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Predicting terrestrial 222 Rn flux

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Predicting terrestrial ²²²Rn flux using gamma dose rate as a proxy^{*}

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Abstract

²²²Rn is commonly used as a natural tracer for validating climate models. To improve such models a better source term for ²²²Rn than currently used is necessary. The aim of this work is to establish a method for mapping this source term by using a commonly measured proxy, the gamma dose rate (GDR). Automatic monitoring of GDR has been networked in 25 European countries by the Institute for Environment and Sustainability at the Joint Research Centre (JRC IES) in Ispra, Italy, using a common data format. We carried out simultaneous measurements of ²²²Rn flux and GDR at 63 locations in Switzerland, Germany, Finland and Hungary in order to cover a wide range of GDR. Spatial variations in GDR resulted from different radionuclide concen-10 trations in soil forming minerals. A relatively stable fraction (20%) of the total terrestrial GDR originates from the ²³⁸U decay series, of which ²²²Rn is a member. Accordingly, spatial variation in terrestrial GDR was found to describe almost 60% of the spatial variation in ²²²Rn flux. Furthermore, temporal variation in GDR and ²²²Rn was found to be correlated. Increasing soil moisture reduces gas diffusivity and the rate of ²²²Rn flux but it also decreases GDR through increased shielding of photons. Prediction of ²²²Rn flux through GDR for individual measurement points is imprecise but un-biased. Verification of larger scale prediction showed that estimates of mean ²²²Rn fluxes were not significantly different from the measured mean values.

20 1 Introduction

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²²²Rn is commonly known as a hazardous radioactive (noble) gas in indoor air. Yet, ²²²Rn is also often used as a natural tracer of air transport. Observations of atmospheric ²²²Rn have been very useful in the evaluation of climate models simulating transport, transformation and removal processes of gases and aerosols (e.g. Rasch, 2000). Used in inverse mode, these models can provide information on location, extent and strength of sources and sinks of greenhouse gases based on the measurement

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of changes in their atmospheric concentrations (Chevillard, 2002; Gupta, 2004). Currently, the effective use of ²²²Rn in this context is limited by the poor accuracy of the ²²²Rn source function (WMO GAW report no. 155, 2004). Current practice is to assume a spatial and temporal uniform flux rate of 1 atom cm⁻² s⁻¹ from all ice-free land ⁵ surfaces. Improvement of the source term was attempted by Schery and Wasiolek (1998), who created a global ²²²Rn flux map based on porous media transport theory and calibrated with experimental radon flux data from Australia and Hawaii. It predicted regional variations of a factor of three not to be uncommon. However, current lack of detailed data on input parameters in large parts of the world results in the proposed map still being preliminary and depending on more data becoming available. Furthermore, additional flux measurements over a greater variety of conditions are needed for robust validation and eventual verification of the model. A different interpretation of

the flux term was proposed by Conen and Robertson (2002), based on atmospheric profile measurements integrating over larger areas and indicating a decline in ²²²Rn flux from ice-free land surface from 1 atom cm⁻² s⁻¹ at 30° N to 0.2 atom cm⁻² s⁻¹ at 70° N. This source term was found to improve predictions but it was speculated that ²²²Rn flux might begin to decline well north of 30° N (Robertson et al., 2005). A more detailed source term is highly desirable to improve validation of atmospheric transport models since the quality of validation is directly proportional to the quality of the ²²²Rn

20 source term used.

Therefore, we are proposing a new method to describe the ²²²Rn source term, initially focusing on the European continent. Our approach is to calibrate direct measurements of ²²²Rn flux against terrestrial gamma dose rate (GDR). We made use of the high density of European GDR measurements, established after the nuclear reactor accident in Chernobyl in 1986, to produce a full description of the European ²²²Rn

accident in Chernobyl in 1986, to produce a full description of the European ²²² I source term.



2 Basic concept

The source of ²²²Rn is ²²⁶Ra, a member of the ²³⁸U decay chain. Gamma spectroscopic analysis of soil surface samples (0-20 cm depth) in geologically diverse regions of Switzerland showed that ²³⁸U contributes an almost constant proportion to the terrestrial GDR (Fig. 1a) and that ²²⁶Ra activity is closely related to the ²³⁸U activity (Fig. 1b). Large radioactive disequilibria of the uranium decay series have been found in the limestone Karst soils in the Jura mountains (Von Gunten et al., 1996). Selective migration of individual members of the ²³⁸U decay chain could lead to an over- or underestimated GDR-based ²²²Rn flux in such locations. However, such cases seem to be rare, as seen in the close correlations in Figs. 1a and b. Therefore, we assume that 10 ²²²Rn flux resulting from the decay of ²²⁶Ra is directly related to terrestrial GDR. This assumption is probably a good first approximation but not entirely correct as indicated by the relatively large scatter in the ratio of ²²²Rn flux to ²²⁶Ra activity (Fig. 1c). Firstly, only part of the produced ²²²Rn emanates into air filled pore space from where it might escape into the atmosphere and the fraction emanating may depend on grain size (Nazaroff, 1992). Secondly, differences in grain size and soil moisture modulate gas diffusivity and thus the fraction of emanated ²²²Rn that may reach the atmosphere before decay. Thus, the proportion of ²²²Rn produced that escapes into the atmosphere is variable and depends on factors other than ²²⁶Ra content.

20 3 Methods

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3.1 ²²²Rn flux measurement techniques

A barely modified closed chamber method as described in Lehmann et al. (2000, 2003) was used to measure the ²²²Rn flux. The main modification consisted in air from the chamber not being pumped through a series of two but only one alpha-decay detector (Alphaguard 2000 Pro, Genitron Instruments Frankfurt, Germany). The flow rate was



0.5 l min⁻¹, a delay volume of 1.5 l was used to remove most of the ²²⁰Rn with its half-life of 56 s (Lehmann et al., 2003 used the second detector, which was installed before the delay volume to evaluate also the ²²⁰Rn flux). From there, the air passed to the detector where only ²²²Rn was measured. The ²²²Rn flux was estimated from the increase in ²²²Rn activity measured in 10 min intervals over about 1.5 h. Remaining ²²⁰Rn may have affected the absolute value of measured ²²²Rn activity but not its increase over time, as ²²⁰Rn concentrations reach a steady state between production and decay after about 7 min and we always rejected the first 10 min measurement interval. Due to radioactive decay of ²²²Rn with a half-life time of 3.82 days the assumption of a linear increase of ²²²Rn in the chamber must be corrected by a factor of +0.38 %. Two types of chambers were used: an automatically closing and opening chamber which measured autonomously the ²²²Rn flux from soil over a longer time period. This flux chamber, a cylindrical box with a diameter of 20 cm and 25 cm height had a flap, which closed automatically 6 times a day for 1.5 hours to accumulate ²²²Rn and was then opened for 2.5 h prior to the next measurement. A second analytical system was a 15 manually closable chamber (a plastic box with the dimensions 35 cm×27 cm and 13 cm

height) which was used for spot measurements.

Long-term measurements of ²²²Rn fluxes were made at 7 different field sites of the Swiss Meteorological Service (MétéoSuisse). Normally, measurements took place for

a duration of 3–4 weeks, except at the field site in Basel-Binningen, where continuous measurements were made over a year in order to estimate seasonal variations. Soil moisture at that location was measured with 4 TDR two-rod probes (rod length: 18 cm), connected with a multiplexer to a Tektronix 1502B (Tektronix, Inc., Wilsonville, USA). The signal was evaluated and logged with a data logger (CR10, Campbell Scientific, Inc., USA).

The manually closing chamber was used for in situ measurements of ²²²Rn flux at 29 sites in Switzerland and South-West Germany, at 8 sites each in Southern (Helsinki region) and Northern Finland (Rovaniemi region) and at 12 sites in Hungary. Supplementary data from Scotland (Robertson, 2005) was included. These measurements



(n=9) were done with the same analytical ²²²Rn system. The difficulty of spot measurements of ²²²Rn flux and GDR is to get representative values for the specific location. Especially precipitation has been found to have significant effects on GDR because of the deposition of Rn daughters associated with aerosols, but also on short-term variations in ²²²Rn flux. Therefore, we avoided spot measurements during or immediately (4–8 h) after precipitation events. Additionally we studied on small scale spatial variability in a woodland in Basel (Lange Erlen) using a nested sampling design with lag distances of 0.5 m, 5 m and 50 m.

3.2 Gamma dose rate

An autonomous gamma probe (Gammatracer, Genitron Instruments Frankfurt, Germany) for continuous surveillance of the environmental gamma radiation was used for measuring GDR (10 H*). The gamma probe was placed 1 m above ground during the measurement. Since most of the measurements took place at locations of the national gamma monitoring networks, where GDR is continuously measured, the gamma probe
 was used as a reference probe. This allowed inter-comparison of different probes at the network sites. The terrestrial component of the gamma dose rate was obtained by subtracting the cosmic part (which depends on altitude above sea level and can be calculated) from the measured total GDR (Murith and Gurtner, 1994). A correction was made for the artificial radiation, which is mainly derived from ¹³⁷Cs from the Chernobyl powerplant accident in 1986, based on the "Atlas of Caesium deposition on Europe after the Chernobyl accident" (De Cort et al, 1998).



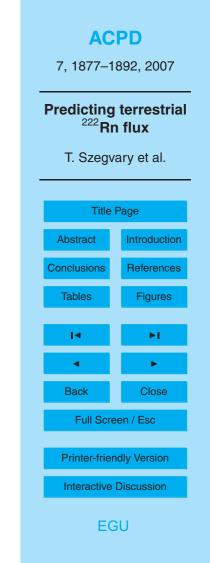
4 Results and discussion

4.1 Correlation of ²²²Rn flux and terrestrial GDR at different locations

There is a linear relationship between ²²²Rn flux and terrestrial GDR (Fig. 2), though the effect of heteroscedasticity is observed, i.e. the variability described by standard deviation depends on the mean value. This means high GDR values are associated with higher variability (an effect, which is often observed in nature). The measured data covers a range from almost 0 to 200 nSv h⁻¹ respectively 0 to 250 Bq m⁻² h⁻¹. Most soils in Europe have gamma dose rates between about 40 to 140 nSv h⁻¹ well within this range. Very low GDR (~40 nSv/h and a ²²²Rn flux less than 15 Bq m⁻² h⁻¹)
10 can be found at locations which have either a high water content and/or low or no mineral content like peat soils. Overall, almost 60% of the variation in ²²²Rn flux can be described by the spatial variation of terrestrial GDR.

Still, there is a lot of variation, which may also be caused by the gamma probe and the ²²²Rn measurement chamber integrating over different soil volumes. The measurement of GDR is mostly influenced by the variability of radionuclides and soil moisture near the soil surface (0 to 0.1 m) within a radius of about 10 m around its location. In contrast, measured ²²²Rn flux is mostly influenced by ²²⁶Ra content and soil moisture in 0 to 1 m soil depth but a three to four orders of magnitude smaller area. Thus, inhomogeneities in radionuclide and moisture distribution on this scale will affect both

- ²⁰ parameters to a different extent. The scatter in Fig. 2 is unlikely to be caused by short-term fluctuations in either parameter. Not only the short-term measurements (triangles) show the scattering effect, but also the long-term measurements (circles), which would smoothen out such short term effects. The nested sampling near Basel revealed that the coefficient of variation between measurements separated by a distance of 0.5 m
- ²⁵ was 19%, increasing to 21% and 36% for 5.0 and 50.0 m distances, respectively. The large coefficient of variation at the smallest distance may to a large part be caused by the error in our ²²²Rn measurement, which we estimate to be around ±15% of the mean. For atmospheric tracer applications, regional information on the ²²²Rn flux is



required. The variability in the correlation between GDR and ²²²Rn flux, which can be found on the local scale, seem to counter balance on the regional scale, as discussed later (see Sect. 4.4).

4.2 Correlation of ²²²Rn flux and terrestrial GDR over time

- Temporal variations in ²²²Rn flux can be observed in GDR at the long-term measurement in Basel (Fig. 3) during the period from June to November 2006, where soil moisture and precipitation was also measured. At the beginning of July a prolonged dry period began without nearly any precipitation and soil moisture decreased almost constantly. During this period the ²²²Rn flux was observed to increase by about 100% until the beginning of August. Simultaneously, GDR increased from 82 nSv/h to 98 nSv/h, which is nearly 20%. Decreasing soil moisture increases the air filled pore volume and with it the diffusivity of soil. Therefore, ²²²Rn flux is larger when soils are dry and less ²²²Rn decavs before it may reach the soil surface (Grasty, 1997). At the same time, low soil moisture leads to reduced shielding of gamma-rays and a larger proportion of them can be detected in the atmosphere above the ground. Diurnal changes in the 15 amplitude of GDR during periods without precipitation are supposed to be influenced by changes in Rn and Rn-progeny concentrations in the near surface air, where they accumulate during atmospherically stable conditions at night (Greenfield, 2002, 2003). At end of September through the beginning of October three intense rain events were recorded (Fig. 3). These were days within a period of otherwise stable weather 20 conditions, where during a short time period between 60 mm and 80 mm of rain fell, approximately the same amount for all three rain events. After each of the three events, the ²²²Rn flux decreased immediately with the beginning of precipitation, probably because of the wet soil surface severely inhibiting ²²²Rn diffusion into the atmosphere. The reaction of GDR was initially to the contrary. It suddenly increased after the first 25
- rain event from 85 nSv/h to 110 nSv/h, an increase of 29%. This effect is caused by outwash of particles from the lower atmosphere, carrying previously absorbed ²²²Rn

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progeny back to the soil surface (Greenfield, 2002, 2003). The cummulative half-life of the short-lived ²²²Rn progeny is about 50 min. Thus, the GDR decreased within a few hours once rain had stopped and was lower than it was before the rain event (~8–10%). The second and third rain event showed the same effect. The only difference ⁵ between the three rain events was the amplitude of the peak at the start of each rain fall, which was smaller for the second and third compared to the first one because the atmosphere was getting increasingly cleaner of particles carrying ²²²Rn progeny.

4.3 Factors affecting ²²²Rn flux but not GDR

Our analysis of the correlation between ²²²Rn and terrestrial GDR showed that both parameters are affected similarly by the radionuclide content of the soil and by soil 10 moisture. However, there are also factors affecting ²²²Rn flux without having a similar effect on GDR which we have not evaluated so far. Total pore space and tortuosity are important variables that affect ²²²Rn flux (Nazaroff, 1992) but not GDR. A larger proportion of ²²²Rn produced within the soil profile will escape to the atmosphere from coarse grained soils with a large total pore volume than from compacted fine grained soils, whereas the escape of gamma rays is unlikely to be affected by this. There already exist models for ²²²Rn flux prediction based on geological and pedological factors, but such models require numerous parameters which are not well known due to the complicated interactions between different geological and pedological units influencing the ²²²Rn flux (lelsch et al., 2002). Temperature differences between air and soil have also 20 been found to be a factor influencing ²²²Rn flux (Nazaroff, 1992), which is driven by diffusion and possibly mass flow.

4.4 Verification on a regional scale

As mentioned in the introduction, our interest in describing the ²²²Rn flux term is because of its application in the validation of atmospheric transport models. We therefore would like to be able to correctly predict regional averages of ²²²Rn flux. To test our ap-



proach of using GDR as a proxy, we split our data in one part to produce the correlation function between ²²²Rn flux and GDR (Switzerland, Germany, Scotland) and another part to verify the correlation (N- and S-Finland, Hungary). The correlation function derived was: y = 0.995x-14.97 (r²=0.66), where y is the ²²²Rn flux in Bq m⁻² h⁻¹ and x is the GDR in nSv h⁻¹. The measured regional means differed by a factor of up to 3, as considered not to be uncommon by Schery and Wasiolek (1998). Still, predicted means were within the error margin of the respective measured mean (Table 1).

5 Conclusions

Most of the spatial variation in ²²²Rn flux may be explained by the variation in radionu clide activity in soils derived from different parent material. Soil moisture has been shown to have similar effects on ²²²Rn flux as it has on GDR, except for short time periods during precipitation events. Considering additional parameters besides GDR, e.g. soil type, might further improve the prediction of ²²²Rn fluxes on the small scale. However, it may also unnecessarily complicate prediction, especially if we are going
 to extend it to areas where required data may not be available. To predict average regional ²²²Rn flux, the empirical correlation with GDR seems to suffice to produce regional means of ²²²Rn flux within the error margin of measurements.

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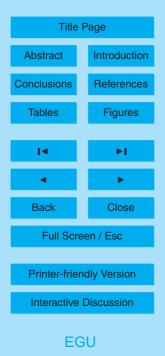
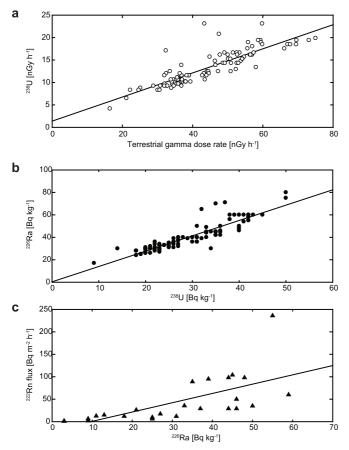


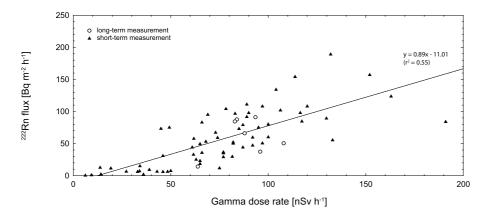
Table 1. Verification of the model in Finland and Hungary for regional mean values of measured and predicted ²²²Rn flux.

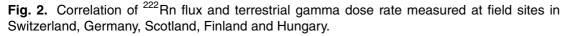
	²²² Rn flux measured	²²² Rn flux predicted	n
	$100 \pm 17 \mathrm{Bq}\mathrm{m}^{-2}\mathrm{h}^{-1}$	$102 \pm 13 \mathrm{Bq}\mathrm{m}^{-2}\mathrm{h}^{-1}$	8
N-Finland		41 ± 6 Bq m ⁻² h ⁻¹	8
Hungary	$60 \pm 9 \mathrm{Bq}\mathrm{m}^{-2}\mathrm{h}^{-1}$	68 ± 3 Bq m ⁻² h ⁻¹	12

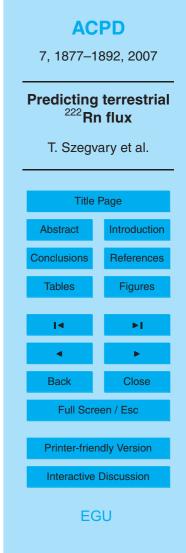


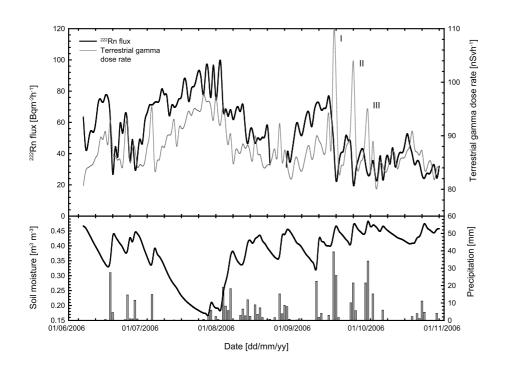












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Fig. 3. ²²²Rn flux, terrestrial gamma dose rate, precipitation and soil moisture time series from June to November 2006 in Basel (Switzerland). Heavy rain events are marked with I, II and III.