_{grass} daily grass consumption rate of cattle ($=65 \text{ kg d}^{-1}$)
 T_{milk} element-specific transfer factor from grass to milk ($=0.003$ for iodine)

Table 2: Comparison of modeled and measured values of Fukushima fallout in Germany. Data for whole Germany were obtained from IMIS [2]

	Regional modeled concentrations	Regional measured data	National measured data (averages)
Deposition	$2.14 \text{ Bq}\cdot\text{m}^{-2} \text{ }^{131}\text{I}$ $0.21 \text{ Bq}\cdot\text{m}^{-2} \text{ }^{137}\text{Cs}$ (Eq. 1)	$1.46 \text{ Bq}\cdot\text{m}^{-2} \text{ }^{131}\text{I}$ (sample 2) $2.72 \text{ Bq}\cdot\text{m}^{-2} \text{ }^{131}\text{I}$ (sample 4) (most of ^{137}Cs "old")	$7.0 \text{ Bq}\cdot\text{m}^{-2} \text{ }^{131}\text{I}$ ($n=67$) $0.20 \text{ Bq}\cdot\text{m}^{-2} \text{ }^{137}\text{Cs}$ ($n=21$)
Rain water	$0.252 \text{ Bq}\cdot\text{l}^{-1} \text{ }^{131}\text{I}$ $0.025 \text{ Bq}\cdot\text{l}^{-1} \text{ }^{137}\text{Cs}$ (Eq. 2)	max. $0.43 \text{ Bq}\cdot\text{l}^{-1} \text{ }^{131}\text{I}$ $< 0.04 \text{ Bq}\cdot\text{l}^{-1} \text{ }^{137}\text{Cs}$	
Grass		max. $3.58 \text{ Bq}\cdot\text{kg}^{-1} \text{ }^{131}\text{I}$ max. $1.59 \text{ Bq}\cdot\text{kg}^{-1} \text{ }^{137}\text{Cs}$ (most of ^{137}Cs "old") max. $0.32 \text{ Bq}\cdot\text{kg}^{-1} \text{ }^{134}\text{Cs}$	$1.56 \text{ Bq}\cdot\text{kg}^{-1} \text{ }^{131}\text{I}$ ($n=10$)
Milk	$0.20 \text{ Bq}\cdot\text{kg}^{-1} \text{ }^{131}\text{I}$ (Eq. 3)	$0.08 \text{ Bq}\cdot\text{kg}^{-1} \text{ }^{131}\text{I}$	$0.008 \text{ Bq}\cdot\text{kg}^{-1} \text{ }^{131}\text{I}$ ($n=4$)

Conclusions

- Despite the large distance between source and deposition area and the usage of standard equipment, it was possible to detect traces of the emissions from Fukushima in NW Germany in various environmental media.
- Isotope ratios could be used to discriminate between "recent" and "old" deposition.
- Values were plausible when compared to reported air concentrations and predictions from simple radioecological models.
- Comparison with nationwide data shows similar values.

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