



EFFECTS OF THE FUKUSHIMA DAIICHI ACCIDENT IN HUNGARY

MEASUREMENTS, RESULTS AND PUBLIC ACCEPTANCE

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Introduction

The contaminated air mass reached Europe from the North in two waves, firstly detected in Reykjavik (Iceland) on 20th of March.

Hungary:

Radiological Monitoring and Data Acquisition Network (RAMDA)

- ¹³¹I isotope in aerosol samples 22nd 24th of March at first time
- Maximum activity concentration in late March: $1.0 1.4 \text{ mBq/m}^3$
- Decreased under the detection limit until late May (1 2 μ Bq/m³)
- The "Frédéric Jouliot-Curie" National Research Institute for Radiobiology and Radiohygenie (the centre of RAMDA) collected aerosol samples, milk, grass and vegetable samples originated from open filed cultures.
- The Environmental Protection Service (EPS) collected aerosol samples daily + whole body counting measurements on tourists and workers arrived from Japan.



Method

- Cesium isotopes: quickly adsorb on the surface of the aerosol molecules (glass fiber filter) lodine isotopes: parlty adsorb, partly remains gaseous (activated carbon containing filter)
- There is a measurable difference between samples collected in Budapest and in the country side in terms of lodine isotopes: Isotope Institute Ltd. located in the KFKI Campus produces radioiodine (5 – 20 μBq/m³ in the air) which is not measurable outside the capital city.
- There are traces of 137 Cs (with 30.1 year of half life) in the soil: typically under 2 μ Bq/m³.
- The Annual Committed Effective Dose for inhaled radioisotopes was calculated for ¹³¹I, ¹³⁴Cs and ¹³⁷Cs referring to the highest activity concentrations measured in Hungary



- Activity concentration maxima in aerosol samples (mBq/m³)
- Food and grass samples (Bq/kg or Bq/L)

in contrast with the measurements in Budapest after the Chernobyl accident

1986 (1 st - 3 th of May)			2011 (11 th of March - 14 th of April)			Sample type	1986 (1 st - 3 th of May)		2011 (11 th of March - 14 th of April)	
¹³¹	¹³⁴ Cs	¹³⁷ Cs	¹³¹	¹³⁴ Cs	¹³⁷ Cs		131	¹³⁴ Cs and ¹³⁷ Cs	¹³¹	¹³⁴ Cs and ¹³⁷ Cs
3190	550	1010	1.47 ± 0.15	0.09 ± 0.01	0.11 ± 0.01	Grass	9700	1180	0.5 – 3.0	0.2 – 0.7
						Vegetables	400	2700	0.2 – 1.0	< 1.13
						Milk	1500	45	< 0.7	< 0.6

• Annual Committed Effective Dose (nSv/h)

20)10	2011 (17 th of Feb 9 th of May)			
¹³¹	¹³⁷ Cs	131	¹³⁴ Cs and ¹³⁷ Cs		
0.7	0.2	3.9	2.7		

Hazai környezetradiológia Fukushima után (Inland environmental radiology after Fukushima), Nukleon, 2012, V., 113



Conclusions

- Due to the huge distance between Budapest, Hungary and Fukushima, Japan (~ 8922 km in a straight line) there where small, but measurable amounts of radioisotopes in the air which could reach Hungary.
- In the biological samples there where also traceable amounts of ¹³¹I, ¹³⁴Cs and ¹³⁷Cs but significantly lower than in the measurements taken after the Chernobyl accident.
- The Annual Committed Effective Dose for the inhalation of the ¹³¹I, ¹³⁴Cs and ¹³⁷Cs isotopes showed elevated values, but **remained far under the dose limit** to the members of the public from artificial source (1 mSv/year).
- Whole body counting measurements on tourists and workers arrived from Japan resulted in **insignificant effects** (in 4 cases only the natural ⁴⁰K was measurable, in one measurement taken on a NPP worker from the Tsukuba Ibaraki Prefecture there was negligible amount of ¹³¹I)
- In a 2009 survey (EUROBAROMETER) 50 60 % of Hungarians advocated nuclear energy in association with safety and maintaining the proportion (40%) in electricity generation. After the Fukushima accident, the supportiveness dropped with about 10 % but only in relation with safety.