- Long-term assessment of airborne radiocesium after the 1 Fukushima nuclear accident: Re-suspension from bare soil 2 and forest ecosystem  $\frac{1}{2}$ 3 4 M. Kajino<sup>1, 2</sup>, M. Ishizuka<sup>3</sup>, Y. Igarashi<sup>1</sup>, K. Kita<sup>4</sup>, C. Yoshikawa<sup>5</sup>, and M. Inatsu<sup>6</sup> 5 6 7 [1] {Meteorological Research Institute (MRI), Japan Meteorological Agency (JMA), Tsukuba, 8 Ibaraki, Japan} 9 [2] {RIKEN Advanced Institute for Computational Science (AICS), Kobe, Hyogo, Japan} 10 [3] {Kagawa University, Takamatsu, Kagawa, Japan} 11 [4] {Ibaraki University, Mito, Ibaraki, Japan} 12 [5] { Japan Agency for Marine-Earth Science and Technology (JAMSTEC), Yokosuka, 13 Kanagawa, Japan} 14 [6] {Hokkaido University, Sapporo, Hokkaido, Japan} Correspondence to: M. Kajino (kajino@mri-jma.go.jp) 15 16 17 Abstract
- The long-term effect of <sup>137</sup>Cs re-suspension from contaminated soil and forests due to the 18 19 Fukushima nuclear accident has been quantitatively assessed by numerical simulation, a field 20 experiment on dust deflation flux in the contaminated area (Namie, Fukushima), and air concentration measurements inside (Namie) apputside (Tsukuba, Ibaraki) the contaminated 21 area. The assessment period is for the year 2013 just after the start of the field experiments, 22 December 14, 2012. The <sup>137</sup>Cs concentrations at Namie and Tsukuba were approximately 10<sup>-1</sup> 23 -1 and  $10^{-2} - 10^{-1}$  mBq/m<sup>3</sup>, respectively. The observed monthly median concentration at 24 Namie was one to two orders of magnitude larger than that at Tsukuba. This observed 25 difference between the two sites was consistent with the simulated difference, indicating 26 successful modeling of <sup>137</sup>Cs re-suspension and atmospheric transport. The estimated re-27 suspension rate was approximately  $10^{-6}$  /d, which was significantly lower than the decreasing 28

rate of the ambient gamma lose rate in Fukushima prefecture  $(10^{-4} - 10^{-3} / d)$  as a result of 1 radioactive decay, land surface processes, and decontamination. Consequently, re-suspension 2 contributed negligib po reducing ground radioactivity. The dust deflation model could 3 account for the air concentration of <sup>137</sup>Cs in winter, whereas the summer air concentration was 4 5 underestimated by one to two orders of magnitude. Re-suspension from forestat a constant rate of  $10^{-7}$  /h, multiplied by the green area fraction, quantitatively accounted for the air 6 concentration of <sup>137</sup>Cs at Namie and its seasonal variation. The simulated contribution of dust 7 8 re-suspension to the air concentration was 0.6 - 0.8 in the cold season and 0.1 - 0.4 in the warm season at both sites; the remainder of the contribution was re-suspension from forest. 9 The re-suspension mechanisms, especially through the forest ecosystems, remain unknown, 10 11 and thus the current study is the first but crude estimation of the long-term assessment of 12 radiocesium re-suspension. Further study will be needed to understand the re-suspension 13 mechanisms and to accurately assess the re-suspension mechanisms through field experiments 14 and numerical simulations.

Keywords Atmospheric radioactivity, Re-suspension, Dust deflation, Unknown re suspension source, Aerosol, Numerical simulation, Budget analysis

17

# 18 **1** Introduction

19 The Fukushima Daiichi Nuclear Power Plant (FDNPP) accidentally released nuclear fission products into the atmosphere and our environment following the catastrophic earthquake 20 and tsunam (سنا March 2011. The accident caused serious contamination of the ground soil 21 over the Tohoku region (northeastern part of Japan, including Fukushima and Miyagi 22 23 prefectures) and the Kanto region (eastern part of Japan, including Ibaraki, Tochigi, Gunma, and Chiba prefectures) (NR 2012). Since then, a number of studies have been conducted, 24 25 particularly in the crisis phase of the disaster. These assessments include primary emission estimations (Chino et al., 2011; Danielache et al., 2012; Stohl et al., 2012; Terada et al., 2012; 26 Katata et al., 2012a, 2012b; Winiarek et al., 2012, 2014; Hirao et al., 2013; Saunier et al. 27 2013; Katata et al., 2015; Yumimoto et al., 2016; Danielache et al., 2016), field observatio 28 29 (Masson et al., 2011, 2013; NRA, 2012; Kaneyasu et al., 2012; Adachi et al., 2013; Tsuruta et al., 2014; Hososhima and Kaneyasu, 2015; Igarashi et al., 2015; Oura et al., 2015), and 30 numerical simulation; deterministic simulation: Chino et al., 2011; Morino et al., 2011; 31 32 Yasunari et al., 2011; Stohl et al., 2012; Terada et al., 2012; Katata et al., 2012a, 2012b;

Winiarek et al., 2012, 2014; Hirao et al., 2013; Saunier et al., 2013; Katata et al., 2015; 1 2 Yumimoto et al., 2016; Danielache et al., 2016, deterministic simulation with sensitivity runs: Morino et al., 2013; Adachi et al., 2013; Groëll et al., 2014; Saito et al., 2015; Sekiyama et al., 3 4 2015; Quérel et al., 2016, uncertainty modeling and probabilistic forecast: Girard et al., 2016; 5 Sekivama et al., 2016, and multi-model inter-comparison and multi-model ensemble analysis: 6 SCJ, 2014; Draxler et al., 2015) The targeted radionuclides were species with both short and long half-lives: <sup>99</sup>Mo-<sup>99m</sup>Tc (half-life 65.9 – 6 h), <sup>129m</sup>Te (33.6 d), <sup>131</sup>I (8.02 d), <sup>132</sup>Te-<sup>132</sup>I (3.2 d 7 -2.3 h),  ${}^{134}$ Cs (2.07 y),  ${}^{136}$ Cs (13.2 d),  ${}^{137}$ Cs (30.1 y),  ${}^{133}$ Xe (5.2 d), and  ${}^{35}$ S (87.5 d). 8

9 In contrast, there have been few studies on the long-term (more than one year) quantitative assessment of radioactivity in the atmosphere associated with the Fukushima accident 10 (Igarashi et al., 2015; Ishizuka et al., 2016; Kinase et al., 2016). More than 100,000 people 11 were evacuated (METI, 2012), most have still not been able to return to their homes and 12 the public remains anxious about the safety of the affected areas. Radionuclides with long 13 half-lives such as <sup>134</sup>Cs (2.07 y) and <sup>137</sup>Cs (30.1 y) are of particular concern. Following the 14 Chernobyl accident there were several studies on the re-suspension and long-term assessment 15 of these radionuclides, such as Holländer and Garger (1996), Garger et al. (1998), Hatano and 16 Hatano (2003) and Garger et al. (2012). For example, Garger et al. (2012) estimated the re-17 suspension "descending trend" as having a half-life of 300 d based on the surface activity 18 concentration of <sup>137</sup>Cs. In the case of the Fukushima accident, Igarashi et al. (2015) estimated 19 the half-reduction time by fitting multi-component exponential functions based on the <sup>137</sup>Cs 20 concentration at the Meteorological Research Institute. Tsukuba, as 5.9 d, 16 d, and 1.1 y. 21 These estimates were based on the trend in the observed surface air concentrations of <sup>137</sup>Cs, 22 23 and thus the contributions from advection, diffusion, emission and deposition terms were quantified. 24

There are thousands of monitoring posts situated in the contaminated area in Fukushima 25 prefecture to measure the ambient gamma dose rate, with the data cannot be used for 26 27 evaluating internal exposure: evaluation of internal exposure requires direct measurement of 28 the surface air activity concentration. There are only a few observation sites that continuously measure the concentration of radiocesium (e.g., Igarashi et al., 2015; Ishizuka et al., 2016; 29 Kinase et al., 2016). To assess the spatial distribution of the impal exposure hazard, 3D 30 numerical simulation is necessary to interpolate values at locations where there are no 31 32 measurements. The numerical simulation requires emission flux as a boundary condition.

1 However, the mechanism and thus the radioactivity flux associated with the re-suspension of  $2^{137}$ Cs were unknown, despite extensive efforts based on field observations (e.g., Igarashi et al.,

3 2015; Ishizuka et al., 2016; Kinase et al., 2016).

4 Garger et al. (2012) summarized the re-suspension sources following the Chernobyl accident 5 as (1) dust deflation, (2) human activity in fields, and on roads and construction sites, (3) 6 forest fires, and (4) emissions from opening of the Chernobyl sarcophagus. Re-suspension 7 sources (1) and (4) were considered in the present study. With respect to source (2), since Namie, Fukushima, is located in the evacuation zone, human activity has been extremely 8 limited except for decontamination-related work. As to source (3), there is a low chance of 9 forest fires in Japan given the high humidity before open biomass burning is estimated by the second 10 Kinase et al. (2016) found no increase in <sup>137</sup>Cs concentration when the concentration of 11 levoglucosan, a marker of biomass burning, was increased, and thus re-suspension due to 12 13 biomass burning was not considered in the present study. In addition to the four sources of <sup>137</sup>Cs from the Chernobyl accident, re-suspension from terrestrial biota was considered as 14 suggested by Kinase et al. (2016). They found substantial amounts of bioaerosols upon 15 16 scanning electron microscopy examination of samples collected in the summer, when <sup>137</sup>Cs 17 concentration was high.

18 In the present study, the long-term effect of radiocesium re-suspension from contaminated 19 soil and terrestrial biota was quantitatively assessed using 3D numerical simulation, a field experiment on dust deflation flux in a contaminated area (Namie, Fukushima), and air 20 concentration measurements tak inside (Namie) and outside (Sukuba, Ibaraki) the 21 contaminated area. The current study is the firs t crude estimation of the spatial budget of 22 23 radiocesium via re-suspension since the re-suspension mechanisms, especially through forest 24 ecosystems, remain unknown. By utilizing observational data collected both inside and outside of the contaminated area, together with 3D numerical simulation, we aimed to provide 25 as robust a budget analysis as possible the re-suspension, transport, and re-deposition of 26 <sup>137</sup>Cs over the Tohoku and Kanto regions of Japan. 27

28

# 29 2 Numerical simulation

30 A brief description of the numerical method, such as processes considered in the model and 31 simulation settings, are presented in this section, and detailed model formulations are described in Appendix A. Because the schemes and assumptions regarding the emissions are
 keys the current study, they are described in detail in the following subsections.

### 3 2.1 Lagrangian Model and simulation settings

4 Figure 1 shows the domain of the Lagrangian Model (LM) with model terrestrial elevations, 5 covering 138 - 143 °E and 34 - 39 °N. The model domain covers the southern part of the Tohoku region (the northern mountainous part of the domain, including Yamagata, Miyagi, 6 and Fukushima prefectures), and includes the FDNPP and highly polluted areas such as the 7 8 Habitation-Restricted Zone (HRZ) (20 - 50 mSv/y) and Difficult-to-Return Zone (DRZ) (> 50 9 mSv/y) (METI, 2012). It also covers the Kanto region (or Kanto Plain, the largest plain in Japan, approximately 120 km  $\times$  120 km), a highly populated region that includes moderately 10 polluted areas such as Tokyo, Gunma, Tochigi, Ibaraki, Saitama, Chiba, and Kanagawa 11 12 prefectures.

13 LM considers horizontal and vertical diffusion and advection, gravitational settling, dry and 14 wet depositions, and radioactive decay. It uses simple parameterizations for dry and wet 15 deposition schemes, and it can be driven by meteorological analysis data sets so that it does 16 not require a meteorological model to predict detailed meteorological fields and variables. 17 The model was designed to be easily handled and computationally efficient so that non-18 specialists of numerical simulations can conduct long-term assessments of atmospheric 19 diffusion problems using their desktop or laptop computers. The LM was designed for rough budget estimates, as presented in the current study, or for sensitivity analyses using a number 20 21 of parameters (e.g., Groëll et al., 2014; Girard et al., 2016; Quérel et al., 2016), rather than for process-oriented analysis (e.g., Morino et al., 2013; Katata et al., 2015) or sensitivity analyses 22 23 of the physical and chemical parameters of aerosols (Adachi et al., 2013). Details of each process and parameter are described in Appendix A. Statistical error of a Lagrangian 24 25 simulation is inversely proportional to the square of the number of Lagrangian particles (LPs). 26 The statistical accuracy of the current simulation setting is discussed in Appendix B.

The Grid Point Value Meso-Scale Model (GPV-MSM) of the Japan Meteorological Agency (JMA) was used for meteorological analysis to calculate the transport of LPs. It covers 120 -150 °E and 23 - 47 °N and provides 3 hourly and 16 pressure levels of 3D meteorological variables, from 1000 hPa to 100 hPa, with a horizontal grid resolution of approximately 11 km ( $\Delta$ longitude = 0.125° and  $\Delta$ latitude = 0.1°) and surface variables at twice the resolution as that for the 3D variables ( $\Delta$ longitude = 0.0625° and  $\Delta$ latitude = 0.05°). In the simulation, the whole model domain where LPs can travel is 138 – 143 °E, 34 – 39 °N and from ground surface to 500 hPa. For output of the model results, LP fields are converted to Eulerian concentration (Bq/m<sup>3</sup>) and deposition (Bq/m<sup>2</sup>) fields in the same horizontal space as the 3D variables but are vertically allocated from the ground surface to an altitude of 1 km at 100 m intervals. The observed surface air concentration was compared with the simulated mean concentration at 0 – 100 m above ground level (AGL).

## 8 2.2 Re-suspension from bare soil

9 Ishizuka et al. (2016) developed a re-suspension scheme for radiocesium from bare soil based
10 on measurements on the ground an Namie High School, Tsushima Campus (denoted as Namie
11 (Tsushima) in Table 1 and Fig. 1) in the DRZ.

12 
$$F_{soil} = p_{20\,\mu m} F_M (1 - f_{forest}) B_{5\,mm}(t),$$
 (1)

where  $F_{soil}$  is the <sup>137</sup>Cs dust re-suspension flux from soil (Bq/m<sup>2</sup>/s),  $p_{20\mu m}$  is the mass fraction 13 of dust smaller than 20 µm in diameter against soil containing a maximum size of 2 mm 14 particles, and varies depending on soil texture  $(2 \times 10^{-9} \text{ for sand}, 0.03 \text{ for loamy sand}, 0.09 \text{ for}$ 15 sandy loam, and 0.32 for silt loam),  $F_{\rm M}$  is the total dust mass flux (kg/m<sup>2</sup>/s),  $f_{\rm forest}$  is the forest 16 area fraction, and  $B_{5mm}(t)$  is the specific radioactivity of surface soil (from the surface to a 17 depth of 5 mm) (Bq/kg) as a function of time since March 2011. The formula is based on the 18 19 assumption that dust particles smaller than 20 µm in diameter originated from the surface soil 20 and to a depth of 5 mm were suspended and transported through the atmosphere.  $F_{\rm M}$  is 21 formulated as being proportional to the cube of the friction velocity  $u_*$  (m/s) as described by 22 Loosmore and Hunt (2002) and was applied to the dust emission:

23 
$$F_M = 3.6 \times 10^{-9} u_*^3$$
. (2)

Since  $u_*$  is not available in GPV-MSM,  $u_*$  was estimated using a wind speed at 10 m AGL by assuming neutral stratification conditions.

26  $B_{5mm}(t)$  was derived from the combination of  $B_{obs}$ , the observed horizontal distribution of 27  $^{137}$ Cs deposition obtained from an airborne radiological survey (NRA, 2012) (Bq/m<sup>2</sup>) and  $r_{5mm}$ , 28 the surface soil activity ratio of 0 – 5 mm to 0 – 5 cm obtained from a vertical profile 29 measurement of  $^{137}$ Cs in the ground soil at Namie High School (= 0.57 Bq/Bq) as

1 
$$B_{5\,\mathrm{mm}}(t) = \frac{B_{obs}r_{5mm}R_{decay}(t)}{5 \times 10^{-3}\rho_{soil}},$$
 (3)

where  $\rho_{\text{soil}}$  is the density of soil particles per unit volume in the ground space (kg/m<sup>3</sup>) obtained 2 from the porosity (0.4 m<sup>3</sup>/m<sup>3</sup>) and the density of dust particles (2650 kg/m<sup>3</sup>). For  $R_{decay}$ , which 3 is the decreasing rate of activity in the ground, only radioactive decay was considered for the 4 re-suspension calculation. The decreasing rate due to other processes such as land surface 5 processes (or migration in the soil and biota) and decontamination were not considered here. 6 A suppression of dust deflation due to soil moisture and snow cover was not considered. 7 8 Therefore, it should be noted here that  $F_{soil}$  in Eq. (1) is considered as a maximum estimate of <sup>137</sup>Cs re-suspension flux from surface soil. Effects such as land surface processes, 9 10 decontamination, and dust deflation suppression due to snow cover are extensively discussed 11 in Appendix C using ambient gamma dose rate measurements obtained by the monitoring 12 posts (Table 1, Fig. 1b, and Fig. C1).

13 Eq. (1) is a function of soil texture. The areal fraction of soil texture of the model grid was 14 obtained from the database of the advanced research Weather Research and Forecasting 15 model version 3 (WRFV3; Skamarock et al., 2008). Sixteen categories of soil texture (Miller 16 and White, 1998) with a 30 s resolution dataset can be obtained from the web after subscription at (http://www2.mmm.ucar.edu/wrf/users/download/get\_sources\_wps\_geog.html-17 last access: 12 February 2016) and were re-categorized into the above-mentioned four 18 19 categories and interpolated to the LM resolution (~11 km) as shown in Fig. 2a-c. Note that the 20 loamy sand fraction is not presented because it is zero for the entire domain. The parameter forest (Fig. 2d) was also obtained from the database of WRFV3 and was calculated based on 21 the 24 United States Geological Survey (USGS) Land Use Categories, which are constant 22 23 over time. The Land Use Category dataset can also be obtained from the above website.

Ishizuka et al. (2016) validated their dust deflation module by using a 1D model and observed 24 the surface air concentration of <sup>137</sup>Cs at Namie in the winter. After applying the module to our 25 3D simulation, we found that the air concentration at Namie was underestimated by about one 26 27 order of magnitude for the same period. The module was formulated based on physical 28 parameters but contains parameters obtained at a single location and under a fixed 29 atmospheric condition, whereas a parameter describing the complexity and differences among 30 locations ideally would have been applied to Eqs (1) - (3). We set the parameter to 10 after adjusting the simulation results against the observed concentration of <sup>137</sup>Cs at Namie in the 31

winter. This is one of the simplest top-down approaches for adjusting the emission flux
 according to the air concentration. The module requires improvement in the future as more
 reliable parameters become available for various conditions and locations.

#### 4 2.3 Re-suspension from the forest ecosystems

5 The re-suspension mechanism of radiocesium from land ecosystems remains unknown. 6 Kinase et al. (2016) found substantial amounts of bioaerosols (rather than mineral dust particles) in samples collected for scanning electron microscopy in the summer, when the 7 <sup>137</sup>Cs concentration was high. This does not prove that the bioaerosol was carrying 8 radiocesium, but that it could be a potential carrier. The behavior of Cs in the environment 9 can be inferred by analogy with K, a congener of Cs. Potassium is a necessary and abundant 10 element in plants and circulates between land ecosystems. The addition of potassium fertilizer 11 12 to a rice field in Fukushima significantly reduced the Cs content of the rice (Ohmori et al., 2014). Substantial amounts of K-salt-rich particles, possibly emitted by active biota such as 13 14 plants and fungi, and coated with secondary organic aerosols, were observed in pristine Amazonian rainforest (Pöhlker et al., 2012). The major areal fraction of the contaminated area 15 16 in Fukushima is covered by biota-rich mountain forests (Figs. 1a, 2d, and 4a). Despite the differences in plant species and locations, it is plausible that water-soluble radiocesium 17 18 circulating in the biota and soil in the forests was somehow re-emitted into the atmosphere 19 and contributed to the surface air concentration. The re-suspension from the forest ecosystem 20 was simply formulated as follows:

21 
$$F_{\text{forest}} = f_{\text{forest}} f_{\text{green}} r_{\text{const}} B_{\text{obs}} R_{\text{decay}}(t), \qquad (4)$$

where  $F_{forest}$  is the <sup>137</sup>Cs re-suspension flux from forest (Bq/m<sup>2</sup>/s),  $f_{green}$  and  $r_{const}$  are the 22 monthly green area fraction and the constant re-suspension coefficient (/s), respectively, and 23  $r_{const}$  is a tunable parameter to adjust the simulated air concentration of <sup>137</sup>Cs to that observed. 24 In the current study,  $r_{const}$  is set to  $10^{-7}$  /h by adjusting the simulation data using the observed 25 <sup>137</sup>Cs concentration at Namie in the summer, when the re-suspension from soil was negligible 26 27 due to the higher soil moisture content (following considerable rain) and lower wind speed. As with re-suspension from soil, only radioactive decay was considered for  $R_{decay}$  and the 28 other processes were not considered. Unlike re-suspension from son, precipitation might not 29 suppress re-suspension from the forest ecosystems since substantial amounts of K-containing 30 particles were observed in the wet season in the Amazon (Pöhlker et al., 2012 and references 31

1 therein). The parameter  $f_{green}$  was obtained from the database of WRFV3 and was originally 2 derived from satellite Advanced Very High Resolution Radiometer (AVHRR)/Normalized 3 Difference Vegetation Index (NDVI) data (Gutman and Ignatov, 1998). Whereas  $f_{forest}$ 4 remains constant, the monthly averaged  $f_{green}$  was used in order to reflect seasonal changes in 5 the activity of the biota.

# 6 2.4 Emission from FDNPP (primary emission, additional emissions from the 7 reactor buildings, and unexpected re-suspension associated with debris 8 removal operations)

9 Katata et al. (2015), the Japan Atomic Energy Agency (JAEA)'s latest estimate of the primary 10 emission from FDNPP, was applied for the emergency situation of March 2011, to evaluate the performance of the LM model against the horizontal distribution of <sup>137</sup>Cs deposition of the 11 airborne radiological survey (NRA, 2012) (Fig. 4a) and surface air concentrations measured at 12 Tsukuba (Fig. 1a). We selected this inventory because it is JAEA's most up-to-date version. 13 14 Based on an integrated understanding of environmental radioactivity, atmospheric dispersion, and the nuclear reactors, the JAEA team has carefully established a series of inventories for 15 about five years, staring with Chino et al. (2011), followed by Katata et al. (2012a), (2012b), 16 Terada et al. (2012), and finally the current inventory (Katata et al., 2015), which is 17 substantially improved compared to its pressors. 18

Ongoing emissions during the study analysis period after the emergency situation, that is, January to December 2013, was obtained from the Tokyo Electric Power Co., Inc. (TEPCO) monthly mean emission flux from the reactor buildings (TEPCO, 2012; 2013; 2014a; 2014b; 2015). Because only the sum of <sup>134</sup>Cs and <sup>137</sup>Cs was provided, the fractions of these two isotopes were calculated based on their half-lives and the assumption that their activities were equal in March 2011 (e.g., Katata et al., 2015), as shown in Fig. 3. The values range from  $10^5$ to  $10^7$  Bq/h,  $\Omega$  for simplicity we set a constant value of  $10^6$  Bq/h in the current simulation.

In August 2013, unexpected re-suspension associated with debris removal operations was reported by TEPCO (2014c) and NRA (2014) and the gross amount was  $10^{10} - 10^{11}$  Bq of <sup>137</sup>Cs (TEPCO, 2014c, NRA, 2014, Steinhauser et al., 2015). The impact of this unexpected re-suspension is briefly discussed in Sect. 5.2 along with an additional finding, but this emission was not considered in the present LM simulation. In this study we focused on the ongoing emission, mostly from the natural environment, that is difficult to control.

# 2 3 Field observation

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3 Details of the surface air activity concentration measurements can be found in Ishizuka et al. 4 (2016) and Kinase et al. (2016) for Namie and Igarashi et al. (2015) for Tsukuba. At both sites, ambient aerosols were collected using a high-volume air sampler and <sup>134</sup>Cs and <sup>137</sup>Cs 5 concentrations were obtained by  $\gamma$ -ray spect, popy using a Ge semiconductor detector. The 6 sampling intervals were 1 - 2 d (sometimes several days) at Namie and 1 w at Tsukuba for the 7 8 analysis period of this study, the year 2013. The observations at Namie started on December 9 14, 2012, while those at Tsukuba started on March 31, 2003, before the FDNPP accident. In March 2011, the sampling interval was increased at Tsukuba to 6 h - 1 d and the data for 10 these periods were used for the validation of LM and its parameters, as presented in Sect. 4.1. 11

The Namie site is located on the ground at Namie High School, Tsushima Campus, in the Tsushima district of Namie town in Fukushima prefecture, as shown in Table 1 and Fig. 1. Namie town extends from the Hamadori coastal area (denoted as C in Fig. 1) to the Abukuma highland area (B in Fig. 1). In order to distinguish the Tsushima site from the monitoring post

16 located in Omaru district in Namie town in the coastal area, the Tsushima site in the mountain

area is sometimes referred to as Namie (Tsushima). Note that, unless specifically referred to
as Namie (Omaru), Namie without brackets indicates Namie (Tsushima) throughout this
manuscript. The Tsukuba site is located on the premises of the Meteorological Research
Institute (Table 1 and Fig. 1a).

Namie (Tsushima) was located in the DRZ (> 50 mSv/y, ~5.71  $\mu$ Sv/h) and the observed <sup>137</sup>Cs deposition amount was 2,300 kBq/m<sup>2</sup> (Fig. 4a). The ambient gamma dose rate was 11.2  $\mu$ Sv/h on April 1, 2012 at the site and had dropped to 4.8  $\mu$ Sv/h on Feb 16, 2016, and at the HRZ level (20 - 50 mSv/y, 2.28 - 5.71  $\mu$ Sv/h). Tsukuba is located approximately 170 km southwest of FDNPP. The observed <sup>137</sup>Cs deposition amount was 21 kBq/m<sup>2</sup> (Fig. 4a), two orders of magnitude lower than at Namie and the dose rate has be below 0.1  $\mu$ Sv/h since 2012.

28

# 29 **4 Results**

30 Section 4.1 presents a validation of the LM model and the optimization of the model 31 deposition parameters by using airborne observations (NRA, 2012) and the emission  $\supset$ 

inventory of Katata et al. (2015) for the emergency situation of March 2011. Using the
 optimized ranges of model parameters validated in Sect. 4.1, the simulated re-suspension of
 <sup>137</sup>Cs from soil and forest, and emission from FDNPP, is presented in Sect. 4.2, and the
 budgets for re-suspension, transport, and re-deposition are presented in Sect. 4.3.

# 5 4.1 Model and parameter validation for the emergency situation (March 2011)

Figures 4 and 5 show the observed and simulated distribution of <sup>137</sup>Cs deposition in March 2011, and the scattergram comparing the observational and simulation results, respectively. In the simulation shown in the figures, the "reference" sets used for dry and wet deposition parameters, namely, the collection efficiency of aerosols using hydrometeors  $E_c$  (Eq. A2) and the dry deposition velocity over land  $v_d$  (Eq. A4), were 0.04 and 0.1 cm/s, respectively.

Since LM employs simple parameterizations for dry and wet deposition, as described in Appendix A, sensitivity tests were conducted for model validation, together with optimization of the deposition parameters. Table 2 summarizes the ranges of the deposition parameters for the sensitivity tests and the results of the <sup>137</sup>Cs budget and statistical measurements between the observations and the simulation.

The parameter  $E_c$  was 0.05 for the JMA dispersion model (JMA, 1998) but the targeted 16 species are different. For example, volcanic ash (particles per than 1 µm in diameter) used 17 for the JMA model were generally larger than the carrier aerosols of  $^{137}$ Cs (around 1  $\mu$ m in 18 19 diameter observed in the downwind area, Tsukuba, Kaneyasu et al. 2012, Adachi et al., 2013). Since the inertia of these smaller <sup>137</sup>Cs particles is likely to be smaller than that for volcanic 20 ash,  $E_c$  could be smaller. The range of  $E_c$  was set as 0.02 - 0.06. The dry deposition velocity 21  $v_d$  was selected as 0.1 cm/s for <sup>137</sup>Cs in Furuno et al. (1999). The range for  $v_d$  was set as 0.05 22 to 0.15 in the present study. 23

The emission inventory of Katata et al. (2015) amounted 14.1 PBq from March 12 to April 1, 24 2013. The simulated deposition over the model domain (138 - 143 °E, 34 - 39 °N) ranged 25 from 3.4 - 4.7 PBq, which is approximately a guarter to one third of the emission from the 26 FDNPP. Sixty percent of the total deposition occurred over land, for a total of 2.0 - 2.8 PBq, 27 28 which is close to the observed value of 2.68 PBq, and the observed value is within the range 29 of the sensitivity runs. Statistical measures such as the fractional bias FB, the correlation 30 coefficient R, and FAx (fraction of the simulated values within a factor of x) are listed in 31 Table 2. To find better combinations of (or to optimize) the dry and wet deposition parameters,

sensitivity runs were screened based on the criteria FA10 > 0.9, FA5 > 0.7, R > 0.75, and an 1 absolute value of FB < 10%. After the screening, only one combination  $(E_c, v_d) = (0.04, 0.1)$ 2 cm/s) was left, and thus this is referred to as the "reference" parameters. To evaluate the 3 4 sensitivity (or uncertainty) of the re-suspension simulation for 2013 due to the deposition 5 parameters, the range of the combination of parameters was set as  $(E_c, v_d) = (0.03 - 0.05, 0.05)$ -0.1 cm/s) around the reference parameters (referred to as the "optimized range") by 6 7 excluding the parameters with the worse performances. The ranges of the statistical measures 8 of the optimized runs are listed in Table 2. FB, R, FA2, FA5, and FA10 after the optimization had the ranges -0.18 - -0.036, 0.74 - 0.77, 0.25 - 0.30, 0.63 - 0.77, and 0.85 - 0.94, 9 10 respectively. These statistical measures were comparable to those reported in previous multi-11 model comparison studies (R: 0.27 - 0.85, FB: -0.84 - 0.56, and FA2: 0.14 - 0.57, in SCJ, 2014 and Draxler et al., 2015). The current model is thus shown to be sufficiently credible for 12 13 the budget analysis in this study, despite the simple parameterization and the low resolution in 14 space ( $\sim$ 11 km) and time (3 h).

Consistent with many previous studies, the simulated contribution of wet deposition was 15 larger than that of dry deposition: the ratio of the amount of dry to wet deposition ranged from 16 17 0.12 - 0.23 for the optimized parameter ranges, indicating that the results were less sensitive to the dry deposition parameter. Generally speaking, R became higher as  $E_c$  became lower, 18 19 whereas FAx became higher as  $E_c$  became higher for the various ranges of the sensitivity tests. Therefore, lower  $E_c$  did not meet the criteria of FAx and higher  $E_c$  did not meet the criteria of 20 R. Consequently, after the optimization, the maximum values of the statistical measures were 21 22 lower but the minimum values became higher, indicating that the optimization was successful 23 in excluding the parameters with the worse performances (rather than selecting the best parameters). It should be noted here that the optimized deposition arameters are not 24 necessarily physically but rather are consistent with the available evidence. The results 25 26 presented in this section indicate that the current LM simulation with these optimized 27 parameters has the potential to reproduce consistent features of the radiocesium budget over 28 the Tohoku and Kanto regions of Japan.

- Figure 6 shows the temporal variation of simulated (red) and observed (black) <sup>137</sup>Cs concentrations at Tsukuba in March 2011. The model successfully reproduced the three major plumes arriving at Tsukuba on March 15-16, 20-21, and 28-30; a plume on March 23 only
- $\frac{32}{32}$  appeared in the simulation. The red shaded areas indicate the range of  $^{137}$ Cs concentrations

obtained when the simulations were run using the optimized parameter ranges  $E_c = 0.03 - 0.05$  and  $v_d = 0.05 - 0.1$  cm/s. Due to differences in the parameters, the surface air concentration could vary by approximately one order of magnitude in transported plumes that experienced wet scavenging along their pathway.

# 5 4.2 Re-suspension in 2013

6 Figure 7 shows the observed and daily mean simulated (with the reference parameters) surface air concentrations at Namie and Tsukuba for the year 2013. The red, green, and blue 7 8 lines indicate re-suspension from soil, re-suspension from forest, and emission from the 9 FDNPP reactor buildings, respectively. Note that the re-suspension flux due to the dust deflation module (Ishizuka et al., 2016) is multiplied by 10 in this study by adium g to the 10 observation level at Namie in the cold season (January to March, October to December). Also 11 note that the re-suspension coefficient  $r_{const}$  in Eq. (4) was set as  $10^{-7}$ /h by adjusting to the 12 observation level at Namie in the warm season (May to September). The emission flux 13 reported by TEPCO varied from  $10^5$  to  $10^7$  Bg/h during the study period but we set it to  $10^6$ 14 Bq/h for simplicity. We did not try to precisely adjust  $r_{const}$  to the observation by, for example, 15 using inverse modeling, and instead we simply multiplied by wer-of-ten values which are 16 constant in time and space be real (1) this aided straightforward interpretation of the 17 18 simulation results by keeping the simulated variation as it was, and therefore (2) this provided 19 simple but useful hints for understanding the re-suspension mechanisms, which remain 20 unknown.

Using the dust deflation module (which has a physical basis), <sup>137</sup>Cs flux re-suspended from 21 soil could account for the level of the observed surface air concentration of <sup>137</sup>Cs at Namie in 22 23 the cold season. Under the influence of the northwesterly winter monsoon, the surface wind speed is high over the contaminated area compared to Che summer (see Fig. 10). Note that 24 the flux might be a maximum estimate since it does not consider land surface processes (such 25 as soil moisture, snow cover, or migration of <sup>137</sup>Cs in the soil) and decontamination, which 26 could reduce the <sup>137</sup>Cs re-suspension flux. In contrast, in the warm season, the estimated flux 27 28 significantly underestimated the observation by one to two orders of magnitude due to the 29 weak surface wind, indicating that the dust deflation process may not be the sole process involved in sustaining the air concentration of <sup>137</sup>Cs during this period. Introducing the <sup>137</sup>Cs 30 re-suspension component from forest with a resuspension coefficient of  $10^{-7}$ /h and a monthly 31 32 variation in the green area fraction (derived from NDVI) could quantitatively account for the

observed air concentration together with its seasonal variation at Namie. Even though both the 1 2 simulated re-suspension from soil and forests reproduced the quantity and seasonal variation of the background concentration (in other words, concentrations originating from constantly 3 presenting emissions) at Namie, sporadic peak events, such as the daily mean <sup>137</sup>Cs 4 concentration exceeding 10 mBq/m<sup>3</sup> as observed in June and August at Namie, were not 5 simulated. Some specific re-suspension events might occur within the premises of FDNPP or 6 very close to this area on these days, as indicated later in Sect. 5.2. The simulated <sup>137</sup>Cs 7 concentrations due to the monthly mean emission from the reactor buildings (= $10^6$  Bq/h) 8 9 significantly underestimated the observed concentration by more than three orders of 10 magnitude at Namie and by two orders of magnitude at Tsukuba. Even the maximum estimate of 10<sup>7</sup> Bq/h does not reach the observed level. The emission from FDNPP may not 11 have been the sole process sustaining the air concentration of <sup>137</sup>Cs in 2013, supporting the 12 13 discussion in Igarashi et al. (2015), which concluded that direct emission from the FDNPP 14 played a minor role in the observed atmospheric radiocesium concentrations over Tsukuba 15 during 2013-2014.

The observed air concentration of <sup>137</sup>Cs at Tsukuba was about one to two orders of magnit lower than that at Namie. The simulated difference between the two sites in and out of the contaminated areas was consistent with the observed difference. This finding indicates that the current LM simulation provided consistent features of re-suspension, transport, and redeposition in the Tohoku and Kanto regions of Japan of <sup>137</sup>Cs originating from Fukushima.

# 21 4.3 Budget analysis

22 Figure 8 illustrates the simulated (with the reference parameters) annual total re-suspension and re-deposition amounts of <sup>137</sup>Cs, together with their ratios to the observed deposition 23 24 amount (Fig. 4a). The simulated areal total re-suspension amount was 1.01 TBq, which was equivalent to 0.037% of the total deposition amount, 2.68 PBq. The areal total re-deposition 25 amount (with the reference parameters) was 0.22 TBq (0.18 - 0.23 TBq for the optimized)26 range of the deposition parameters), corresponding to approximately 21.7 (17.8 - 22.8) % of 27 the re-suspended amount deposited mainly in the Tohoku region, with the remainder being 28 29 transported out of the region. Therefore, the regional mean rate in the decrease of the land surface <sup>137</sup>Cs concentration due to re-suspension was estimated to be 0.029 (0.029 -30

1 0.031) %/y<sup>1</sup>, equivalent to 7.9  $(7.9 - 8.2) \times 10^{-7}$  /d. The spatial distribution of the resuspension and re-deposition ratio to the primary deposition ranged from 0.01 – 0.3% and 0.001 – 0.03%, respectively. The spatial distribution of the land surface <sup>137</sup>Cs deposition decay due to re-suspension ranged from  $2.2 \times 10^{-7} - 6.6 \times 10^{-6}$  /d. Re-suspension therefore had a negligible effect on reducing land surface radioactive contamination.

6

# 7 5 Discussion

8 Seasonal variation of the surface activity concentration and its source contributions are 9 extensively discussed in Sect. 5.1. A possible source of the observed sporadic peak events, 10 which could not be reproduced by the simulation, is discussed in Sect. 5.2. Future issues are 11 summarized in Sect. 5.3. The effects of other processes that were not considered in the model, 12 such as land surface processes and decontamination, are discussed based on the dose rate 13 measurements from the monitoring posts in Fukushima in Appendix C.

# 14 **5.1** Seasonal variation and source contribution

The discussion in this section expands on that in Sect. 4.2. Figure 9 shows the same temporal variation as Fig. 7 but for simulated (using the optimized ranges of parameters) results for <sup>137</sup>Cs from dust and FDNPP in winter (January to March) and from forest and FDNPP in summer (June to August).

19 In the winter, the simulated trend for dust agreed well with the observed trend (Fig. 9a), and the surface air concentration during this period was positively correlated with the surface 20 21 wind speed in both the simulation and the observations. There was a sporadic peak in the observational data of 6.7 mBq/m<sup>3</sup> from the March 17 at 13:00 local time (LT) to March 18 at 22 23 13:00 LT that could not be reproduced by the dust module, and this peak coincided with a plume arriving from FDNPP, as shown in Fig. 9c. The discrepancy between the observed 24 peak and the dust simulation is likely due to underestimation of the simulation because t 25 simulated dust peak reached an intensity of  $4 - 5 \text{ mBg/m}^3$  in the winter, which is of the same 26

27 order of magnitude as that of the observed peak. The observed peak could also be accounted

<sup>&</sup>lt;sup>1</sup> The amount re-suspended, excluding re-deposition (1.01 TBq minus 0.22 (0.18 - 0.23) TBq) for the year 2013, divided by the total deposition amount of 2.68 PBq.

for by specific re-suspension events on the order of  $10^9$  Bq/h (the left axis divided by the right axis multiplied by  $10^6$  Bq/h in Fig. 9c) if they occurred on the premises of FDNPP or close to the area. There are also two events exceeding 2 mBq/m<sup>3</sup>, one in January and another in February. It is unlikely that the two peaks originated from the direct emission from FDNPP and likely that they originated from the dust deflation because the observed peaks coincided with the simulated dust peaks (Fig. 9a) and not with the simulated peaks due to the FDNPP emission (Fig. 9c).

8 In the summer, the simulated quantity as well as the variation in the forest data agreed well 9 with the observed data (Fig. 9b). Because there is only monthly variation in the simulated emission, the simulated daily trend solely originated from variations in the meteorological 10 parameters (wind field, turbulent mixing, and wet scavenging). A significant peak of 60.4 11 mBq/m<sup>3</sup> is observed from August 14 at 13:00 LT to August 15 at 13:00 LT. This observed 12 level was approximately two orders of magnitude larger than the simulated level and one to 13 14 two orders of magnitude larger than the observed level for the other days in this period. Therefore, constant emission such as re-suspension from forest is less likely to be the origin 15 of the peak. Because the observed peak and the simulated peak of <sup>137</sup>Cs from FDNPP 16 coincided (Fig. 9d), the observed level could be accounted for by specific re-suspension 17 18 events on the order of 10<sup>10</sup> Bq/h either on the premises of FDNPP or close to the area. There have been several arguments that the observed peaks in August 2013 were associated with 19 debris removal operations at FDNPP and this is discussed separately in Sect. 5.2. 20

Figure 10 illustrates the seasonal mean surface wind field and surface air <sup>137</sup>Cs concentration 21 (simulated using the reference parameters) due to (a) dust re-suspension in the winter and (b). 22 forest re-suspension in the summer. Due to the prevailing northwesterly winter monsoon 23 <sup>137</sup>Cs was carried southeastward in the winter. In the summer, under the influence of the 24 Pacific high pressure system, <sup>137</sup>Cs was carried inland. The monthly mean wind speed is high 25 26 in winter and low in summer. The upper panels of Fig. 11 illustrate the observed and 27 simulated (using the optimized ranges of parameters) total (from soil, forest, and FDNPP) <sup>137</sup>Cs concentration at Namie and Tsukuba. The time resolutions of the simulation are daily 28 29 for Namie and weekly for Tsukuba to be consistent with the sampling intervals of the two 30 respective sites. The simulation successfully reproduced the quantity and variation in the observed background concentration at Namie and Tsukuba but could not reproduce the 31 32 sporadic peak events observed at Namie, as discussed above. The simulation also significantly

1 underestimated the observations at Tsukuba from January to March, 2013. Due to the

2 northwesterly monsoon (Fig. 10a), there was less air mass transported from FDNPP to

3 Tsukuba in the winter (Fig. 12b) and therefore this underestimation is probably due

4 specifically to underestimation of the simulated re-suspension. The lower panels of Fig. 11

5 show the relative contributions of  $^{137}$ Cs from soil and forests at Namie and Tsukuba. The

6 contribution from FDNPP was negligible throughout the year. At both sites, the contribution

7 from dust was high (0.6 - 0.8) in the cold season and low (0.1 - 0.4) in the warm season.

8 Figure 12a shows the observed and simulated (with the reference parameters) monthly Namie to Tsukuba <sup>137</sup>Cs concentration ratios. The mean concentration ratio exceeded 100 in June and 9 10 200 in August due to the sporadic peak events. The monthly median would be relevant for comparing the background observation with the simulation results by considering only 11 constant emission he values of the simulated concentration ratio and its seasonal variation 12 agreed fairly well with the observed monthly median ratio: the observed and simulated annual 13 14 means were 38.9 and 30.3, respectively. Fig. 12b shows the monthly mean simulated resuspension source area contributions to the <sup>137</sup>Cs air concentration at Namie and Tsukuba. 15 16 The re-suspension source area is defined as the model grid where the observed deposition amount exceeded 300 kBg/m<sup>2</sup> (Fig. 4a) and includes the Namie grid (2.300 kBg/m<sup>2</sup>). Eighty 17 to 90% of the <sup>137</sup>Cs air concentration at Namie originated from the source region, and there 18 was no clear seasonal variation in the value. In contrast, Tsukuba is characterized as a 19 20 downwind region and there was clear seasonal variation in the source contribution ratio: high in summer and low in winter, due to the summer and winter monsoons, as discussed above. 21 Nonetheless, the highest value at Tsukuba was 0.4 in July, and so more than half of the <sup>137</sup>Cs 22 23 concentration at Tsukuba originated locally or from areas other than the contaminated regions throughout the year. As shown in Figs. 11a and 11b, the variability in the simulated 24 25 concentration at Tsukuba due to uncertainty in the deposition parameters was much larger 26 than that at Namie. The differences in the variability indicated that the Namie and Tsukuba 27 sites can be characterized as the source area and the downwind area, respectively: as the time 28 required for the plume to move from the emission site to the observation site increases, the 29 variability becomes larger due to the increased chance for the plume to experience dry and wet scavenging. 30

 $\mathcal{O}$ 

## **5.2** Possible source of sporadic peak events

There have been several scientific studies and governmental reports on the unexpected re-2 suspension from FDNPP in August 2013. The high dose rate alarm was activated on August 3 4 19 within the premises of FDNPP associated with the debris removal operation. Matsunami et al. (2016) related the radiocesium contamination of brown rice in Fukushima in 2013 to this 5 operation, whereas MAFF (2015) denied any association. The NRA estimated the <sup>137</sup>Cs 6 emission rate during the debris removal operation as  $6.7 \times 10^{10}$  Bq/h and the total amount as 7 1.1×10<sup>11</sup> Bq (NRA, 2014). TEPCO (2014c) estimated the emission rate during the operation 8 as  $5.8 \times 10^{10} - 1.2 \times 10^{11}$  Bg/h and the total amount as  $1.3 - 2.6 \times 10^{11}$  Bg. Steinhauser et al. 9 (2015) estimated the gross amount as  $2.8 \times 10^{11}$  Bq using measurements of weekly air filter 10 sampling and monthly deposition, and a numerical simulation. Their estimates are of the same 11 order of magnitude as our estimate  $(10^{10} \text{ Bg/h}, \text{ see Sect. 5.1})$  but the dates are different: our 12 observed peak was earlier than the reported removal operation. 13

Our daily sampling showed a peak concentration ( $60.4 \text{ mBg/m}^3$ ) from August 14 at 13:00 LT 14 to August 15 at 13:00 LT before the reported operation, but did not detect high concentrations 15 16 in the August 19 (0.33 mBq/h for August 18 at 13:00 LT to August 19 at 13:00 LT and 1.2 mBq/h for August 19 at 13:00 to August 20 at 13:00 LT). Figure 13 shows the forward 17 18 trajectories predicted by the LM (statistical locations of LPs) starting from FDNPP on August 19 14 (left) and August 19 (right). The sky-blue lines and red dashed circles indicate areas 20 containing approximately two-thirds of the LPs within 1 km AGL: the extent of the area 21 reflects horizontal and vertical atmospheric diffusion. The highest dose rate peaks were 22 observed from 13:50 LT to 14:10 LT on August 19 at 2.8 - 8.3 km north and north-northwest the leeward side. 23 of **FDNPP** on as reported by Fukushima prefecture 24 (https://www.pref.fukushima.lg.jp/download/1/20130827moni.pdf.pdf, last access: March 11, 2016). The forward trajectories on August 19 indicated that plumes during the debris removal 25 26 operation traveled north-northwest to north of FDNPP (Figs 13b and 13d), rather than toward 27 the west-northwest where the Namie site is located. On August 14, on the other hand, plumes 28 were transported toward the west (starting at 12 LT, Fig 13a), and then to the north (starting at 15 LT, Fig. 13c) due to fast changes in wind direction, resulting in the simulated peak 29 30 concentration shown in Fig. 9d during this period. Our simulation and observations together indicated that the same order of magnitude of  $^{137}$ Cs emission occurred on August 14 – 15 and 31

on August 19. Alarm activation was not reported on August 14 – 15 but debris removal
operation was also conducted on August 14 and 16 (MAFF, 2015).

# 3 **5.3 Future issues**

4 Issues that remain to be resolved in future research are summarized as follows:

Re-suspension from the biota could be predominant in the warm season but the re suspension sources as well as mechanisms remain essentially unknown. Further study is
 needed to understand the mechanism based on field experiments and numerical
 simulations.

9 2. The current estimation could account for the measured background concentration  $(0.1 - 1 \text{ mBq/m}^3)$  but could not reproduce the observed sporadic peak concentration  $(1 - 10 \text{ mBq/m}^3)$  at the Namie site. Further study is needed to identify the cause.

12 3. The flux module has been validated at a single location. The module could be
extended to be applicable to various land use and soil texture conditions.

- 4. The current estimation was based on a single model simulation. Variability in multi-model
  simulations is rather large (SCJ, 2014; Draxler et al., 2015) and therefore multi-model
  assessment will be indispensable for long-term re-suspension analysis.
- 17

# 18 6 Conclusions

The long-term effect of <sup>137</sup>Cs re-suspension from contaminated soil and biota due to the 19 20 Fukushima nuclear accident has been quantitatively assessed using a numerical simulation, a field experiment on dust deflation in the contaminated area (Namie, Fukushima), and air 21 concentration measurements inside (Namie) and outside (Tsukuba, Ibaraki) of torrate. The 22 re-suspension mechanism remains unknown. We therefore utilized observational data 23 obtained both inside and outside the contaminated area, together with 3D numerical 24 25 simulation, to provide a robust budget analysis of the re-suspension, transport, and redeposition of <sup>137</sup>Cs in the eastern part (the Tohoku and Kanto regions) of Japan. Our findings 26 27 are summarized as follows:

Optimization of the deposition parameters of the LM for simulating the emergency
 situation of March 2011, using aircraft observation data (NRA, 2012) and the prescribed

emission inventory (Katata et al., 2015), provided 0.1 (0.05 - 0.1) cm/s for a dry
deposition velocity over land and 0.04 (0.03 - 0.05) for a hydrometeor collection
efficiency for aerosols. The optimized (or validated) ranges of the deposition parameters
were applied to long-term re-suspension assessment for the year 2013.

5 Using the dust deflation module (Ishizuka et al., 2016), which was developed based on 2. physical parameters, <sup>137</sup>Cs re-suspension from soil could account for the observed <sup>137</sup>Cs 6 surface air concentration measured at Namie Only the cold season; the module 7 underestimated the <sup>137</sup>Cs concentration by one to two orders of magnitude in the warm 8 9 season. Introducing re-suspension from forest using a constant re-suspension coefficient of  $10^{-7}$  /h and monthly green area fraction could quantitatively account for the observed 10 concentration together with its seasonal variation. The contribution from additional 11 emission from the reactor buildings of FDNPP  $(10^6 \text{ Bg/h})$  was negligible throughout the 12 year and underestimated the observed air concentration by two to three orders of 13 14 magnitude at both observation sites.

15 3. At Namie and Tsukuba, the simulated contribution of re-suspension from soil was high 16 (0.6 - 0.8) in the cold season and low (0.1 - 0.4) in the warm season; the remaining 17 contribution was from forest and was low in winter and high in summer. The contribution 18 of the re-suspension source area (where the aircraft-observed deposition exceeded 300 19 kBq/m<sup>2</sup>) to the air concentration at Namie was 0.8 - 0.9 throughout the year, while that at 20 Tsukuba varied from 0.1 to 0.4, and was high in the summer and low in the winter.

21 The simulated total re-suspended amount for the whole region was 1.01 TBq, equivalent 4. to 0.037% of the aircraft-observed total deposition amount of 2.68 PBq. The total re-22 23 deposition was 0.18 - 0.23 TBg, equivalent to 17.8 - 22.8% of the total re-suspended amount: the rest of the <sup>137</sup>Cs was transported out of the model domain. The spatial 24 distribution of the decreasing rate of land surface <sup>137</sup>Cs due to re-suspension ranged from 25  $2.2 \times 10^{-7} - 6.6 \times 10^{-6}$  /d. The first order decrease rate of the ambient gamma dose rate in 26 Fukushima ranged from  $5.2 - 12.1 \times 10^{-4}$  /d. By subtracting the radioactive decay rate of 27  $3.0 - 4.2 \times 10^{-4}$  /d, the ground radioactivity decay due to land surface processes, 28 decontamination, and re-suspension from air was found to range from  $1.0 - 7.9 \times 10^{-4}$  /d. 29 30 The estimated re-suspension rate was two to three orders of magnitude lower than the decrease in rate due to the other processes, showing that re-suspension contributed 31 32 negligibly toward reducing ground radioactivity.

1

## 2 Appendix A: Model description

3 The current study employs a Lagrangian type model for the simulation of emission (either 4 point sources or areal sources), horizontal and vertical diffusion and advection, gravitational 5 settling, dry and wet depositions, and radioactive decay in the air. As described in Sect. 2.1, 6 the current Lagrangian Model (LM) uses simple parameterizations for dry and wet deposition schemes for computational efficiency, so long-term assessment and parameter sweep 7 8 experiments are easily feasible. The source code for the model is open with the BSD 3-Clause 9 License and is available on the web (http://157.82.240.167/~dl3/, in Japanese, last access: 12 10 February 2016).

11 The coordinate system of the model is horizontal for longitude and latitude and vertical for 12 pressure level, consistent with meteorological analysis data commonly used. The model can 13 be driven only by fundamental meteorological parameters such as temperature, humidity, 3D 14 wind field, geopotential height, and surface precipitation provided by meteorological analysis 15 data such as GPV-MSM. The model does not need to drive meteorological models to predict 16 detailed meteorological variables such as cloud microphysics, turbulence quantities, and 17 surface variables. Since the temporal and spatial resolution of the meteorological analysis is 18 not very high (e.g., 3 h and ~11 km for GPV-MSM, respectively), linear interpolation is 19 conducted in time and space. Alternatively, higher temporal and spatial resolution can be 20 achieved by using a meteorological model. Furthermore, although currently not implemented, 21 detailed variables predicted by a meteorological model can be used for more accurate 22 predictions of turbulent diffusion, surface flux, and dry and wet deposition.

23 In the LM model, LPs are released constantly in time but the initial activity of LPs (Bq/LP) 24 differs accordingly to the emission flux (Bg/h). The initial positions of LPs were randomly 25 distributed within a fixed volume (or line) of plume centered at a point emission source such 26 as FDNPP for the primary emission case simulation, or randomly distributed within a 27 horizontal model grid for the areal emission cases (such as re-suspension from soil and forest). 28 LPs do not disappear unless transported across lateral and upper boundaries or if they reach 29 the surface layer due to gravitational settling (technically, gravitational settling velocity in the 30 surface layer is included in the dry deposition velocity). The other processes, such as dry 31 deposition, wet deposition and radioactive decay, do not decrease the number of LPs but do 32 decrease the radioactivity carried by LPs because LPs represent an air mass rather than an

actual particle, except in the case of gravitational settling. The lowest level permitted for the 1 2 position of LPs is set as 2 m AGL and LPs going down across the level due to vertical turbulent motion will rebound at the level and go up. An LP whose radioactivity is smaller 3 than a preset value, i.e., 10<sup>-10</sup> Bq, due to deposition or radioactive decay will disappear from 4 5 the computation to maintain computational efficiency, since the cost of the computation is proportional to the number of LPs in the model domain. To output the model results, the LP 6 fields are converted to Eulerian concentration  $(Bq/m^3)$  and deposition  $(Bq/m^2)$  fields on a 7 8 prescribed coordinate system of grids. In Lagrangian type models, the spatial resolution of 9 tracer emission, concentration, and deposition fields can be set independent of each other and 10 with the spatial resolution of meteorological fields. In the current implementation of the LM, 11 the coordinate system of meteorological fields and radioactivity fields is horizontal for common (longitude and latitude) but vertical for different (pressure level and meters AGL, 12 13 respectively).

14 The horizontal and vertical diffusion calculation followed JMA (2008), using the horizontal 15 diffusion scheme of Uliasz (1990) with a constant horizontal diffusivity of  $5.864 \times 10^4 \text{ m}^2/\text{s}$ 16 and using a vertical diffusivity calculated based on Louis et al. (1982) (see Eqs. 8.1.8 through 17 8.1.15 of JMA (2008) for details). The incremental change in location of an LP  $\delta x$  (*y*, *z*) after 18 a time step  $\delta t$  was defined as

19 
$$\delta x = \frac{\mathrm{d}x}{\mathrm{d}t} G \,\delta t$$
, (A1)

where *G* is the normalized Gaussian random number (average = 0, standard deviation = 1).  $\delta t$ is set large enough for computational efficiency but without violating the Courant-Friedrichs-Lewy (CFL) condition of  $\delta t < 0.5 U/\Delta x$ , where *U* (or dx/dt) is the typical wind speed and  $\Delta x$  is the grid size in the direction of *U*. However, the selection of  $\delta t$  is not critical because every time step prior to applying Eq. (A1) time splitting is made so that the split step always satisfies the CFL condition.

26 The wet scavenging rate  $\Lambda_{wet}$  (/s) is simply parameterized as a function of the surface 27 precipitation rate *P* (mm/s) as

28 
$$\Lambda_{wet} = \frac{3}{4} \frac{E_c(a_m, r_m)}{a_m} P, \qquad (A2)$$

1 where  $E_c$  is the collection efficiency of aerosols by the hydrometeor, and  $a_m$  and  $r_m$  are the 2 mean radii of the hydrometeor and aerosols, respectively (JMA, 2008). Empirically,  $a_m$  is 3 characterized by *P* as

4 
$$a_m = 0.35 P^{0.25}$$
. (A3)

5 JMA (2008) uses 0.05 for  $E_c$ . In the current study, instead of explicitly predicting  $E_c$ , its range 6 was set for the sensitivity tests as listed in Table 2.

7 Conceptually, Eq. (A2) is the formulation for the washout process, i.e., the collection of 8 aerosols by the settling hydrometeor particles such as rain and snow.  $a_{\rm m}$  and  $E_{\rm c}$  should differ 9 for rain and snow, but common parameters are used in the current simulation. Also, Eq. (A2) 10 is not applicable for the rainout process, since this process-cloud condensation nuclei or ice 11 nuclei activation and deposition via subsequent cloud microphysical processes- is totally 12 different from the washout process. Because meteorological models were not utilized in this study and thus only relative humidity and surface precipitation rate are available and no cloud 13 14 microphysical information (such as hydrometeors mixing ratio in each model grid) is 15 available, Eq. (A2) is applied for all the LPs located above the grid with P. In order to partly 16 account for the rainout process, Eq. (A2) is not applied to LPs in a grid, where the relative 17 humidity is lower than the minimum value, set as 95% in the simulation.

18 The dry deposition velocity  $v_d$  (m/s) of aerosols (or gases) is conventionally formulated, using 19 an electrical analogy, as an inverse of the summation of resistances (s/m) representing 20 turbulent diffusion in the surface layer, Brownian diffusion (or molecular diffusion for gases), 21 interaction with the land surface (soil, water and vegetation), and gravitational settling for 22 aerosols (e.g., Wesely and Hicks, 2000). Therefore, vd is a function of height as well as of 23 turbulent flux and surface conditions. Nevertheless, vd is set as constant in the simulation, but the height dependency of  $v_d$  is considered in the dry scavenging rate  $\Lambda_{dry}$  (/s), following 24 25 Furuno et al. (1999) as

$$26 \qquad \Lambda_{\rm dry} = \frac{2}{z_{\rm srf}} \left( 1 - \frac{z}{z_{\rm srf}} \right) v_d \,, \tag{A4}$$

where z is the height of the LP (m AGL) and  $z_{srf}$  is the surface layer height set as 100 m AGL in the study. Instead of explicitly predicting v<sub>d</sub>, its range was set at around 0.1 cm/s, a typical speed for a range of aerosols around 1 µm in diameter, for the sensitivity tests, as listed in Table 2. The value of v<sub>d</sub> is applied over land, whereas v<sub>d</sub> over the ocean is multiplied by 0.1, because v<sub>d</sub> over a flat surface is approximately one and two orders of magnitude smaller than
 v<sub>d</sub> over short vegetation such as grass and tall vegetation such as forest, respectively (e.g.,
 Petroff and Zhang, 2010).

4

# 5 Appendix B: Statistical accuracy of the current simulation setting

Because the statistical error of Lagrangian simulation is inversely proportional to the square
of the number of LPs, the statistical accuracy of the current simulation setting was evaluated
using the following measures (relative errors of quantities of the sensitivity runs to those of
the reference run):

10 
$$E_{con}(x,y) = \frac{\left|C_{sens}(x,y) - C_{ref}(x,y)\right|}{C_{ref}(x,y)},$$
 (B1)

11 
$$E_{dep}(x,y) = \frac{\left| D_{sens}(x,y) - D_{ref}(x,y) \right|}{D_{ref}(x,y)},$$
 (B2)

where x and y indicate grid points on the longitudinal and latitudinal axes, respectively.  $C_{sens}$ and  $D_{sens}$  indicate temporal mean surface concentrations (Bq/m<sup>3</sup>) and temporal cumulative depositions (Bq/m<sup>2</sup>) of the sensitivity runs, respectively.  $C_{ref}$  and  $D_{ref}$  are the same as  $C_{sens}$  and  $D_{sens}$  but for the reference run.  $E_{con}$  and  $E_{dep}$  were sampled only at grids where  $C_{ref}(x,y)$  and  $D_{ref}(x,y)$  are greater than their areal mean values, respectively.

#### 17 **B.1** Point source case

18 The number emission rate of LPs,  $N_{LP}$ , was set as 32,000 /h (= $N_{LP ref}$ ) for a point source 19 emission case such as the primary emission in March 2011 and additional emission from the 20 reactor buildings in 2013. The median values together with the 25th and 75th percentile 21 values of  $E_{con}$  and  $E_{dep}$  of the sensitivity runs (sensitivity to deposition parameters and sensitivity to  $N_{LP}$ ) against the reference run are listed on the top half rows of Table B1. Both 22 23  $E_{con}$  and  $E_{dep}$  of  $N_{LP\_ref} \times 4$  were significantly lower than those for the deposition parameters sensitivity run. This result indicates that 32,000 /h for NLP was sufficient to allow a 24 25 statistically significant simulation for the purpose of this study, as shown in Figs. 4 and 5: the 26 difference in concentration and deposition due to the deposition parameters was much larger 27 than the difference due to model uncertainty in  $N_{\text{LP}}$ .  $E_{con}$  and  $E_{dep}$  of  $N_{\text{LP ref}} \times 0.25$  (which are 1 also smaller than those of the deposition parameters sensitivity run) are larger than those of 2  $N_{\text{LP ref}} \times 4$ , indicating fairly well accuracy convergence of the LM model.

#### 3 B.2 Areal emission case

4  $N_{\rm LP \ ref}$  was 16 /h/grid for the areal emission case simulating re-suspension from soil and forests in 2013. The lower half of Table B1 is the same as the upper half except for the areal 5 6 emission case (re-suspension from forest). Both  $E_{con}$  and  $E_{dep}$  of  $N_{LP ref} \times 4$  were much lower 7 than those for the deposition parameters sensitivity run, indicating that 16 /h/grid supports a 8 statistically significant simulation for the purpose of this study, as shown in Figs. 7 - 12. 9 Usually, Eulerian-type models are appropriate for solving areal emission 10 problems-Lagrangian-type models require many more LPs for areal emission cases 11 compared to point source cases and thus become computationally too expensive. In the case 12 of this simulation, especially for the concentration, sensitivity to deposition parameters was 13 much more significant than sensitivity to model uncertainty in  $N_{\rm LP}$  using the sufficiently small 14 number of  $N_{\text{LP ref}} = 16$  /h/grid.  $E_{con}$  and  $E_{dep}$  of  $N_{\text{LP ref}} \times 4$  are smaller than those of  $N_{\text{LP ref}} \times$ 0.25, indicating fairly accuracy convergence of the LM model. 15

16

# Appendix C: Land surface processes, decontamination, and dust deflation suppression due to snow cover

19 Figure C1 presents the time series of ambient gamma dose rates measured at the monitoring 20 posts in Fukushima prefecture indicated in Fig. 1b. The data were obtained from the Nuclear Regulation Authority (NRA), Japan website (http://radioactivity.nsr.go.jp/map/ja/index.html, 21 22 last access: Feb 16, 2016). A total of six, two of the monitoring posts from three geographical areas (Hamadori coastal area, Abukuma highland area, and Nakadori valley area), were 23 24 selected. There are tens to hundreds of monitoring posts in each municipality (village, town, and city) in Fukushima prefecture. We selected a monitoring post in each municipality  $\mathbb{D}^{+}$ 25 applying the following conditions: a post showing the highest dose rate of all posts in the 26 27 municipality at the time of downloading (around 11:00 a.m., December 28, 2015), data are 28 available since April 1, 2012, and the instruments are situated 100 cm above the ground.

The government of Japan designated the evacuation-directed zones as a Difficult-to-Return zone (DRZ) (> 50 mSv/y;  $5.71\mu$ Sv/h), a Habitation-Restricted Zone (HRZ) (20 - 50 mSv/y;

- 31  $2.28 5.71 \,\mu$ Sv/h), and a zone being prepared to have the evacuation directive lifted (< 20
  - 25

1 mSv/y; 2.28  $\mu$ Sv/h), in April 2012 (METI, 2012). The two sites in the Hamadori area, Okuma 2 and Namie (Omaru), have been designated DRZ (13.6 and 11.8  $\mu$ Sv/h on December 28, 2015). 3 The dose rates at the two sites in the Abukuma area, Iitate and Kawamata, dropped below the 4 HRZ level during the analysis period in this study (1.17 and 0.521  $\mu$ Sv/h on December 28, 5 2015). The dose rates in the Nakadori area are below 1  $\mu$ Sv/h (0.242 and 0.201  $\mu$ Sv/h on 6 December 28, 2015).

7 The dose rate significantly dropped when the ground was covered with snow, in January and 8 December 2013 in Hamadori and Nakadori, and from January to early March and December 9 2013 in Abukuma (the elevation of Abukuma is 500 - 1,000 m and higher than Hamadori and Nakadori). Snow cover suppresses re-suspension due to dust deflation. Namie (Tsushima) is 10 located in Abukuma and the ground was covered with snow until early March (Ishizuka et al., 11 2016). The observed air concentration of <sup>137</sup>Cs at Namie (Tsushima) was correlated with the 12 wind speed in the winter from January to March, indicating that re-suspension during the 13 period was mechanically induced. In the winter, dust re-suspension from outside Abukuma, 14 such as from Hamadori and Nakadori, or from land surface where the snow cover was partly 15 melted due to solar radiation, might be the dominant source contributing to the <sup>137</sup>Cs surface 16 air concentration at Namie (Tsushima) when the ground was covered with snow. 17

18 The first order decreasing rates fitted by the least-square approximation for the period without 19 snow cover, May to October 2012 and 2013, are presented in Fig. C1. The rates ranged from  $5.2 - 12.1 \times 10^{-4}$  /d. The monthly mean radioactive decay rates of total radiocesium ( $^{134}$ Cs + 20  $^{137}$ Cs), determined by assuming that the activities of  $^{134}$ Cs and  $^{137}$ Cs were equivalent in March 21 2011, were  $4.2 \times 10^{-4}$  /d and  $3.0 \times 10^{-4}$  /d in April 2012 and March 2014, respectively, due to the 22 difference in half-life of <sup>134</sup>Cs and <sup>137</sup>Cs (2.07 v and 30.1 v, respectively). By assuming that 23 24 the gamma dose rate primarily originated to radiation from the land surface radiocesium, the radioactive decay accounted for 35 - 50% of the decreasing rate of total ground radioactivity; 25 the exception was Shirakawa, where radioactive decay accounted for 55 - 80% of the 26 decrease. In other words, 50 - 65% of the ground radioactivity decrease was likely due to land 27 28 surface processes, decontamination, and re-suspension to air. As discussed in Sect. 4.3, the estimated decreasing rate due to re-suspension was  $2.2 \times 10^{-7} - 6.6 \times 10^{-6}$  /d, which is two to 29 three orders of magnitude smaller than the decreasing rates due to the other processes  $(10^{-4} -$ 30  $10^{-3}$  /d). 31

It is difficult to distinguish the contributions of land surface processes and decontamination. 1 By subtracting the radioactive decay rate  $(3.0 - 4.2 \times 10^{-4} / d)$  and the decreasing rate due to re-2 suspension  $(2.2 \times 10^{-7} - 6.6 \times 10^{-6} / d)$  from the gross decreasing rate  $(5.2 - 12 ) \times 10^{-4} / d)$ , the 3 estimated decreasing rates due to land surface processes (migration of radiocesium in soil) 4 and decontamination ranged from  $1.0 - 7.9 \times 10^{-4}$  /d. Matsuda et al. (2015) summarized the 5 depth profiles of radiocesium in soil at more than 80 locations in Fukushima, including 6 7 Hamadori, Abukuma, and Nakadori. They found that the radiocesium levels have been slowly migrating downward with rates ranging from  $1.7 - 9.6 \text{ kg/m}^2/\text{y}$  (equivalent to 1.1 - 6.0 mm/y8 for a dust particle density of 2650 kg/m<sup>3</sup> and a porosity of 0.4 m<sup>3</sup>/m<sup>3</sup>, for example). The 9 downward migration resulted in decreasing the air dose rate due to incremental soil layers 10 11 blocking radiation, but the downward migration rate has not been quantitatively related to the 12 decrease in the air dose rate. This quantitative relationship needs to be assessed for the 13 quantitative and individual assessment of land surface processes and decontamination effects.

14

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### 27 References

Adachi, K., Kajino, M., Zaizen, Y., and Igarashi, Y.: Emission of spherical cesium-bearing particles from an early stage of the Fukushima nuclear accident, Scientific Report, 3, 2554,

30 doi:10.1038/srep02554, 2013.

- 1 Chino, M., Nakayama, H., Nagai, H., Terada, H., Katata, G., and Yamazawa, H.: Preliminary
- 2 estimation of release amounts of <sup>131</sup>I and <sup>137</sup>Cs accidentally discharged from the Fukushima
- 3 Daiichi nuclear power plant into atmosphere, J. Nucl. Sci. Technol., 48, 1129-1134, 2011.
- 4 Danielache, S. O., Yoshikawa, C., Priyadarshi, A., Takemura, T., Ueno, Y., Thiemens, M.
- 5 H.,and Yoshida, N.: An estimation of the radioactive <sup>35</sup>S emitted into the atmospheric from
- 6 the Fukushima Daiichi Nuclear Power Plant by using a numerical simulation global transport,
- 7 Geochemical Journal, 46, 355-339, 2012.
- 8 Danielache, S. O., Yoshikawa, C., Kajino, M., Itou, S., Kakeya, W., Yoshida, N., Igarashi, Y.:
- 9 Radioactive <sup>35</sup>S emitted from the Fukushima Nuclear Power Plan studied by a modified
- 10 regional model, 2016, manuscript in preparation.
- 11 Draxler, R., Arnold, D., Chino, M., Galmarini, S., Hort, M., Jones, A., Leadbetter, S., Malo,
- 12 A., Maurer, C., Rolph, G., Saito, K., Servranckx, R., Shimbori, T., Solazzo, E., Wotawa, G.:
- 13 World Meteorological Organization's model simulations of the radionuclide dispersion and
- 14 deposition from the Fukushima Daiichi nuclear power plant accident, J. Environ. Radioact.,
- 15 139, 172-184, 2015.
- 16 Furuno, A., Chino, M., and Yamazawa, H.: Worldwide version of System of Prediction of
- 17 Environmental Emergency Dose Information (WSPEEDI) Model Code (II) Three-
- 18 dimensional Atmospheric Dispersion Model for Synoptic Scale: GEARN, JAERI-Data/Code
- 19 99-044, Japan Atomic Energy Research Institute, Japan (in Japanese), 68 pp., 1999.
- Garger, E.K., Paretzke, H.G., Tschiersch, J.: Measurement of resuspended aerosol in the
  Chernobyl area. Part III: size distribution and dry deposition velocity of radioactive particles
  during anthropogenic enhanced resuspension, Radiation and Environmental Biophysics 37,
  201-208, 1998.
- Garger, E. K., Kuzmenko, Y. I., Sickinger, S., Tschiersch, J.: Prediction of the <sup>137</sup>Cs activity
  concentration in the atmospheric surface layer of the Chernobyl exclusion zone, J. Environ.
  Radioact., 110, 53-8, 2012.
- 27 Girard, S., Mallet, V., Korsakissok, I., and Mathieu, A.: Emulation and Sobl' sensitivity
- analysis of an atmospheric dispersion model applied to the Fukushima nuclear accident, J.
- 29 Geophys. Res., 2016, in press.

- 1 Groëll J., Quélo, D., and Mathieu, A.: Sensitivity analysis of the modelled deposition of <sup>137</sup>Cs
- on the Japanese land following the Fukushima accident, Int. J. of Environment and Pollution,
  55, 67-75, 2014.
- Gutman, G., and Ignatov, A.: The derivation of green vegetation fraction from
  NOAA/AVHRR data for use in numerical weather prediction models, Int. J. Remote Sens., 19,
  1533-1543, 1998.
- Hatano, Y., and Hatano N.: Formula for the resuspension factor and estimation of the date of
  surface contamination, Atmos. Environ., 37, 3475-3480, 2003.
- 9 Hirao, S., Yamazawa, H., Nagae, T.: Estimation of release rate of iodine-131 and cesium-137
- 10 from the Fukushima Daiichi nuclear power plant, J. Nucl. Sci. Technol., 50, 139-147,
  11 doi:10.1080/00223131.2013.757454, 2013.
- 12 Holländer, W., Garger, E.: Contamination of Surface by Resuspended Material. ECP-1, Final
- Report, Rep. EUR 16527. Office for Official Publications of the European Communities,
  Luxembourg, 1996.
- Hososhima, M. and Kaneyasu, N.: Altitude-dependent distribution of ambient gamma dose
  rates in mountainous area of Japan caused by the Fukushima nuclear accident. Environ. Sci.
  Technolo., 49(6), 3341-3348, 2015.
- 18 Igarashi, Y., Kajino, M., Zaizen, Y., Adachi, K., and Mikami, M.: Atmospheric radioactivity
- 19 over Tsukuba, Japan: a summary of three years of observations after the FDNPP accident,
- 20 Progress in Earth and Planetary Science, 2:44, doi:10.1186/s40645-015-0066-1, 2015.
- Ishizuka, K, Mikami, M., Tanaka, T. Y., Igarashi, Y., Kita, K., Yamada, Y., Yoshida, N.,
  Toyoda, S., Satou, Y., Kinase, T., Ninomiya, K., Shinohara, A.: Use of a size-resolved 1-D
  resuspension scheme to evaluation resuspended radioactive material associated with mineral
- 24 dust particles from the ground surface, J. Environ. Radioact. 2016, in press.
- JMA: Japan Meteorological Agency's Non-Hydrostatic Model II, *Suuchi Yohoka Hokoku Bessatsu* (Additional Volume to Report of Numerical Prediction Division) 54, 265 pp., 2008
  (in Japanese).
- 28 Kaneyasu, N., Ohashi, H., Suzuki, F., Okada, T., and Ikemori, F.: Sulfate aerosol as a
- 29 potential transport medium of radiocesium from the Fukushima nuclear accident, Environ. Sci.
- 30 Technol. 46, 5720-5726, 2012.

- 1 Katata, G., Terada, H., Nagai, H., and Chino, M.: Numerical reconstruction of high dose rate
- zones due to the Fukushima Daiichi Nuclear Power Plant accident, J. Environ. Radioact., 111,
  2-12, 2012a.
- Katata, G., Ota, M., Terada, H., Chino, M., and Nagai, H.: Atmospheric discharge and
  dispersion of radionuclides during the Fukushima Dai-ichi Nuclear Power Plant accident, Part
  I: Source term estimation and local-scale atmospheric dispersion in early phase of the
- 7 accident, J. Environ. Radioact, 109, 103-113, doi:10.1016/j.jenvrad.2012.02.006, 2012b.
- 8 Katata, G., Chino, M., Kobayashi, T., Terada, H., Ota, M., Nagai, H., Kajino, M., Draxler, R.,
- 9 Hort, M. C., Malo, A., Torii, T., and Sanada, Y.: Detailed source term estimation of the
- 10 atmospheric release for the Fukushima Daiichi Nuclear Power Station accident by coupling
- 11 simulations of an atmospheric dispersion model with an improved deposition scheme and
- 12 oceanic dispersion model. Atmos. Chem. Phys., 15, 1029-1070, 2015.
- 13 Kinase, T., Kita, K. Igarashi, Y., Adachi, K., Ninomiya, K., Shinohara, A., Okochi, H., Ogata,
- 14 Y., Ishizuka, M., Toyoda, S., Yamada, K., Yoshida, N., Zaizen, Y., Mikami, M., Demizu, H.,
- 15 and Onda, Y.: Seasonal variation in activity concentration of atmospheric <sup>134,137</sup>Cs and their
- 16 possibile resuspension host aerosol observed at Tsuahima and Yamakiya, Fukushima, 2016,
- 17 in preparation.
- 18 Loosmore, G. A., Hunt, J. R.: Dust resuspension without saltation, J. Geophys. Res.,
  19 105(D16), 20,663-20,671, doi:10.1029/2000JD900271, 2002.
- Louis, J. F., Tiedtke, M., Geleyn, J. F.: A short history of the PBL parameterization at ECMWF, *Workshop on planetary boundary layer parameterization*. *ECMWF*, 59-80, 1982.
- MAFF (Ministry of Agriculture, Forestry and Fisheries), Fukushima Prefecture, Tohoku
  Agricultural Research Center, National Institute for Agro-Environmental Sciences:
- 24 Investigation of the cause for brown rice exceeding the allowable radiocesium concentration,
- $25 \qquad available \ at \ http://www.maff.go.jp/j/kanbo/joho/saigai/fukusima/pdf/150526\_youin\_chosa.pdf$
- 26 (last access, 10 March, 2016), 2015 (in Japanese).
- 27 Masson, O., Baeza, A., Bieringer, J., Brudecki., Bucci, S., Cappai, M., Carvalho, F. P.,
- 28 Connan, O., Cosma, C., Dalheimer, A., Didier, D., Depuydt, G., De Geer, L. E., De Vismes,
- 29 A., Gini, L., Groppi, F., Gudnason, K., Gurriaran, R., Hainz, D., Halldórsson, O., Hammond,
- 30 D., Hanley, O., Holey, K., Homoki, Zs., Ioannidou, A., Isajenko, K., Jankovic, M.,
- 31 Katzlberger, C., Kettunen, M., Kierepko, R., Kontro, R., Kwakman, P. J. M., Lecomte, M.,

- 1 Leon Vintro, L., Leppänen, A.-P., Lind, B., Lujaniene, G., Mc Ginnity, P., Mc Mahon, C.,
- 2 Malá, H., Manenti, S., Manolopoulou, M., Mattila., A., Mauring., A., Mietelski, J. W., Moller,
- 3 B., Nielsen, S. P., Nikolic, J., Overwater, R. M. W., Pálsson, S. E., Papastefanou, C., Penev, I.,
- 4 Pham, M. K., Povinec, P. P., Ramebäck, H., Reis, M. C., Ringer, W., Rodriguez, A., Rulik, P.,
- 5 Saey, P. R. J., Samsonov, V., Schlosser, C., Sgorbati, G., Silobritiene, B. V., Söderström, C.,
- 6 Sogni, R., Solier, L., Sonck, M., Steinhauser, G., Steinkopff, T., Steinmann, P., Stoulos, S.,
- 7 Sýkora, I., Todorovic, D., Tooloutalaie, N., Tositti, L., Tschiersch, J., Ugron, A., Vagena, E.,
- 8 Vargas, A., Wershofen, H., Zhukova, O. Tracking of Airborne Radionuclides from the
- 9 Damaged Fukushima Dai-Ichi Nuclear Reactors by European Networks. Environ. Sci.
- 10 Technol. 2011, 45 (18), 7670-7677, 2011.
- 11 Masson, O., Ringer, W., Malá, H., Rulik, P., Dlugosz-Lisiecka, M., Eleftheriadis, K.,
- 13 radionuclides from the Fukushima Nuclear Accident at several places in Europe, Environ. Sci.

Meisenberg, O., De Vismes-Ott, A., and Gensdarmes, F.: Size distribution of airborne

14 Technol., 47, 10995-11003, 2013.

12

- Matsunami H., Murakami, T., Fujiwara, H., and Shinano, T.: Evaluation of the cause of
  unexplained radiocaesium contamination of brown rice in Fukushima in 2013 using
  autoradiography and gamma-ray spectrometry, Scientific Report, 6, 20386,
  doi:10.1038/srep20386, 2016.
- METI (Ministry of Economy, Trade and Industry): Convention on Nuclear Safety National
  Report of Japan for the Second Extraordinary Meeting, July 2012, Government of Japan,
  available at www.meti.go.jp/english/press/2012/pdf/0705\_01b.pdf (last access: 19 February,
  2016), 2012.
- Miller, D. A. and White, R. A.: A conterminous United States multi-layer soil characteristics
  data set for regional climate and hydrology modeling, Earth Interactions, 2, 1-26, 1998.
- Matsuda, N., Mikami, S., Shimoura, S., Takahashi, J., Nakano, M., Shimada, K., Uno, K.,
  Hagiwara, S., and Saito, K.: Depth profiles of radioactive cesium in soil using a scraper plate
- 27 over a wide area surrounding the Fukushima Dai-ichi Nuclear Power Plant, Japan., 139, J.
- 28 Environ. Radioact., 427-434, 2015.
- 29 Morino, Y., Ohara, T., and Nishizawa, M.: Atmospheric behavior, deposition, and budget of
- 30 radioactive materials from the Fukushima Daiichi nuclear power plant in March 2011,
- 31 Geophys. Res. Lett. 38, doi:10.1029/2011GL048689, 2011.

- 1 Morino, Y., Ohara, T., Watanabe, S., Hayashi, S., and Nishizawa, M.: Episode analysis of
- 2 deposition of radiocesium from the Fukushima Daiichi Nuclear Power Plant accident, Environ.
- 3 Sci. Technol., 47, 2314-2322, 2013.
- 4 NRA (Nuclear Regulation Authority): Airborne Monitoring Results in each prefecture,
  5 available at: http://radioactivity.nsr.go.jp/en/list/203/list-1.html (last access: 25 December
  6 2015), 2012.
- NRA: Assessment of radionuclide emission involved in the debris removal operations for
  reactor 3; Handout for the 28th meeting on the supervision of a specific nuclear facility,
  available at http://www.nsr.go.jp/data/000051154.pdf (last access, 10 March 2016), 2014 (in
  Japanese).
- 11 Ohmori, Y., Kajikawa, M., Nishida, S., Tanaka, N., Kobayashi, N. I., Tanoi, K., Furukawa,
- 12 and J., Fujiwara, T.: The effect of fertilization on cesium concentration of rice grown in a
- paddy field in Fukushima Prefecture in 2011 and 2012, J. Plant Res., 127, 1, 67-71, 2014.
- Oura, Y., Ebihara, M., Tsuruta, H., Nakajima, T., Ohara, T., Ishimoto, M., Sawahata, H.,
  Katsumura, Y., and Nitta, W.: A database of hourly atmospheric concentrations of
  radiocesium (<sup>134</sup>Cs and <sup>137</sup>Cs) in suspended particulate matter collected in March 2011 at 99
  air pollution monitoring stations in Eastern Japan, Journal of Nuclear and Radiochemical
  Sciences, 15(2), 15-26, 2015.
- Petroff, A. and Zhang L.: Development and validation of a size-resolved particle dry
  deposition scheme for application in aerosol transport models, Geosci. Model Dev. 3, 753-769,
  2010.
- Pöhlker C., Wiedemann, K. T., Sinha, B., Shiraiwa, M., Gunthe, S. S., Smith, M., Su, H.,
  Artaxo, P., Chen, Q., Cheng, Y., Elbert, W., Gilles, M. K., Kilcoyne, A. L. D., Moffet, R. C>,
  Weigand, M., Matin, S. T., Pöschl, U., and Andreae, M. O.: Biogenic potassium salt particles
- as seeds for secondary organic aerosol in the Amazon, Science, 337, 1075-1078, 2012.
- 26 Quérel, A., Roustan, Y., Quélo, D., and Benoit, J. -P.: Hints to discriminate the choice of wet
- 27 deposition models applied to an accidental radioactive release, Int. J. of Environment and
- 28 Pollution, in review.
- 29 Saito, K., Shimbori, T., and Draxler, R.: JMA's regional atmospheric transport model
- 30 calculations for the WMO technical task team on meteorological analyses for Fukushima
- 31 Daiichi Nuclear Power Plant accident, J. Environ. Raioact., 139, 185-199, 2015.

- Saunier, O., Mathieu, A., Didier, D., Tombette, M., Quelo, D., Winiarek, V., and Bocquet,
   M.: An inverse modeling method to assess the source term of the Fukushima Nuclear Power
   Plant accident using gamma dose rate observations, Atmos. Chem. Phys., 13, 11403-11421,
- 4 doi:10.5194/acp-13-11403-2013, 2013.
- 5 SCJ (Science Council of Japan): A review of the model comparison of transportation and 6 deposition of radioactive materials released to the environment as a result of the Tokyo 7 Electric Power Company's Fukushima Daiichi Nuclear Power Plant accident, available at 8 http://www.scj.go.jp/ja/info/kohyo/pdf/kohyo-22-h140902-e1.pdf (last access: 22 February 2016),
- 9 2014.
- 10 Sekiyama, T. T., Kunii, M., Kajino, M., and Shimbori, T.: Horizontal resolution dependence
- 11 of atmospheric simulations of the Fukushima nuclear accident using 15-km, 3-km, and 500-m
- 12 grid models, J. Meteor. Soc. Japan, 93, 49-64, 2015.
- 13 Sekiyama, T. T., Kajino, M., and Kunii, M.: Ensemble dispersion simulation of a point-source
- 14 radioactive aerosol using a square root Kalman filter, J. Geophys. Res., 2016, in review.
- 15 Skamarock, W. C., Klemp, J. B., Dudhia, J., Grill, D. O., Barker, D. M., Duda, M. G., Huang,
- 16 X. Y., Wang, W., and Powers, J. G.: A description of the advanced research WRF version 3,
- 17 Tech. Note, NCAR/TN~475+STR, 125 pp., Natl. Cent. For Atmos. Res., Boulder, Colo.,18 2008.
- 19 Steinhauser, G., Niisoe, T., Harada, K. H., Shozugawa, K., Schneider, S., Synal, H.-A,
- 20 Walther, C., Christl, M., Nanba, K., Ishikawa, H., and Koizumi, A: Post-accident sporadic
- 21 releases of airborne radionuclides from the Fukushima Daiichi Nuclear Power Plant Site,
- 22 Environ. Sci. Technol., 49, 14028-14035, doi:10.1021/acs.est.5b03155, 2015.
- Stohl, A. Seibert, P., Wotawa, G., Arnold, D., Burkhart, J. F., Eckhardt, S., Tapia, C., Vargas,
  A., and Yasunari, T. J.: Xenon-133 and caesium-137 releases into the atmosphere from the
  Fukushima Dai-ichi nuclear power plant: determination of the source term, atmospheric
- dispersion, and deposition. Atmos. Chem. Phys., 12, 2313-2343, 2012.
- TEPCO (Tokyo Electric Power Co., Inc.): Estimation of additional emissions from nuclear
  reactor buildings (30 July 2012), available at: http://www.tepco.co.jp/nu/fukushima-
- 29 np/roadmap/images/m120730\_05-j.pdf (last access: 25 December 2015), 2012 (in Japanese).

- 1 TEPCO: Estimation of the additional emission from nuclear reactor buildings (30 May 2013),
- 2 available at: http://www.tepco.co.jp/nu/fukushima-np/roadmap/images/d130530\_05-j.pdf (last
- 3 access: 25 December 2015), 2013 (in Japanese).
- 4 TEPCO: Estimation of the additional emission from nuclear reactor buildings (27 March
- 5 2014), available at: http://www.tepco.co.jp/nu/fukushima-np/roadmap/images/d140327 06-
- 6 j.pdf (last access: 25 December 2015), 2014a (in Japanese).
- 7 TEPCO: Estimation of the additional emission from nuclear reactor buildings (25 December
- 8 2014), available at: http://www.tepco.co.jp/nu/fukushima-np/roadmap/images/d141225\_08-
- 9 j.pdf (last access: 25 December 2015), 2014b (in Japanese).
- 10 TEPCO: Estimation of radioactive material released during debris removal operations at

11 reactor unit 3 of the Fukushima Daiichi Nuclear Power Plant in August 2013, available at

- 12 http://www.nsr.go.jp/data/000051136.pdf (last access: 10 March, 2016), 2014c (in Japanese).
- 13 TEPCO: Estimation of the additional emission from nuclear reactor buildings (9 October
- 14 2015), available at: http://www.tepco.co.jp/life/custom/faq/images/d151109\_11-j.pdf (last
- 15 access: 25 December 2015), 2015 (in Japanese).
- Terada, H., Katata, G., Chino, M., and Nagai, H.: Atmospheric discharge and dispersion of
  radionuclides during the Fukushima Dai-ichi Nuclear Power Plant accident, Part II:
  Verification of the source term and analysis of regional-scale atmospheric dispersion, J.
  Environ. Radioact., 112, 141-154, 2012.
- 20 Tsuruta, H., Oura, Y., Ebihara, M., Ohara, T., and Nakajima, T.: First retrieval of hourly 21 atmospheric radionuclides just after the Fukushima accident by analysing filter-tapes of 22 monitoring stations, Scientific operational air pollution Reports, 4, 6717, doi:10.1038/srep06717, 2014. 23
- Uliasz, M.: Development of the mesoscale dispersion modelling system using personal
  computers. Part I: Models and computer implementation. Z. Meteor., 40, 110-120, 1990.
- Wesely, M. L. and Hicks, B. B.: A review of the current status of knowledge on dry
  deposition, Atmos. Environ., 34, 2261-2282, 2000.
- 28 Winiarek, V., Bocquet, M., Saunier, O., and Mathieu, A.: Estimation of errors in the inverse
- 29 modeling of accidental release of atmospheric pollutant: Application to the reconstruction of

- 1 the cesium-137 and iodine-131 source terms from the Fukushima Daiichi power plant, J.
- 2 Geophys. Res., 117, D05122, 16 pp, doi:10.1029/2011JD016932, 2012.
- 3 Winiarek, V., Bocquet, M., Duhanyan, N., Roustan, Y., Saunier, O., and Mathieu, A.:
- 4 Estimation of the caesium-137 source term from the Fukushima Daiichi nuclear power plant
- 5 using a consistent joint assimilation of air concentration and deposition observations, Atmos.
- 6 Environ., 82, 268-279, 2014.
- 7 Yasunari, T. J., Stohl A., Hayano R. S., Burkhart J. F., Eckhardt S., and Yasunari T.: Cesium-
- 8 137 deposition and contamination of Japanese soils due to the Fukushima nuclear accident.
- 9 Proc. Natl. Acad. Sci., 108, 19530-19534, 2011.
- 10 Yumimoto, K., Mirno, Y., Ohara, T., Oura, Y., Ebihara, M., Tsuruta, H., and Nakajima, T.:
- 11 Inverse modeling of the 137Cs source term of the Fukushima Dai-ichi Nuclear Power Plant
- 12 accident constrained by a deposition map monitored by aircraft, J. Environ. Radioact., 2016,
- 13 in review.
- 14
- 15

N	lame	Location	Description
0	bservation sites		
N	amie (Tsushima)	140.7683 °E, 37.5621 °N	Namie High School, Tsushima Campus <sup>1</sup>
Т	sukuba	140.1254 °E, 36.0551 °N	Meteorological Research Institute
M	lonitoring posts		
0	kuma	140.9969 °E, 37.4163  °N	Ottozawa 3 Community Center <sup>2</sup>
N	amie (Omaru)	140.9296 °E, 37.4665 °N	Omaru Multipurpose Community Center
Ii	tate	140.7385 °E, 37.6772 °N	litate Junior High School <sup>4</sup>
K	awamata	140.6979 °E, 37.5836 °N	Yamakiya Otsu 8 Community Firehouse <sup>5</sup>
F	ukushima	140.4765 °E, 37.6870 °N	Fukushima-Minami Fire Department
S	hirakawa	140.1904 °E, 37.1241 °N	Takayama-Kita Park
1.	Original location (now moved to Nihonmatsu city).		
2.	Ottozawa San-ku Chiku Shukaijo (in Japanese)		
3.	Omaru Tamokuteki Shukaijo (in Japanese)		
4.	Original location (now moved to Fukushima city).		
5.	. Yamakiya Ots	u Hachi-ku Community Shou	ıbou Center (in Japanese)

1 Table 1. The observation sites and monitoring posts used to provide data for this study.
	$E_c^{a}$	$v_d^{b}$	$D_{\rm all}{}^{\rm c}$	$D_{\text{land}}^{d}$	$FB^{e}$	$R^{\mathrm{f}}$	FA2 <sup>g</sup>	FA5 <sup>h</sup>
	(-)	(cm/s)	(PBq)	(PBq)				
Sensitivity test								
Range	0.02 -	0.05 -	3.4 -	2.0 -	-0.25 -	0.73 –	0.25 -	0.63 -
	0.06	0.15	4.7	2.8	0.00050	0.78	0.30	0.77
Optimization use	ed for the re	e-suspension	n analysis	for 2013				
Reference	0.04	0.10	4.2	2.5	-6.2	0.75	0.28	0.74
Optimized	0.03 -	0.05 -	3.7 -	2.2 -	-0.18 -	0.74 –	0.26 -	0.68 -
range	0.05	0.10	4.3	2.6	-0.036	0.77	0.30	0.74
Reference value.	5							
Observed depos	ition over la	and $D_{\rm obs}$ (N	RA, 2012)	2.68	PBq			
Emission amount (Katata et al., 2015)				14.1 PBq				

Table 2. <sup>137</sup>Cs budget and statistical analysis for the comparison of observed and simulated
deposition data for March 2011.

<sup>a</sup> Correction efficiency, see Eq. (A2) <sup>b</sup> Dry deposition velocity over land, see Eq. (A4). <sup>c</sup> Simulated deposition amount over the whole model domain. <sup>d</sup> Simulated deposition amount only over land. <sup>e</sup> Fractional bias between  $D_{\text{land}}$  and  $D_{\text{obs.}}$  <sup>f</sup> Correlation coefficient between each grid cell of the observed and simulated deposition (linear vs. linear). <sup>g</sup> Fraction of simulated values within a factor of 2 of the observed values. <sup>h</sup> Fraction of simulated values within a factor of 5 of the observed values. <sup>f, g, h</sup> Compared only at grids where the observed values are greater than 10 kBq/m<sup>2</sup>.

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1 Table B1. Statistical measures of temporal mean <sup>137</sup>Cs surface concentration ( $E_{con}$ ) and 2 cumulative deposition ( $E_{dep}$ ) of the sensitivity runs against the reference run for (top) the point 3 source case and (bottom) the areal emission case.

	Number emission	$E_c^{\ a}$	$v_d^{\ b}$	Median (25th –	Median (25th –	
	rate of LP, $N_{\rm LP}$	(-)	(cm/s)	75th percentile)	75th percentile)	
	(/h/grid)			of $E_{con}^{c}$ (%)	of $E_{dep}^{d}$ (%)	
Point source case,	, March 2011					
Reference run	32,000	0.04	0.01	-	-	
Sensitivity runs						
Deposition	32,000	0.03 -	0.05 -	2.7	7.5	
parameters		0.05	0.01	(0.37 – 5.0)	(3.6 – 13)	
$N_{\rm LP\_ref} \times 4$	128,000	0.04	0.01	0.51	0.72	
				(0.21 – 0.96)	(0.32 – 1.5)	
$N_{\rm LP\_ref} \times 0.25$	8,000	0.04	0.01	0.95	1.6	
				(0.39-1.8)	(0.73 – 2.9)	
Areal emission ca	se, 2013 (re-suspensi	on from fo	rest)			
Reference run	16	0.04	0.10	-	-	
Sensitivity runs						
Deposition	16	0.03 -	0.05 -	7.3	7.7	
parameters		0.05	0.01	(2.6 – 13)	(3.6 – 17)	
$N_{\rm LP\_ref} \times 4$	64	0.04	0.10	0.39	2.0	
				(0.17 – 0.78)	(1.3 – 2.6)	
$N_{\rm LP\_ref} \times 0.25$	4	0.04	0.10	0.78	2.3	
				(0.34-1.5)	(1.5 – 3.1)	

<sup>a</sup> Correction efficiency, see Eq. (A2). <sup>b</sup> Dry deposition velocity over land, see Eq. (A4). <sup>c</sup>
Relative errors of temporal mean surface concentration at each grid cell of the sensitivity run
to that of the reference run, see Eq. (B1). <sup>d</sup> same as *E<sub>con</sub>* but for cumulative deposition, see Eq. (B2).





Name of prefectures: 1. Miyagi, 2. Fukushima, 3. Ibaraki, 4. Tochigi, 5. Gunma, 6. Saitama, 7. Chiba, 8. Tokyo

3 Figure 1. (a) The model domain showing the model terrestrial elevation, observation sites, and other locations described in the study. The linear distances from FDNPP to Namie (Tsushima 4 5 district, Namie town), Tsukuba and Tokyo are approximately 30 km, 170 km, and 220 km, respectively. The numbers denote prefectures: 1. Miyagi, 2. Fukushima, 3. Ibaraki, 4. Tochigi, 6 7 5. Gunma, 6. Saitama, 7. Chiba, and 8. Tokyo. (b) Fukushima prefecture and (colored circles) 8 the locations (village, town, or city name) of monitoring posts used in this study (see Fig. C1). 9 The letters in both (a) and (b) denote the name of the area based on geographical features: A. 10 Nakadori valley, B. Abukuma highland, and C. Hamadori coastal area.



Figure 2. The areal fractions of (a)-(c) soil texture and (d)-(f) land use category used for the
boundary conditions of the simulation.



Figure 3. Monthly mean emission flux of radiocesium released from the reactor buildings of
FDNPP from October 2011 to August 2015 as estimated by TEPCO (TEPCO, 2012-2015).



Figure 4. (a) Aircraft observation and (b), (c) simulation of <sup>137</sup>Cs deposition depicted (b) only 2 over land and (c) for the whole domain  $(kBq/m^2)$ . The observation was interpolated to the 3 4 model grid ( $\Delta$ longitude = 0.125° and  $\Delta$ latitude = 0.1°). A decay correction for the observation 5 was made for March - May 2012, varied depending on the region. The simulation provided a 6 cumulative value from March 12 to April 1, 2011. The total activities are presented as numbers. The color scales are the same for (a) - (c) but observed values are not depicted 7 below the detection limit, 10 kBq/m<sup>2</sup>. The deposition parameters of the simulation in the 8 figure are  $E_c = 0.04$  and  $v_d = 0.1$  cm/s. 9



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Figure 5. Scattergram between the observational data and the simulation results for <sup>137</sup>Cs deposition (kBq/m<sup>2</sup>). The deposition parameters of the simulation in the figure are  $E_c = 0.04$ and  $v_d = 0.1$  cm/s. *N* indicates the number of samples, and the statistical measures *R*, *FA2*, and *FA5* are described in Table 2.



Figure 6. Time series of the surface activity concentration of (black) observed and (red) simulated <sup>137</sup>Cs concentrations at Tsukuba from March 12 to April 1, 2011 (Bq/m<sup>3</sup>). The red shaded areas indicate the range of <sup>137</sup>Cs concentrations obtained when the simulations were run using the optimized parameter ranges  $E_c = 0.03 - 0.05$  and  $v_d = 0.05 - 0.1$  cm/s.



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Figure 7. Time series of the surface air concentration of (black) observed <sup>137</sup>Cs and (colors) simulated daily <sup>137</sup>Cs levels at (a) Namie and (b) Tsukuba. The colored lines indicate simulated <sup>137</sup>Cs concentrations due to (red) re-suspension from soil using the scheme given in Ishizuka et al. (2016) (10 times), (green) re-suspension from forest with a re-suspension rate of 10<sup>-7</sup> /h, and (blue) emission from the FDNPP reactor buildings with a constant emission rate of 10<sup>6</sup> Bq/h. The deposition parameters of the simulation in the figure are  $E_c = 0.04$  and  $v_d = 0.1$  cm/s.



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Figure 8. (a) Annual total re-suspension amounts of <sup>137</sup>Cs and (b) the re-deposition amounts. The total activities are presented as numbers. Ratios of the re-suspension and the redeposition to the observed <sup>137</sup>Cs deposition amounts are also shown in (c) and (d), respectively. The deposition parameters of the simulation in the figure are  $E_c = 0.04$  and  $V_d = 0.1$  cm/s.



Figure 9. Time series of (black) the observed and (colors) the simulated <sup>137</sup>Cs surface air concentration at Namie (a, c) in winter from January to March and (b, d) in summer from June to August, 2013. The colors of the lines are the same as in Fig. 7 but the time variation is hourly. The shaded areas indicate the range of <sup>137</sup>Cs concentrations obtained when the simulations were run using the optimized parameter ranges  $E_c = 0.03 - 0.05$  and  $v_d = 0.05 - 0.1$  cm/s.



9 Figure 10. Seasonal mean surface (10 m above ground level) wind field and <sup>137</sup>Cs surface 10 concentration (a) due to dust re-suspension in winter from January to March and (b) due to 11 forest re-suspension in summer from June to August. The deposition parameters of the 12 simulation in the figure are  $E_c = 0.04$  and  $v_d = 0.1$  cm/s.



Figure 11. Time series of (a, b) (black) the observed and (purple) the simulated surface <sup>137</sup>Cs concentration due to total re-suspension and (c, d) the relative contribution of (red) dust and (green) forest re-suspension to the <sup>137</sup>Cs concentration at (a, c) Namie and (b,d) Tsukuba. The shaded areas of (1) and (b) indicate the range of <sup>137</sup>Cs concentrations obtained when the simulations were run using the optimized parameter ranges  $E_c = 0.03 - 0.05$  and  $v_d = 0.05 - 0.1$  cm/s.



9 Figure 12. Monthly mean (a) observed mean, observed median and simulated Namie to 10 Tsukuba <sup>137</sup>Cs concentration ratio and (b) simulated re-suspension source area (where the 11 observed deposition amount > 300 kBq/m<sup>2</sup>) contributions to <sup>137</sup>Cs air concentration at Namie 12 and Tsukuba. The deposition parameters of the simulation in the figure are  $E_c = 0.04$  and  $v_d =$ 13 0.1 cm/s.



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Figure 13. 48 h forward trajectory (statistical locations of LPs within 1 km AGL) predicted by the LM starting at (a) 12 LT and (c) 15 LT of August 14 and (b) 9 LT and (d) 15 LT of August 19. Blue lines indicate median locations of LPs at 1 h and 1 d intervals as blue and red dots, respectively. Sky-blue lines indicate 17th and 83rd percentile locations of LPs and red dashed circles indicate areas containing 66% of the LPs.



Figure C1. Daily mean gamma dose rate at the six monitoring sites: (a) (red) Okuma town and 1 2 (orange) Namie town (Omaru district) in the coastal area, (b) (blue) litate village and (sky blue) Kawamata town in the Abukuma highland area, and (c) (green) Fukushima city and 3 4 (greenish yellow) Shirakawa city in the Nakadori valley area of Fukushima prefecture as presented in Fig. 1b for two fiscal years (from April 2012 to March 2014). The first order 5 decreasing rates of the least-square approximation over the period of no snow cover (May to 6 October, 2012 and 2013) is also presented, along with the radioactive decay (<sup>134</sup>Cs plus <sup>137</sup>Cs, 7 by assuming the same amount of activity on March 12, 2011) in April 2012 and March 2013. 8 9