

Indian Journal of Pure & Applied Physics Vol. 61, October 2023, pp. 869-873 DOI: 10.56042/ijpap.v61i10.2800

Radon and Thoron Exhalation Rate Measurements in Soil Samples Collected from the Vicinity of a Thermal Power Plant

Pankaj Kumar & Mukesh Kumar^{*}

Department of Physics, S.V. College, Aligarh, Uttar Pradesh 202 001, India

Received 20 June 2023; accepted 31 July 2023

Radon mass and thoron surface exhalation rate of soil samples collected from the surrounding area of Harduaganj thermal power plant (HTPP), Aligarh is measured using an active radon/thoron monitor (SMART RnDuo). The radon mass exhalation rate values range from 27±1 to 100±3 mBq kg⁻¹ h⁻¹ with a mean value of 63±17 mBq kg⁻¹ h⁻¹. The thoron surface exhalation rate values vary from 1.0 ± 0.2 to 10 ± 1 kBq m⁻²h⁻¹ with a mean value of 5.7 ± 2.2 kBq m⁻²h⁻¹. The mean values of radon mass and thoron surface exhalation rate are found to be higher than the worldwide mean values.

Keywords: Soil samples; SMART RnDuo; Exhalation rates

1 Introduction

Radon exhalation measurements from soil and building material samples are important to know the radon risk to human populations¹. Radon and its progeny are the significant contributors to the average dose from natural background sources of radiation and constitute approximately half of the total dose to the general population. Epidemiologic studies in Europe and other case-control investigations in China and North America show that exposure to radon increases the risk of lung cancer²⁻⁴. Radon, a radioactive inert gas, escapes from grain as a result of recoil and if the recoil terminates in open pore space, radon becomes available to migrate⁵. Radon which is available in the interstitial space may be transported to the surface by diffusion and adjective flow and released into the atmosphere, referred to as radon exhalation⁶. Diffusion, which is usually the dominant transport mechanism, is caused by the radon concentration gradient

While advection takes place if there is a pressure difference between the air of the pore space and the ground surface⁷. The infiltration of radon gas from soil has been identified as one of the main mechanisms influencing indoor radon levels in buildings. It is reported that more than 90% of the contribution to indoor radon comes from the ground and surrounding soil of a building⁸. Under normal conditions, thoron concentrations in soil gas are roughly comparable to

the radon concentrations because of the similar production rates in rocks and soils and their similar behavior in the ground⁹. However, high thoron entry rates from the ground are rarely encountered. Because of its short half-life (55 sec), most of the thoron decays before reaching the surface.

Radon and thoron gases are members of the 238 U and 232 Th decay series, respectively, widely distributed in the earth's crust. Since both gases are members of different decay chains, their concentrations in the atmosphere depend on the 238 U and 232Th contents, respectively, in local soil and building materials. Various factors such as radium content, soil morphology, grain size, porosity, and radon flux density directly influence radon exhalation dynamics while meteorological parameters like temperature, pressure, and humidity indirectly affect the process of exhalation in soil $1^{10,11}$.

In the present study radon mass (J_m) and thoron surface (J_S) exhalation rates in soil samples collected from the surrounding area of about 90 km^2 around HTPP situated are measured using a continuous radon monitor (SMART RnDuo)¹².

2 Material and Methods

The installed capacity of HTPP (28°01'03''N 78° 07´48´´E) is 665 MW having seven generation units. Longitudes and latitudes of the study area around HTPP vary from 78.0925° to78.1810° and 27.9722 to 28.0510 , respectively. The soil of this area is sandy loam soil with brown or radish color.

^{*}Corresponding author: (E-mail: mksvc@rediffmail.com)

Fig. 1 — Samples locations around HTPP, Aligarh.

A total of 75 samples were collected from twenty-five villages. The locations of the soil samples taken for exhalation rate studies are shown in Fig. 1.

Technical specifications and experimental procedures for the measurement of J_m and J_s are discussed in detail elsewhere¹³. The SMART RnDuo is operated in diffusion mode with 1-hour cycles for the measurement of *Jm*.

The radon concentration C_{Rn} (t) buildup in the accumulation chamber at time *t* after closing the chamber can be calculated as per the following relation¹³

$$
C_{Rn}(t) = \frac{J_m M}{V \lambda_e} \left[1 - e^{-\lambda_e t} \right] + C_{Rn}(0) e^{-\lambda_e t} \qquad \dots (1)
$$

where $C_{Rn}(0)$ represents the initial radon concentration (Bq m^{-3}) present in the chamber volume at $t = 0$, J_m is the radon mass exhalation rate, measured in Bqkg⁻¹h⁻¹, *M* is the total mass of the dry sample, measured in kg, V is the residual air volume in the measurement system, which includes the detector volume of 150cc, the residual air volume of the exhalation chamber, and porous volume of the sample measured in m^3 , λ_e is the effective decay constant, which is the sum of the leak rate (if any) and the radon decay constant. The measurement time, *t* is measured in hours. If measurement time is limited to 12 hours, a linear approximation can be used and the Eq. (1) can be written as:

$$
C_{Rn}(t) = \frac{J_m M}{V} t + C_{Rn}(0) \qquad ...(2)
$$

Fig. 2 — Set up for the measurement of *Jm.*

By performing a least-square fitting of the data to Equation (2) mentioned above, the value of J_m can be determined using the slope obtained from the fitting, along with the mass (M) of the sample and the residual air volume (V) of the setup.

However, for the measurement of J_S , the SMART RnDuo is operated in a flow mode with 15-minute cycles. To calculate the thoron surface exhalation J_s (Bqm⁻²h⁻¹) rate of the sample, the following Equation (3) is used¹³

$$
J_S = \frac{c_{Th} V \lambda}{A} \qquad \qquad \dots (3)
$$

where C_{Th} is the equilibrium thoron concentration (Bqm⁻³). *V* is the residual air volume of the setup (m^3) . *A* is the surface area of the sample (m^2) , and λ is the decay constant $(44.87h^{-1})$ for thoron.

The accuracy in the estimation of J_m and J_s with this device is under 10% and 20% respectively. Figs 2 & 3 show the SMART RnDuo setup for the measurement of J_m and J_s , respectively.

Fig. 3 — Set up for the measurement of *JS.*

3 Results and Discussion

Global Positioning System (GPS) coordinates of soil samples and experimentally determined values of J_m and J_s along with their statistical values of parameters are given in Table 1.

The measured values of J_m are found to vary from 27 \pm 1 to 100 \pm 3 mBq kg⁻¹ h⁻¹. The median and the mean values with one slandered deviation are 60 mBq kg^{-1} h⁻¹ and 63±17mBq kg^{-1} h⁻¹, respectively. The observed wide variation in J_m may be due to the

radon emanation factor, and porosity of soil samples¹⁴. Sample H 26 of village Luhara has a maximum value of J_m (100 \pm 3 mBq kg⁻¹ h⁻¹) while the sample H 29 (27±1 mBq kg⁻¹ h⁻¹) of village Bhootpura has a minimum value of J_m as shown in Table 1.

The excess kurtosis is less than zero (-0.5) , showing platykurtic distribution. The skewness (0.5) is between -5 and $+5$, indicating that the distribution

is approximately symmetric. The values of J_S vary from 1.0 ± 0.2 to 10 ± 1 kBq m⁻²h⁻¹ with a mean value of 5.7 ± 2.2 kBq m⁻²h⁻¹. The median value is 5.8 kBq m⁻²h⁻¹. In the soil sample (H 53) collected from village Pilonathe value of J_S is found to be highest and is 10 ± 1 kBq m⁻²h⁻¹. The lowest level is 1.0 ± 0.2 kBq m⁻²h⁻¹ found in a soil sample (H 22) of village Kasimpur (Table 1).The value of excess kurtosis is found to be -0.4 which is less than zero, showing that

Fig. 4 — Distribution curve of J_m in the surrounding region of HTPP, Aligarh.

Fig. 5 — Distribution curve of J_S in the surrounding region of HTPP, Aligarh.

the nature of distributions is platykurtic. The value of skewness (-0.4) shows that the distribution is approximately symmetric.

The frequency distributions of J_m and J_s are given in Figs. 4 & 5, respectively. Notably, about 69% of the samples have a value of J_m , and about 79% of the samples have a value of J_S higher than worldwide mean values of 57 mBqkg⁻¹ h⁻¹ and 3.6 kBq m⁻² h⁻¹ for radon mass and thoron surface exhalation rate, respectively¹⁵.

Conclusions

The average value (63 mBq $kg^{-1} h^{-1}$) of J_m in the soil of the surrounding region of HTPP, Aligarh is found to be higher than the worldwide mean value of 57 mBqkg⁻¹ h⁻¹. The average value of J_S (5.7 kBqm⁻¹) ${}^{2}h^{-1}$) is also found to be higher than the worldwide mean value of 3.6 kBq m⁻² h⁻¹. The distribution of J_m and *Js* in the study area is found to be approximately symmetric.

Acknowledgment

The authors are thankful to the Board of Research in Nuclear Science (BRNS), Department of Atomic Energy, Government of India (Project Ref. No.: 2013/36/59-BRNS/2468) for providing financial assistance to carry out this study.

References

- 1 Nazaroff WW & Nero AV, *Radon and its decay product in indoor*, (John Wiley and Sons, New York), 1988.
- 2 Lubin J H, Wang Z Y, Boice J D, Xu Z Y, Bolt J W, De Wang L & Kleinerman A R, *Int J Cancer*, 109 (2004) 132.
- 3 Kerwski D, Lubin J H, Zielinski J M, Alavanja M, Catalan V S, Field R W, Klotz J B, Letourneau E G, Schoenberg J B, Steck D J, Stolwijk J N, Weinberg C & Wilcox H B, *Epidemiology*, 16 (2005) 137.
- 4 Darby S C, Hill D, Deo H, Auvinen A, Barros-Dios J M, Baysson H, Bochicchio F, Falk R, Farchi S, Figueiras A, Hakama M, Heid I, Hunter N, Kreienbrock L, Kreuzer M, Lagarde F, Makelainen I, Muirhead C, Oberaigner W, Pershagen G, Ruosteenoja E, Rosario AS, Tirmarche M, Tomasek L, Whitely E, Wichmann H E & Doll R, *Scand J Work Environ Health*, 32 (2006) 1.
- 5 UNSCEAR, United Nations Scientific Committee on the Effects of Atomic Radiation: Sources and effects ionizing radiation: Report to the General Assembly with Scientific Annexes, Annex B: Exposure from natural sources of radiation, United Nations, New York, 2000.
- 6 Ishimori Y, Lange K & Martin P, Technical reports series no. 474, International Atomic energy Agency, Vienna, 2013.
- 7 Sun K, Guo Q & Zhuo W, *J Nucl Sci Technol*, 41 (2004) 86.
- 8 Nero A V, Gadgil A J, Nazaroff W W & Revzan K L, Technical Report, U.S. Department of Energy, Office of Health and Environmental Research Washington, D C, 20545 (1990).
- 9 Hutter A R, *Environ Int J*, 22 (1996) 455.
- 10 Khan M S, Srivastava D S & Azam A, *Environ Earth Sci*, 67 (2012) 1363.
- 11 Sharma S, Kumar A, Mehra R & Mishra R, *J Soils Sediments*, 19 (2019) 1441.
- 12 Gaware J J, Sahoo B K, Sapra B K & Mayya Y S, *BARC News Lett*, 318 (2011) 45.
- 13 Sahoo B K, Sapra B K, Kanse S D, Gaware J J & Mayya Y S, *Radiat Meas*, 58 (2013) 52.
- 14 Sahoo B K, Agarwal T K, Gaware J J & Sapra B K, *J Radioanal Nucl Chem*, 302 (2014) 1417.
- 15 Prajith R, Rout R P, Kumbhar D, Mishra R, Sahoo B K & Sapra B K, *Environ Earth Sci*, 78 (2019) 35.