Dosimetry of radon progeny deposited on skin in air and thermal water

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ABSTRACT

It is held that the skin dose from radon progeny is not negligibly small and that introducing cancer is a possible consequence under normal circumstances as there are a number of uncertainties in terms of related parameters such as activity concentrations in air and water, target cells in skin, skin covering materials, and deposition velocities. An interesting proposal has emerged in that skin exposure to natural radon-rich thermal water as part of balneotherapy can produce an immune response to induce beneficial health effects. The goal of this study was to obtain generic dose coefficients with a focus on the radon progeny deposited on the skin in air or water in relation to risk or treatment assessments. We thus first estimated the skin deposition velocities of radon progeny in air and thermal water based on data from the latest human studies. Skin dosimetry was then performed under different assumptions regarding alpha-emitting source position and target cell (i.e. basal cells or Langerhans cells). Furthermore, the impact of the radon progeny deposition on effective doses from all exposure pathways relating to ‘radon exposure’ was assessed using various possible scenarios. It was found that in both exposure media, effective doses from radon progeny inhalation are one to four orders of magnitude higher than those from the other pathways. In addition, absorbed doses on the skin can be the highest among all pathways when the radon activity concentrations in water are two or more orders of magnitude higher than those in air.

Keywords: radon progeny; skin; deposition velocity; air; thermal water; alpha particle

INTRODUCTION

Radon (222Rn) is a natural radioactive gas that can exist anywhere in the environment and is well known to present a risk of lung cancer. In general, the pathway of so-called ‘radon exposure’ is the inhalation of its short-lived progeny rather than radon itself [1,2]. Arguments have also been put forward regarding other exposure pathways such as the inhalation and ingestion of radon gas [3–5], the skin permeability of radon gas [6,7], the skin deposition of radon progeny [8–12], and external exposure to radon progeny existing in air [13]. Most of these studies were focused on radiation protection, while others have been dedicated to promoting radon spa therapy research.

A fraction of the short-lived radon progeny (i.e. 218Po, 214 Pb, 214 Bi, and 214 Po) are attached to aerosol particles in the air (known as ‘attached fraction’), with the ‘unattached fraction’ remaining free ions. Both fractions can plate out onto any surface in the environment, including human lung airways and skin. A previous report indicated that in addition to the lungs, the deposition of the radon progeny may, under certain circumstances, lead to significant doses from alpha emitters to sensitive cells in the skin, that is, basal cell layers of between 50 and 100 μm in the skin surface [14]. Here, it has also been noted that if 70 μm is simply adopted as a nominal depth of the cell layer in accordance with the ICRP’s recommendation for practically assessing
equivalent doses to locally exposed skin [15], the skin doses calculated will be negligible due to the shorter ranges of the alpha particles in skin (ca. 45 μm for $^{218}$Po [6.002 MeV] and 65 μm for $^{214}$Po [7.687 MeV]).

Natural radon-rich water, air, and vapor in hot springs and thermal galleys have been traditionally and successfully used for the treatment of various diseases, including rheumatism and asthma [16,17]. However, the role played by radon and/or its progeny remains controversial, with the biological mechanisms triggered by very low radiation doses resulting from such radon exposure yet to be fully elucidated. Generally, therapeutic responses are supposed to be attributed to the radiation irritation to certain organs, tissues, and cells; however, an alternative explanation was proposed, that is, an immune response caused by irradiation to a deeper layer of the skin (e.g. Langerhans cells in the epidermis) with alpha particles from the radon progeny attached to the skin contributes to the effectiveness of that treatment [8].

For the assessment of skin doses under any conditions, the deposition velocity of the radon progeny on the skin is a key parameter that allows for predicting the skin surface activities. Two main papers have quantitatively represented the skin surface activities under given conditions, the first of which was compiled by Eatough et al. [12] and the second by Tempfer et al. [8]. In the former paper, 41 volunteers spent approximately one month wearing wristwatch alpha-particle detectors (apart from when showering or bathing) for individually monitoring the ambient activity concentrations of radon (a few to 400 Bq m$^{-2}$) as well as the skin surface activities of $^{224}$Po and $^{214}$Po. Meanwhile, in the latter paper, the skin surface activities of $^{218}$Po, $^{214}$Pb, $^{214}$Bi, and $^{214}$Po were repeatedly obtained from six volunteers during and after a 20 minute or 60 minute exposure to radon-rich thermal water (around 950 Bq l$^{-1}$ = 950,000 Bq m$^{-2}$). While both studies attempted to calculate the skin doses, there was some difficulty with arbitrary exposure conditions due to the lack of information on the deposition velocity, which means that it is unfeasible to evaluate the skin dose in cases where the individual radon progeny concentrations, the attached/unattached fraction or the exposure time are different from those in these studies.

To address this issue, we first devised a methodology to estimate and optimize the deposition velocities in both air and thermal water using published experimental data [8,12]. Following this, an integrated analysis of the skin deposition was conducted and is comprehensively discussed herein. The optimized values were then used for formulating skin doses for any exposure conditions. The doses were calculated in terms of various possible scenarios for comparisons between the exposure in air and water and with doses from other exposure pathways.

**MATERIALS AND METHODS**

Deposition model

The change in skin surface activity of the radon progeny $S_i$ (Bq m$^{-2}$) per time $t$ (s) can be expressed as follows [8,12,18]:

$$\frac{dS_{Po-214}}{dt} = C_{Po-214} \cdot P_{Po-214} - S_{Po-214} \lambda_{Po-214}$$

$$\frac{dS_{Bi-214}}{dt} = C_{Bi-214} \cdot P_{Bi-214} - S_{Bi-214} \lambda_{Bi-214} + (1 - k_\beta) S_{Bi-214} \lambda_{Po-214}$$

where $C_i$ (Bq m$^{-2}$) is the concentration in air or water, $v_i$ (m s$^{-1}$) is the deposition velocity, $\lambda_i$ (s$^{-1}$) is the decay constant, $k_\alpha$ or $k_\beta$ (–) is the fraction of desorption occurring after the alpha or beta decay of a parent nuclide, respectively and the subscript stands for a radionuclide ($i = 218$Po, $^{214}$Pb, $^{214}$Bi or $^{214}$Po) (see the explanation below regarding $v_i$, $k_\alpha$ and $k_\beta$). The fact that radioactive equilibrium is instantly established between $^{214}$Bi and $^{214}$Po due to the short half-life of $^{214}$Po (164.3 μs) is the basis of equation (4) and $C_{Bi-214} = C_{Po-214}$.

Since the parameter $v_i$ can be expected to be the same among all radon progeny [18], the relation of $v_{Po-218} = v_{Po-214} = v_{Bi-214}$ was assumed in the case of water; however, special attention was paid to the case of air. Here, the parameter $v_i$ was divided into two components: $v_{in}$ for the unattached fraction and $v_{it}$ for the attached fraction. Based on a previous work [18], the $v_{in}$ was assumed to be 100 times higher than the $v_{it}$. The parameters $C_{in}$ and $C_{it}$ were then defined accordingly: $C_i = C_{in} + C_{it}$. Thus, $v_i$ can be written as:

$$v_i = \frac{C_{in} v_{in} + C_{it} v_{it}}{C_{in} + C_{it}} = \frac{v_{in}}{C_{in}} \left( C_{in} + C_{it} \frac{100}{100} \right).$$

The parameter $k_\alpha$ or $k_\beta$, which can be in the range of 0–1, were incorporated into equations (2–3), based on the experimental findings: for example, if $k_\alpha$ or $k_\beta = 1$, $S_i$ is not influenced by $S_{Po-214}$ and is built up independently in every nuclide. In the case of air, the radon progeny ($i = ^{214}$Pb) can be desorbed from the material surface due to the alpha-recoil energy (much higher than the beta recoil energy) immediately after its generation by the alpha decay of the parent nuclide ($i = ^{218}$Po) [19]. In the case of water, the radon progeny ($i = ^{218}$Po or $^{214}$Bi) can be readily desorbed from the material surface immediately following its generation via the decay of the parent nuclide ($i = ^{218}$Po or $^{214}$Pb, respectively), irrespective of the type of decay [8,20,21]. In the present modeling, a value of 0, 0.5 or 1 was given to $k_\alpha$ or $k_\beta$ to evaluate the range of resulting doses. Based on the above information on desorption, different combinations of ($k_\alpha$, $k_\beta$) were adopted for the different media: i.e. (0, 0) or (0.5, 0) for air and (0, 0), (0.5, 0.5) or (1, 1) for water.

**Application of the deposition model to the measured data**

The data used for our model application were taken from Eatough et al. [12] for the exposure in air and from Tempfer et al. [8] for the thermal water. Both the previous experiments and our application methods are described below.

(1) In air:

A total of 41 individuals participated in the one-month monitoring test. The activity concentration of radon in air ($C_{Po-222}$) was also measured in parallel with $S_{Po-218}$ and $S_{Po-214}$ monitoring at the wrist part. The results indicated a linear relationship between $C_{Po-222}$ and $S_{Po-218}$.
Fig. 1. Buildup and decay of the surface activities ($S_i$) on the skin exposed to radon-rich water ($C_{\text{w,Rn-222}} = 950 \text{ Bq l}^{-1}$). The first 60 minutes was the exposure time via bathing, whereas the remaining time was simply the resting time without exposure. The approximate relative uncertainties of the measurement were in the range of 40–90% for $^{218}$Po, 10–20% for $^{214}$Pb, and 10–25% for $^{214}$Bi and $^{214}$Po. The curves were fitted to the empirical data from Ref. [8], assuming three desorption fractions: (a) $k_\alpha = k_\beta = 0$, (b) $k_\alpha = k_\beta = 0.5$, and (c) $k_\alpha = k_\beta = 1$. 

- **(a) Fitted with $k_\alpha = k_\beta = 0$**
- **(b) Fitted with $k_\alpha = k_\beta = 0.5$**
- **(c) Fitted with $k_\alpha = k_\beta = 1$**
or \( S_{\text{Po-214}} \). Only the data from 13 individuals were used here to obtain more appropriate values of \( S_i \) (Bq m\(^{-2}\)) per \( C_{\text{Ra-222}} \) (Bq m\(^{-1}\)), i.e. and 
\[ S_{\text{Po-218}} = 0.25 \pm 0.19 \text{ m and } S_{\text{Po-214}} = 0.22 \pm 0.11 \text{ m}. \]

Given that these measured values corresponded to the saturated values, the parameters \( \nu_a \), \( \nu_u \), and \( \nu_p (= \nu_u / 100) \) can be calculated under the steady state of 
\[ \frac{dC_{\text{Po-218}}}{dt} = \frac{dC_{\text{Po-214}}}{dt} = \frac{dC_{\text{Po-222}}}{dt} = 0 \text{ in equations } (1-3). \]

This should be performed for a variety of possible indoor activity concentrations of the radon progeny [22]. It is reasonable to suppose indoor conditions, since all the voluntary participants were medical staff and are expected to have high occupancy factors. Thus, individual activity concentrations of the radon progeny (\( C_{\text{iu}} \) and \( C_{\text{ui}} \): \( i = \text{Po-218, Po-214, Pb-214, Bi or Po-214} \)) that satisfy a certain equilibrium factor \( F \) of between 0.2 and 0.7 and an unattached fraction \( f_p \) of between 0.04 and 0.2, which were concluded as the parameter ranges of general indoor conditions in a recent international report [22], were randomly determined with 10,000 patterns under the following constraints: (a) \( C_{\text{Po-218,a}} > C_{\text{Po-214,a}} > C_{\text{Po-214,u}} \), (b) \( C_{\text{Po-218,u}} > C_{\text{Po-218,a}} > C_{\text{Po-214,a}} \), and (c) \( C_{\text{Po-214,u}} = C_{\text{Po-214,a}} = 0 \). Accordingly, 10,000 patterns of \( \nu_a \) and \( \nu_u \) were computed and summarized to ascertain their dependences on the assumption of \( F \) and \( f_p \).

In addition, the variability analysis of \( \nu_a \) or \( \nu_u \) was also performed based on the Monte Carlo simulation while considering uniform distributions of \( F \) (0.2–0.7) and \( f_p \) (0.04–0.2) and normal distributions of 
\[ 10 \pm 5 \text{ m and } 50 \pm 5 \text{ m}. \]

A total of six individuals underwent the exposure test in a thermal water bath. The activity concentration of radon in thermal water was relatively stable at \( C_{\text{Ra-222}} = 950 \pm 73 \text{ Bq l}^{-1} \), and the radon progeny was empirically verified to be in equilibrium with radon, that is, \( C_{\text{Ra-222}} = C_{\text{Po-218}} = C_{\text{Po-214}} = C_{\text{Po-222}} \) for each participant in several tests with different exposure times (10, 20, 30, 40, and 60 minutes). In the present work, the time-series data on \( S_{\text{Po-218}}, S_{\text{Po-214}}, S_{\text{Po-222}}, \) and \( S_{\text{Po-214}} \) at the forearm part over 60 minutes were utilized for better evaluation. That is, the model of equations (1–3) was fitted to the plots in Fig. 1 to estimate \( \nu_v \).

Dosimetry

The depth-dose distribution was first computed for alpha particles isotropically emitted from \( ^{218}\text{Po} \) or \( ^{214}\text{Po} \) using the particle and heavy ion transport system (PHITS) code [23]. The transport of alpha particles in a tissue-equivalent cube (\( 10 \times 10 \times 10 \text{ cm}^3 \)) was simulated, and the absorbed dose was calculated by dividing the energy imparted between the depth \( d \) and \( d + 1 \mu m \) by the mass corresponding to the volume of \( 1 \text{ cm} 	imes 1 \text{ cm} 	imes 1 \mu m \). The elemental composition of the skin with a density of \( 1.1 \text{ g cm}^{-3} \) was taken from ICRP [24]. The source was assumed to be positioned on the skin surface, whereas in the case of the water, the skin penetration of radon progeny, which was previously indicated in Tempfer et al. [8], was also considered (Fig. 2).

Radiation doses to given targets that had to be set depending on the assessment purpose were then calculated with the following depth-dose distributions: (i) basal cell layer from the viewpoint of radiation risk, and (ii) Langerhans cell layer from the treatment viewpoint. The basal cells are deemed to be potentially at risk from radiation and are assumed to be positioned at the bottom layer of the epidermis [24], whereas Langerhans cells are immune cells that could play a role in indicating a positive response in the radon spa therapy [8] and are assumed to be distributed uniformly in the epidermis. Two forms of probability distribution of the basal cell layer in depth were taken from the existing literature [12,25,26] (see Fig. 2), the first of which was a relatively realistic model developed on an empirical basis (epidermis model 1), while the second was a simple model proposed for the practice of radiation protection (epidermis model 2).

Finally, absorbed dose rates to the target cells, \( D (\text{mGy h}^{-1} \text{ (Bq m}^{-3} \text{,}^{220}\text{Rn})^{-1}) \), were determined as follows:

\[ D = \sum_{d=1}^{r} P_j D_d \text{ for basal cells} \quad (6) \]

\[ D = \sum_{j=1}^{r} \left( \frac{P_j}{j} \sum_{d=1}^{r} D_d \right) \text{ for Langerhans cells} \quad (7) \]

where \( r (\mu m) \) is the range of alpha particles in the skin, \( P (-) \) is the probability of the target cells existing at the depth \( d \) or \( j (\mu m) \), and \( D_d \) is the absorbed dose rate at the depth \( d (\mu m) \) that is available from the depth-dose distribution as calculated above.

RESULTS

Skin deposition velocity in air and water

Figure 3 shows the variation in \( \nu_v \) or \( \nu_u \) (100\( \nu_v \)) for the exposure in air as a function of \( f_p \) between \( F = 0.2 \) and 0.7. Under the same \( F \) and \( f_p \),
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Fig. 3. Variation of the calculated skin deposition velocity of the radon progeny in air ($v_{ia}$ or $v_{iu}$) as a function of the assumed unattached fractions ($f_p$). The calculation was performed for $F = 0.2$ to $0.7$, based on the skin surface activity ($S_{Po-218}$ or $S_{Po-214}$) measured by Eatough et al. [12].

conditions, the plots of $v_a$ and $v_u$ calculated from the measured $S_{Po-218}$ (Fig. 3a) were generally two to four times higher than those from the measured $S_{Po-214}$ (Figs 3b,c). In addition, the plots of $v_a$ and $v_u$ calculated from the measured $S_{Po-214}$ at $k_\alpha = 0$ (Fig. 3c) were around 20% greater than those at $k_\alpha = 0.5$ (Fig. 3b), representing relatively similar values of $v_a$ and $v_u$ regardless of $k_\alpha$. This finding confirmed that in addition to $F$ and $f_p$, the selection of the measured $S_{Po-218}$ or $S_{Po-214}$ was also crucial to assessing the $v_a$ and $v_u$. Figure 4 shows the comparison of the variability in $v_a$ or $v_u$ calculated from the measured $S_{Po-218}$ in relation to that from $S_{Po-214}$. In general, the explanation for the results shown in Fig. 3 also holds for those shown in Fig. 4.

Figure 1 shows the fitted curves of $S_i$ to the measured data for the exposure in thermal water. The $S_i$ values increased over time during the bathing with radon exposure for 1 h, with only $S_{Po-218}$ saturated within a shorter time due to the short half-life of $^{218}$Po (3.10 minutes), before the $S_i$ values then decreased during the resting without exposure. Given its plot trend, one could suggest that $S_{Po-218}$ reaches a maximum followed by decrease during the bathing; however, we accepted the idea of the saturation of $S_{Po-218}$ as the relative uncertainties of the $^{218}$Po measurement were high (ca. 90% at 10 minutes, 40% at 20 and 30, 75% at 40 and 60% at 60) and a plausible mechanism producing such trend was unknown and difficult to be incorporated to the deposition model. Given the residual sum of squares, the best fit of $S_{Po-218}$, $S_{Pb-214}$ and $S_{Bi-214}$ was the case of $k_\alpha = k_\beta = 0$, and the next was that of $k_\alpha = k_\beta = 0.5$. The estimated $v_i$ was in the range of 0.021–0.028 m h$^{-1}$, indicating that $k_\alpha$ and $k_\beta$ did not significantly influence the determination of $v_i$.

The estimated values of $v_i$ for both exposure in air and in water are summarized in Table 1.
Fig. 4. Comparison of the variability in the skin deposition velocity of the radon progeny in air (\(v_{sa}\) or \(v_{su}\)) calculated from the measured \(S_{Po-218}\) and \(S_{Po-214}\). It should be noted that the bin ranges were different among the curves.

Table 1. Estimated skin deposition velocity of radon progeny

| Exposure medium | Desorption fraction (\(-\)) | Deposition velocity (m h\(^{-1}\)) | Deposition velocity (m h\(^{-1}\)) | Deposition velocity (m h\(^{-1}\)) |
|----------------|----------------------------|---------------------------------|---------------------------------|---------------------------------|
| Air\(^{b}\)    | \(k_{\alpha} = 0; k_{\beta} = 0\) | \(v_{sa} = 0.18 (0.27 \pm 0.29)\) | \(v_{su} = 18 (27 \pm 29)\) | Estimated from the measured \(S_{Po-214}\) \(v_{sa} = 0.040 (0.052 \pm 0.042)\) | \(v_{su} = 4.0 (5.2 \pm 4.2)\) | Estimated from both of the measured \(S_{Po-218}\) and \(S_{Po-214}\) \(v_{sa} = 0.075 (0.16 \pm 0.24)\) | \(v_{su} = 7.5 (16 \pm 24)\) |
| Water          | \(k_{\alpha} = k_{\beta} = 0\) | \(v_{sa} = 0.028\) | \(v_{su} = 0.024\) | \(v_{su} = 0.024\) | \(v_{su} = 8\) |
| \(k_{\alpha} = k_{\beta} = 0.5\) | \(v_{sa} = 0.024\) | \(v_{su} = 0.021\) | \(v_{su} = 0.024\) |
| \(k_{\alpha} = k_{\beta} = 1\) | \(v_{sa} = 0.08\) | | |

\(^{a}\)The value of \(v_{sa}\) or \(v_{su}\) for the medium of air corresponds to a median, with an arithmetic mean ± standard deviation in parentheses (see Fig. 4).

\(^{b}\)The estimation of \(v_{sa}\) and \(v_{su}\) based on the Monte Carlo simulation was performed under the conditions of \(F = 0.45 \pm 0.14\) and \(f_p = 0.12 \pm 0.05\).

\(^{c}\)The single value of \(v_{sa}\) or \(v_{su}\) for both cases of \(k_{\alpha} = 1\), and 0.5 was obtained since its calculation did not rely on the parameter \(k_{\alpha}\).
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DISCUSSION

Comparison of skin deposition velocities

The analysis derived $v_{sa} = 0.08$ and $v_{mu} = 8 \text{ m h}^{-1}$ for air and $v_i = 0.024 \text{ m h}^{-1}$ for water as representative values. The deposition velocity in water was closer to that for the attached fraction in air. It should be noted here that the conditions of the subjects in the two human studies [8,12], the data of which were used to calculate the present deposition velocities, can be expected to be quite different.

Fig. 5. Depth-dose distribution computed for alpha particles from $^{218}\text{Po}$ or $^{214}\text{Po}$ existing (a) on the skin surface or (b) from the surface to a depth of around 20 $\mu$m. The probability distribution of the deposited radon progeny in depth in terms of the case of exposure in water is presented in Fig. 2.

Fig. 6. Absorbed dose to a specific target cell layer as a function of time exposed in water. The epidermis models, i.e. the probability distribution of epidermis in depth, are presented in Fig. 2. The source was assumed to exist (a) on the skin surface, and (b) from the surface to 20 $\mu$m (Fig. 2). The fitted lines are numerically expressed as dose coefficients in Table 2.
Table 2. Evaluated absorbed dose rate to a specific target cell layer in the skin

| Exposure medium | Deposition velocity (m h$^{-1}$) | Desorption fraction (−) | Source position | Target cell | Absorbed dose rate (nGy (Bq m$^{-3}$ h Rn$^{222}$)$^{-1}$) |
|-----------------|----------------------------------|-------------------------|-----------------|-------------|----------------------------------------------------------|
| Air$^a$         | $v_{u} = 0.08; v_{a} = 8$ | $k_{u} = 0; k_{\beta} = 0$ | Surface | Langerhans cell | Model 1: 14.341, Model 2: 9.716 |
|                 |                                  |                         |                 | Basal cell   | Model 1: 4.811, Model 2: 0.536 |
| Water$^b$       | $v_{i} = 0.024$                | $k_{u} = 0; k_{\beta} = 0$ | Surface | Langerhans cell | Model 1: 1.033, Model 2: 0.718 |
|                 |                                  |                         |                 | Basal cell   | Model 1: 0.394, Model 2: 0.0514 |
|                 | Surface to 20 μm                |                         |                 | Langerhans cell | Model 1: 1.259, Model 2: 0.908 |
|                 |                                  |                         |                 | Basal cell   | Model 1: 0.560, Model 2: 0.122 |

$^a$In accordance with the ICRP [27] and ICRU [22], the ratios of activity concentrations were assumed to meet the representative indoor conditions ($F = 0.4$ and $f_{p} = 0.08$). For the attached fraction, $C_{Bi-214,a} = C_{Bi-214,u}; C_{Pb-214,a} = C_{Pb-214,u}; C_{Po-214,a} = C_{Po-214,u} = 1; 0.51: 0.3825: 0.306: 0.306$; for the unattached fraction, $C_{Bi-214,a} = C_{Bi-214,u}; C_{Pb-214,a} = C_{Pb-214,u}; C_{Po-214,a} = C_{Po-214,u} = 1: 0.2: 0.02: 0.0$. $^b$The slopes of the fitted lines in Fig. 6 were equivalent to the absorbed dose rates.

For example, maybe the subjects for the bathing exposure test were sitting calmly for 60 minutes, while those for the exposure test in air were working and living as usual for a month. In short, fluid or human movement can influence the friction velocity on a surface, resulting in an increase in the chance of deposition by a factor of 3–10 [28]. Despite this influence, our findings may remain practically reasonable with regard to the similarity between $v_{u}$ in air and $v_{i}$ in water; however, the attendant reason and mechanism cannot be explained from previously obtained experimental data.

It should also be noted that the present study assumed that $v_{u} = 100 v_{u,i}$, which was based on Porstendörfer [18]. This assumption was useful and practical for the modeling but was unclear in terms of accuracy. The uncertainty of this assumption, in addition to certain other environmental parameters, could have influenced our estimation, leading to a discrepancy in the estimated $v_{u}$ or $v_{i}$ (by a factor of around 2) depending on the data source (i.e., $S_{Po-218}$, $S_{Po-214}$, or both $S_{Po-214}$ and $S_{Po-214}$) (Table 1). Further research on this topic is required to quantify the relationship between $v_{u}$ and $v_{i}$ and to improve the estimation of the skin deposition velocity of radon progeny. In addition, the impacts of aerosol particle size and air movement on the skin deposition of the radon progeny must also be better understood since the deposition velocity depends on such parameters [18,29]. The evaluation of these impacts will, in turn, contribute to improved skin dose quantification with some consideration of the environmental conditions.

The values of $v_{u} = 0.08$ and $v_{a} = 8$ m h$^{-1}$ for the skin exposed to air were compared with those reported for specific materials (e.g., grass, filter paper and metal). Porstendörfer [18] comprehensively collated and analyzed experimental data on the deposition of radon and thoron progenies in a general room, estimating the $v_{u,i}$ value to be 0.02 and the $v_{a}$ value to be 2 m h$^{-1}$ (Table 1). This analysis was performed by considering the deposition velocities as a function of particle size, for a room with low ventilation (<0.3 h$^{-1}$), with the assumption of the surface roughness of filter paper. Knutson [29] also provided a summary of the deposition velocity results, which presented representative average values of 0.08 and 8 m h$^{-1}$ for $v_{u}$ and $v_{a}$ respectively. To make a reasonable comparison between these earlier reports and this article, some attention must be given to the movement of fluid as described above. Nevertheless, it can be stated that the estimated deposition velocity to the skin will not differ greatly from that to materials such as paper and grass, when taking into account that air motion, equivalent to human motion, can enhance the deposition velocity by, at most, one order of magnitude [29].

**How significantly does skin-permeating radon affect the activity measurement of the radon progeny directly deposited on the skin?**

Since the intake of radon through skin from thermal water was numerically computed in a previous work [7], it was important to guarantee that there was no significant interference from the skin-permeating radon, followed by its decay, with the activity measurement of radon progeny directly deposited on the skin (Fig. 1). Without this guarantee, an implicit agreement could not be reached on how the measured data resulted solely from the skin deposition of radon progeny in water.

By means of biokinetic modeling dedicated to the skin absorption of radon [7], we quantified the buildup of radon and its progeny activity concentrations (Bq kg$^{-1}$) in the skin compartment during bathing in thermal water (Fig. 7). This model calculation was performed under the following conditions: $C_{Rn,222} = 950$ Bq l$^{-1}$ and skin permeability coefficient $K = 1.5 \times 10^{-6}$ m s$^{-1}$ for an adult male subject. The value of K used here corresponded to the 95th percentile of K estimated for adult males, meaning that the calculation was on the overestimated side. The calculated activity concentrations were converted to the skin surface activities (Bq cm$^{-2}$) by considering the approximated total (epidermis + dermis) skin mass thickness at the upper/lower arms and legs of adult males (i.e. $1.4 \times 10^{-3}$ kg cm$^{-2}$) [24].

Finally, a comparison between Figs. 1 and 7 allowed us to conclude that in the previous experiment, the skin-permeating radon did not significantly influence the activity measurement of radon progeny deposited on the skin. Thus, the earlier measurement data taken for the present study were valid for use without correction with regard to radon permeability.

**Dose assessment**

Table 3 presents the effective doses for radon and its progeny exposure calculated using various given scenarios and parameter conditions. To
### Table 3. Calculated effective doses from pathways relating to radon and its progeny exposures under given scenarios and parameter conditions

| Scenario and condition | Effective dose (nSv (Bq m\(^{-3}\))\(^{-1}\)) |
|------------------------|---------------------------------|
|                        | Radon exposure                  | Radon progeny exposure          |
|                        | Inhalation, \(E_{Rn,inha}\)    | Skin permeability, \(E_{Rn,skin}\) | Inhalation, \(E_{Rn-prog,inha}\) | Skin deposition, \(E_{Rn-prog,skin}\) |
| Living environment     | \(9.7 \times 10^{-2}\)         | N.A.                            | \(3.6\)                          | \(1.1 \times 10^{-1}\)                  |
| Thermal bath           | \(3.2 \times 10^{-2}\)         | \(3.2 \times 10^{-4}\)          | \(1.2\)                          | \(3.4 \times 10^{-3}\)                  |

- **Medium**: Air
- **1-h exposure**
- \(F = 0.4; f_p = 0.08\) (See the footnote of Table 2)
- \(v_{sa} = 0.08 \text{ m h}^{-1}; v_{sa} = 8 \text{ m h}^{-1}\)
- \(k_\alpha = 0; k_\beta = 0\)
- \(K = 2.1 \times 10^{-7} \text{ ms}^{-1} \text{ (median)} \) [7]

### Fig. 7
Biokinetic modeling calculation of the buildup of the calculated surface activities of radon and its progeny originating from the skin permeability of radon during bathing in water (\(C_{Rn,Rn-222} = 950 \text{ Bq l}^{-1}\)).

![Graph](https://example.com/graph.png)

Calculate an effective dose from the absorbed dose, we used a radiation weighting factor of 20 for alpha particles and a tissue weighting factor of 0.01 for skin or 0.12 for lung [15]. Common values were used for \(F, f_p,\) and \(K,\) with the representative values concluded in this article used for \(v_i\) and the values that conservatively provide the doses used for \(k_\alpha\) and \(k_\beta.\) The basal cells were set as the target and were assumed to be distributed as epidermis model 2 (Fig. 2), since this calculation targeted an effective dose that is meant to be used in relation to the risks of stochastic effects (e.g. cancer). The results clearly indicated that, for both exposure media, the effective doses from the radon progeny inhalation were much higher than those from the other pathways, i.e., by an order of magnitude in the case of air, and by two orders of magnitude (radon inhalation and radon progeny skin deposition) or four orders of magnitude (radon skin permeability) in the case of water.

Here, it should be noted that if the skin is covered with clothing or other materials, the skin deposition of the radon progeny would be suppressed so that the effective dose is reduced. As such, the doses presented in Table 3 can be qualitatively regarded as overestimated. At the same time, attention must be given to the case of exposure in water due to the possibility of a significant difference in radon activity concentration in air and water, e.g., radon spas may have a few hundred or thousands of Bq m\(^{-3}\) in air and a few hundred or thousands of kBq m\(^{-3}\) (= a few hundred or thousands of Bq l\(^{-1}\)) in water [7]. If the radon activity concentrations in a radon spa location are plausibly assumed to be 1,000 Bq m\(^{-3}\) for air and 100 kBq m\(^{-3}\) (= 100 Bq l\(^{-1}\)) for water, the ratio of \(E_{Rn,inha}: E_{Rn,skin}: E_{Rn-prog,inha}: E_{Rn-prog,skin}\) is 0.032: 0.032: 1.2: 0.34. Thus, it can be concluded that the skin deposition of
radon progeny may induce significant effective doses in certain specific cases and radon progeny inhalation is the dominant exposure pathway in daily living environments.

Thermal bathing (balneotherapy) and the resulting absorbed doses to the skin from the therapeutic viewpoint should also be considered. If the exposure time is 20 minutes and the activity concentrations of radon in air and water are 1,000 Bq m$^{-3}$ and 100 kBq m$^{-3}$ (=100 Bq l$^{-1}$), the absorbed dose to Langerhans cells can, according to Table 2, be estimated to be in the range of 24–42 μGy, while, according to Table 3, the absorbed dose to the lung (i.e. target cells relating to lung cancer) resulting from radon progeny inhalation can be estimated to be 0.5 μGy. The skin dose was found to be higher than the lung dose, while the aim of selecting the target cells is different in skin dosimetry than in lung dosimetry. This information could be valuable in scrutinizing biological/positive responses from the radon spa therapies that remain controversial and the mechanisms of which have yet to be clarified.

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CONFLICT OF INTEREST

The authors declare no conflicts of interest.

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