

An Overview of the Current Distribution of Radionuclides Released to the Environment Following the Fukushima Accident

Kimiaki Saito

**Fukushima Environmental Safety Center
Japan Atomic Energy Agency (JAEA)**

Estimated radioactive releases into the atmosphere from the Fukushima accident (Bq)

Nuclide	Half life	Reactor 1	Reactor 2	Reactor 3	Total
Xe-133	5.2 d	3.4×10^{18}	3.5×10^{18}	4.4×10^{18}	1.1×10^{19}
Cs-134	2.1 y	7.1×10^{14}	1.6×10^{16}	8.2×10^{14}	1.8×10^{16}
Cs-137	30 y	5.9×10^{14}	1.4×10^{16}	7.1×10^{14}	1.5×10^{16}
Sr-89	50.5 d	8.2×10^{13}	6.8×10^{14}	1.2×10^{15}	2.0×10^{15}
Sr-90	29.1 y	6.1×10^{12}	4.8×10^{13}	8.5×10^{13}	1.4×10^{14}
Te-129m	33.6 d	7.2×10^{14}	2.4×10^{15}	2.1×10^{14}	3.3×10^{15}
Pu-238	87.7 y	5.8×10^{08}	1.8×10^{10}	2.5×10^{08}	1.9×10^{10}
Pu-239	24065 y	8.6×10^{07}	3.1×10^{09}	4.0×10^{07}	3.2×10^{09}
Pu-240	6537 y	8.8×10^{07}	3.0×10^{09}	4.0×10^{07}	3.2×10^{09}
Pu-241	14.4 y	3.5×10^{10}	1.2×10^{12}	1.6×10^{10}	1.2×10^{12}
I-131	8 d	1.2×10^{16}	1.4×10^{17}	7.0×10^{15}	1.6×10^{17}

Contamination mapping projects

- **In order to estimate the impact of the Fukushima accident and take appropriate countermeasures, it has been necessary to obtain precise information on the contamination conditions.**
- **The Ministry of Education, Culture, Sports, Science and Technology (MEXT) commissioned JAEA to construct detailed contamination maps based on reliable environmental monitoring.**
- **JAEA has completed three series of mapping projects in collaboration with many organizations.**
- **The 4th project is in progress being commissioned by the Nuclear Regulation Authority (NRA).**

Tasks of mapping projects

- 1. Mapping of radionuclide deposition and dose rates in air**
- 2. Studies on radionuclide migration in natural environment**
- 3. Construction of a database**
- 4. Prediction of contamination conditions in future**

Contents

- 1. Distribution of radionuclide ground deposition densities (Bq/m^2)**
- 2. Distribution of air dose rates ($\mu\text{Sv/h}$)**
- 3. Depth profiles of radiocesium in ground (Bq/kg , relaxation depth β)**

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Locations

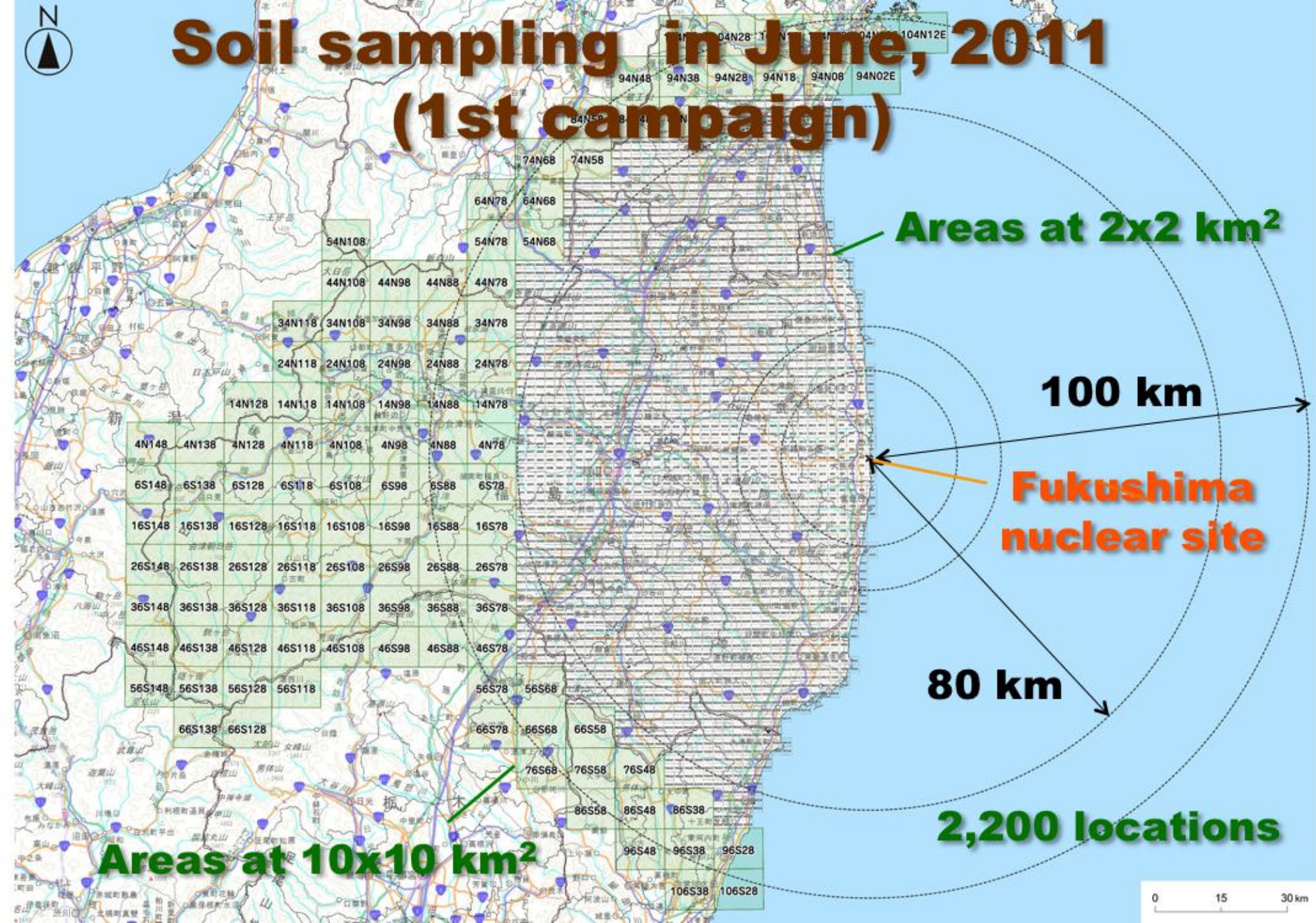
- Undisturbed flat fields for 1, 2, 3**
- Car-borne survey on roads over wide areas for 2**

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- 4. Overview of current contamination conditions in Fukushima region**



Soil sampling in June, 2011 (1st campaign)



Collection procedures of a soil sample



- **Top 5 cm soil**
- **Sufficient mixing**
- **U8 plastic container**



- **11,000 samples at 2,200 locations in 1st campaign**
- **Analyzed at 21 laboratories**

In-situ measurement using a portable Ge detector



In-situ measurements have been used to determine deposition density of radionuclides since December 2011.

Radionuclide deposition maps

1. Gamma-ray emitting nuclides

- Cs-137 (30.2 y) •Cs-134 (2.06 y)
- I-131 (8.02 d)
- Te-129m (33.6 d) •Ag-110m (250 d)

2. Alpha-ray emitting nuclides

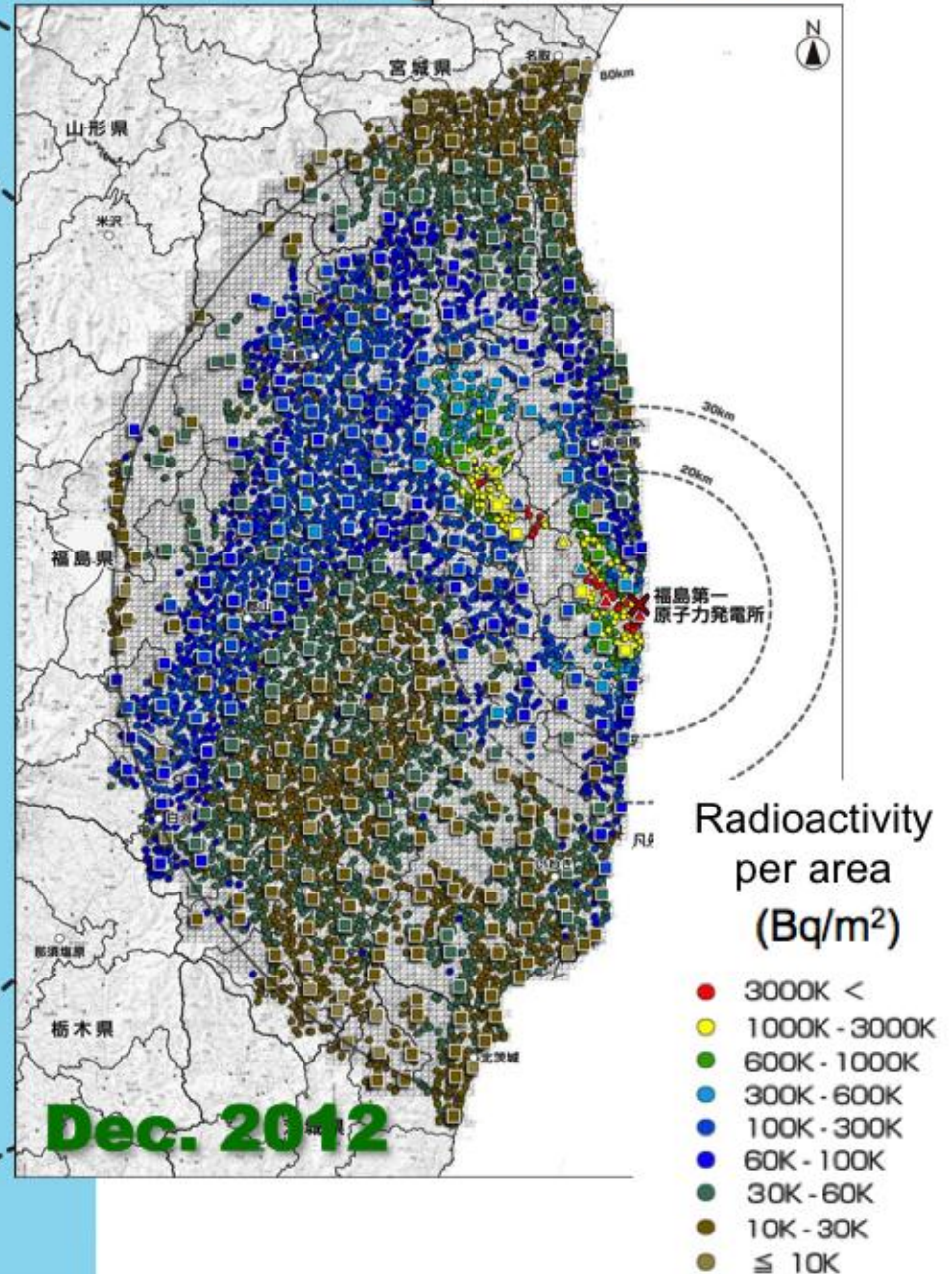
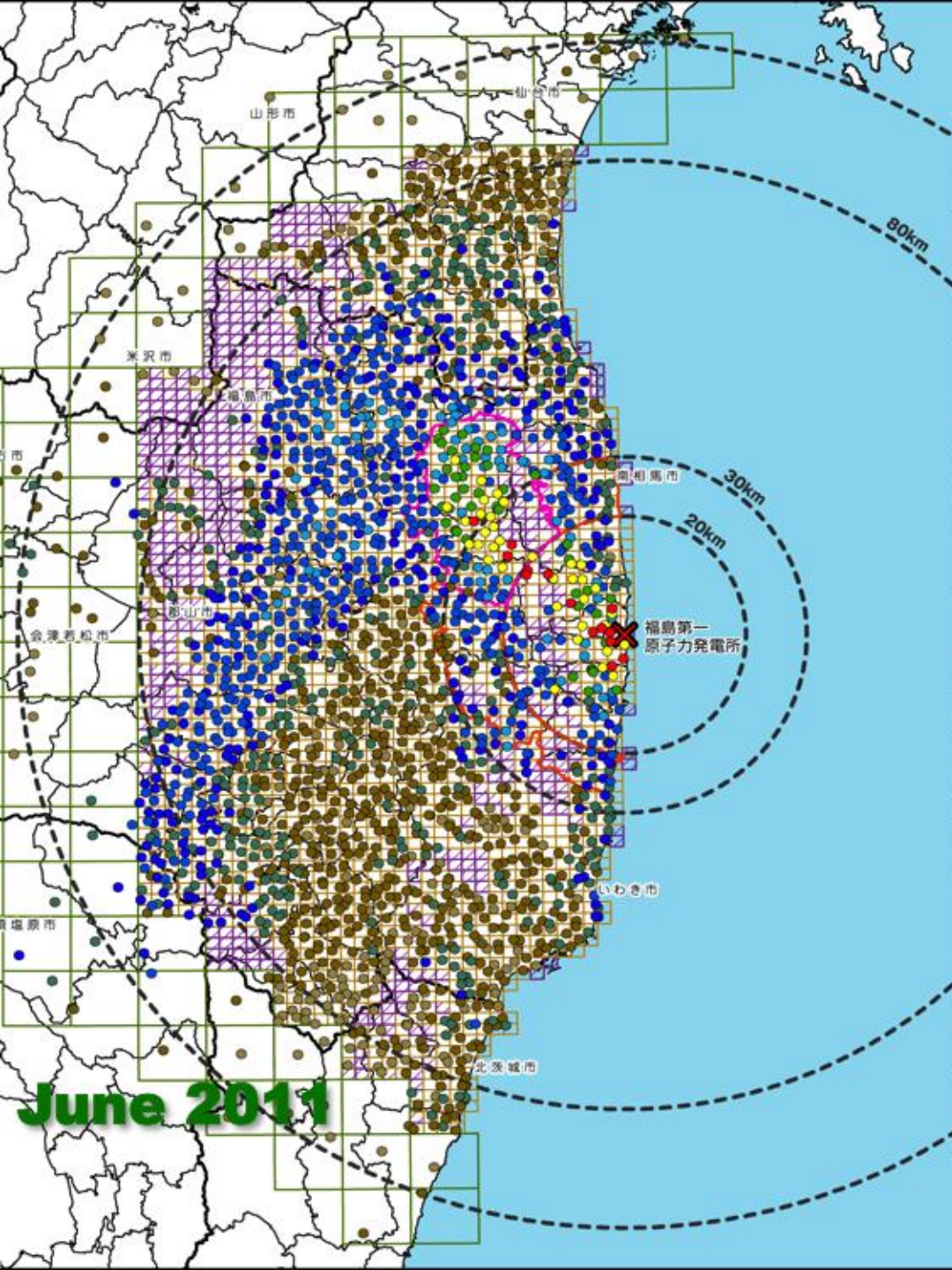
- Pu-238 (87.7 y)
- Pu-239 (24,100 y) + Pu-240 (6,564 y)

3. Beta-ray emitting nuclides

- Sr-89 (50.5 d)
- Sr-90 (28.8 y)

(half life)

¹³⁷Cs deposition

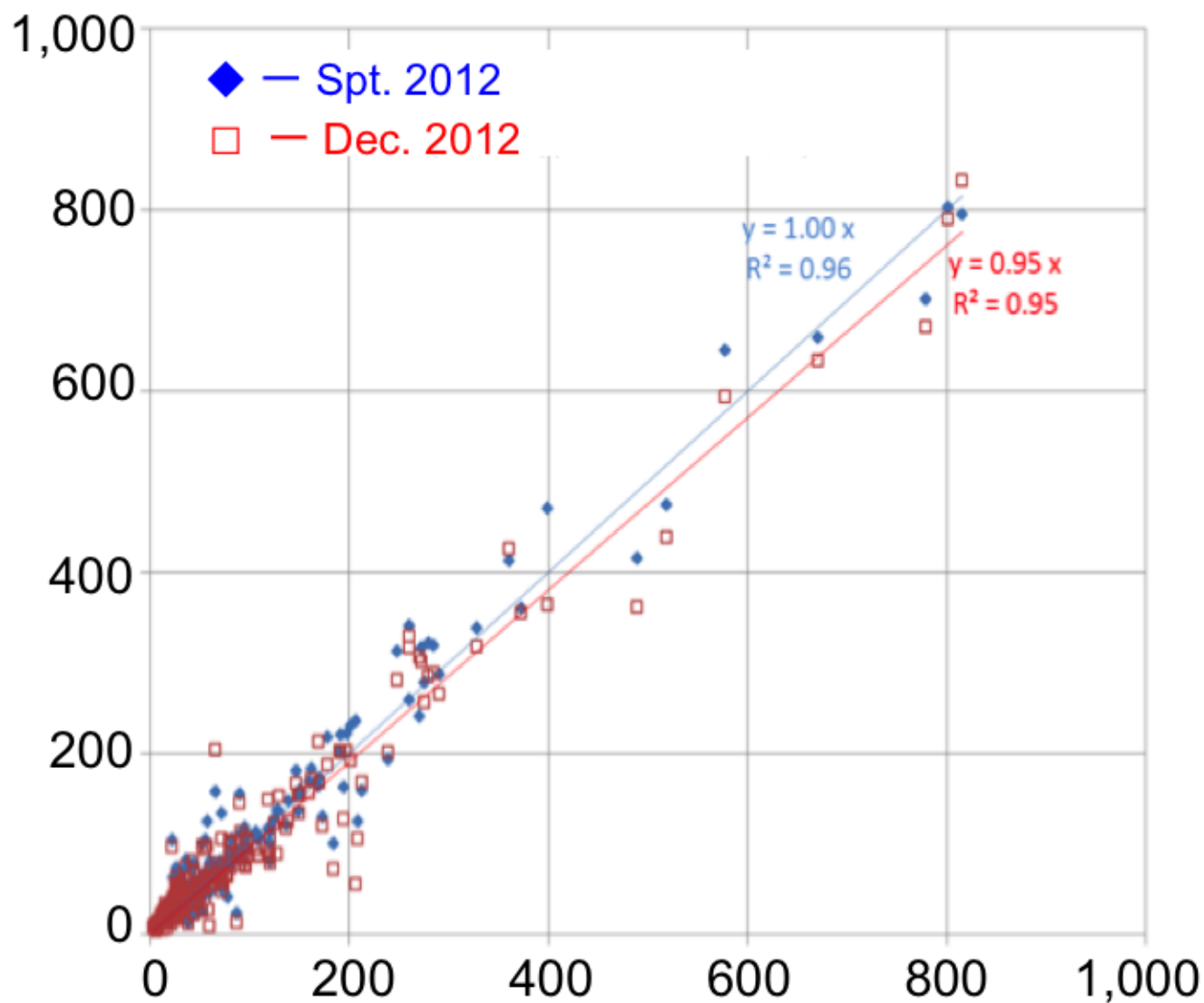


Radioactivity
per area
(Bq/m²)

- 3000K <
- 1000K - 3000K
- 600K - 1000K
- 300K - 600K
- 100K - 300K
- 60K - 100K
- 30K - 60K
- 10K - 30K
- ≤ 10K

Temporal change in Cs-137 deposition density

Deposition density in Sept. 2012 (kBq/m²)

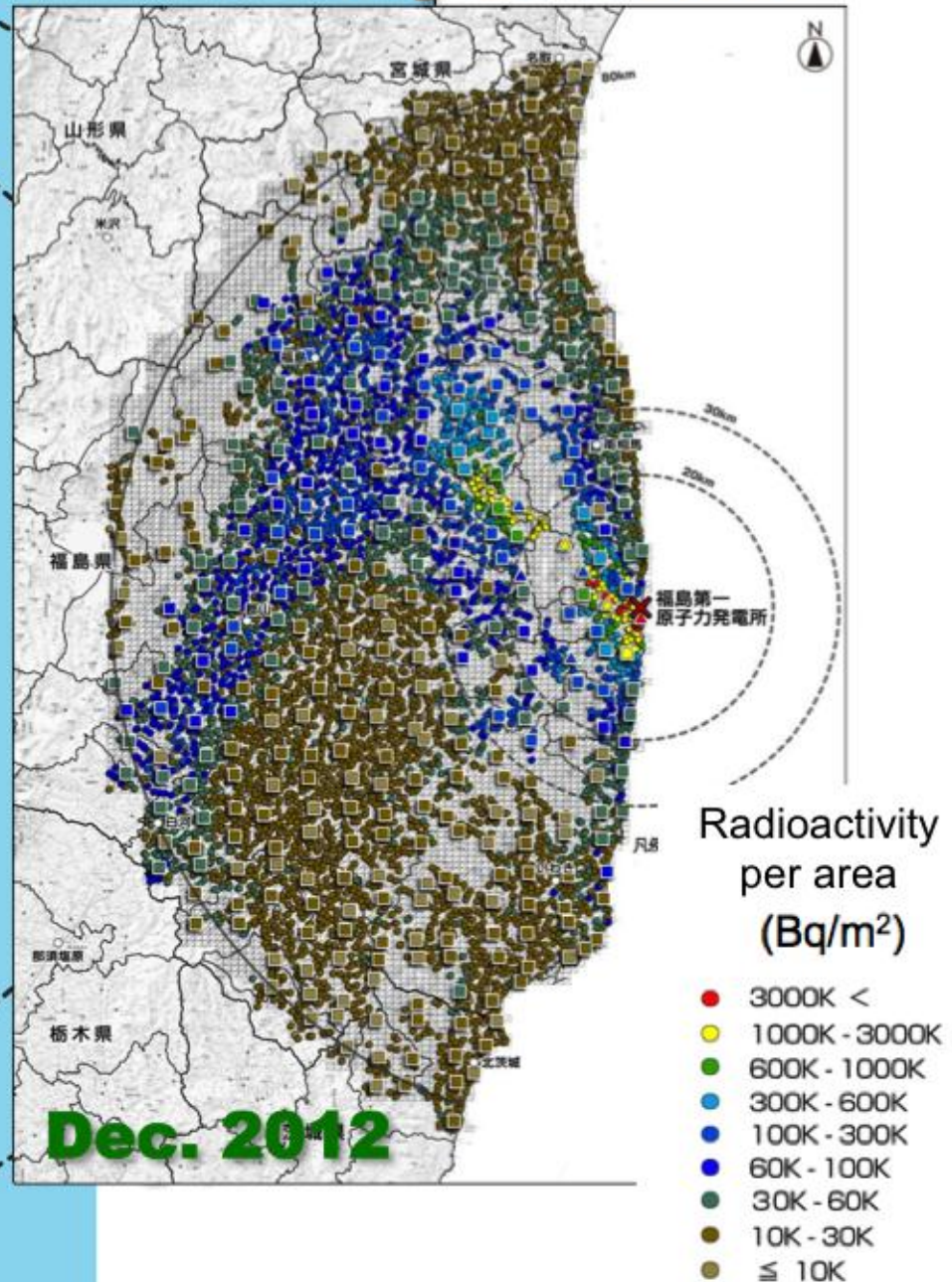
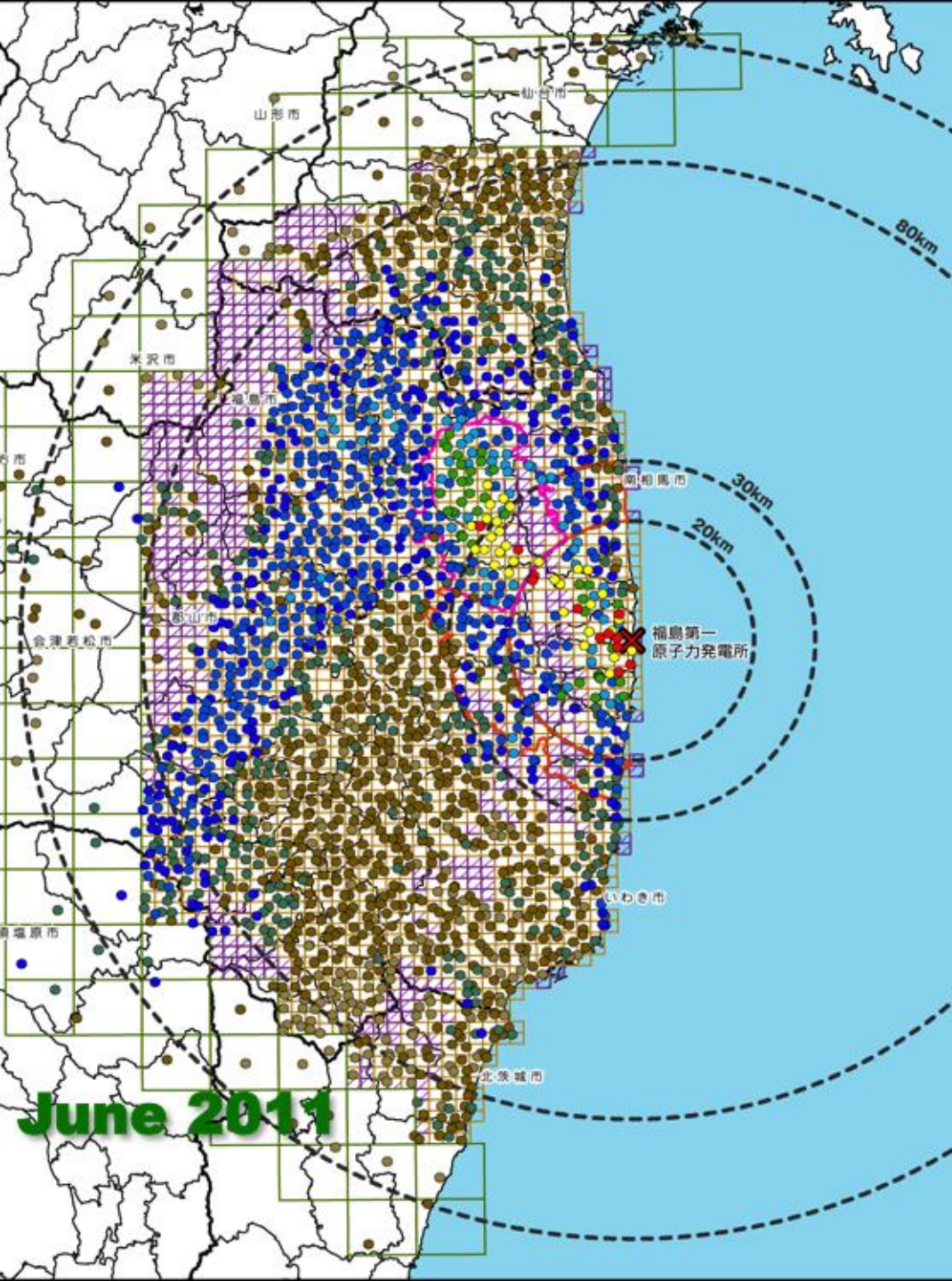


Deposition density in Dec. 2012 (kBq/m²)

● **Cs-137 deposition densities have not changed much in undisturbed flat fields.**

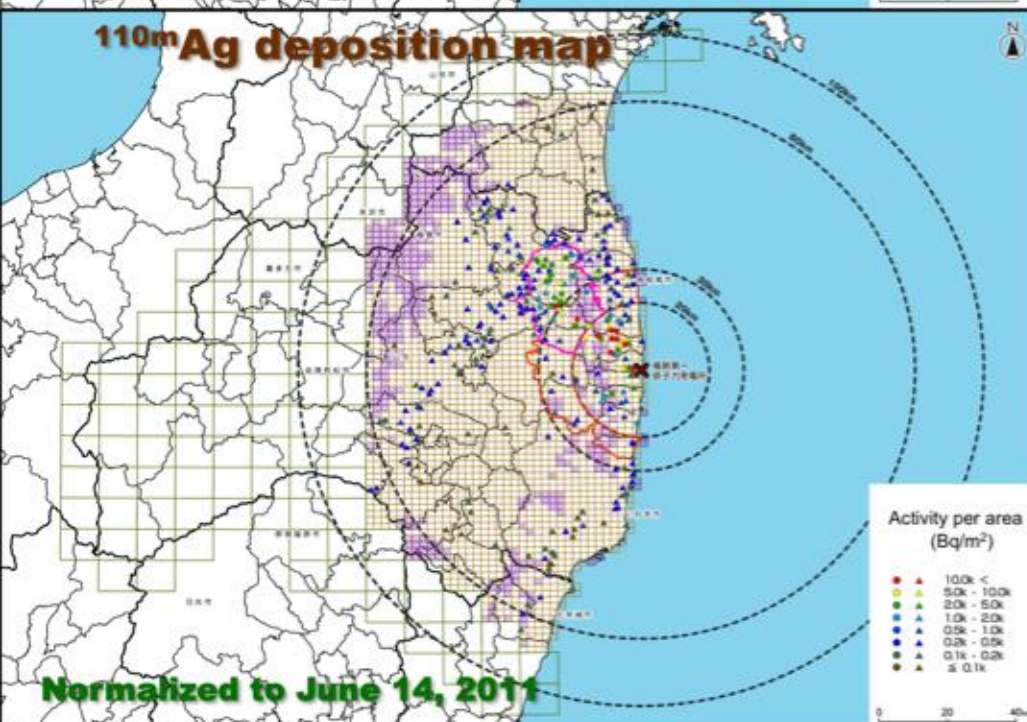
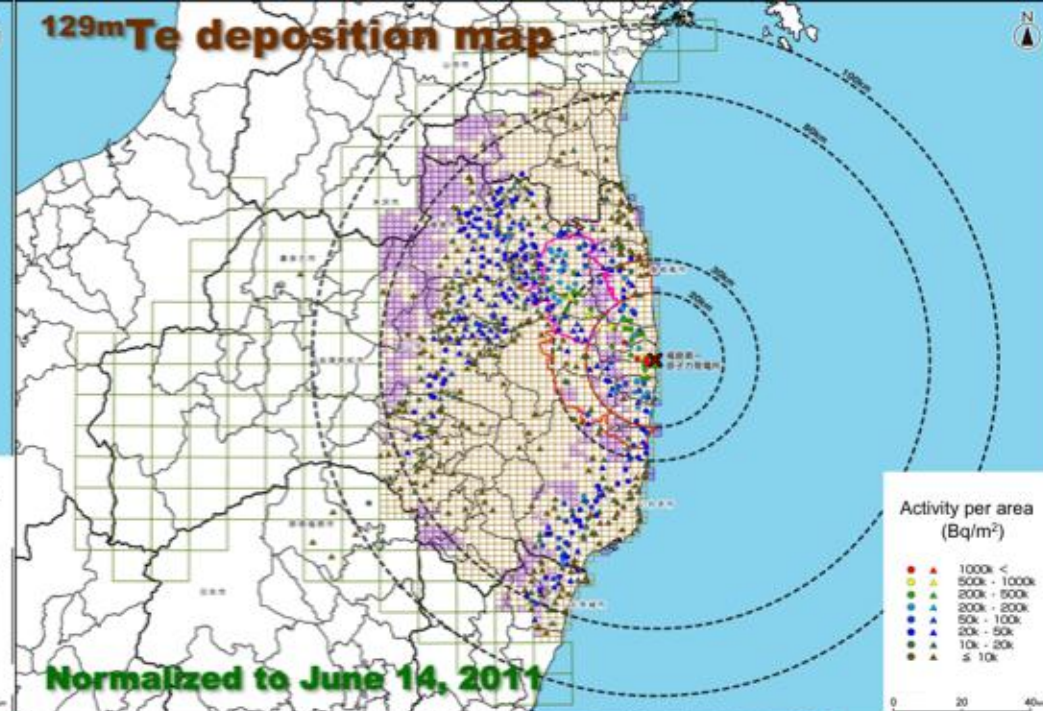
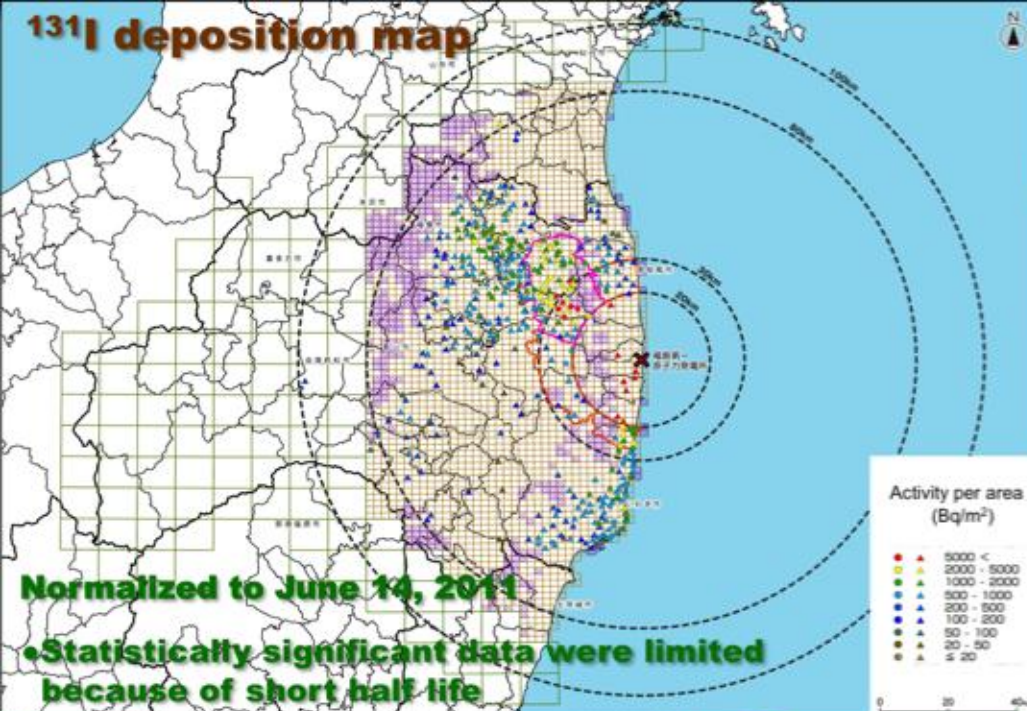
Deposition density in March 2012 (kBq/m²)

¹³⁴Cs deposition



Radioactivity
per area
(Bq/m²)

- 3000K <
- 1000K - 3000K
- 600K - 1000K
- 300K - 600K
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- 30K - 60K
- 10K - 30K
- ≤ 10K

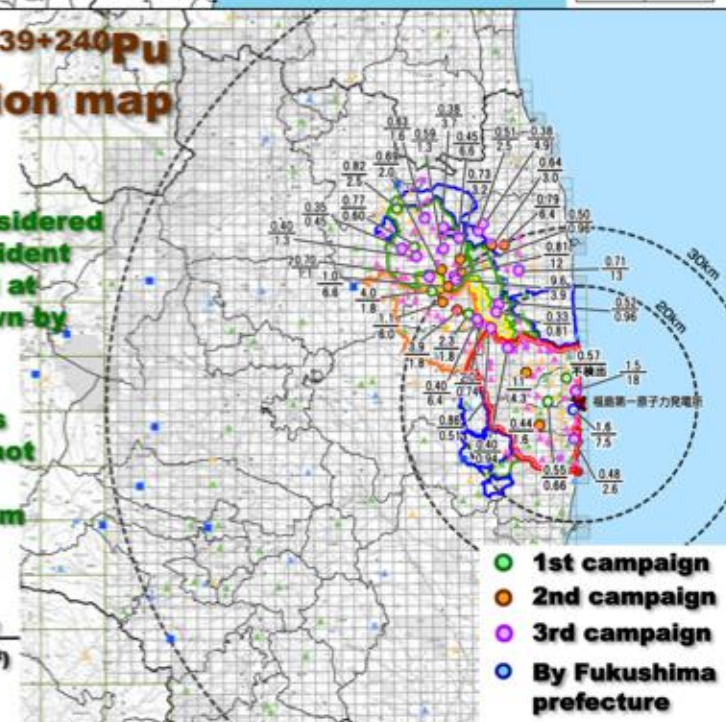


^{238}Pu , $^{239}+^{240}\text{Pu}$ deposition map

○ Plutonium considered
due to the accident
were detected at
locations shown by
circles.

■ Plutonium was
detected but not
identified as
originating from
the accident.

^{238}Pu (Bq/m^2)
 $^{239}+^{240}\text{Pu}$ (Bq/m^2)



Evaluation of accumulated effective doses for 50 years from June 2011

- Maximum nuclide deposition densities (Bq/m²) were used.
- External exposures and inhalation due to re-suspension were evaluated.

Nuclide	Half life	Maximum concentration (Bq/m ²)	Effective dose for 50 years	
			Conversion coef. (μSv/(Bq/m ²))	Dose (mSv)
Cs-134	2.065 y	1.4×10 ⁷	5.1×10 ⁻²	710
Cs-137	30.167 y	1.5×10 ⁷	1.3×10 ⁻¹	2000(2.0Sv)
I-131	8.02 d	5.5×10 ⁴	2.7×10 ⁻⁴	0.015
Sr-89	50.53 d	2.2×10 ⁴	2.8×10 ⁻⁵	0.00061 (0.61 μSv)
Sr-90	28.79 y	5.7×10 ³	2.1×10 ⁻²	0.12
Pu-238	87.7 y	4	6.6	0.027
Pu-239+240	2.411×10 ⁴ y	15	8.5	0.12
Ag-110m	249.95 d	8.3×10 ⁴	3.9×10 ⁻²	3.2
Te-129m	33.6 d	2.7×10 ⁶	2.2×10 ⁻⁴	0.6

(Dose coefficients from TECDOC-1162)

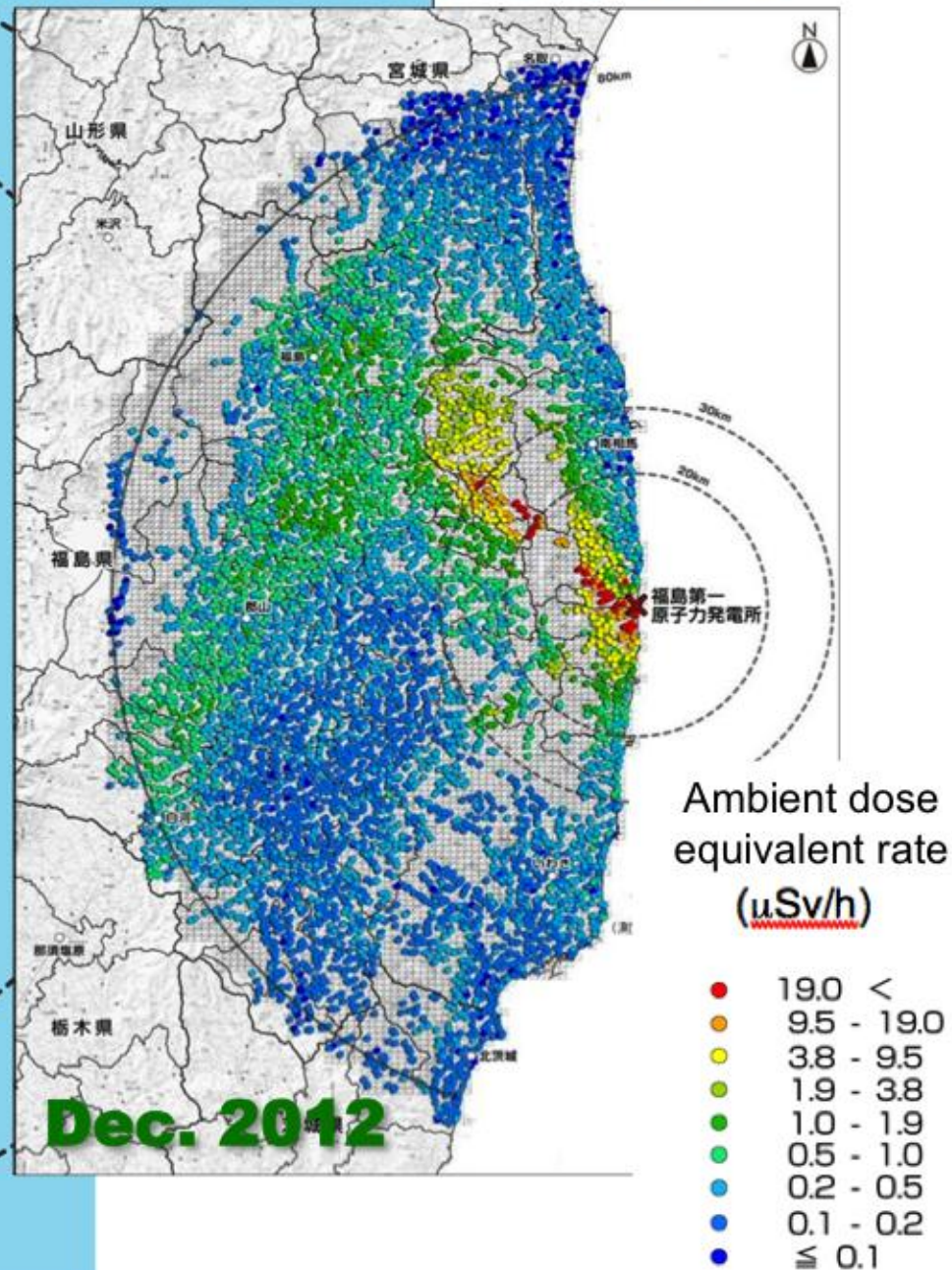
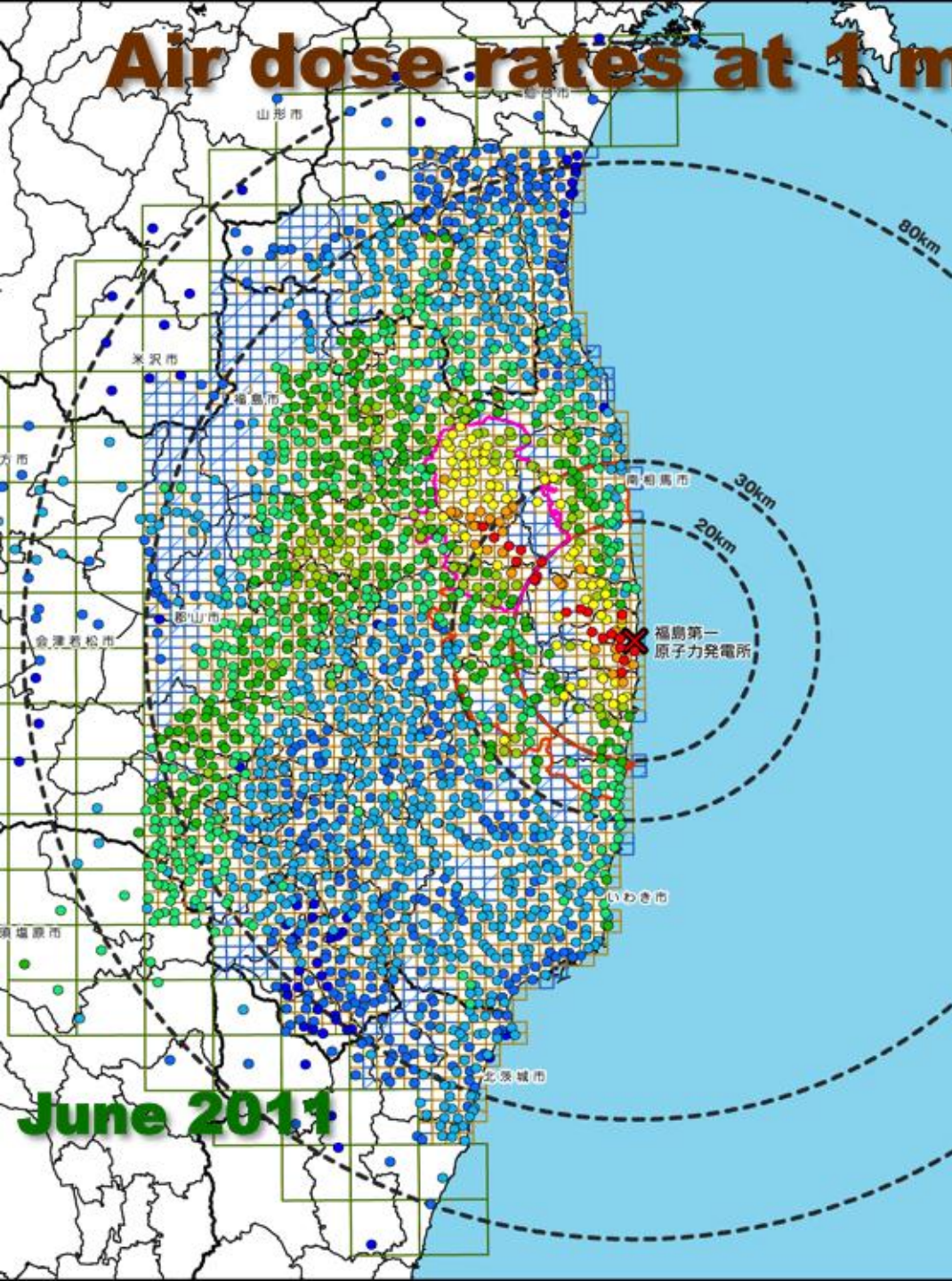
Summary on radionuclide deposition

- 1. Cesium is much more important than other nuclides from the viewpoint of exposure doses in future.**
- 2. Plutonium and Strontium originating from the accident were detected; the radioactivities were not large.**
- 3. Cs-137 deposition densities have not decreased much; while, Cs-134 deposition densities have certainly decreased due to physical decay.**
- 4. Radioactivity ratios of I-131, Te-129m and Ag-110m to Cs-137 have regional dependency.**

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- 4. Overview of current contamination conditions in Fukushima region**

Air dose rates at 1 m above flat fields

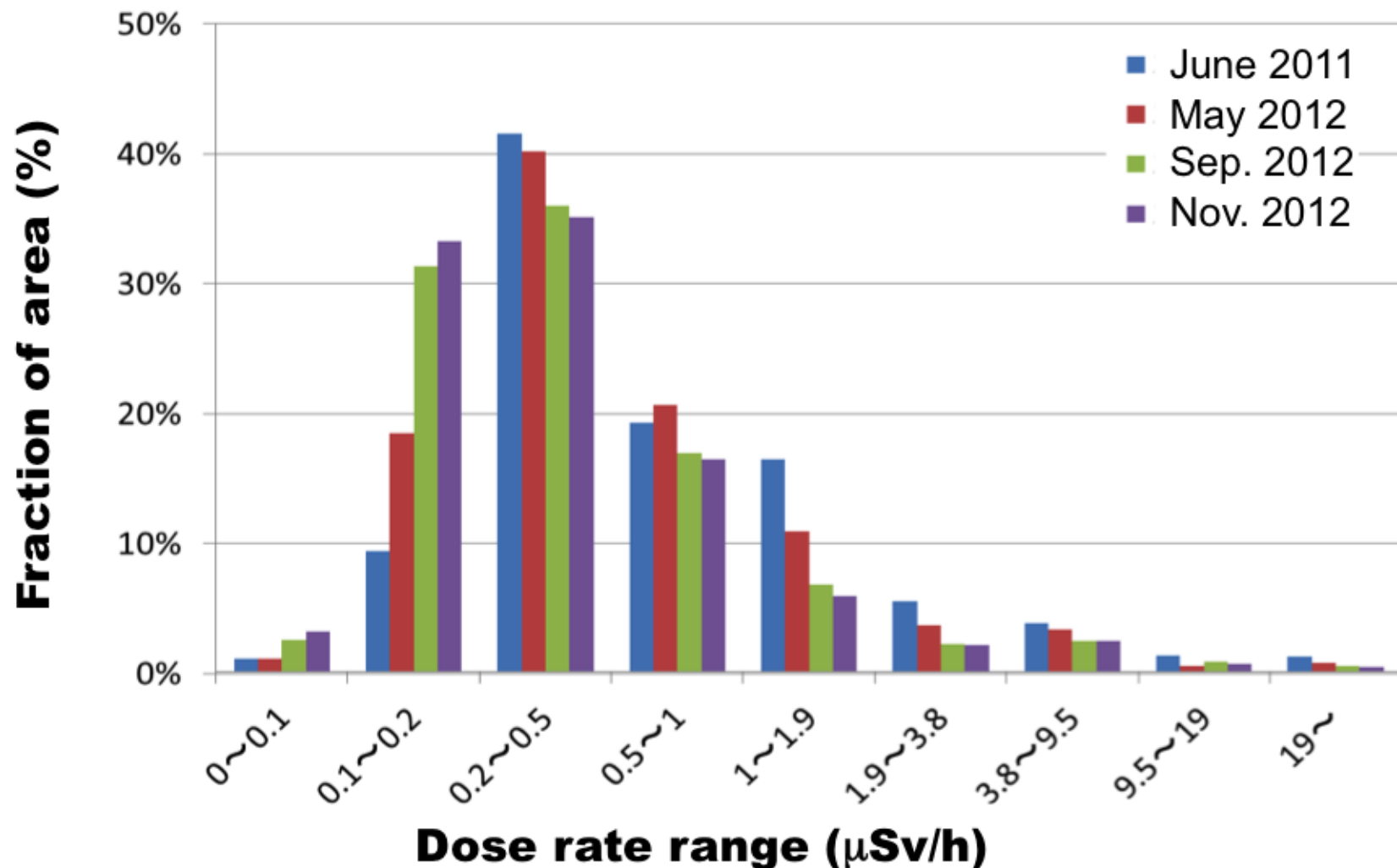


Ambient dose
equivalent rate
($\mu\text{Sv/h}$)

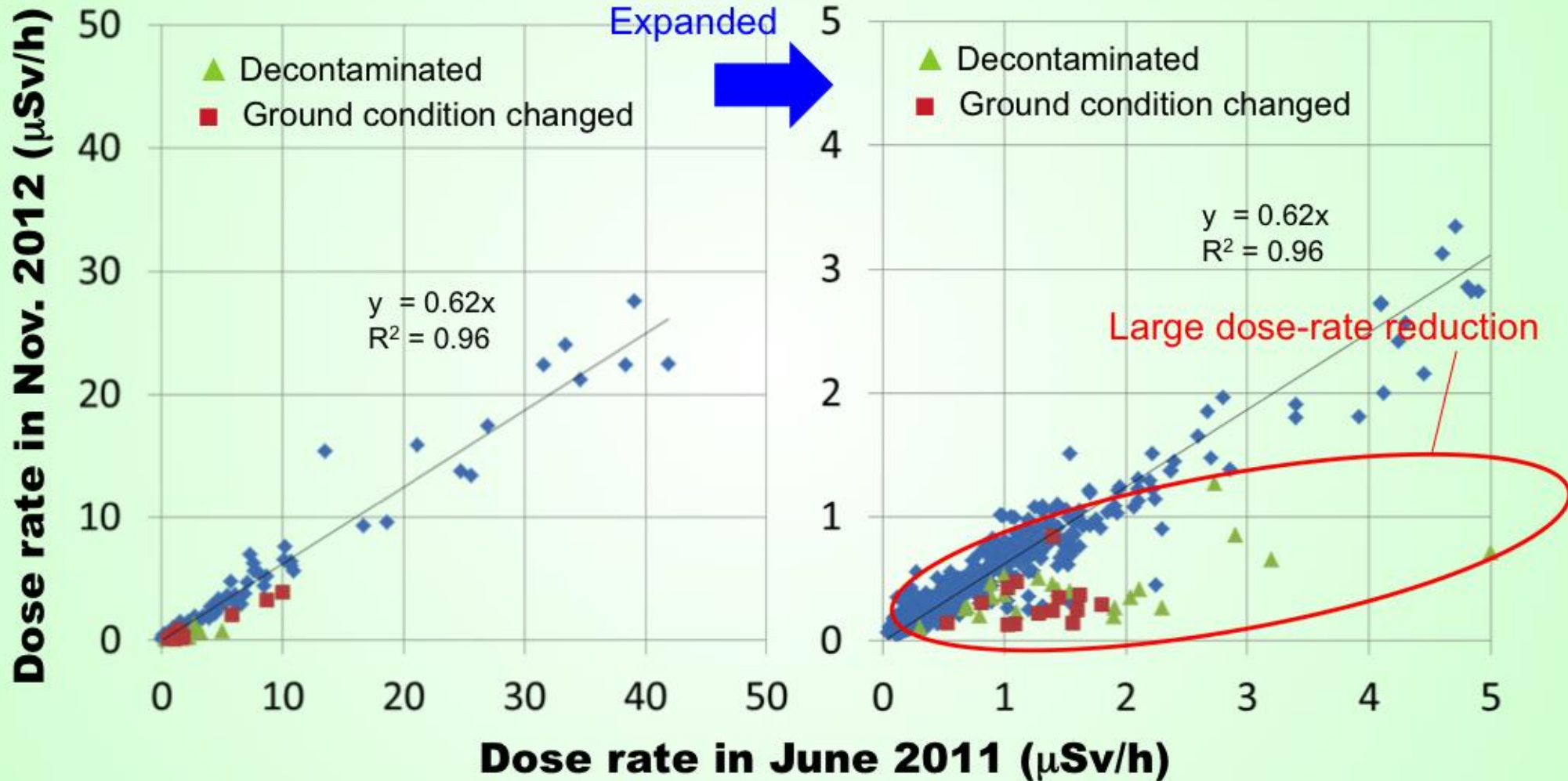
- 19.0 <
- 9.5 - 19.0
- 3.8 - 9.5
- 1.9 - 3.8
- 1.0 - 1.9
- 0.5 - 1.0
- 0.2 - 0.5
- 0.1 - 0.2
- ≤ 0.1

Distribution of areas having different dose rate ranges within the 80 km zone

- Areas more than $0.2 \mu\text{Sv/h}$ are decreasing, less than $0.2 \mu\text{Sv/h}$ increasing.
- Nearly 70% of the total area has dose rates below $0.5 \mu\text{Sv/h}$.

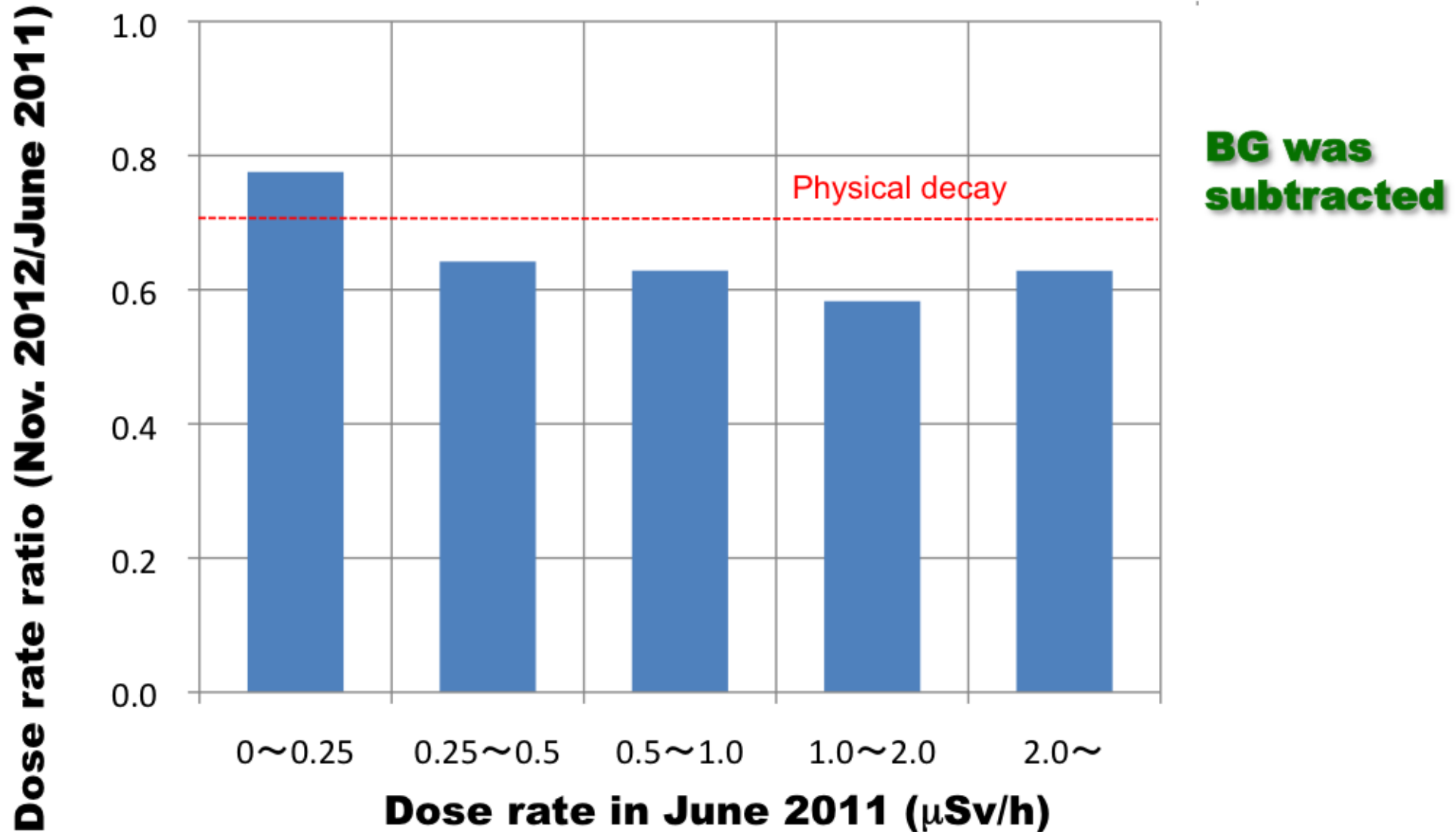


Comparison of dose rates in air at 1 m between June 2011 and Nov. 2012



- Dose-rate reduction was smaller than 40%. (Physical decay : 29%)
- There exist locations showing large dose-rate reduction.

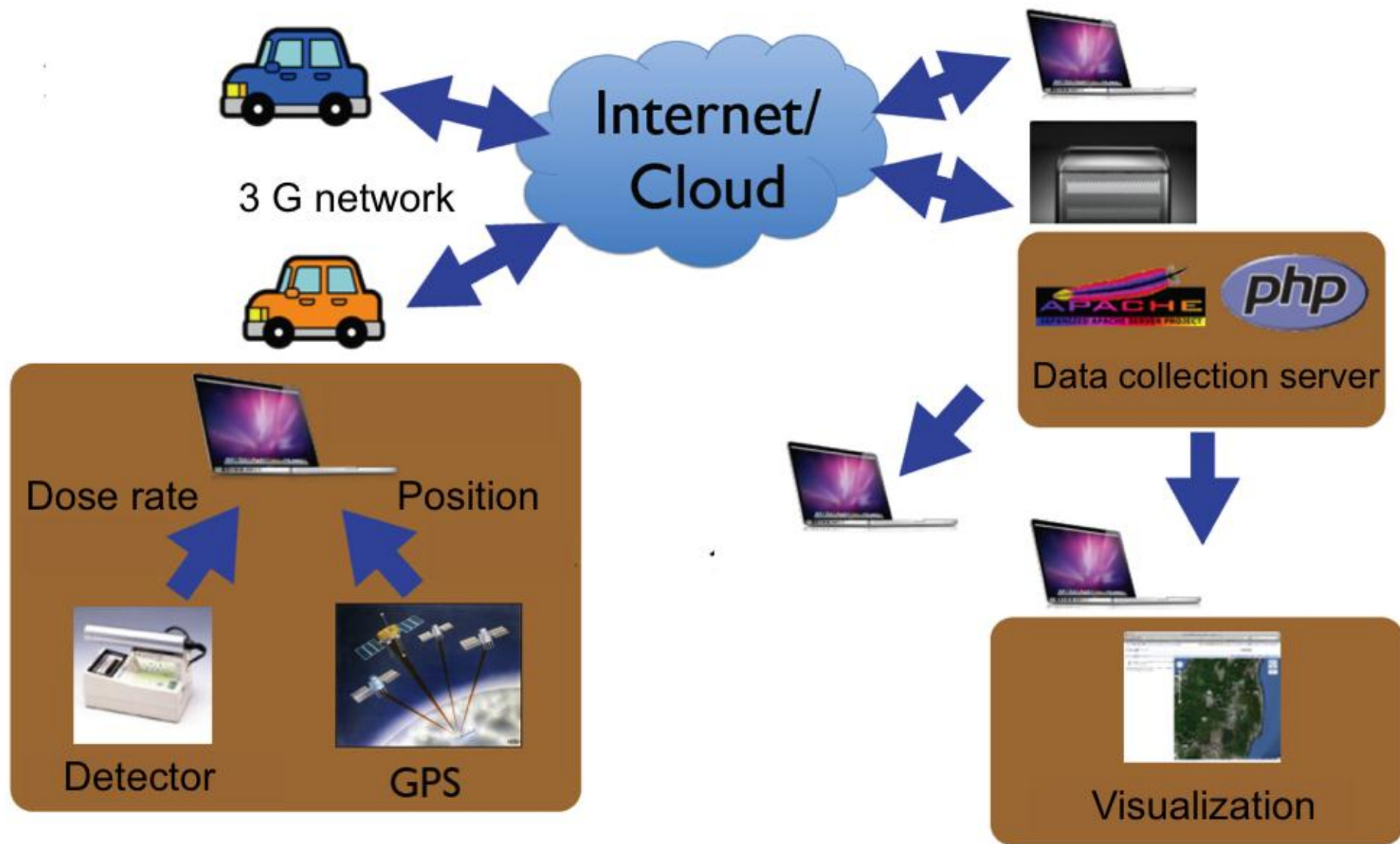
Comparison of dose rates in air at 1 m between June 2011 and Nov. 2012



● **Dose-rate reduction has an initial dose rate dependency.**

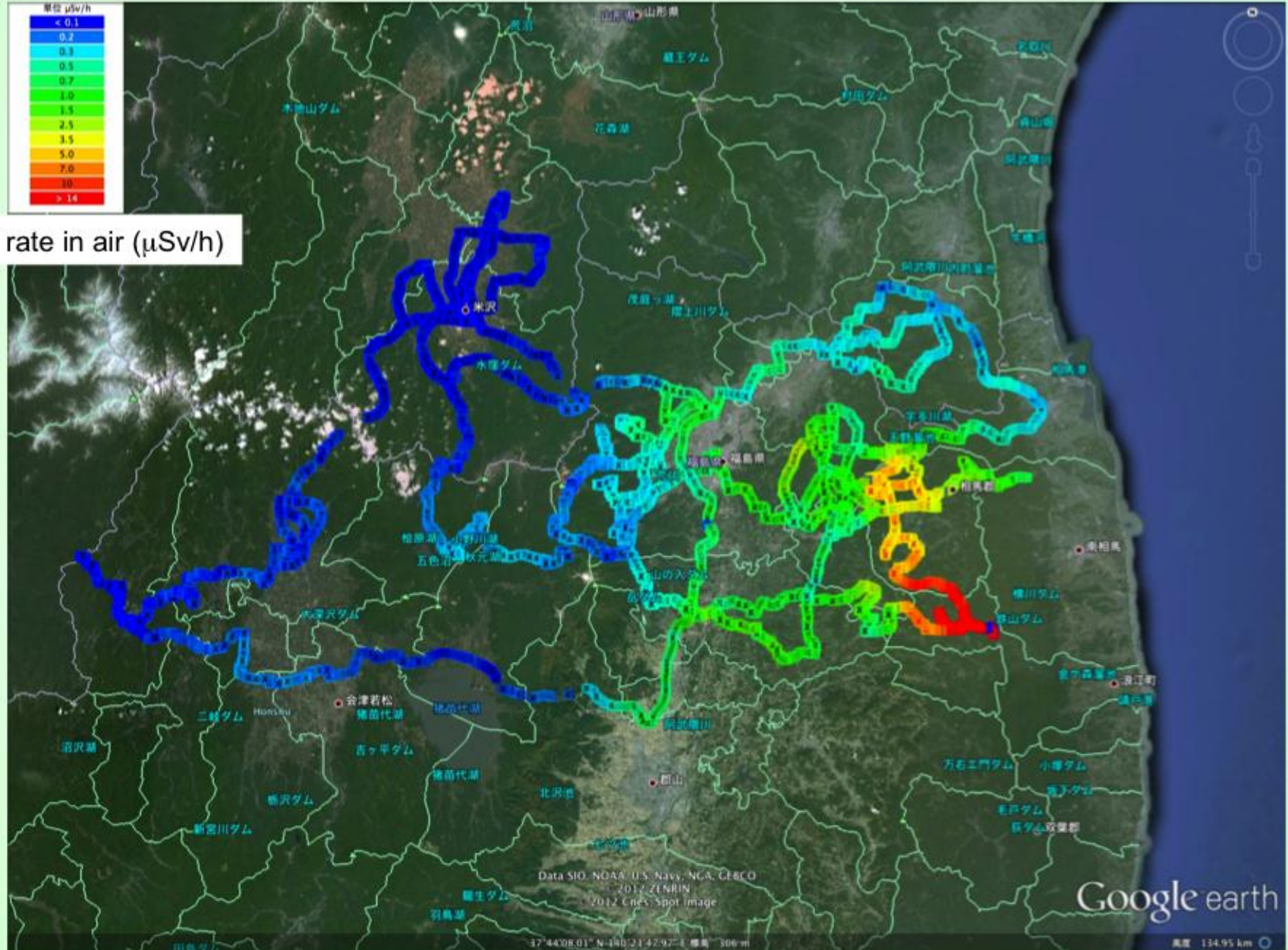
Car-borne survey using KURAMA system

(developed at Kyoto Univ.)



● **Data transfer through a cellular phone network**

Real-time display of car-borne survey data

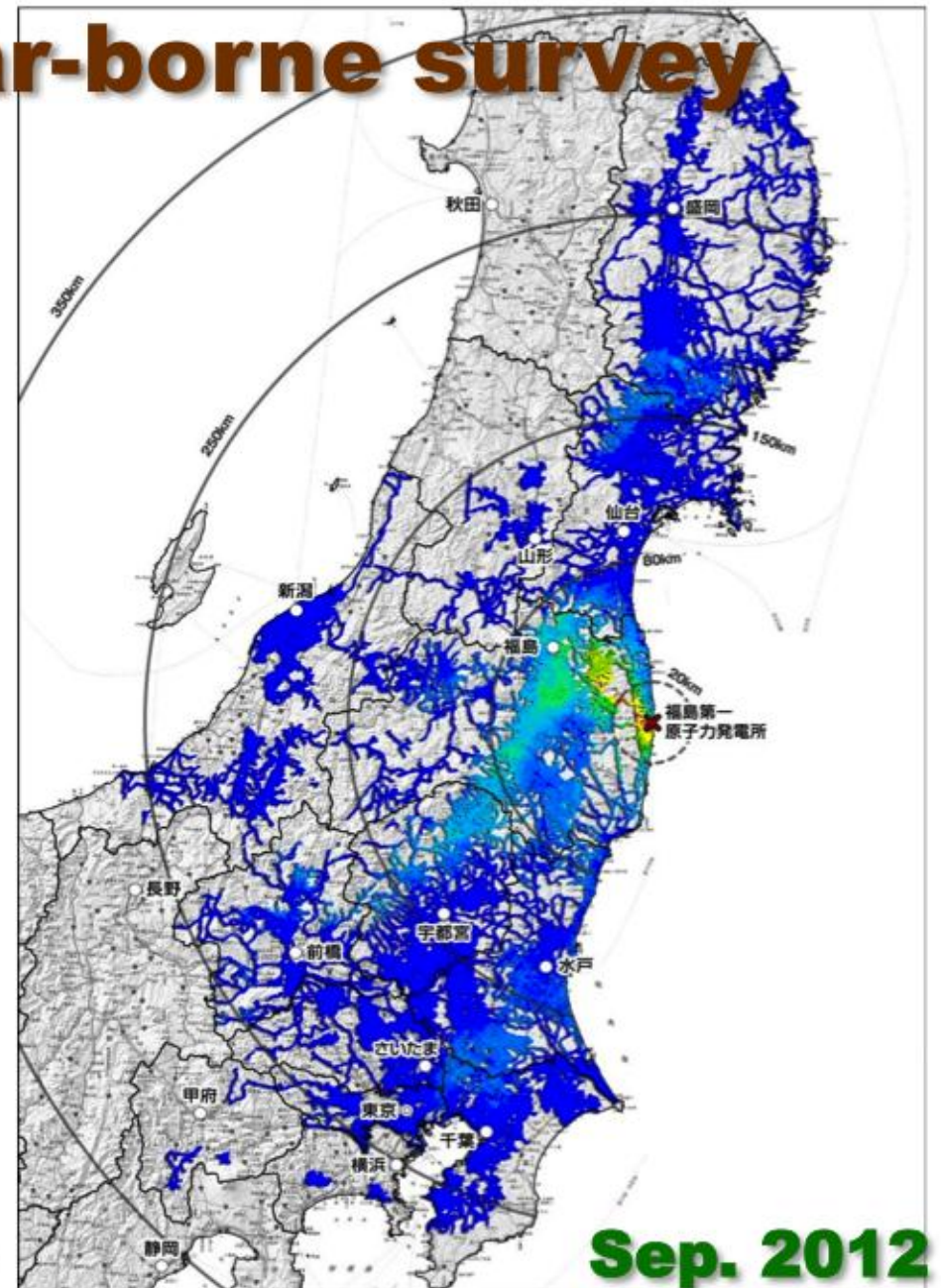
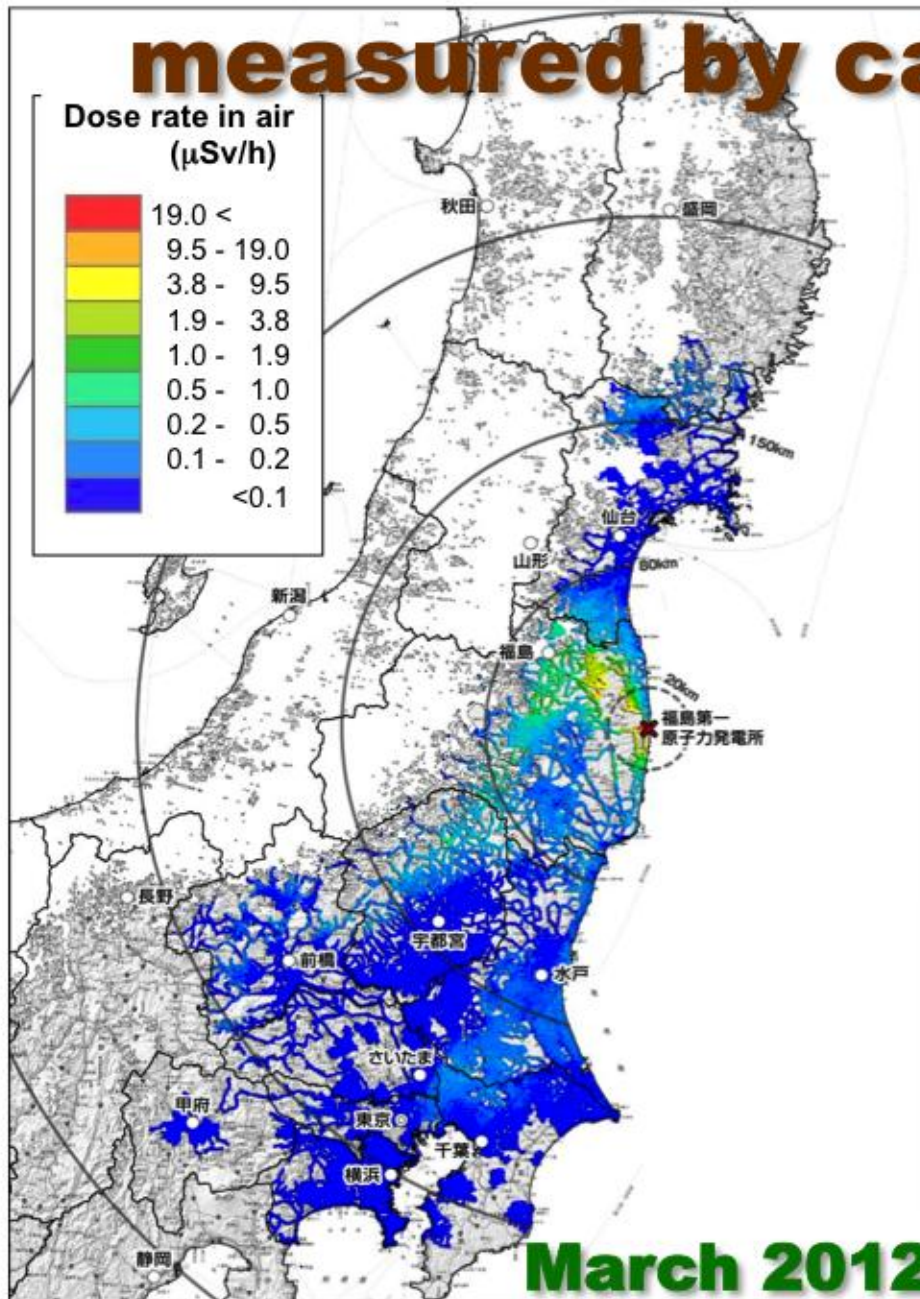


KURAMA-II system

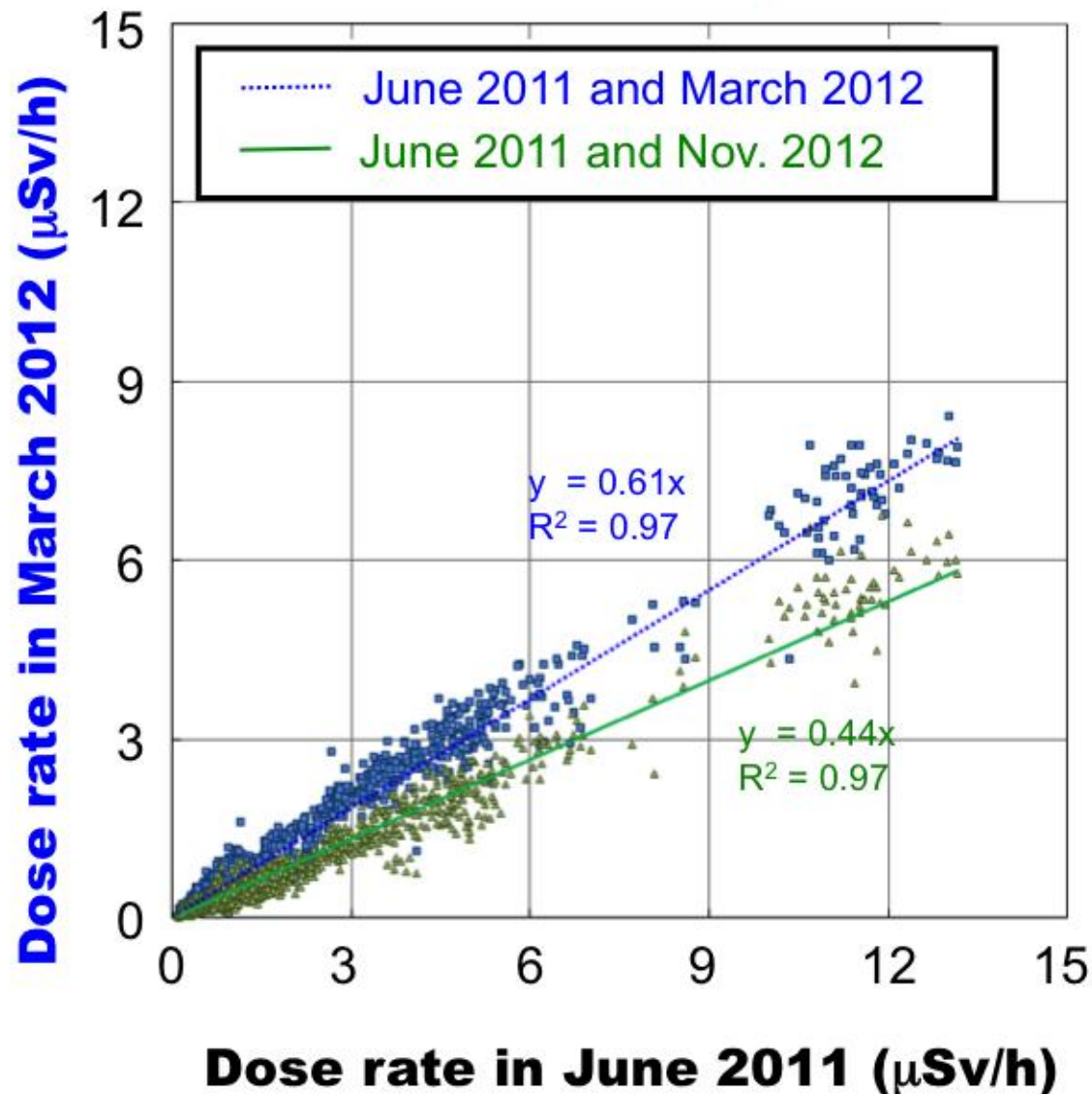
- **Compact**
- **Easy to operate**
- **Distribute 100 systems to about 200 local governments**
- **Each local government makes a survey as it would like.**



Distribution of dose rates in air measured by car-borne survey



Comparison of dose rates in air measured by car-borne survey

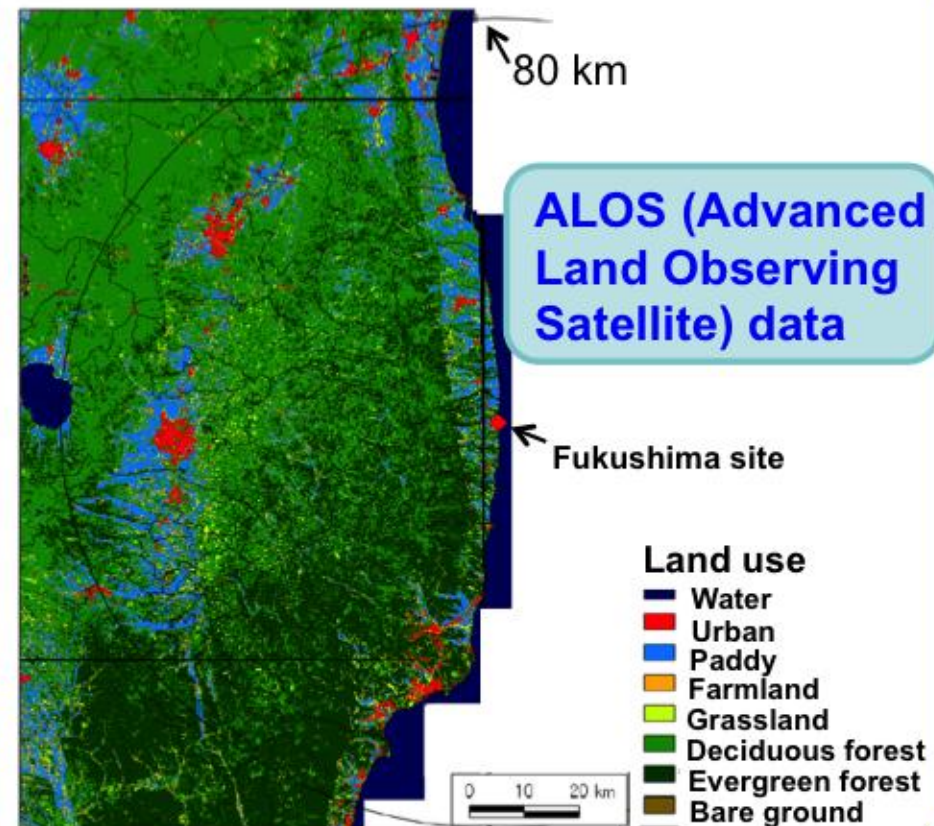
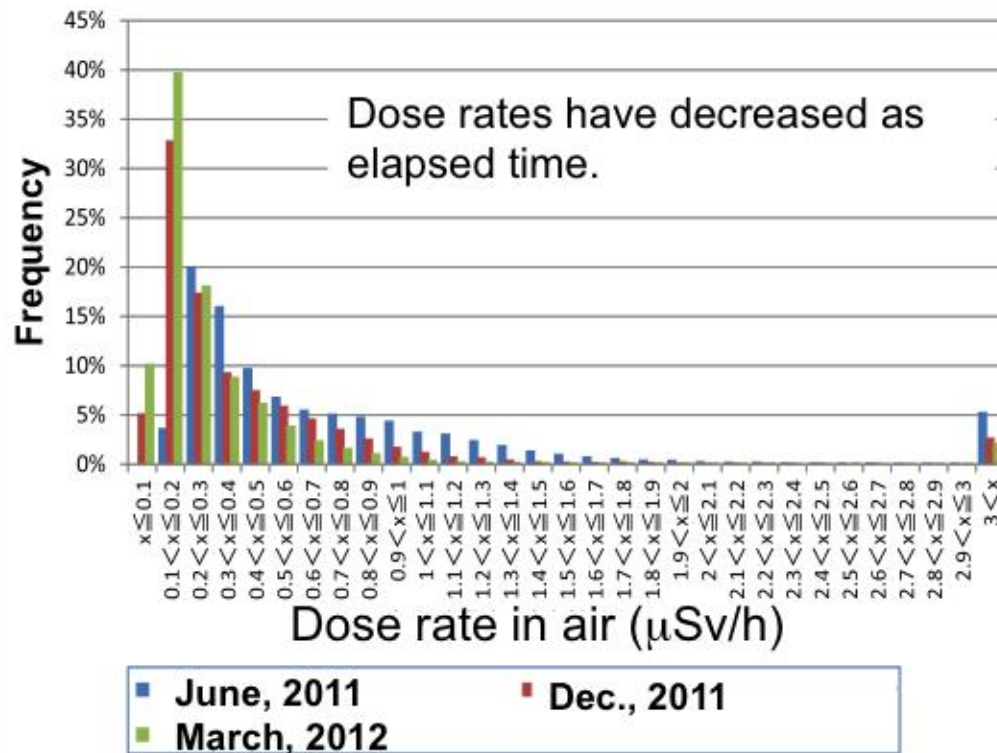


● Dose rates decreased by about 40% from June 2011 to March 2012. (Physical decay 20%)

● Dose rates decreased by more than 55% from June 2011 to Nov. 2012. (Physical decay 29%)

Analyses based on land uses

Car-bone survey data

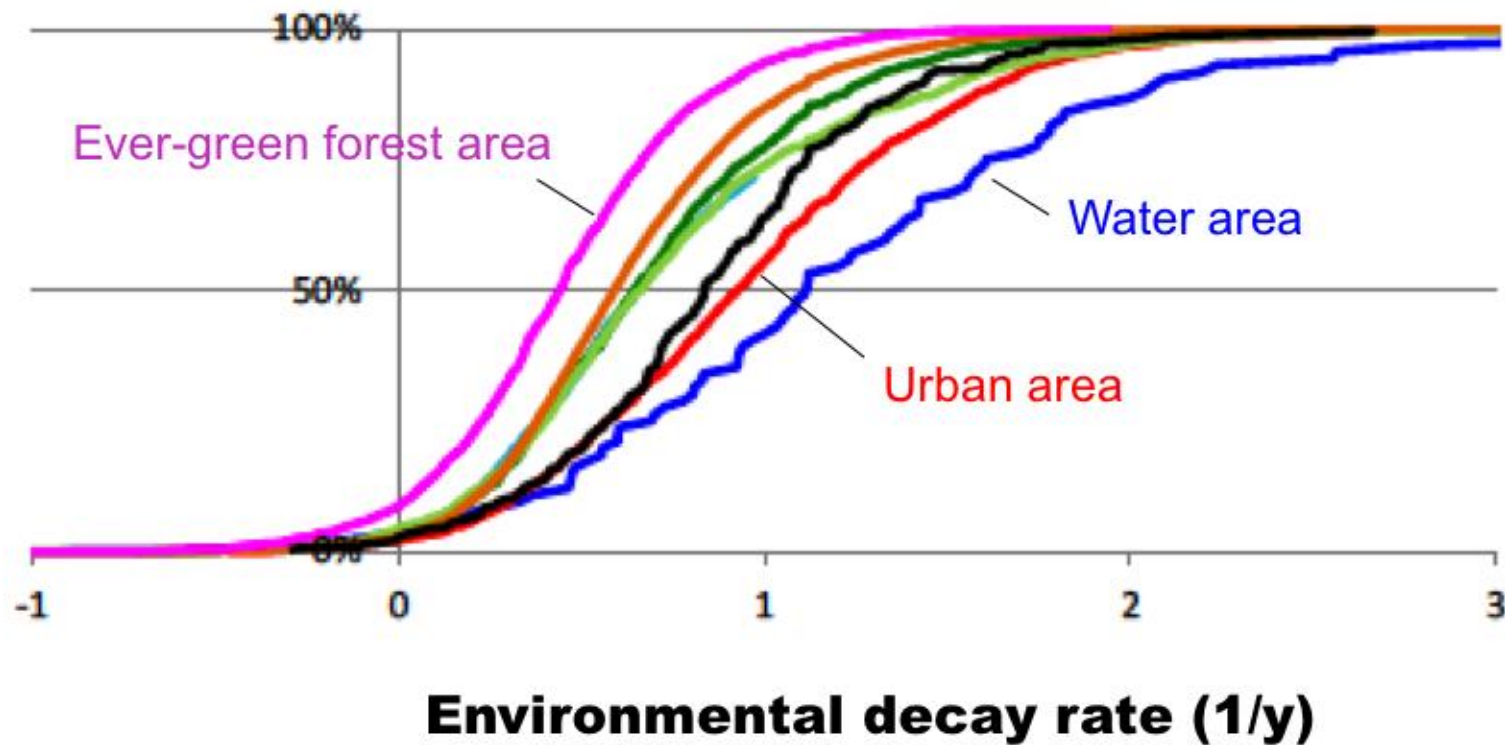


• Dose rate reduction tendency was analyzed in connection with different land uses

Dose rate reduction tendency for different land uses

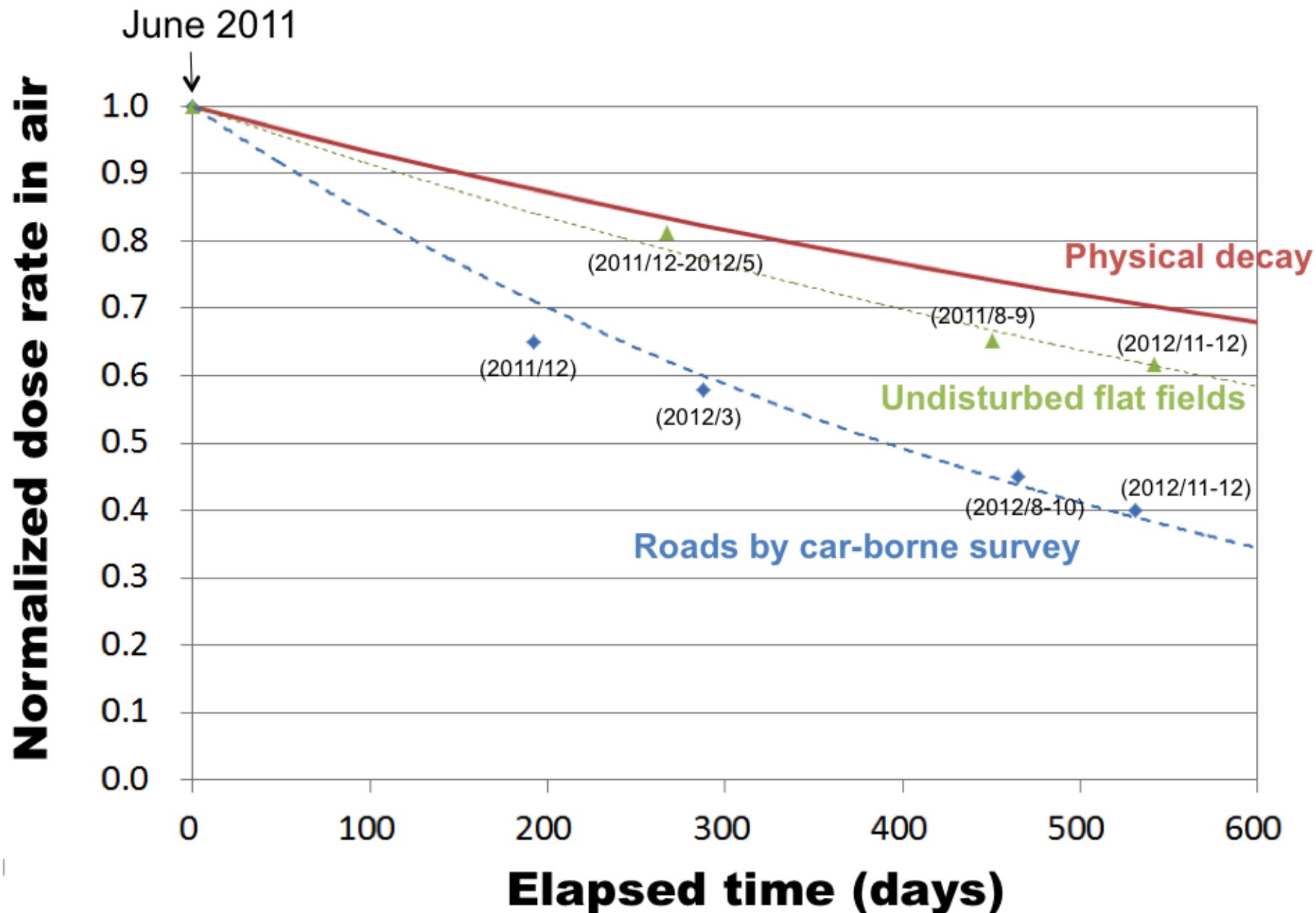
- Slow in ever-green forest area
- Fast in urban and water areas

Cumulative Distribution Frequency of environmental decay rate



Excluding physical decay

Temporal change of dose rates in air



Summary on dose rates in air

- 1. Dose rates in air above roads have decreased much faster than those at undisturbed flat fields.**
- 2. Dose-rate reduction tendency depends on**
 - a) land-use: fast in urban, slow in forest,**
 - b) magnitude of initial air dose rate,**
 - c) local distribution of contamination.**
- 3. Decontamination is considered to reduce dose rates in air by a factor of 2-5**

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Soil sampling for investigating depth profiles

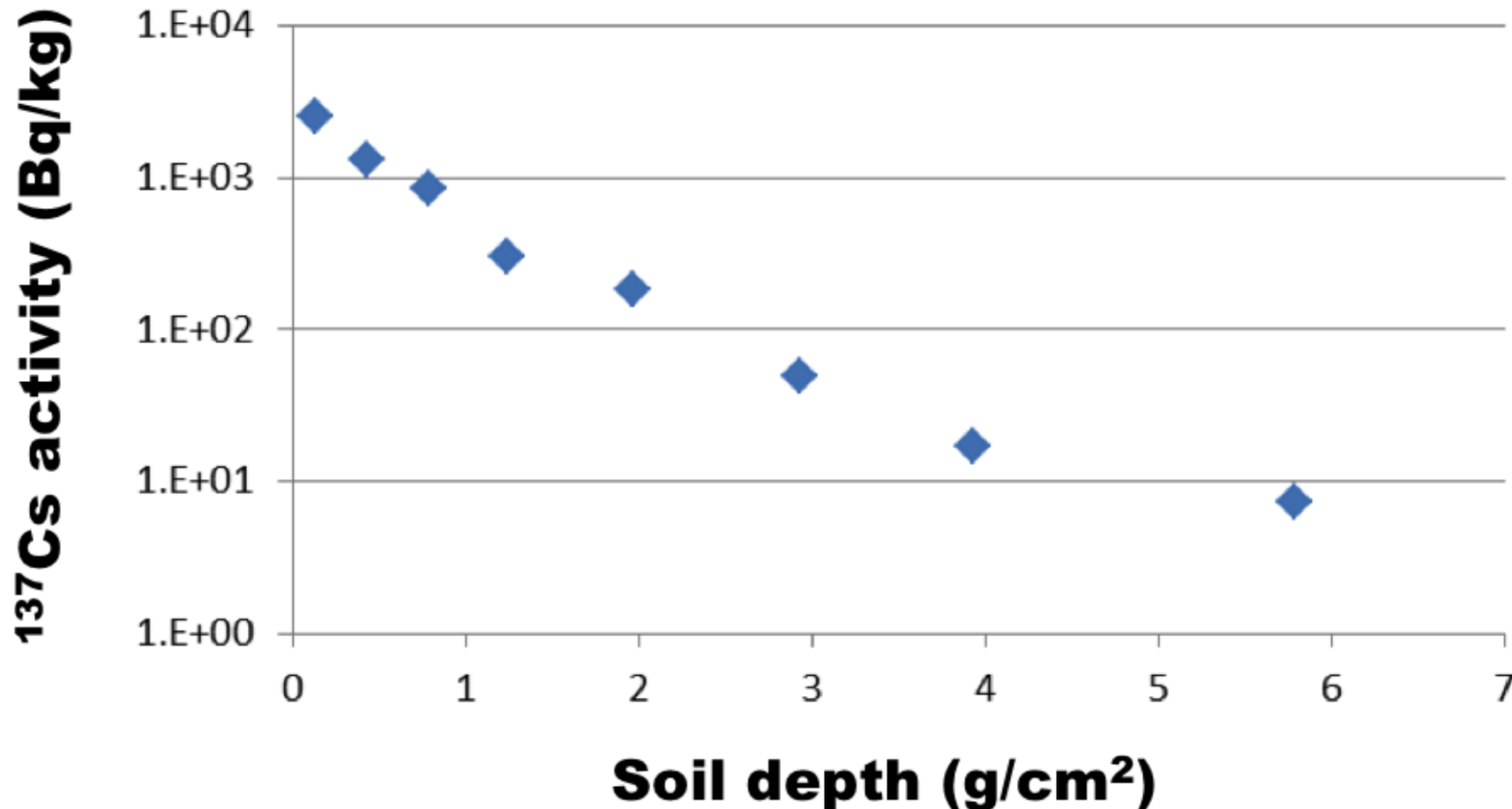
● Soil was sampled at 85 locations using scraper plates in:

- 1) Dec. 2011,
- 2) Aug. 2012,
- 3) Dec. 2012.



Example of ^{137}Cs depth profile in soil

- Most depth profiles could be approximated by exponential distribution

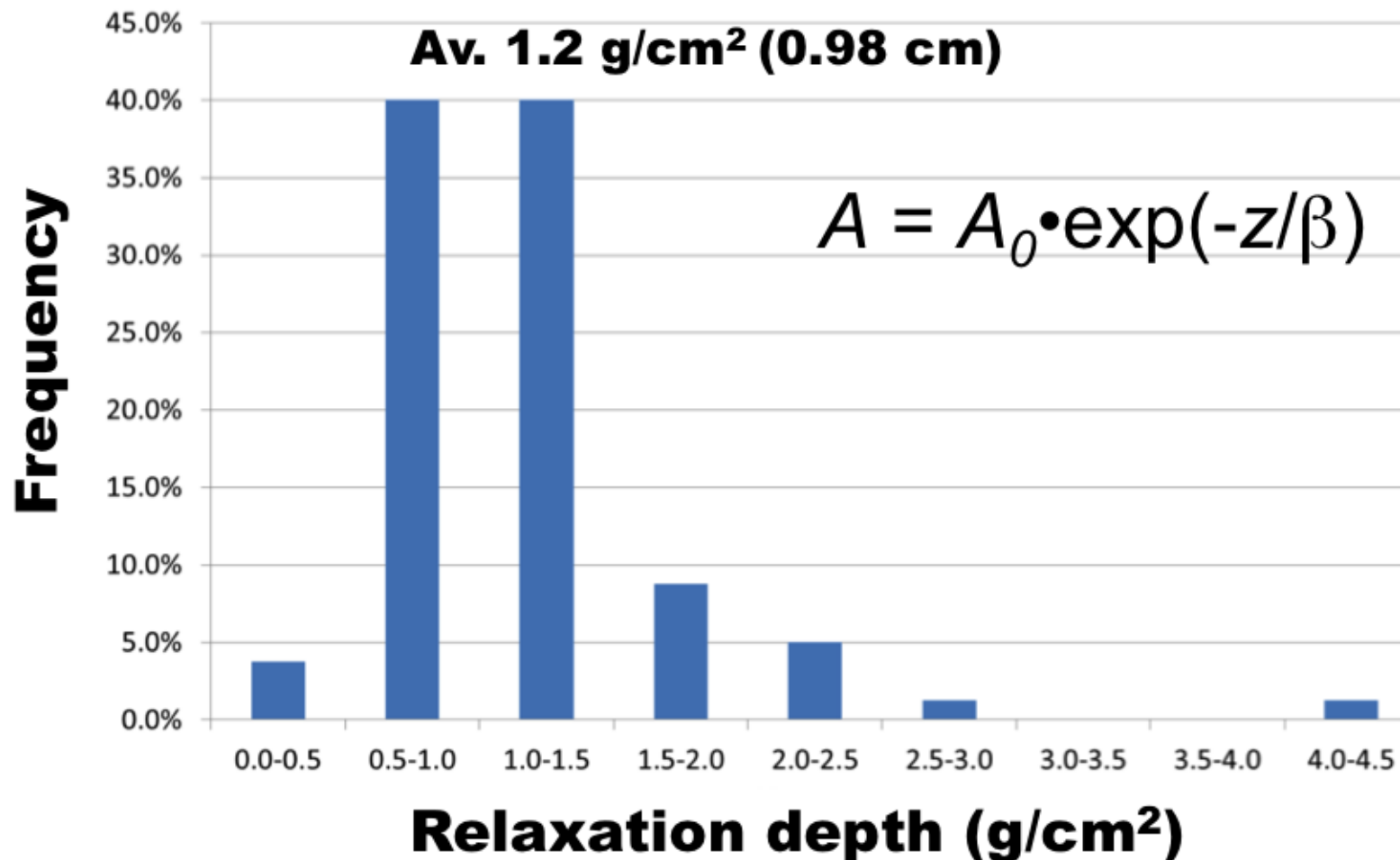


(December, 2011)

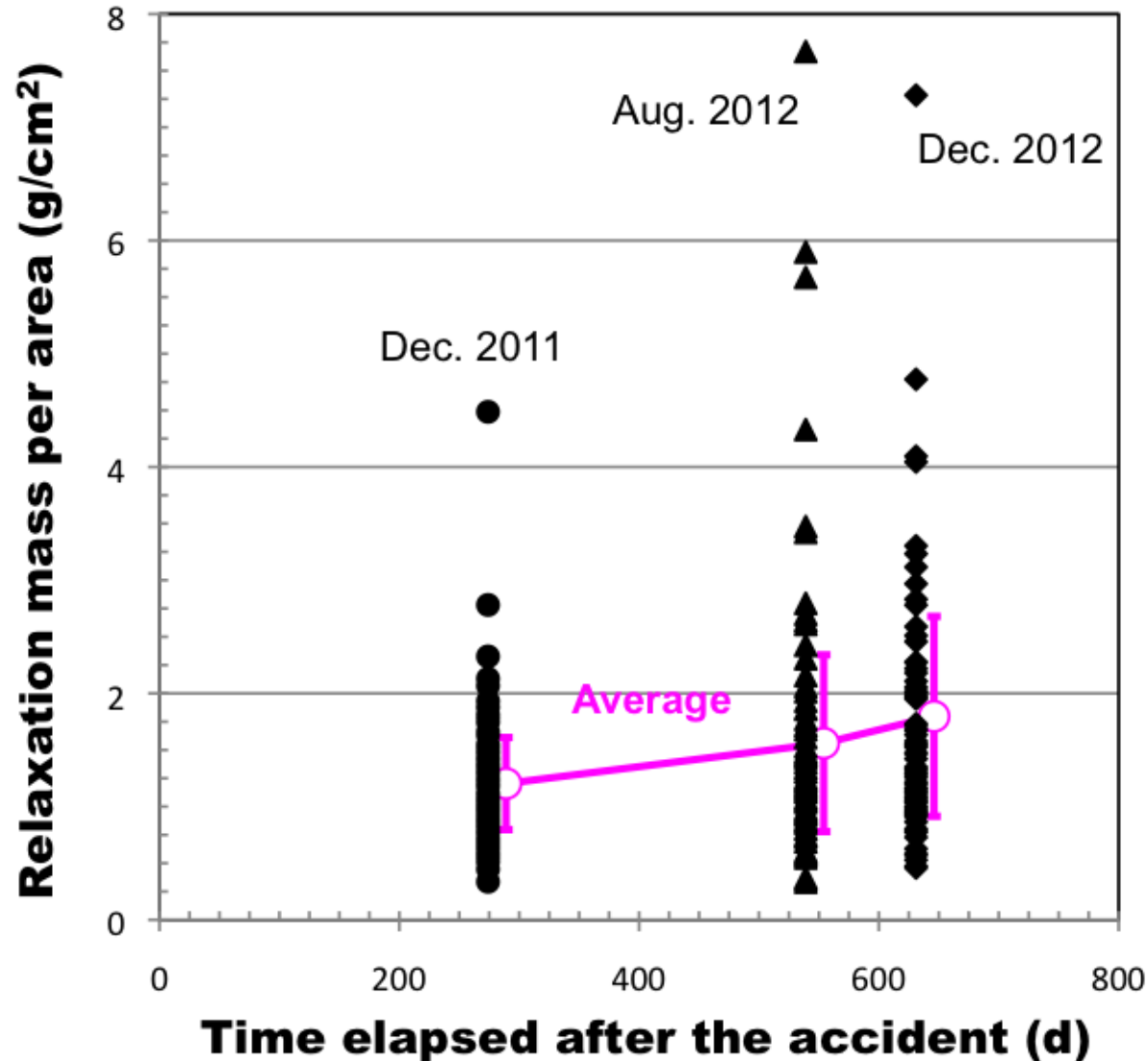
Distribution of relaxation depth β

- Indicator of radionuclide migration into soil

(Dec. 2011)



Temporal change of relaxation depth β



$$A = A_0 \cdot \exp(-z/\beta)$$

Average β

● **1.2 ± 0.4 g/cm²**
(Dec. 2011)

▲ **1.6 ± 0.8 g/cm²**
(Aug. 2012)

◆ **1.8 ± 0.9 g/cm²**
(Dec. 2012)

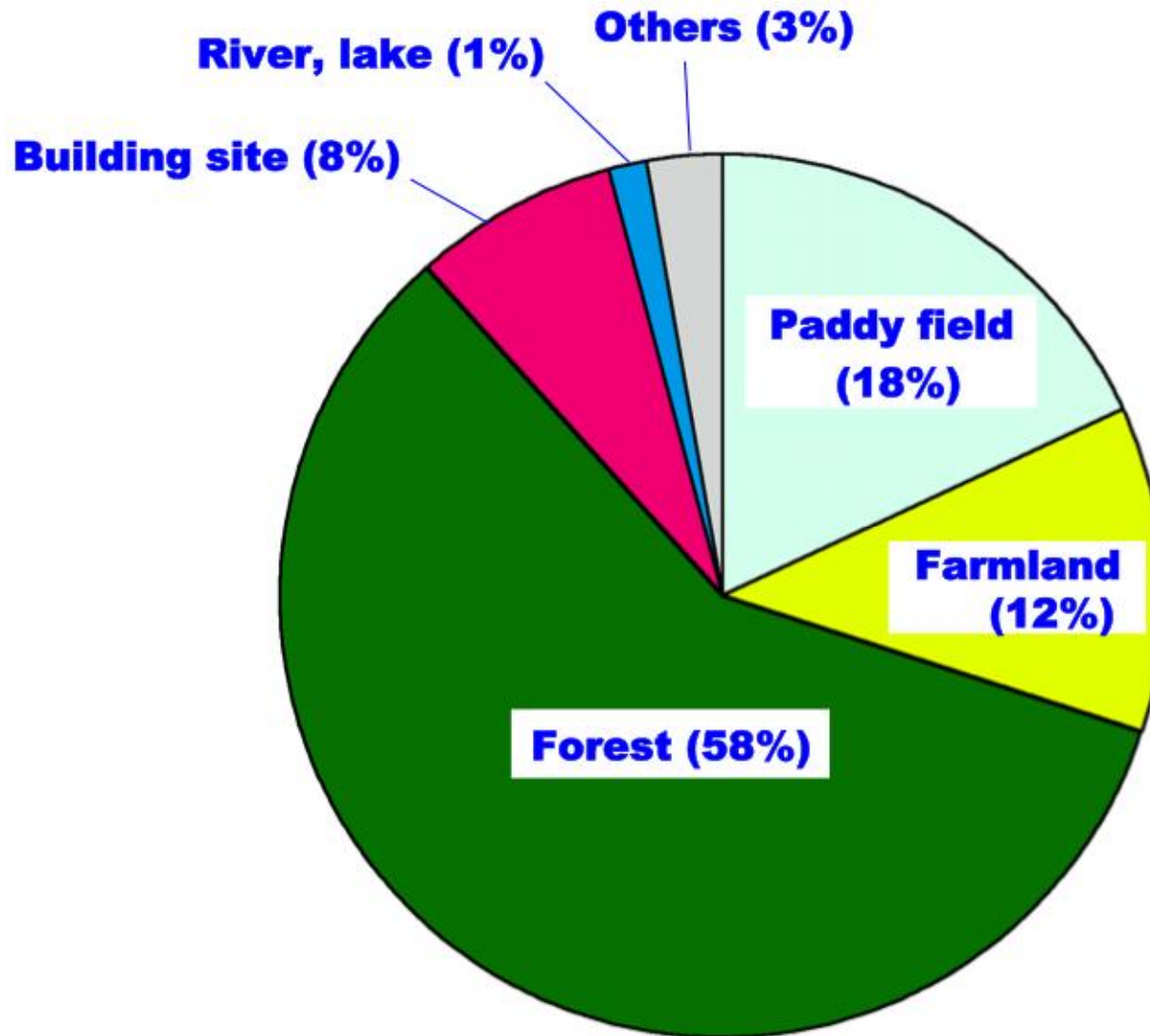
● **β has gradually increased with time.**

● **Most cesium exists within 5 cm depth**

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Fractions of Cs-137 deposition for different land uses within the 80 km zone

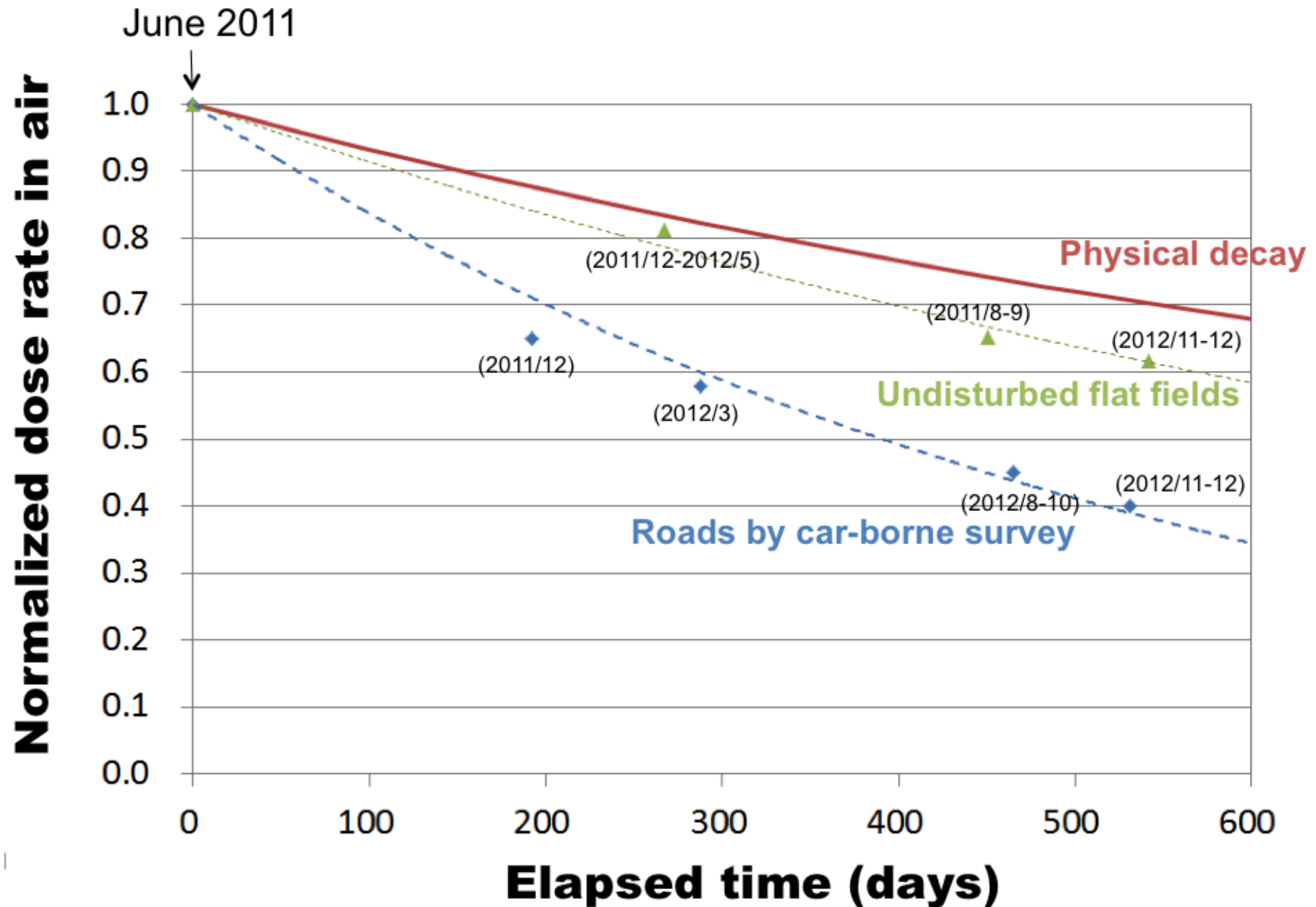


● Evaluated based on deposition densities measured in undisturbed fields in Dec. 2012.

● Assumed that the density does not change within 1 km square.

(Dec. 2012)

Temporal change of dose rates in air

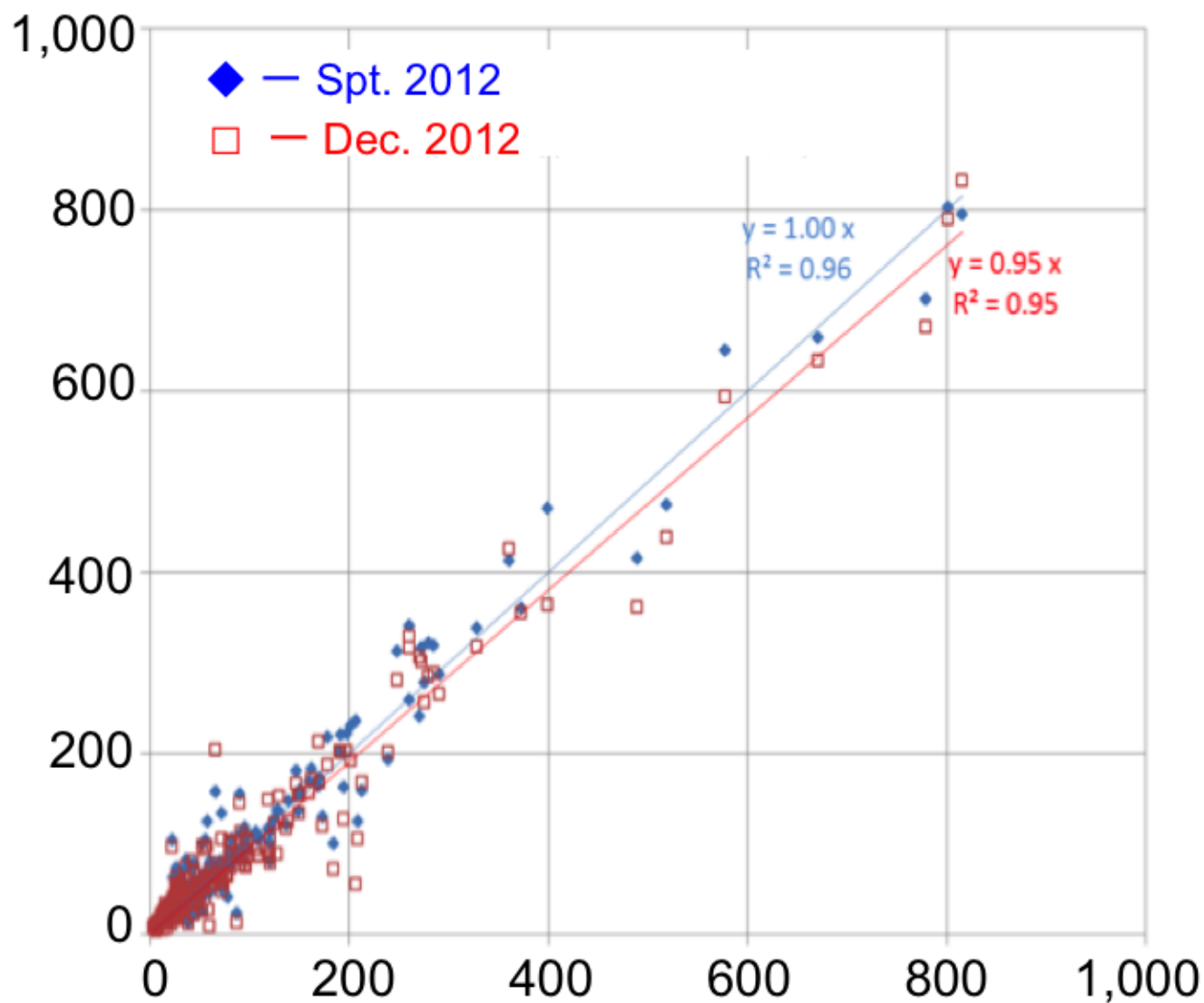


Undisturbed fields

- 1. Dose rates in air have decreased a little faster than the physical decay.**
- 2. Movement of cesium in horizontal directions seems small.**
- 3. Cesium has gradually penetrated into deeper parts in the ground: the excess decrease in dose rates can be explained by this penetration.**
- 4. Yet small amount of cesium is considered to have moved in horizontal directions into low-contaminated areas.**

Temporal change in Cs-137 deposition density

Deposition density in Sept. 2012 (kBq/m²)



Deposition density in Dec. 2012 (kBq/m²)

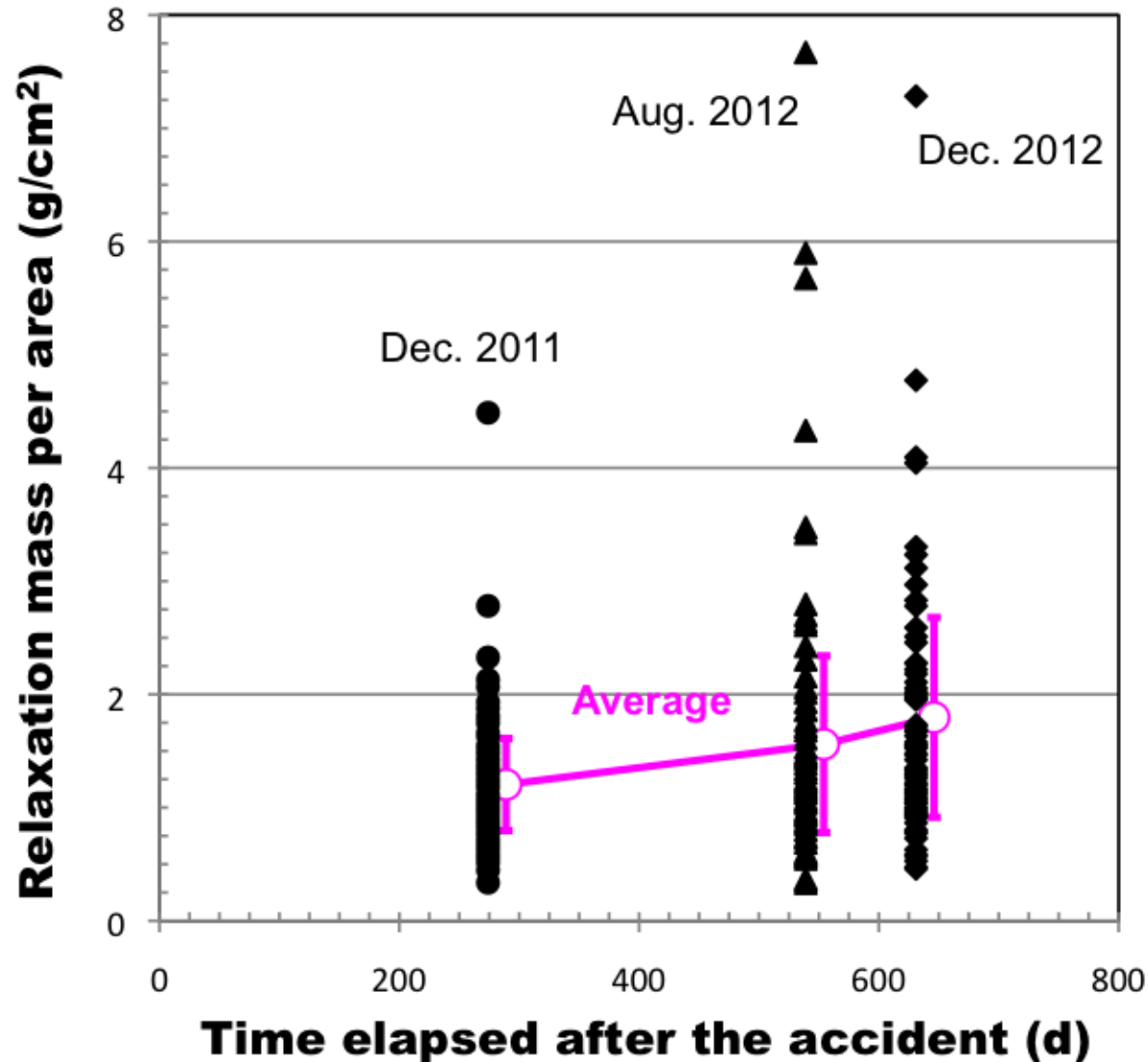
● **Cs-137 deposition densities have not changed much in undisturbed flat fields.**

Deposition density in March 2012 (kBq/m²)

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Temporal change of relaxation depth β



$$A = A_0 \cdot \exp(-z/\beta)$$

Average β

● **1.2 ± 0.4 g/cm²**
(Dec. 2011)

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(Aug. 2012)

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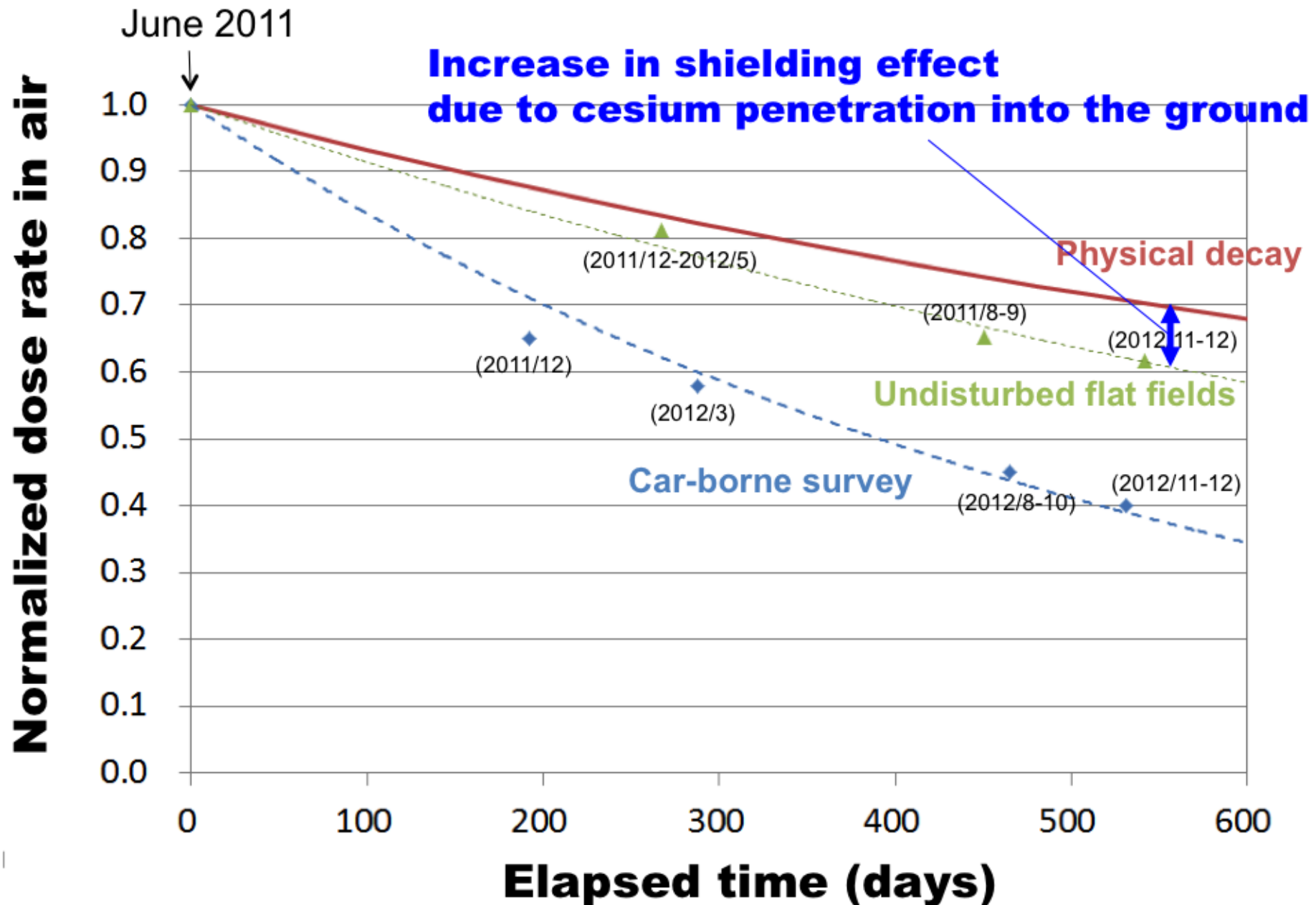
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● **Most cesium existed within 5 cm depth**

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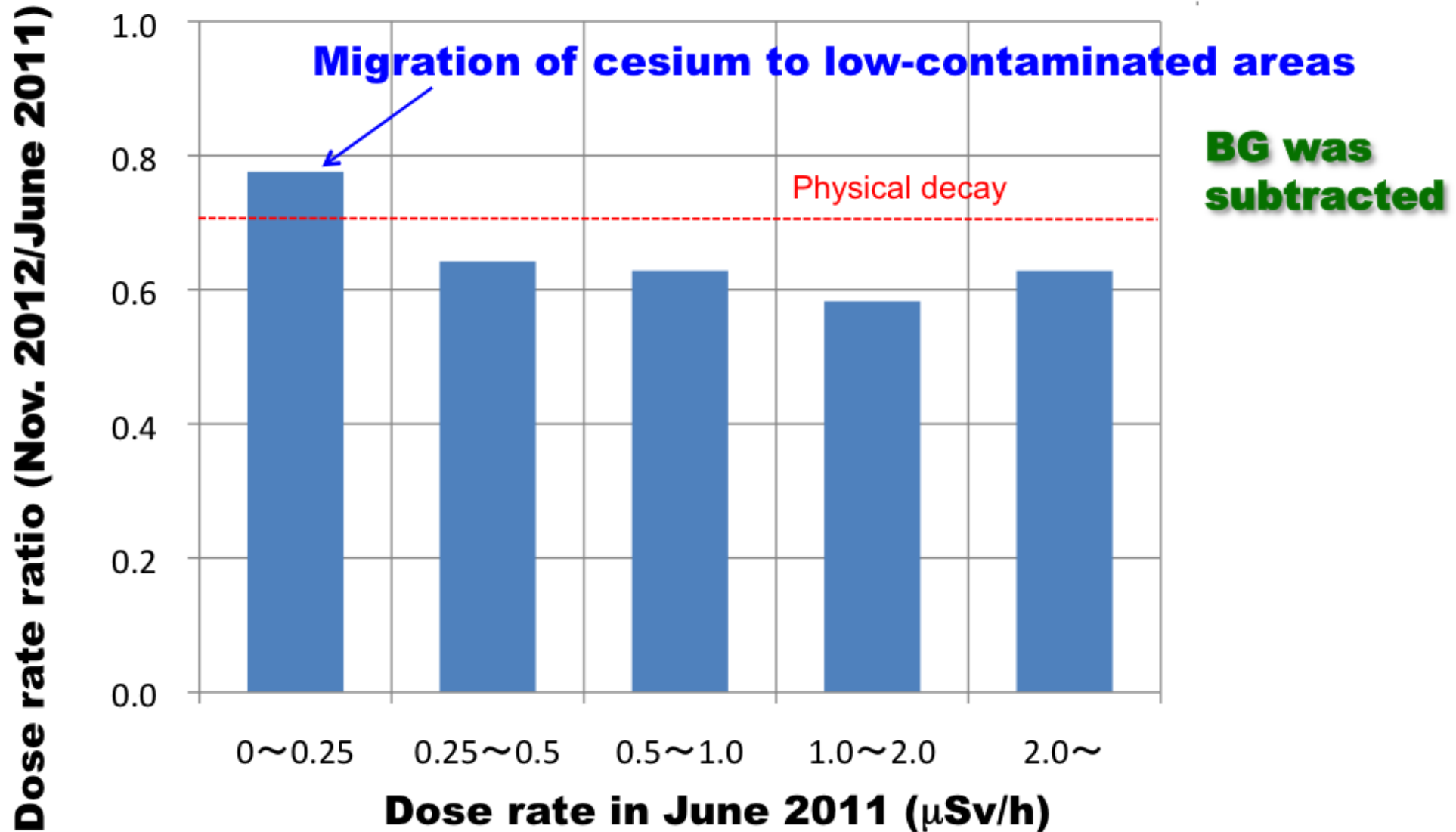
Temporal change of dose rates in air



Undisturbed fields

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Comparison of dose rates in air at 1 m between June 2011 and Nov. 2012

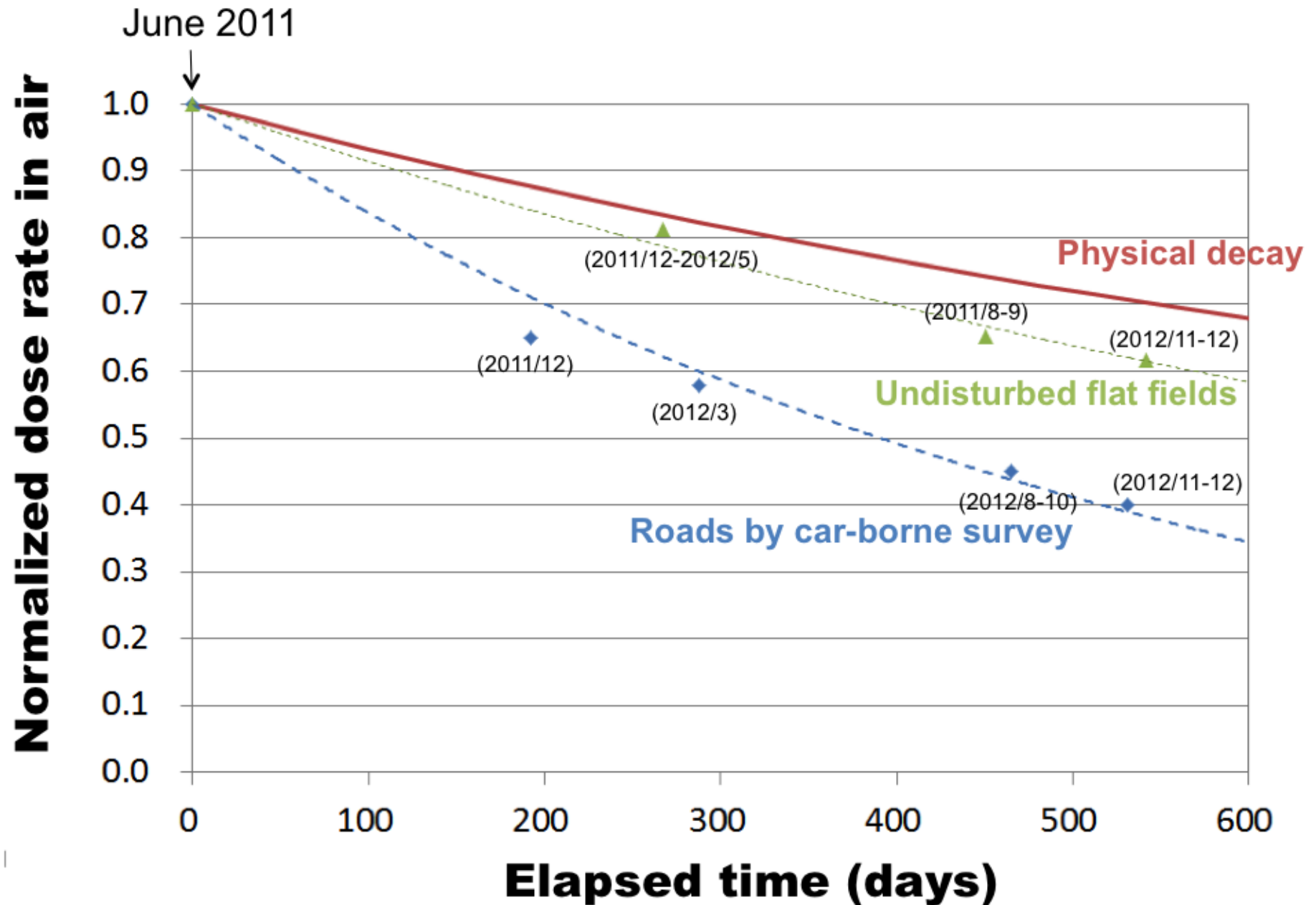


● **Dose-rate reduction has an initial dose rate dependency.**

Car-borne survey on roads

- 1. Air dose rates above roads have decreased much faster than the physical decay.**
 - a) Cesium on roads is easily washed away.**
 - b) Cesium around roads is inferred to have been removed somehow.**
- 2. Air dose rate reduction is slow in forest area, and fast in urban and water areas.**
 - a) Cesium is being kept in the forest system.**
 - b) Flooding plays an important role in removing cesium.**

Temporal change of dose rates in air



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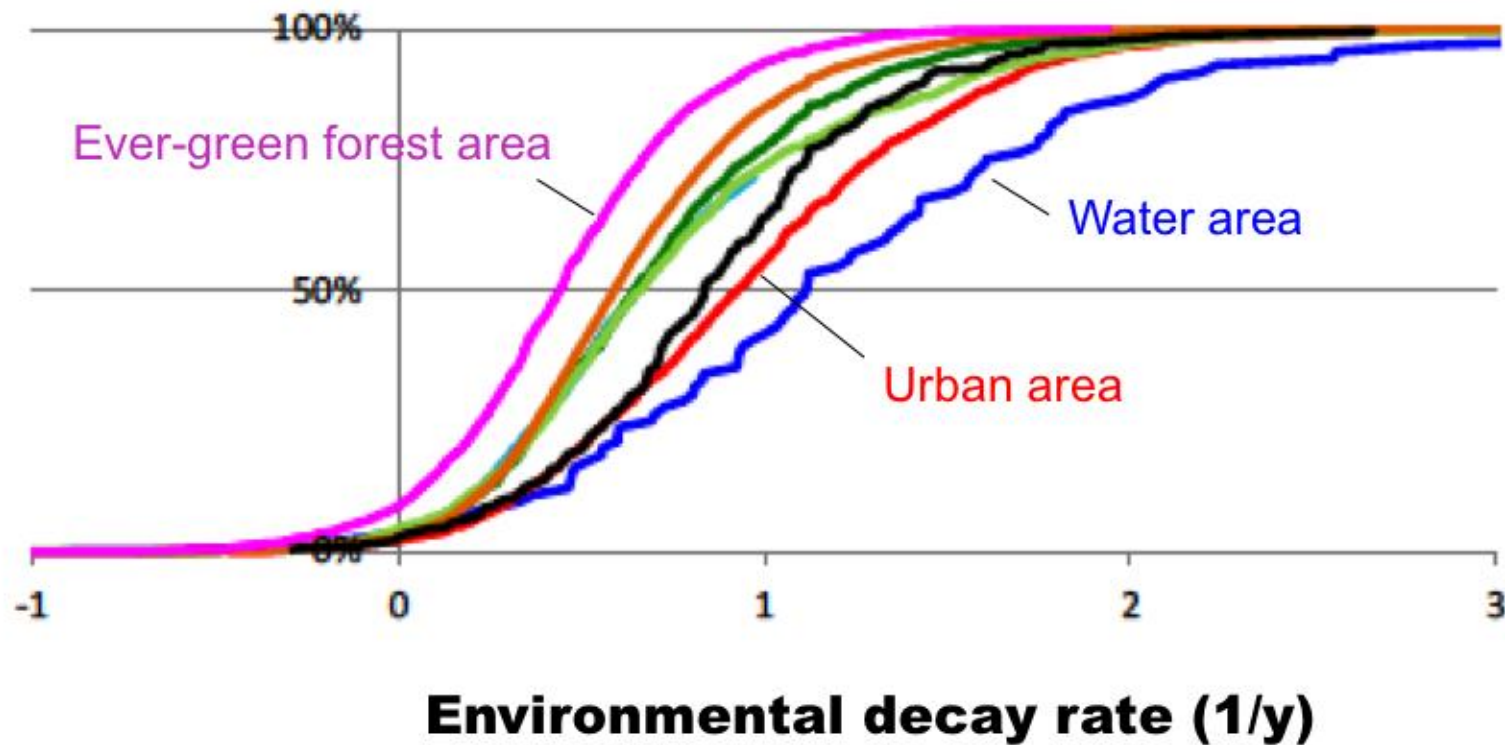
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Dose rate reduction tendency for different land uses

- Slow in ever-green forest area
- Fast in urban and water areas

Cumulative Distribution Frequency of environmental decay rate



Excluding physical decay

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Air dose rate reduction

Slow

Fast



Forest



**Undisturbed
field**



Near road

Air dose rate reduction

Slow

Fast



Forest



**Undisturbed
field**



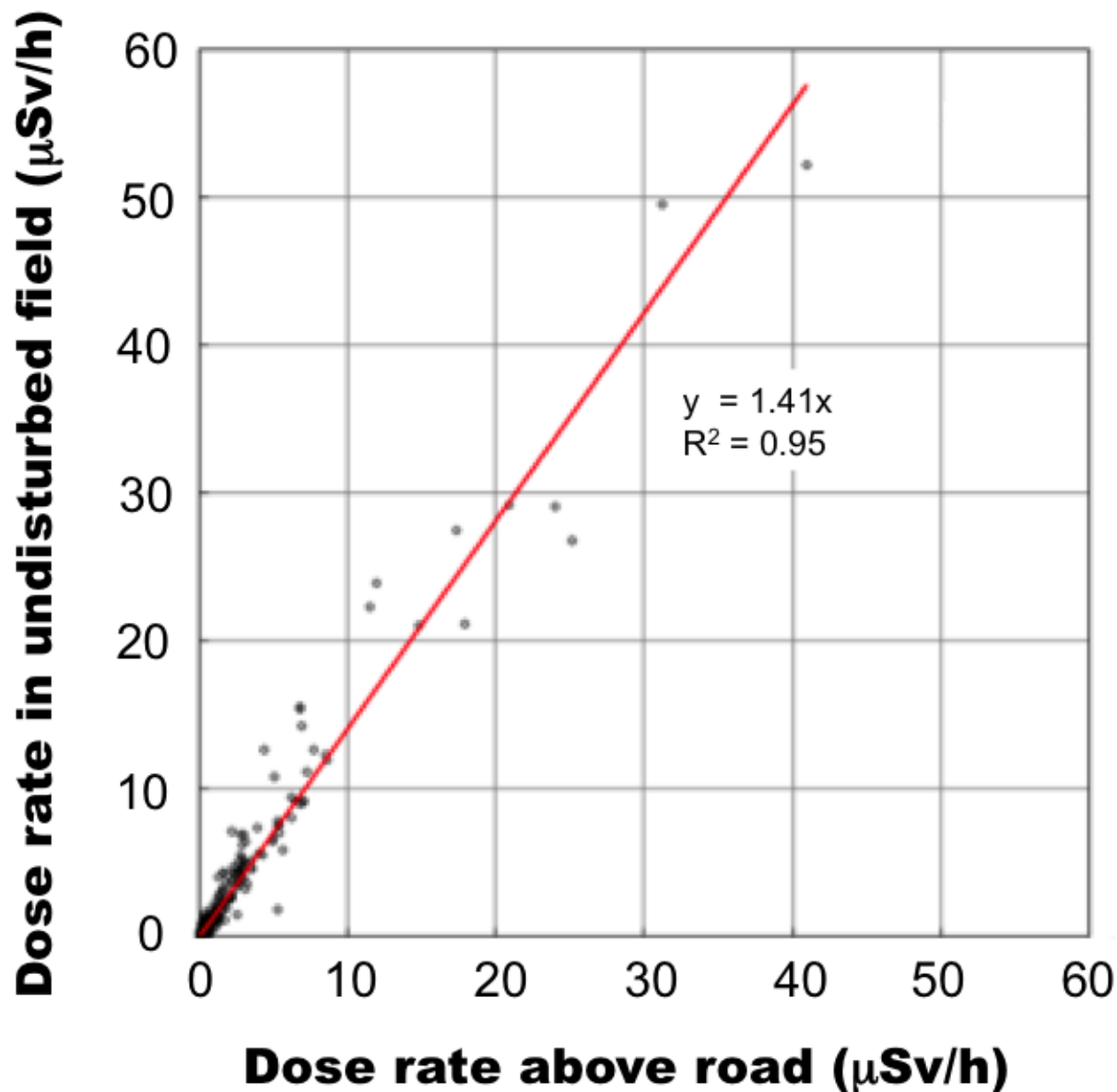
Near road

High

Low

Air dose rates

Comparison of dose rates between roads and undisturbed field



Dose rates in undisturbed fields are several tens percent higher than those by car-borne survey.

(Nov. 2012)

Air dose rate reduction

Slow

Fast



Forest



**Undisturbed
field**



Near road

High

Low

Air dose rate

How about in intermediate areas?



Farmland

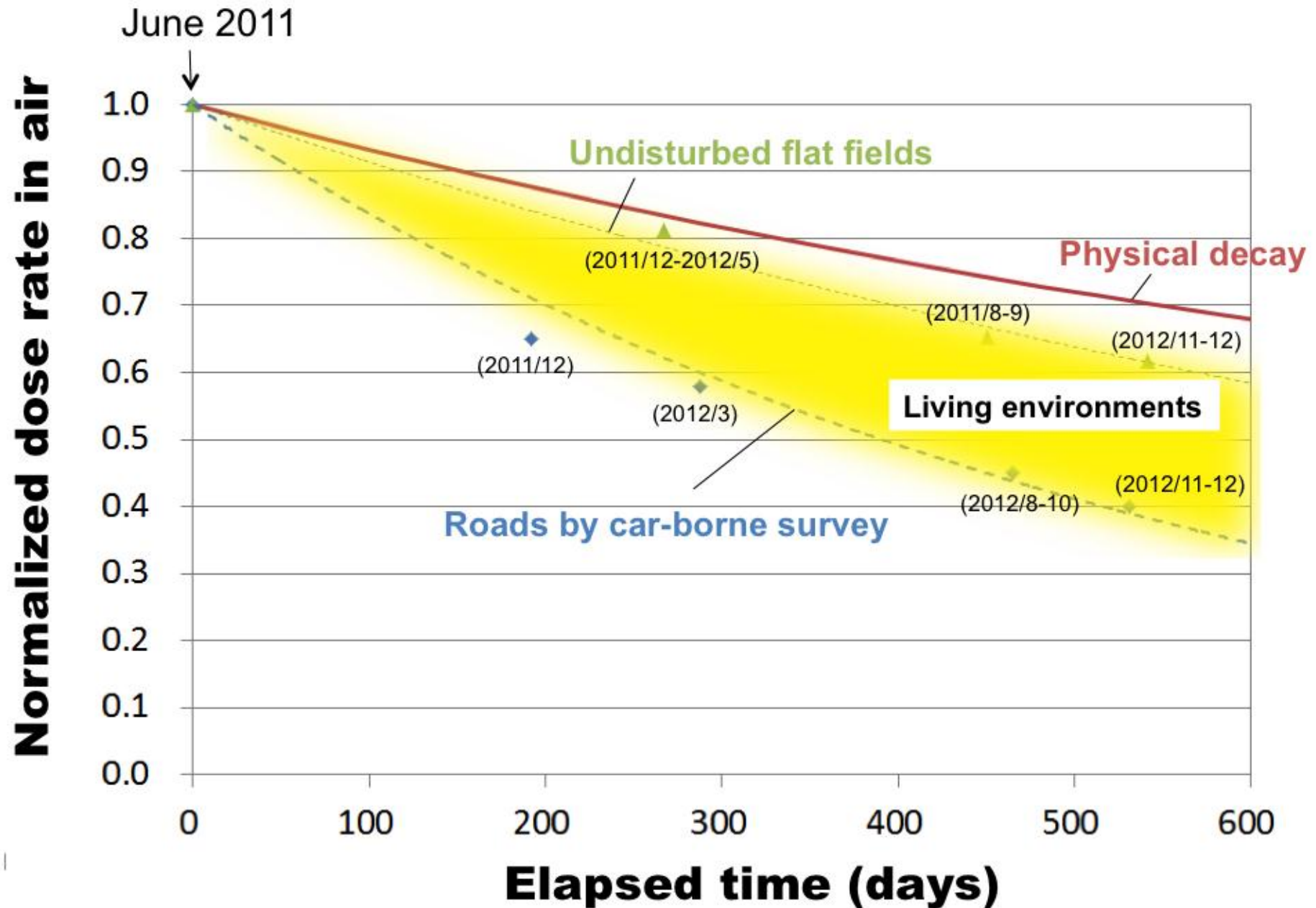


Residence



Town

Temporal change of dose rates in air



Example of man-borne survey



Example of man-borne survey



Ongoing program

1. Man-borne survey

- a) Examine detailed distributions of air dose rates in living environment**
- b) Find out the relation of the dose rates to those in undisturbed fields and above roads**

2. Prediction model

- a) Semi-empirical model assuming two components indicating time-dependent reduction (fast and slow) in air dose rates**
- b) Determine the parameters on the basis of statistical analysis of large amount of air dose data**



Thank you for your attention