



HAL
open science

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

T. Szegvary, M. C. Leuenberger, F. Conen

► **To cite this version:**

T. Szegvary, M. C. Leuenberger, F. Conen. Predicting terrestrial ^{222}Rn flux using gamma dose rate as a proxy. *Atmospheric Chemistry and Physics*, 2007, 7 (11), pp.2789-2795. hal-00296241

HAL Id: hal-00296241

<https://hal.science/hal-00296241>

Submitted on 18 Jun 2008

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

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

T. Szegvary¹, M. C. Leuenberger², and F. Conen¹

¹Environmental Geosciences, Department of Geosciences, University of Basel, Switzerland

²Climate and Environmental Physics, Physics Institute, University of Bern, Switzerland

*Invited contribution by T. Szegvary, one of the EGU Outstanding Young Scientist Award winners 2006.

Received: 14 December 2006 – Published in Atmos. Chem. Phys. Discuss.: 5 January 2007

Revised: 23 April 2007 – Accepted: 23 May 2007 – Published: 6 June 2007

Abstract. ^{222}Rn is commonly used as a natural tracer for validating climate models. To improve such models a better source term for ^{222}Rn 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 ^{222}Rn 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 concentrations in soil forming minerals. A relatively stable fraction (20%) of the total terrestrial GDR originates from the ^{238}U decay series, of which ^{222}Rn is a member. Accordingly, spatial variation in terrestrial GDR was found to describe almost 60% of the spatial variation in ^{222}Rn flux. Furthermore, temporal variation in GDR and ^{222}Rn was found to be correlated. Increasing soil moisture reduces gas diffusivity and the rate of ^{222}Rn flux but it also decreases GDR through increased shielding of photons. Prediction of ^{222}Rn flux through GDR for individual measurement points is imprecise but un-biased. Verification of larger scale prediction showed that estimates of mean ^{222}Rn fluxes were not significantly different from the measured mean values.

1 Introduction

^{222}Rn is commonly known as a hazardous radioactive (noble) gas in indoor air. Yet, ^{222}Rn is also often used as a natural tracer of air transport. Observations of atmospheric ^{222}Rn have been very useful in the evaluation of climate models

Correspondence to: T. Szegvary
(t.szegvary@unibas.ch)

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 of changes in their atmospheric concentrations (Chevallard, 2002; Gupta et al., 2004). Currently, the effective use of ^{222}Rn in this context is limited by the poor accuracy of the ^{222}Rn source function (WMO GAW report no. 155, 2004). Current practice is to assume a spatial and temporal uniform flux rate of $1 \text{ atom cm}^{-2} \text{ s}^{-1}$ from all ice-free land surfaces. Improvement of the source term was attempted by Schery and Wasiolek (1998), who created a global ^{222}Rn 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 ^{222}Rn flux from ice-free land surface from $1 \text{ atom cm}^{-2} \text{ s}^{-1}$ at 30° N to $0.2 \text{ atom cm}^{-2} \text{ s}^{-1}$ at 70° N . This source term was found to improve predictions but it was speculated that ^{222}Rn 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 ^{222}Rn source term used.

Therefore, we are proposing a new method to describe the ^{222}Rn source term, initially focusing on the European continent. Our approach is to calibrate direct measurements of ^{222}Rn flux against terrestrial gamma dose rate (GDR). We

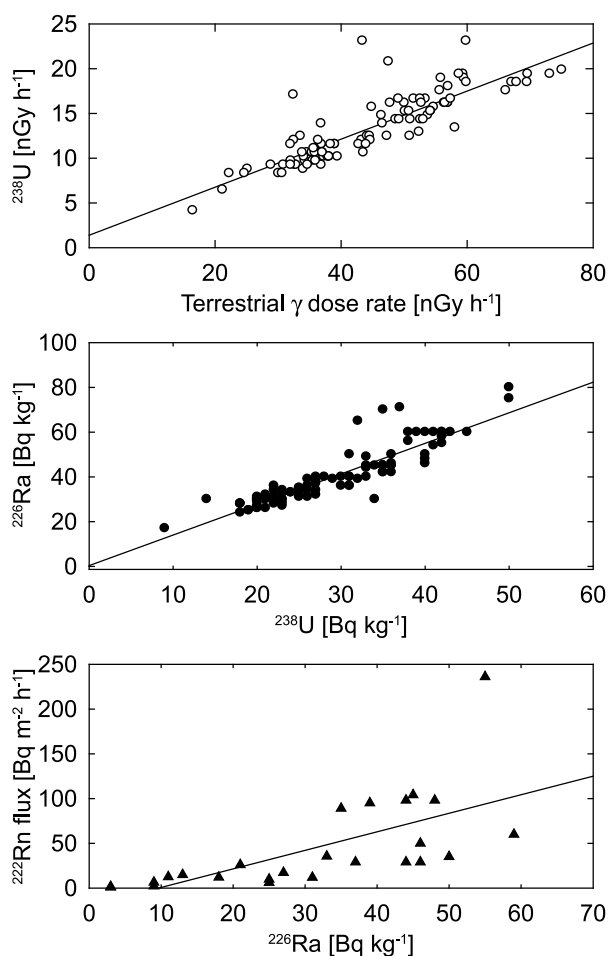


Fig. 1. Correlations between the contribution of GDR originating from the ^{238}U decay series and total terrestrial gamma dose rate (top), ^{226}Ra activity and ^{238}U activity (middle) and ^{222}Rn flux at the soil surface and soil ^{226}Ra activity (bottom). Data from the first two diagrams was kindly provided by SUER (Section of Surveillance of Radioactivity, Switzerland).

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 ^{222}Rn source term.

2 Basic concept

The source of ^{222}Rn is ^{226}Ra , a member of the ^{238}U decay chain. Gamma spectroscopic analysis of soil surface samples (0–20 cm depth) in geologically diverse regions of Switzerland showed that ^{238}U contributes an almost constant proportion to the terrestrial GDR (Fig. 1a) and that ^{226}Ra activity is closely related to the ^{238}U 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 mem-

bers of the ^{238}U decay chain could lead to an over- or underestimated GDR-based ^{222}Rn flux in such locations. However, such cases seem to be rare, as seen in the close correlations in Figs. 1a and b.

The proportion of the contribution of the ^{238}U series to total gamma dose rate is also reported for North-West Italy in Chiozzi et al. (2002), for Spain in Quindos et al. (2004) and for Cyprus in Tzortzis et al. (2003). Contributions of the ^{238}U series for individual types of rocks reported in these four studies range from 12% to 90%. However, the average for each country or region ranges from 27% (Spain), 29% (North-West Italy) to 30% (Cyprus, Switzerland). Thus, in the context of our objective to predict larger scale averages for radon flux, it seems justified to assume a constant contribution of the ^{238}U series to the total gamma dose rate.

Therefore, we assume that ^{222}Rn flux resulting from the decay of ^{226}Ra 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 ^{222}Rn flux to ^{226}Ra activity (Fig. 1c). Firstly, only part of the produced ^{222}Rn 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 ^{222}Rn that may reach the atmosphere before decay. Thus, the proportion of ^{222}Rn produced that escapes into the atmosphere is variable and depends on factors other than ^{226}Ra content. Indeed, the emanation coefficient for radon can vary by a factor of 10. The magnitude of this variation is a question of scale. Greeman and Rose (1996) determined emanation coefficients for each horizon in 12 contrasting soil profiles in the North-East of the United States. Emanation coefficients ranged from 5.5% to 33% for individual horizons. However, average emanation coefficients for entire soil profiles only ranged from 13% to 29% and two-thirds of the soil profiles were in the narrow range between 18% and 22%. Hence, despite large differences at the small scale, emanation factors at larger scales seem to be within a narrow range.

3 Methods

3.1 ^{222}Rn flux measurement techniques

A barely modified closed chamber method as described in Lehmann et al. (2000, 2003) was used to measure the ^{222}Rn 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 ^{220}Rn 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 ^{220}Rn flux). From there, the air passed to

the detector where only ^{222}Rn was measured. The ^{222}Rn flux was estimated from the increase in ^{222}Rn activity measured in 10 min intervals over about 1.5 h. Remaining ^{220}Rn may have affected the absolute value of measured ^{222}Rn activity but not its increase over time, as ^{220}Rn 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 ^{222}Rn with a half-life time of 3.82 days the assumption of a linear increase of ^{222}Rn 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 ^{222}Rn 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 h to accumulate ^{222}Rn and was then opened for 2.5 h prior to the next measurement. A second analytical system was a manually closable chamber (a plastic box with the dimensions 35 cm \times 27 cm and 13 cm height) which was used for spot measurements. The instrument we used in our study was compared in 2003 (Lynette Robertson, PhD Thesis, University of Edinburgh, 2005) to an instrument which has been widely used in East-Asia. The mean flux determined at six locations was $52 \text{ Bq m}^{-2} \text{ h}^{-1}$ (standard error $9 \text{ Bq m}^{-2} \text{ h}^{-1}$) with our instrument and compared well with the mean flux of $49 \text{ Bq m}^{-2} \text{ h}^{-1}$ (standard error $8 \text{ Bq m}^{-2} \text{ h}^{-1}$) measured with the instrument described in Iida et al. (1996).

Long-term measurements of ^{222}Rn 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 ^{222}Rn 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 ^{222}Rn system. The difficulty of spot measurements of ^{222}Rn 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 ^{222}Rn 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.

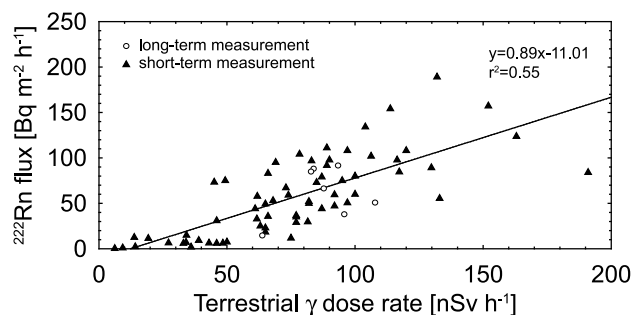


Fig. 2. Correlation of ^{222}Rn flux and terrestrial gamma dose rate measured at field sites in Switzerland, Germany, Scotland, Finland and Hungary.

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 (10H^*). 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 ^{137}Cs 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 ^{222}Rn flux and terrestrial GDR at different locations

The results of the measurement campaign are shown in Table 1, containing field site information and soil properties (all data concerning this research can also be found on the website <http://radon.unibas.ch>). There is a linear relationship between ^{222}Rn 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 $250 \text{ Bq m}^{-2} \text{ h}^{-1}$. Most soils in Europe have gamma dose rates between about 40 to 140 nSv h^{-1} well within this range. Very low GDR ($\sim 40 \text{ nSv h}^{-1}$ and a ^{222}Rn flux less than $15 \text{ Bq m}^{-2} \text{ h}^{-1}$) can be found at locations

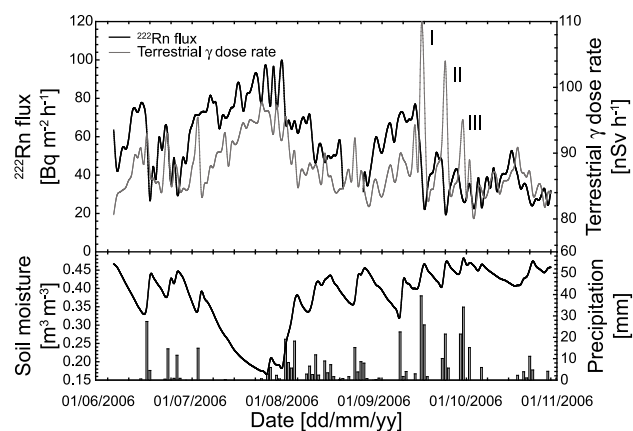


Fig. 3. ^{222}Rn flux, terrestrial gamma dose rate, precipitation and soil moisture time series from June to November 2006 in Basel (Switzerland). Heavy rain events are marked with I, II and III.

which have either a high water content and/or low or no mineral content like peat soils. Overall, almost 60% of the variation in ^{222}Rn 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 ^{222}Rn 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 ^{222}Rn flux is mostly influenced by ^{226}Ra 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 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 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 ^{222}Rn measurement, which we estimate to be around $\pm 15\%$ of the mean. For atmospheric tracer applications, regional information on the ^{222}Rn flux is required. The variability in the correlation between GDR and ^{222}Rn 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 ^{222}Rn flux and terrestrial GDR over time

Temporal variations in ^{222}Rn flux can be observed in GDR at the long-term measurement in Basel (Fig. 3) during the pe-

riod 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 ^{222}Rn 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, ^{222}Rn flux is larger when soils are dry and less ^{222}Rn decays 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 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 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 ^{222}Rn flux decreased immediately with the beginning of precipitation, probably because of the wet soil surface severely inhibiting ^{222}Rn diffusion into the atmosphere. The reaction of GDR was initially to the contrary. It suddenly increased after the first 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 ^{222}Rn progeny back to the soil surface (Greenfield, 2002, 2003). The cumulative half-life of the short-lived ^{222}Rn 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 ($\sim 8\text{--}10\%$). The second and third rain event showed the same effect. The only difference 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 ^{222}Rn progeny.

4.3 Factors affecting ^{222}Rn flux but not GDR

Our analysis of the correlation between ^{222}Rn and terrestrial GDR showed that both parameters are affected similarly by the radionuclide content of the soil and by soil moisture. However, there are also factors affecting ^{222}Rn 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 ^{222}Rn flux (Nazaroff, 1992) but not GDR. A larger proportion of ^{222}Rn 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,

Table 1. Measurement results of field sites in Switzerland, Germany, Hungary and Finland.

Field site information			^{222}Rn flux and GDR		Particle size fractions [%]			Moisture [wt%]
Long. [° E]	Lat. [° N]	Elevation asl [m]	^{222}Rn flux [Bq m ⁻² h ⁻¹]	Terr. GDR [nSv h ⁻¹]	Sand (63–2000 μm)	Silt (2–63 μm)	Clay (0–2 μm)	
Switzerland								
7.58	47.54	316	66*	88	5.9	71.4	22.7	21.3
7.88	47.43	610	14	57	25.7	55.4	18.9	17.6
7.88	47.43	610	14*	64	25.7	55.4	18.9	17.6
6.67	46.51	461	91*	93	30.9	54.0	15.1	
7.74	47.29	453	87*	84				
7.42	46.93	565	84*	83	45.2	37.1	17.7	
7.42	46.93	565	66	63	45.2	37.1	17.7	
6.92	46.33	381	37	96	5.5	82.4	12.1	
6.92	46.33	381	13	92	5.5	82.4	12.1	
7.84	46.30	640	50*	108	44.3	52.6	3.1	
6.58	46.84	1202	49	65	37.2	47.2	15.6	21.8
6.79	47.08	1018	67	73	6.3	70.3	23.4	25.8
6.23	46.40	430	92	89	25.2	49.3	25.6	15.3
9.84	46.81	1590	18	65	54.2	33.8	12.0	29.1
9.88	46.53	1705	37	77	39.6	46.5	13.9	25.5
10.07	46.34	1201	98	92	48.5	34.4	17.1	23.4
7.79	47.26	422	83	66	33.0	41.8	8.8	23.9
8.31	46.50	1345	61	100	62.1	34.3	3.6	18.4
8.90	47.48	536	96	69	28.6	49.4	22.0	24.9
9.40	47.43	779	44	61	33.5	45.7	20.8	34.1
9.07	47.03	515	74	59	8.7	52.8	38.5	41.3
9.52	47.13	460	39	66	20.5	70.1	9.4	26.8
8.46	47.06	1040	7	33	17.3	39.5	43.3	
7.64	47.59	268	109	105				
Germany								
7.81	47.76	850	157	155				
8.00	47.66	700	30	82				
8.14	47.59	300	33	61				
7.95	47.56	280	58	62				
7.78	47.56	350	98	116				
7.82	47.65	500	89	127				
Finland								
25.29	60.39	11	189	132	2.6	39.6	57.8	20.3
26.22	60.46	30	84	191	70.8	23.4	5.8	8.2
26.05	60.44	6	124	166	87.3	7.4	5.3	12.5
23.79	61.51	112	55	135	9.1	48.9	42.0	24.7
24.04	61.27	94	60	94	23.0	36.2	40.9	16.2
24.29	60.89	110	51	100	25.9	61.6	12.5	17.4
22.37	60.45	37	134	104	17.4	40.0	42.7	23.9
23.98	60.47	37	108	124	1.5	42.5	56.0	16.9
28.14	66.14	250	12	80	67.5	25.1	7.5	53.0
26.76	66.37	118	95	82	8.5	69.2	22.3	18.8
25.79	66.51	61	49	81	58.2	37.2	4.5	11.1
26.91	65.40	118	6	53	33.9	41.0	25.2	76.4
26.47	65.95	160	6	50	42.0	49.5	8.6	48.9
26.64	67.41	173	35	45	52.9	42.7	4.4	26.7
27.33	66.72	162	2	43	35.4	58.1	6.5	54.9
24.85	66.12	27	53	73	38.4	51.5	10.1	8.4
Hungary								
17.67	47.71	121	29	86	54.5	37.1	8.4	3.2
18.41	47.56	182	108	100	60.1	25.0	14.9	4.3
19.14	47.94	227	80	108	12.9	60.9	26.2	9.6
19.54	48.10	163	44	89	52.5	31.7	15.9	9.6
19.79	48.05	225	47	94	30.7	44.4	24.9	2.4
17.89	47.10	260	79	93	25.5	57.0	17.6	5.2
17.47	47.35	144	25	70	48.6	31.2	20.2	13.8
20.27	47.73	127	23	70	82.6	9.5	7.9	3.6
20.77	48.10	230	111	91				11.8
20.26	48.23	176	73	92	27.7	47.2	25.1	7.0
18.61	47.76	112	50	88	44.9	45.4	9.6	8.0
18.80	47.55	203	52	92	12.6	68.1	19.3	21.0

* Longterm measurements in Switzerland

Table 2. Verification of the model in Finland and Hungary for regional mean values of measured and predicted ^{222}Rn flux.

	^{222}Rn flux measured	^{222}Rn flux predicted	n
S-Finland	$100 \pm 17 \text{ Bq m}^{-2} \text{ h}^{-1}$	$102 \pm 13 \text{ Bq m}^{-2} \text{ h}^{-1}$	8
N-Finland	$32 \pm 11 \text{ Bq m}^{-2} \text{ h}^{-1}$	$41 \pm 06 \text{ Bq m}^{-2} \text{ h}^{-1}$	8
Hungary	$60 \pm 09 \text{ Bq m}^{-2} \text{ h}^{-1}$	$68 \pm 03 \text{ Bq m}^{-2} \text{ h}^{-1}$	12

whereas the escape of gamma rays is unlikely to be affected by this. There already exist models for ^{222}Rn 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 ^{222}Rn flux (Ielsch et al., 2002). Temperature differences between air and soil have also been found to be a factor influencing ^{222}Rn flux (Nazaroff, 1992), which is driven by diffusion and possibly mass flow. As for other possible correlations between environmental parameters and radon flux, we have tested for correlations with air temperature, atmospheric pressure, soil temperature and difference between air and soil temperature. If one of these parameters was correlated with radon flux, it was very weak. We do not think these parameters have a strong direct effect on radon flux but rather coincide with precipitation events or dry spells. In principle, diurnal pressure variations may cause mass flow through periodic expansion and contraction of the soil gas volume and influence the otherwise mainly diffusion-driven exchange of radon between soil pore space and atmosphere. We would expect this to be a major factor in deeply weathered dry soils with large air volumes. In the commonly humid regions in Europe we studied, it might not be a major issue.

4.4 Verification on a regional scale

As mentioned in the introduction, our interest in describing the ^{222}Rn 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 ^{222}Rn flux. To test our approach of using GDR as a proxy, we split our data in one part to produce the correlation function between ^{222}Rn flux and GDR (Switzerland, Germany, Scotland) and another part to verify the correlation (N- and S-Finland, Hungary). The correlation function derived was: $y = 0.995 (\pm 0.10)x + 14.97 (\pm 8.11)$ ($r^2 = 0.66$), where y is the ^{222}Rn flux in $\text{Bq m}^{-2} \text{ h}^{-1}$ 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 2), strongly supporting the effectiveness of our approach.

5 Conclusion

Most of the spatial variation in ^{222}Rn flux may be explained by the variation in radionuclide activity in soils derived from different parent material. Soil moisture has been shown to have similar effects on ^{222}Rn 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 ^{222}Rn 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 ^{222}Rn flux, the empirical correlation with GDR seems to suffice to produce regional means of ^{222}Rn flux within the error margin of measurements.

Acknowledgements. This project was funded by the Swiss National Science Foundation (project no. 200021-105123). We would like to thank the Swiss National Emergency Operations Center providing us with data for gamma dose rates, SUER (Section of Surveillance of the Radioactivity, Switzerland) for radionuclide databases in Switzerland and MétéoSuisse for meteorological data.

Edited by: M. Ammann

References

- Chiozzi, P., Pasquale, V., and Verdoya, M.: Naturally occurring radioactivity at the Alps-Apennines transition, *Radiation Measurements*, 35, 147–154, 2002.
- Conen, F. and Robertson, L. B.: Latitudinal distribution of radon-222 flux from continents, *Tellus*, 54B, 127–133, 2002.
- Chevillard, A., Ciais, P., Karstens, U., Heimann, M., Schmidt, M., Levin, I., Jacob, D., Podzun, R., Kazan, V., Sartorius, H., and Weingartner, E.: Transport of ^{222}Rn using the regional model REMO: a detailed comparison with measurements over Europe, *Tellus*, 54B, 850–871, 2002.
- De Cort, M. and de Vries, G.: The European Union Radiological Data Exchange Platform: Reference Manual and European Automatic Monitoring Systems, EUR 16415 EN, EC, Office for Official Publications of the European Communities, Luxembourg, EUR (2005), Automatic mapping algorithms for routine and emergency monitoring data, EUR 21595 EN, EC, edited by: Dubois, G., Office for Official Publications of the European Communities, Luxembourg, 1996.
- De Cort, M., Dubois, G., Fridman, Sh. D., Germenchuk, M. G., Izrael, Yu. A., Janssens, A., Jones, A. R., Kelly, G. N., Kvasnikova, E. V., Matveenko, I. I., Nazarov, I. M., Pokumeiko, Yu. M., Sitak, V. A., Stukin, E. D., Tabachnyi, L. Ya., Tsaturov, Yu. S., and Avdyushin, S. I.: Atlas of caesium deposition on Europe after the Chernobyl accident, EUR Report 16733, EC, Office for Official Publications of the European Communities, Luxembourg, 1998.
- Grasty, R. L.: Radon emanation and soil moisture effects on airborne gamma-ray Measurements, *Soc. Geophys.*, 62(5), 1379–1385, 1997.

- Greeman, D. J. and Rose, A. W.: Factors controlling the emanation of radon and thoron in soils of the eastern U.S.A., *Chemical Geology*, 129, 1–14, 1996.
- Greenfield, M. B., Domondon, A. T., Okamoto, N., and Watanabe, I.: Variation in γ -ray count rates as a monitor of precipitation rates, radon concentrations, and tectonic activity, *J. Appl. Phys.*, 91(3), 1628–1633, 2002.
- Greenfield, M. B., Domondon, A., Tsuchiya, S., and Tomiyama, G.: Monitoring precipitation and lightning via changes in atmospheric gamma radiation, application of accelerators in research and industry: 17th Int. conference, American Institute of Physics, 2003.
- Gupta, M. L., Douglass A. R., Kawa, R., and Pawson, S.: Use of radon for evaluation of atmospheric transport models: sensitivity to emissions, *Tellus Series B-Chemical and Physical Meteorology*, 56(5), 404–412, 2004.
- Ielsch, G., Ferry, C., Tymen, G., and Robe, M.-C.: Study of a predictive methodology for quantification and mapping of the radon-222 exhalation rate, *J. Environ. Radioactivity*, 63, 15–33, 2002.
- Iida, T., Ikebe, Y., Suzuki, K., Ueno, K., Wang, Z., and Jin, Y.: Continuous measurements of outdoor radon concentrations at various locations in East Asia, *Environ. Int.*, 22, 139–147, 1996.
- Lebedyte, M., Butkus, D., and Morkunas, G.: Variations of the ambient dose equivalent rate in the ground level air, *J. Environ. Radioactivity*, 64, 45–57, 2002.
- Lehmann, B. E., Lehmann, M., Neftel, A., and Tarakanov, S. V.: Radon-222 monitoring of soil diffusivity, *Geophys. Res. Lett.*, 27, 3917–3920, 2000.
- Lehmann, B. E., Ihly, B., Salzmann, S., Conen, F., and Simon, E.: An automatic static chamber for continuous ^{220}Rn and ^{222}Rn flux measurements from soil, *Radiation Measurement*, 38, 43–50, 2003.
- Murith, C. and Gurtner, A.: Mesures in situ et irradiation externe (French), in: BAG, Environmental radioactivity and radiation exposure in Switzerland 1993, Swiss Federal Office of Public Health, Berne, 1994.
- Nazaroff, W.: Radon transport from soil to air, *Rev. Geophys.*, 30(2), 137–160, 1992.
- Quindos Poncela, L. S., Fernández, P. L., Gómez Arozamena, J., Sainz, C., Fernández, J. A., Suarez Mahou, E., Martin Mataranz, J. L., and Cascón, M. C.: Natural gamma radiation map (MARNA) and indoor radon levels in Spain, *Environ. Int.*, 29, 1091–1096, 2004.
- Rasch, P. J., Feichter, J., Law, K., Mahowald, N., Penner, J., Benkovitz, C., Genthon, C., Giannakopoulos, C., Kasibhatla, P., Koch, D., Levy, H., Maki, T., Prather, M., Roberts, D. L., Roelofs, G.-J., Stevenson, D., Stockwell, Z., Taguchi, S., Kritz, M., Chipperfield, M., Baldocchi, D., McMurry, P., Barrie, L., Balkanski, Y., Chatfield, R., Kjellström, E., Lawrence, M., Lee, H. N., Lelieveld, J., Noone, K. J., Seinfeld, J., Stenchikov, G., Schwartz, S., Walcek, C., and Williamson, D.: A comparison of scavenging and deposition processes in global models: results from the WCRP Cambridge Workshop of 1995, *Tellus*, 52B, 1025–1056, 2000.
- Robertson, L. B.: Radon Emissions to the Atmosphere and their use as Atmospheric Tracers, University of Edinburgh, Ph.D. thesis, 2005.
- Robertson, L. B., Stevenson, D. S., and Conen, F.: Test of a northwards-decreasing Rn-222 source term by comparison of modelled and observed atmospheric Rn-222 concentrations, *Tellus*, 57B, 116–123, 2005.
- Schery, S. D. and Wasiolek, M.: Modeling radon flux from the earth's surface, in: Radon and Thoron in the Human Environment, edited by: Katase, A. and Shimo, M., World Scientific, Singapore, 207–217, 1998.
- Schery, S. D., Whittlestone, S., Hart, K. P., and Hill, S. E.: The flux of radon and thoron from Australian soils, *J. Geophys. Res.*, 94(D6), 8567–8576, 1989.
- Tzortzis, M., Tsertos, H., Christofides, S., and Christodoulides, G.: Gamma-ray measurements of naturally occurring radioactive samples from Cyprus characteristic geological rocks, *Radiation Measurements*, 37, 221–229, 2003.
- Von Gunten, H. R., Surbeck, H., and Rössler, E.: Uranium series disequilibrium and high thorium and radium enrichments in Karst formations, *Environ. Sci. Technol.*, 30(4), 1268–1274, 1996.
- World Meteorological Organization Global Atmosphere Watch: 1st International expert meeting on sources and measurements of natural radionuclides applied to climate and air quality studies, no. 155, 2004.