



Bias-corrected $H_p(10)$ -to-Organ-Absorbed Dose Conversion Coefficients for the Epidemiological Study of Korean Radiation Workers

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ABSTRACT

Background: The effects of radiation on the health of radiation workers who are constantly susceptible to occupational exposure must be assessed based on an accurate and reliable reconstruction of organ-absorbed doses that can be calculated using personal dosimeter readings measured as $H_p(10)$ and dose conversion coefficients. However, the data used in the dose reconstruction contain significant biases arising from the lack of reality and could result in an inaccurate measure of organ-absorbed doses. Therefore, this study quantified the biases involved in organ dose reconstruction and calculated the bias-corrected $H_p(10)$ -to-organ-absorbed dose coefficients for the use in epidemiological studies of Korean radiation workers.

Materials and Methods: Two major biases were considered: (a) the bias in $H_p(10)$ arising from the difference between the dosimeter calibration geometry and the actual exposure geometry, and (b) the bias in air kerma-to- $H_p(10)$ conversion coefficients resulting from geometric differences between the human body and slab phantom. The biases were quantified by implementing personal dosimeters on the slab and human phantoms coupled with a Monte Carlo method and considered to calculate the bias-corrected $H_p(10)$ -to-organ-absorbed dose conversion coefficients.

Results and Discussion: The bias in $H_p(10)$ was significant for large incident angles and low energies (e.g., 0.32 for right lateral at 218 keV), whereas the bias in dose coefficients was significant for the posteroanterior (PA) geometry only (e.g., 0.79 at 218 keV). The bias-corrected $H_p(10)$ -to-organ-absorbed dose conversion coefficients derived in this study were up to 3.09-fold greater than those from the International Commission on Radiological Protection publications without considering the biases.

Conclusion: The obtained results will aid future studies in assessing the health effects of occupational exposure of Korean radiation workers. The bias-corrected dose coefficients of this study can be used to calculate organ doses for Korean radiation workers based on personal dose records.

Keywords: Conversion Coefficient, Organ-Absorbed Dose, Bias Quantification, Monte Carlo Method, Epidemiological Study

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Introduction

Radiation workers are constantly susceptible to occupational exposure. Thus, the assessment of the radiation effect on health has become important because of increasing

concerns about protracted exposure. Previous studies generally used the personal dose equivalent $H_p(10)$ to evaluate the health effects associated with occupational exposure. Similar to other countries, the occupational exposure of Korean radiation workers is managed in terms of $H_p(10)$, measured by personal dosimeters such as the thermoluminescent dosimeter (TLD), and the measured $H_p(10)$ values are reported every 3 months to the national dose registry.

However, to evaluate cancer morbidity and mortality, the International Commission on Radiological Protection (ICRP) strongly recommends the use of an organ-absorbed dose rather than $H_p(10)$ [1]. Unfortunately, it is impossible to directly measure organ-absorbed doses because dosimeters should not be inserted into the human body. For this reason, organ-absorbed doses for radiation workers should be derived from $H_p(10)$ values using dose conversion coefficients. The ICRP has provided the dose conversion coefficients for air kerma-to- $H_p(10)$ and air kerma-to-organ-absorbed doses calculated based on a 30 cm × 30 cm × 15 cm slab phantom (mass composition: 76.2% oxygen, 11.1% carbon, 10.1% hydrogen, and 2.6% nitrogen) and anthropomorphic phantoms coupled with Monte Carlo methods [2, 3], respectively. Accordingly, the organ-absorbed dose can be calculated by converting $H_p(10)$ using the aforementioned dose conversion coefficients.

However, several biases affect the reliability and validity of the risk assessment of occupational exposure in the reconstruction of the organ-absorbed dose. First, a bias in the recorded $H_p(10)$ occurs during the calibration and reading of personal dosimeters. Although personal dosimeters are practically exposed to radiation from various directions in addition to having different angular dependencies depending on the incident angle, the exposure geometry, except the anteroposterior (AP), is ignored in the existing calibration and reading procedure. In addition, the slab phantom used for calibration is not similar to the human body, where an actual personal dosimeter is positioned. Therefore, the accuracy of the recorded $H_p(10)$ varies with exposure conditions and calibration geometry. The calibration procedure of a personal dosimeter can introduce bias in the recorded $H_p(10)$ because the angular dependency and the differences between the slab phantom and the human body are not considered.

Furthermore, an important bias is introduced while deriving organ-absorbed doses from recorded $H_p(10)$ through the ICRP air kerma-to- $H_p(10)$ conversion coefficients. The ICRP

air kerma-to- $H_p(10)$ conversion coefficients are derived from a cuboid slab phantom. However, in actual practice, $H_p(10)$ is measured using a personal dosimeter worn on a radiation worker's body, not a slab phantom. Therefore, differences exist between the actual measurement geometry of $H_p(10)$ and the slab phantom-based calibration geometry that are used to calculate the conversion coefficients. These differences introduce bias in the estimation of air kerma from recorded $H_p(10)$ and consequently affect the derivation of organ-absorbed doses. For example, for exposure from the postero-anterior (PA) direction, the attenuation thickness for the human body is typically expected to be higher than 200 mm, whereas that for a slab phantom is only 140 mm, according to the definition of $H_p(10)$. In this case, although the recorded $H_p(10)$ is measured after being attenuated by the human body, only the attenuation by the slab phantom is considered in the air kerma calculations. Therefore, the calculated air kerma is underestimated, which means that the organ-absorbed dose is underestimated. For these reasons, to improve the validity of the estimated organ-absorbed doses, it is necessary to analyze biases that affect dose conversions.

In this study, we attempted to quantify the biases introduced into the recorded $H_p(10)$ and subsequently into the slab phantom-based conversion coefficients (air kerma-to- $H_p(10)$) to finally derive the organ-absorbed dose for radiation workers. For this purpose, a Monte Carlo simulation of the dosimeter measurement coupled with the slab and the anthropomorphic phantoms was performed.

Materials and Methods

1. Background of Organ-Absorbed Dose Reconstruction

An organ-absorbed dose can be reconstructed from recorded $H_p(10)$ as follows:

$$D_T = H_p(10) \left[\frac{H_p(10)}{K_a} \right]^{-1} \left[\frac{D_T}{K_a} \right], \quad (1)$$

where D_T is the tissue or organ-absorbed dose (Gy); $H_p(10)$ is the personal dose equivalent at a depth of 10 mm (Sv); $H_p(10)/K_a$ is the air kerma-to- $H_p(10)$ conversion coefficient (Sv/Gy); and D_T/K_a is the air kerma-to-organ-absorbed dose conversion coefficient (Gy/Gy).

The two conversion coefficients (i.e., $H_p(10)/K_a$ and D_T/K_a) are determined according to the exposure scenarios (energy and geometry) to which radiation workers are exposed. ICRP $H_p(10)/K_a$ values calculated based on a slab phantom according to the photon energy and the incident angle are pro-

vided in Table A.24 of ICRP Publication 74 [2]. D_T/K_a can be calculated based on simulations using computational human phantoms. ICRP has provided reference D_T/K_a values calculated based on the ICRP reference voxel phantom [3]. Computation was performed according to idealized geometries: AP, PA, right lateral (RLAT), left lateral (LLAT), rotational (ROT), and isotropic (ISO) [3]. The cranial-caudal and caudal-cranial geometries introduced in the US National Council on Radiation Protection and Measurements (NCRP) report were also considered in this study [4].

2. Bias in Organ Dose Reconstruction

Two types of biases were considered in this study: the bias B_1 in the recorded $H_p(10)$ and the bias B_2 in $H_p(10)/K_a$. B_1 arises because the response of the personal dosimeter under the current calibration conditions (i.e., the irradiation in the AP direction and the use of the slab phantom) may differ that under the actual exposure conditions. Therefore, B_1 can be quantified using the ratio of the dosimeter response under the irradiation on the anthropomorphic phantom at the incident angle α to that under the irradiation on a slab phantom at 0° . In addition, the difference in reference $H_p(10)$ (i.e., the value used as the correction factor for calibration) introduced by the differences between the calibration geometry (AP) and actual exposure geometry is also considered.

$$B_1 = \left[\frac{R(E, \alpha)_{\text{human}}}{R(E, 0^\circ)_{\text{slab}}} \right] \left[\frac{H_p(10; E, \alpha)_{\text{human}}}{H_p(10; E, 0^\circ)_{\text{slab}}} \right]^{-1}, \quad (2)$$

where $R(E, \alpha)_{\text{human}}$ denotes the dosimeter response on the anthropomorphic phantom irradiated at an incident angle α ; $R(E, 0^\circ)_{\text{slab}}$ denotes the dosimeter response on the slab phantom irradiated at 0° (AP); $H_p(10; E, \alpha)_{\text{human}}$ denotes the $H_p(10)$ values depending on the actual exposure energy and geometry; and $H_p(10; E, 0^\circ)_{\text{slab}}$ denotes the $H_p(10)$ values for AP geometry, which have been provided by ICRP. More specifically, to quantify the response R with Monte Carlo simulation, R was defined as the absorbed dose delivered to the element of the personal dosimeter per unit of air kerma.

Another bias, B_2 , can also be quantified by comparing the conversion coefficients based on the anthropomorphic and slab phantoms. Because this bias involved in the dose conversion coefficients results from the geometric differences between the human body and the slab phantom, B_2 is defined as follows:

$$B_2 = \left[\left(\frac{H_p(10)}{K_a} \right)_{\text{human}} \right] \left[\left(\frac{H_p(10)}{K_a} \right)_{\text{slab}} \right]^{-1}, \quad (3)$$

where $(H_p(10)/K_a)_{\text{human}}$ is the anthropomorphic phantom-based $H_p(10)/K_a$, and $(H_p(10)/K_a)_{\text{slab}}$ is the slab phantom-based $H_p(10)/K_a$, which have been provided by ICRP.

3. Monte Carlo Simulation for Bias Estimation

In the quantification of the biases, we used the Korean Typical Man-2 (KTMAN-2) computational phantom constructed from high-resolution anatomical images of Koreans to minimize the physical differences between Korean radiation workers and the anthropomorphic phantom. KTMAN-2 is a computed tomography image-based voxel phantom consisting of $300 \times 150 \times 344$ voxels with a resolution of $2 \text{ mm} \times 2 \text{ mm} \times 5 \text{ mm}$ [5]. KTMAN-2 was implemented based on the Monte Carlo N-Particle 6 (MCNP 6) code. The field size was defined as $50 \text{ cm} \times 60 \text{ cm}$ for AP, PA, RLAT, and LLAT geometries and $60 \text{ cm} \times 80 \text{ cm}$ for the cranial-caudal and caudal-cranial geometries; the size was considered sufficient to approximately model whole-body irradiation. In the LAT geometry (i.e., an incident angle of 90°), because a depth of 10 mm cannot be defined, the $H_p(10)$ value cannot be defined. Therefore, the response was calculated with an incident angle of 75° [4]. For the same reason, the cranial-caudal and caudal-cranial geometries were also exposed at 75° . The responses for ROT and ISO geometries were calculated by averaging those for four geometries (AP, PA, LLAT, and RLAT) and six geometries (AP, PA, LLAT, RLAT, cranial-caudal, and caudal-cranial), respectively.

To calculate B_1 from the relative response of a TLD, personal dosimeters on the slab and KTMAN-2 phantom (Fig. 1) were modeled after the Panasonic TLD UD-802 model (Panasonic Corporation, Tokyo, Japan), which is commonly used in radiation facilities in Korea. TLD UD-802 measured $H_p(10)$ using a 0.05-mm thick thermoluminescent material (CaSO_4) and a $1 \text{ g} \cdot \text{cm}^{-2}$ filter made of lead and plastic. The absorbed doses in the CaSO_4 element were calculated using the energy deposition (type F6) tally for AP, PA, RLAT, LLAT, cranial-caudal, and caudal-cranial geometries. Thus, the relative response was calculated as the ratio of the absorbed dose in the CaSO_4 element on the KTMAN-2 to that on the slab phantom. To confine the statistical error of the simulation result within 5%, 10^8 – 10^9 histories were used. The position of the personal dosimeter worn on the front was selected as 14.4 cm below the thyroid gland and 4 cm left from the midline, where it is commonly worn [6].

The bias estimation was performed for energies of 218, 397, and 662 keV, suggested as representative exposure ener-

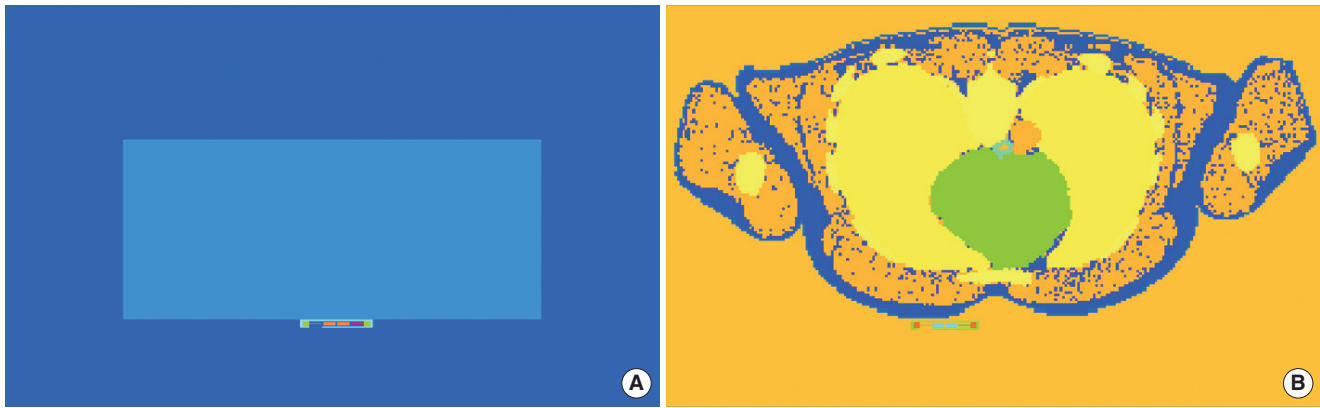


Fig. 1. Transverse plane images of personal dosimeter implemented in Monte Carlo N-Particle 6 (MCNP6) code for the calculation of B_1 with the slab phantom (A) and with Korean Typical Man-2 (KTMAN-2) (B).

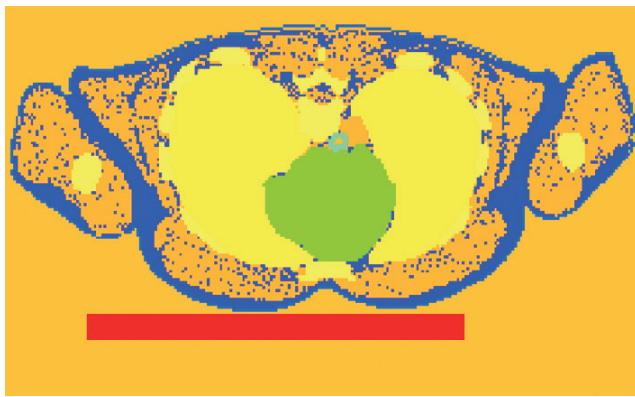


Fig. 2. Transverse plane image of cuboid tissue equivalent material (red) and Korean Typical Man-2 (KTMAN-2) implemented in Monte Carlo N-Particle 6 (MCNP6) code for the calculation of $H_p(10; E, \alpha)_{\text{human}}$.

gies for Korean radiation workers [7]. The use of these energies facilitates proper interpolation of the biases over a wide energy range.

$H_p(10; E, \alpha)_{\text{human}}$ in equation (2) was calculated based on a tissue equivalent material of $30 \text{ cm} \times 30 \text{ cm} \times 2 \text{ cm}$ positioned on the chest of the KTMAN-2. The cuboidal material is the same as that of the slab phantom, and its thickness was selected in consideration of the filter thickness of TLD. The absorbed dose was tallied in a cylindrical cell (thickness of 0.01 cm; diameter of 1 cm) defined at a depth of 10 mm (Fig. 2).

To estimate B_2 , $H_p(10)$ was calculated in the same manner as B_1 (Fig. 2), and the air kerma values based on energy were calculated using the fluence-to-air kerma conversion coefficients provided in Table A.1 of the ICRU Report 47 [8]. B_2 was calculated as the ratio of the $H_p(10)/K_a$ obtained in this simulation to that provided in the ICRP Publication 74 [2].

Table 1. Results of Verification for the Bias Estimation Method

Energy (MeV)	ICRP 74 (A)	This study (B)	Ratio (B/A)
0.03	1.11	1.13	1.01
0.04	1.49	1.51	1.01
0.05	1.77	1.79	1.01
0.06	1.89	1.90	1.00
0.08	1.90	1.89	0.99
0.2	1.49	1.51	1.01
0.4	1.30	1.30	1.00
0.6	1.23	1.23	1.00
0.8	1.19	1.19	1.00
1	1.17	1.16	1.00
10	1.11	1.07	0.96

ICRP, International Commission on Radiological Protection.

4. Calculation of Bias-Corrected $H_p(10)$ -to-Organ-Absorbed Dose Conversion Coefficients

For use in epidemiological studies, bias-corrected $H_p(10)$ -to-organ-absorbed dose conversion coefficients are derived by considering the two types of biases in this study and are expressed as follows:

$$\frac{D_T}{H_p(10)} = \frac{1}{B_1 B_2} \left[\left(\frac{H_p(10)}{K_a} \right)_{\text{slab}} \right]^{-1} \left[\frac{D_T}{K_a} \right], \quad (4)$$

where $D_T/H_p(10)$ is the $H_p(10)$ -to-organ-absorbed dose conversion coefficient and D_T/K_a is the air kerma-to-organ-absorbed dose conversion coefficient. We used $(H_p(10)/K_a)_{\text{slab}}$ values provided in the ICRP Publication 74 and D_T/K_a values calculated using mesh-type reference computational phantoms (MRCPs) developed based on the ICRP reference anatomical data [2, 9].

Table 2. Biases Calculated by Monte Carlo Simulation Depending on Energy and Geometry

	Energy (keV)	Bias							
		AP	PA	RLAT	LLAT	Cranial-caudal	Caudal-cranial	ROT	ISO
B_1	218	1.04	0.68	0.32	0.35	0.31	0.31	0.60	0.50
	397	1.01	0.79	0.79	0.85	0.81	0.82	0.86	0.84
	662	1.02	0.85	0.97	1.03	0.97	0.97	0.97	0.97
B_2	218	0.98	0.79	1.00	0.98	0.99	0.95	0.94	0.95
	397	0.99	0.83	1.01	0.99	1.00	0.98	0.96	0.97
	662	1.00	0.88	1.01	0.98	1.01	1.00	0.96	0.98

AP, anteroposterior; PA, posteroanterior; RLAT, right lateral; LLAT, left lateral; ROT, rotational; ISO, isotropic.

Results and Discussion

1. Simulation Verification

To verify the bias estimation method, the $H_p(10)$ values on the slab phantom were calculated using MCNP 6, and $H_p(10)/K_a$ values were compared with those of ICRP Publication 74 [2]. The $H_p(10)$ values were calculated in a cylindrical cell with a thickness of 0.01 cm and diameter of 1 cm, in the same manner as the bias calculations. The radiation field size was selected as 50 cm × 60 cm, and the fluence was 3.3×10^{-4} cm⁻². The photon energies ranged from 0.03 keV to 10 MeV, and 10^8 histories were considered. The $H_p(10)/K_a$ from this study and ICRP are presented in Table 1, and the differences were within 5% for all energies.

2. Quantified Biases

The quantified biases are shown in Table 2. The B_1 values, depending on the incident angle, were calculated considering the angular dependency of the personal dosimeter and differences between the slab and KTMAN-2 phantom. In AP geometry, B_1 for all energies is approximately 1, which means that the geometric differences between the two phantoms are not significant. For the AP geometry (i.e., 0°), the angular dependency is not considered in the quantification of B_1 . However, although $H_p(10; E, \alpha)_{\text{human}}$ values are typically smaller than $H_p(10; E, 0^\circ)_{\text{slab}}$ values for all energies and geometries, the B_1 values for lower energies were observed to be noticeably smaller than 1, except for the AP geometry. In addition, the magnitude of bias (i.e., the difference from 1) is larger at low energies; for example, B_1 for RLAT was 0.32 and 0.97 at 218 and 662 keV, respectively. This is because the attenuation of the incident radiation in TLD is larger for longer attenuation distances and lower energies, resulting in a relatively low response. The radiation entering the lead and plastic filters should travel longer distances as the incident angle in-

creases. In the case of ROT and ISO geometries, the B_1 values, calculated by weighting those of other geometries, were considerably larger than those for the 75° directions (i.e., RLAT, LLAT, cranial-caudal, and caudal-cranial) because of the effects of the AP geometries.

B_2 is close to 1 for all energies and geometries except for the PA geometry, i.e., 0.79 and 0.88 at 218 keV and 662 keV, respectively. As expected, the attenuation thickness of KTMAN-2 was higher than 200 mm, whereas that of the slab phantom was only 140 mm. Therefore, the absorbed dose per unit of air kerma calculated based on KTMAN-2 was smaller than that calculated based on the slab phantom. Consequently, the calculated B_2 was also smaller in the PA geometry.

3. $H_p(10)$ -to-Organ-Absorbed Dose Conversion Coefficients

For use in epidemiological studies, a library of bias-corrected $H_p(10)$ -to-organ-absorbed dose conversion coefficients, $D_T/H_p(10)$, was created for three energy points, six geometries, and 30 organs (28 organs for each sex) using the biases derived in the current study and the dose coefficients provided by ICRP. The full data are tabulated in Appendices A and B. $D_T/H_p(10)$ values for energy points between the energies considered in this study can be derived by interpolation. Tables 3 and 4 show example dose conversion coefficients for adult male and female phantoms for five major organs with a tissue weighting factor of 0.12 (i.e., red bone marrow, colon, lungs, stomach, and breasts). Although most organs are exposed to the highest doses in the AP direction, the bias-corrected $D_T/H_p(10)$ values were greatest in the PA direction due to the significantly greater biases.

The practical impact of the biases on dose reconstruction can be assessed by $D_T/H_p(10)$. Ratios of $D_T/H_p(10)$ derived in the current study to those from ICRP publications without

Table 3. $D_T/H_p(10)$ Conversion Coefficients with Biases Considered: Male

Organs	Energy (keV)	$D_T/H_p(10)$ (Gy/Sv)					
		AP	PA	RLAT	LLAT	ROT	ISO
Bone marrow	218	0.598	3.474	1.496	1.388	1.342	1.270
	397	0.654	2.390	0.640	0.594	0.926	0.759
	662	0.688	1.882	0.557	0.534	0.833	0.685
Colon	218	0.809	2.515	1.755	1.948	1.339	1.269
	397	0.830	1.762	0.721	0.790	0.932	0.746
	662	0.832	1.517	0.629	0.675	0.830	0.676
Lungs	218	0.704	3.171	1.249	1.273	1.284	1.284
	397	0.753	2.244	0.567	0.568	0.932	0.797
	662	0.798	1.807	0.519	0.533	0.840	0.725
Stomach	218	0.742	2.301	0.934	2.620	1.329	1.220
	397	0.793	1.683	0.455	1.047	0.888	0.727
	662	0.807	1.383	0.426	0.876	0.799	0.644
Breasts	218	0.936	1.371	2.085	2.020	1.491	1.615
	397	0.972	1.218	0.853	0.832	1.022	0.968
	662	0.976	1.144	0.716	0.713	0.912	0.842

AP, anteroposterior; PA, posteroanterior; RLAT, right lateral; LLAT, left lateral; ROT, rotational; ISO, isotropic.

Table 4. $D_T/H_p(10)$ Conversion Coefficients with Biases Considered: Female

Organs	Energy (keV)	$D_T/H_p(10)$ (Gy/Sv)					
		AP	PA	RLAT	LLAT	ROT	ISO
Bone marrow	218	0.632	3.499	1.606	1.492	1.389	1.324
	397	0.687	2.403	0.684	0.641	0.951	0.791
	662	0.715	1.890	0.596	0.570	0.855	0.706
Colon	218	0.829	2.694	1.841	1.524	1.391	1.276
	397	0.863	1.957	0.771	0.665	0.951	0.778
	662	0.859	1.569	0.658	0.599	0.851	0.682
Lungs	218	0.678	3.412	1.383	1.380	1.372	1.333
	397	0.735	2.468	0.615	0.611	0.964	0.828
	662	0.751	1.943	0.555	0.563	0.889	0.758
Stomach	218	0.782	2.805	1.117	2.903	1.440	1.342
	397	0.810	2.038	0.504	1.147	0.984	0.791
	662	0.831	1.654	0.483	0.926	0.882	0.721
Breasts	218	0.934	1.872	2.193	2.117	1.509	1.592
	397	0.964	1.551	0.928	0.891	1.047	0.949
	662	0.964	1.376	0.784	0.770	0.939	0.835

AP, anteroposterior; PA, posteroanterior; RLAT, right lateral; LLAT, left lateral; ROT, rotational; ISO, isotropic.

Table 5. Ratios of $H_p(10)$ -to-Organ-Absorbed Dose Conversion Coefficients, $D_T/H_p(10)$, Derived in This Study to Those from ICRP Publications

Geometry	Ratio (this study/ICRP)		
	218 keV	397 keV	662 keV
AP	0.98	1.00	0.98
PA	1.86	1.53	1.34
RLAT	3.09	1.26	1.03
LLAT	2.93	1.19	1.00
ROT	1.78	1.22	1.07
ISO	2.10	1.23	1.06

ICRP, International Commission on Radiological Protection; AP, anteroposterior; PA, posteroanterior; RLAT, right lateral; LLAT, left lateral; ROT, rotational; ISO, isotropic.

considering the biases are shown in Table 5. Note that the differences in dose conversion coefficients arise only from the biases, and thus the ratios are equal in all organ dose coefficients. For the AP direction, the biases in the dose coefficients do not have a significant impact on the dose reconstruction. Nevertheless, the ratios (this study/ICRP) vary by exposure scenario and range from 0.98 to 3.09 for three energies and six geometries. The greatest difference in $D_T/H_p(10)$ was observed in the case of the exposure energy of 218 keV and RLAT geometry (3.09), indicating that the consideration of the biases derived in this study can result in 3.09-fold higher organ doses than those calculated using ICRP refer-

ence dose coefficients without considering the biases.

The differences in $D_T/H_p(10)$ have a significant implication for dose reconstructions for Korean industrial radiographers, whose cumulative $H_p(10)$ is highest among all Korean radiation workers [10]. In practice, an industrial radiographer who places an image film can be exposed with PA direction because the acquisition of radiographic images usually begins on the way back to the shielded area (this information was gathered in a personal conversation). In this case (i.e., 397 keV and PA direction), organ doses calculated considering the biases can be 1.53-fold higher than those without considering the biases.

Conclusion

In this study, two types of biases in organ-absorbed dose reconstruction were quantified using Monte Carlo simulations. The study demonstrated that the bias B_1 , caused by personal dosimeters and calibration geometries, increases when the incident angle is large or the exposure energy is low, and B_2 is approximately 1, except for the PA geometry, which has a large difference in attenuation thickness between anthropomorphic and slab phantoms. The findings of this study indicate that the biases in $H_p(10)$ and dose coefficients ($H_p(10)/K_a$) can result in a significant underestimation of a radiation worker's dose and thus should be considered in the dose reconstruction. Moreover, this study might be used as a framework for the detailed analysis and reliable assessment of the organ-absorbed dose from occupational exposure. The application of the quantified biases to other epidemiological studies, based on new scenarios and methods, will widen the prospects of this study and is therefore of significant interest for future research.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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Author Contribution

Conceptualization: Jeong A, Kwon TE. Methodology: Jeong A, Kwon TE, Lee W. Formal analysis: Jeong A, Kwon TE. Project administration: Park S. Visualization: Jeong A. Writing - original draft: Jeong A. Writing - review and editing: Kwon TE. Approval of final manuscript: all authors.

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Appendix A. Bias-Corrected $D_r/H_p(10)$ Conversion Coefficients for Male

Organs	Energy (keV)	$D_r/H_p(10)$ (Gy/Sv)						Organs	Energy (keV)	$D_r/H_p(10)$ (Gy/Sv)					
		AP	PA	RLAT	LLAT	ROT	ISO			AP	PA	RLAT	LLAT	ROT	ISO
Bone marrow	218	0.598	3.474	1.496	1.388	1.342	1.270	Extrathoracic region	218	0.697	2.165	3.039	2.622	1.646	1.474
	397	0.654	2.390	0.640	0.594	0.926	0.759		397	0.782	1.615	1.173	1.143	1.115	0.848
	662	0.688	1.882	0.557	0.534	0.833	0.685		662	0.789	1.407	0.981	0.927	1.000	0.826
Colon	218	0.809	2.515	1.755	1.948	1.339	1.269	Gallbladder	218	0.699	2.382	2.785	0.951	1.285	1.180
	397	0.830	1.762	0.721	0.790	0.932	0.746		397	0.728	1.749	1.105	0.452	0.882	0.702
	662	0.832	1.517	0.629	0.675	0.830	0.676		662	0.742	1.468	0.893	0.442	0.806	0.636
Lungs	218	0.704	3.171	1.249	1.273	1.284	1.284	Heart	218	0.775	2.645	1.306	1.803	1.319	1.261
	397	0.753	2.244	0.567	0.568	0.932	0.797		397	0.805	1.906	0.597	0.766	0.907	0.753
	662	0.798	1.807	0.519	0.533	0.840	0.725		662	0.817	1.563	0.548	0.676	0.821	0.682
Stomach	218	0.742	2.301	0.934	2.620	1.329	1.220	Kidneys	218	0.443	4.081	1.481	1.347	1.355	1.202
	397	0.793	1.683	0.455	1.047	0.888	0.727		397	0.500	2.731	0.627	0.572	0.920	0.715
	662	0.807	1.383	0.426	0.876	0.799	0.644		662	0.544	2.094	0.551	0.515	0.825	0.646
Breasts	218	0.936	1.371	2.085	2.020	1.491	1.615	Lymph nodes	218	0.773	2.954	1.603	1.766	1.426	1.324
	397	0.972	1.218	0.853	0.832	1.022	0.968		397	0.813	2.102	0.696	0.754	0.977	0.791
	662	0.976	1.144	0.716	0.713	0.912	0.842		662	0.823	1.698	0.613	0.663	0.874	0.706
Urinary bladder	218	0.838	2.833	1.156	1.204	1.325	1.176	Muscle	218	0.656	3.509	1.710	1.615	1.493	1.472
	397	0.862	1.944	0.542	0.542	0.888	0.715		397	0.720	2.442	0.721	0.677	1.022	0.879
	662	0.855	1.622	0.498	0.514	0.803	0.647		662	0.749	1.927	0.614	0.596	0.910	0.778
Esophagus	218	0.713	3.090	1.321	1.373	1.353	1.191	Oral mucosa	218	0.684	1.813	2.761	2.593	1.521	1.468
	397	0.740	2.115	0.608	0.635	0.901	0.760		397	0.768	1.396	1.137	1.077	1.049	0.879
	662	0.743	1.764	0.514	0.555	0.837	0.677		662	0.755	1.209	0.965	0.912	0.965	0.777
Liver	218	0.700	2.631	2.542	0.941	1.311	1.241	Pancreas	218	0.687	2.706	1.702	1.681	1.246	1.115
	397	0.732	1.893	1.017	0.441	0.894	0.740		397	0.720	1.945	0.734	0.718	0.856	0.670
	662	0.750	1.547	0.831	0.426	0.804	0.667		662	0.737	1.578	0.646	0.640	0.776	0.609
Thyroid	218	0.965	2.479	2.071	1.736	1.588	1.449	Small intestine	218	0.800	2.617	1.534	1.901	1.329	1.198
	397	0.985	1.840	0.853	0.725	1.067	0.854		397	0.818	1.893	0.671	0.802	0.932	0.715
	662	0.970	1.530	0.722	0.634	0.938	0.751		662	0.820	1.551	0.599	0.688	0.827	0.642
Endosteum	218	0.684	3.398	2.011	1.884	1.563	1.549	Spleen	218	0.470	3.895	0.473	2.034	1.332	1.253
	397	0.716	2.299	0.803	0.755	1.029	0.886		397	0.524	2.626	0.243	0.814	0.907	0.746
	662	0.740	1.825	0.673	0.646	0.909	0.778		662	0.572	2.026	0.250	0.692	0.813	0.674
Brain	218	0.481	2.725	2.655	2.500	1.456	1.554	Thymus	218	0.946	2.234	1.084	1.104	1.400	1.365
	397	0.561	1.997	1.067	1.010	1.015	0.930		397	0.972	1.709	0.483	0.494	0.958	0.822
	662	0.619	1.650	0.879	0.848	0.917	0.826		662	0.953	1.420	0.445	0.457	0.861	0.739
Salivary glands	218	0.701	3.356	2.736	2.599	1.642	1.566	Lens of the eye	218	0.897	0.808	2.901	2.859	1.654	1.751
	397	0.780	2.454	1.092	1.046	1.131	0.936		397	0.952	0.778	1.168	1.159	1.142	1.018
	662	0.819	1.992	0.894	0.868	1.009	0.825		662	0.945	0.803	0.991	0.949	0.997	0.897
Skin	218	0.694	3.346	2.104	2.009	1.560	1.682	Testis	218	0.899	3.219	1.179	1.357	1.472	1.465
	397	0.604	1.873	0.705	0.679	0.852	0.800		397	0.915	2.388	0.569	0.629	1.015	0.867
	662	0.508	1.213	0.488	0.477	0.619	0.568		662	0.910	1.959	0.534	0.586	0.897	0.771
Adrenals	218	0.404	4.083	1.086	1.096	1.276	1.150	Prostate	218	0.712	3.717	1.061	1.031	1.230	1.142
	397	0.463	2.732	0.480	0.475	0.875	0.702		397	0.752	2.586	0.502	0.486	0.850	0.696
	662	0.511	2.101	0.437	0.431	0.794	0.639		662	0.771	2.021	0.474	0.462	0.770	0.633

AP, anteroposterior; PA, posteroanterior; RLAT, right lateral; LLAT, left lateral; ROT, rotational; ISO, isotropic.

Appendix B. Bias-Corrected $D_T/H_p(10)$ Conversion Coefficients for Female

Organs	Energy (keV)	$D_T/H_p(10)$ (Gy/Sv)						Organs	Energy (keV)	$D_T/H_p(10)$ (Gy/Sv)					
		AP	PA	RLAT	LLAT	ROT	ISO			AP	PA	RLAT	LLAT	ROT	ISO
Bone marrow	218	0.632	3.499	1.606	1.492	1.389	1.324	Extrathoracic region	218	0.598	2.685	2.935	2.690	1.648	1.480
	397	0.687	2.403	0.684	0.641	0.951	0.791		397	0.671	2.142	1.199	1.200	1.092	0.898
	662	0.715	1.890	0.596	0.570	0.855	0.706		662	0.697	1.738	0.946	0.949	0.986	0.796
Colon	218	0.829	2.694	1.841	1.524	1.391	1.276	Gallbladder	218	0.743	2.865	2.232	1.439	1.358	1.210
	397	0.863	1.957	0.771	0.665	0.951	0.778		397	0.777	2.062	0.929	0.647	0.939	0.727
	662	0.859	1.569	0.658	0.599	0.851	0.682		662	0.788	1.657	0.778	0.600	0.840	0.657
Lungs	218	0.678	3.412	1.383	1.380	1.372	1.333	Heart	218	218	0.792	2.935	1.480	2.211	1.428
	397	0.735	2.468	0.615	0.611	0.964	0.828		397	397	0.822	2.089	0.671	0.915	0.977
	662	0.751	1.943	0.555	0.563	0.889	0.758		662	662	0.830	1.700	0.602	0.782	0.877
Stomach	218	0.782	2.805	1.117	2.903	1.440	1.342	Kidneys	218	218	0.580	4.049	1.706	1.598	1.441
	397	0.810	2.038	0.504	1.147	0.984	0.791		397	397	0.638	2.705	0.721	0.683	0.984
	662	0.831	1.654	0.483	0.926	0.882	0.721		662	662	0.672	2.080	0.626	0.607	0.873
Breasts	218	0.934	1.872	2.193	2.117	1.509	1.592	Lymph nodes	218	0.757	3.065	1.708	1.647	1.433	1.331
	397	0.964	1.551	0.928	0.891	1.047	0.949		397	0.797	2.180	0.740	0.713	0.990	0.797
	662	0.964	1.376	0.784	0.770	0.939	0.835		662	0.811	1.752	0.643	0.634	0.881	0.715
Urinary bladder	218	0.956	2.089	1.278	1.195	1.318	1.302	Muscle	218	0.670	3.467	1.759	1.633	1.499	1.473
	397	0.943	1.552	0.599	0.545	0.913	0.765		397	0.728	2.416	0.740	0.689	1.028	0.879
	662	0.913	1.290	0.531	0.519	0.831	0.660		662	0.757	1.916	0.635	0.605	0.916	0.778
Esophagus	218	0.714	3.306	1.473	1.564	1.411	1.332	Oral mucosa	218	0.684	2.334	3.050	2.773	1.691	1.500
	397	0.753	2.311	0.622	0.629	0.966	0.790		397	0.757	1.889	1.211	1.129	1.124	0.891
	662	0.805	1.788	0.552	0.543	0.880	0.709		662	0.763	1.479	0.973	0.918	1.034	0.783
Liver	218	0.770	2.950	2.549	1.209	1.439	1.333	Pancreas	218	0.799	2.773	2.035	1.810	1.409	1.238
	397	0.801	2.089	1.023	0.547	0.977	0.791		397	0.822	1.997	0.853	0.772	0.971	0.740
	662	0.813	1.684	0.842	0.511	0.870	0.707		662	0.829	1.632	0.731	0.683	0.859	0.661
Thyroid	218	0.982	2.601	1.490	1.072	1.593	1.493	Small intestine	218	0.781	2.824	1.428	1.814	1.356	1.221
	397	1.005	1.918	0.633	0.462	1.067	0.879		397	0.810	2.063	0.639	0.760	0.927	0.734
	662	0.987	1.572	0.560	0.437	0.939	0.786		662	0.823	1.646	0.575	0.664	0.840	0.668
Endosteum	218	0.705	3.414	2.062	1.936	1.587	1.568	Spleen	218	0.532	4.313	0.589	2.327	1.450	1.353
	397	0.736	2.312	0.822	0.772	1.048	0.898		397	0.593	2.875	0.294	0.927	0.984	0.803
	662	0.757	1.833	0.687	0.667	0.920	0.786		662	0.636	2.199	0.297	0.780	0.878	0.714
Brain	218	0.503	2.748	2.699	2.526	1.486	1.583	Thymus	218	0.961	2.475	1.099	1.194	1.411	1.408
	397	0.581	2.010	1.086	1.022	1.034	0.949		397	0.985	1.840	0.491	0.541	0.958	0.841
	662	0.632	1.661	0.892	0.859	0.930	0.840		662	0.971	1.536	0.453	0.509	0.865	0.752
Salivary glands	218	0.639	3.382	2.875	2.732	1.697	1.543	Lens of the eye	218	0.899	0.921	2.926	2.862	1.641	1.734
	397	0.715	2.467	1.155	1.100	1.163	0.923		397	0.927	0.894	1.211	1.153	1.124	1.038
	662	0.755	1.979	0.938	0.915	1.031	0.820		662	0.949	0.884	0.958	0.985	1.014	0.930
Skin	218	0.698	3.389	2.158	2.036	1.579	1.697	Ovaries	218	0.669	3.201	1.299	1.104	1.214	1.192
	397	0.608	1.900	0.730	0.685	0.865	0.806		397	0.720	2.194	0.570	0.504	0.837	0.702
	662	0.510	1.231	0.500	0.480	0.623	0.570		662	0.741	1.728	0.520	0.473	0.758	0.648
Adrenals	218	0.559	3.805	1.657	1.403	1.356	1.204	Uterus	218	0.730	3.055	1.151	1.148	1.204	1.161
	397	0.618	2.574	0.721	0.617	0.932	0.727		397	0.769	2.168	0.533	0.529	0.830	0.702
	662	0.651	1.997	0.629	0.564	0.833	0.662		662	0.786	1.737	0.491	0.495	0.762	0.637

AP, anteroposterior; PA, posteroanterior; RLAT, right lateral; LLAT, left lateral; ROT, rotational; ISO, isotropic.