

A size-specific effective dose for patients undergoing CT examinations

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A size-specific effective dose for patients undergoing CT examinations

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Abstract. This study aims to develop a simple method for estimating the size-specific effective dose using the ImPACT software. The size-specific effective dose was calculated from the images of patients who underwent CT examinations of the thorax and abdomen regions. Volume computed tomography dose index (CTDI_{vol}) and mAs data were obtained from the dose report. The average of mAs and CTDI_{vol} were used to determine the normalized CTDI_{vol} (nCTDI_{vol}). Patient size was expressed in effective diameter (D_{eff}), and was measured at nine slices along the z-axis. The normalized size-specific dose estimate (nSSDE) was then calculated. The normalized size-specific effective dose was obtained in the “effective dose” text-box by inputting an arbitrary value in the “CTDI (air)” text-box so that the “CTDI_{vol}” text-box showed a similar value to the nSSDE. The results show that the normalized size-specific effective dose decreases exponentially with increasing patient size. The results are consistent with results of Sahbaee formula to within 20%.

1. Introduction

There are several dosimetric quantities in CT scanning, including volume computed tomography dose index (CTDI_{vol}) [1, 2], dose-length product (DLP) [3, 4], size-specific dose estimate (SSDE) [5, 6], organ dose [7, 8], and effective dose (E) [9]. Among them, the effective dose is considered as the best quantity to relate cancer risk to X-ray radiation from CT [10]. It is known that cancer risk is influenced not only by the magnitude of the average patient dose or organ dose, but is also determined by the sensitivity of each exposed organ to the radiation [11]. With this effective dose, the magnitude of CT dose can be compared to other radiological modalities, such as fluoroscopy, mammography, dental radiology, and conventional radiology [12], and it can also be compared with the natural radiation dose [13].

Up to now, there were several techniques for calculating the effective dose. One way was by using a software package such as ImPACT [14], CT-Expo [15] or Waza-ari [16], another was by measurement using an anthropomorphic phantom and a small detector such as photoluminescence dosimeter (PLD) [17], or by calculation using Monte Carlo simulation [18], or it could be estimated from DLP using a conversion value of DLP/E or the so-called *k*-factor [19]. However, all of these approaches are still limited to a phantom of a standard size, so the results may differ from an actual patient.



Several investigators have reported that estimates of patient dose, organ dose, and effective dose using output CT machine only is not sufficient, and that the size of the patient should be taken into account [20-23]. This study introduced a simple method for estimating the size-specific effective dose using ImPACT.

2. Method

2.1. Overall steps of this study

The overall steps in this study are shown in Figure 1. The size-specific effective dose was calculated from patients who underwent thoracic and abdominal CT examinations. The effective diameter (D_{eff}) was determined from the image of the patient. $CTDI_{vol}$ and mAs values were derived from the dose report. From the average of the mAs and $CTDI_{vol}$ values, the normalized $CTDI_{vol}$ (in units of mGy /100 mAs) and normalized effective dose (nSSDE, taking into account the patient's size) were determined. Finally, based on this nSSDE value, the normalized size-specific effective dose was calculated using ImPACT software. The results obtained in this approach were compared with the results calculated by the Sahbaee formula [21].

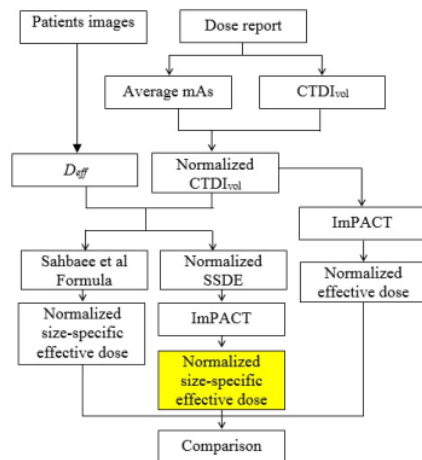


Figure 1. Overall steps in this study.

2.2. Patient Images

We calculated the size-specific effective dose from 24 patients who underwent thoracic CT examination and 27 patients who underwent abdominal CT examinations. The age averages were $57.8 (\pm 13.8)$ and $48.7 (\pm 14.1)$ years for the patients underwent thoracic and abdominal examinations respectively. All patients were scanned using a MDCT Siemens Somatom 6. Scan parameters were shown in Table 1. The MDCT was equipped with tube current modulation (TCM) with 70 mAs reference for thoracic examinations and 95 mAs for abdominal examinations.

Table 1. Scan parameters of CT examinations in this study.

Scanning parameter	Values
Tube voltage	130 kVp
Rotation time	0.6 s
Pitch	0.8
Slice thickness	2 mm

2.3. D_{eff} calculation

In this study, the patient size was characterized by the effective diameter (D_{eff}) [24]. The D_{eff} of each patient was calculated from images of the patient at nine slices (locations) along the z-axis [5] as shown in Figure 2(a). The maximum diameters in the lateral (D_{LAT}) and antero-posterior (D_{AP}) directions were determined, as shown in Figure 2(b). The value of D_{eff} was calculated from:

$$D_{eff} = \sqrt{D_{AP} \times D_{LAT}} \quad (1)$$

The average of D_{eff} was then calculated from these nine values of D_{eff} .

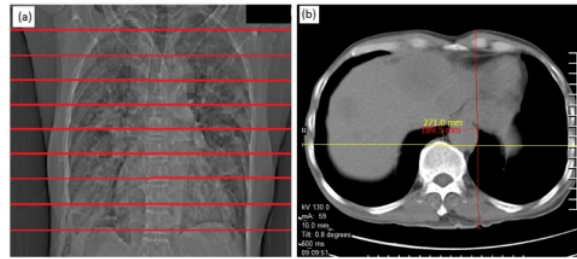


Figure 2. (a) Nine slices along the z-axis for determining D_{eff} for every patient. (b) D_{eff} for each slice was determined by the square root of the product of the maximum diameters in the lateral direction (D_{LAT}) and in the antero-posterior direction (D_{AP}).

2.4. $nCTDI_{vol}$ and $nSSDE$

$CTDI_{vol}$ and mAs values were taken from the CT scanner dose report. The mAs value is the product of tube current and rotation time. In the tube current modulation (TCM) technique, the tube current is not constant throughout the scan process, so that the average mAs should be calculated. The normalized $CTDI_{vol}$ ($nCTDI_{vol}$) was calculated in units of mGy/100 mAs [5]. The quantity used for describing patient dose is the size-specific dose estimate (SSDE). Normalized SSDE ($nSSDE$) was calculated by the equation:

$$nSSDE = nCTDI_{vol} \times a \times e^{-b \times D_{eff}} \quad (2)$$

where a is 3.704369 and b is 0.03671937 for thoracic and abdominal examinations [25].

2.5. ImPACT and scan protocol

ImPACT is software developed by Im-PACT group (UK National Health Service CT Evaluation Centre, London, UK) which can be used to estimate organ doses and effective doses for many scanners with different examination parameters [12]. In this study, we used ImPACT 1.0.1a (Figure 3). The input of "scan region" for the thoracic and abdominal examination was "body", and the "scan range" inputs are shown in Table 2. Selection of the scan range refers to the protocol and a previous study by Sahbaee [21]. In addition, the effective dose is greatly influenced by the organ weighting factors. In this study we used the ICRP 103 data.

All acquisition parameters, such as spiral pitch and collimation, were inserted into the ImPACT software. The product of tube current and rotation time was set to be 100 mAs. The "CTDI (air)" was filled by trial and error, resulting in expected $nCTDI_{vol}$ (equal to the value of the $nCTDI_{vol}$ as in the dose report). It should be noted that this effective dose does not take patient size into consideration. The "CTDI (air)" text-box was filled again by trial and error with a value such that the "CTDI_{vol}" box showed the $nSSDE$ value. The effective dose value shown was the normalized size-specific effective dose.

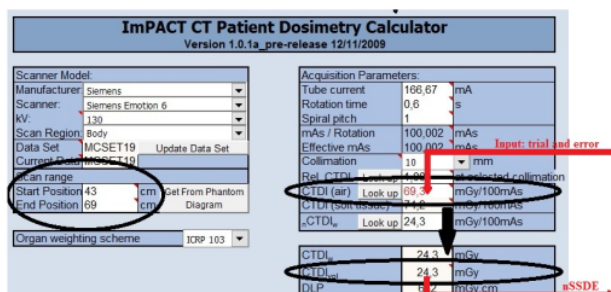


Figure 3. Screen capture of ImPACT software for calculating normalized effective dose and normalized size-specific effective dose. The text box of “CTDI (air)” was filled by trial and error, so that the “CTDI_{vol}” box indicated the value of nSSDE.

Table 2. Standard head and body CT examinations [19].

Examination	Start and End z coordinates (cm)	Scan Length (cm)	Scan Coverage
Thorax	43 and 69	26	Start from 1 cm above the lung down to 1 cm below the lung base
Abdomen	20 and 44	24	Start from 1 cm above the superior liver down to 1 cm below the superior iliac crest

2.6. Sahbaee formula

The result of this calculation was compared with the calculated results using the Sahbaee formula [21]. The formula is a patient-based estimation of effective dose for adult protocols across 13 categories, including thoracic and abdominal protocols. It was derived using a validated Monte Carlo program on 58 adult cardiac-torso extended computational phantoms. The correlation between the normalized effective dose (E) and patient size inside the scan coverage is described by an exponential fit:

$$E(D_{eff, avg}) = DLP \times e^{(\alpha \times D_{eff, avg} + \beta)} \quad (3)$$

The α values are -0.04 and -0.05 and the β values are -2.52 and -2.49, for thorax and abdomen respectively [21].

3. Results

3.1. CTDI_{vol} and nCTDI_{vol}

The patient size was characterized by the average D_{eff} from nine points along the z-axis. The average D_{eff} for the thoracic examinations was $22.4 (\pm 2.4)$ cm; and for the abdominal examinations was $23.5 (\pm 2.0)$ cm. The average D_{eff} of our patients was lower by about 30% for thorax and 28% for abdomen than the size of the standard 32-cm phantom.

The correlations between CTDI_{vol} and D_{eff} for thoracic and abdominal examinations are shown in Figure 4. The value of CTDI_{vol} increased linearly with D_{eff} , because the average tube current increases with D_{eff} . The average CTDI_{vol} was $4.5 (\pm 1.2)$ mGy for thoracic examinations, and the average CTDI_{vol} was $5.1 (\pm 1.1)$ mGy for abdominal examinations. The CTDI_{vol} value was 3.5 mGy for D_{eff} about 20 cm, and 7.0 mGy (2-fold increase) for D_{eff} about 28 cm. R^2 values were 0.514 and 0.728 for the thoracic and abdominal examinations respectively.

After normalization, $CTDI_{vol}$ was independent of D_{eff} . The $nCTDI_{vol}$ was constant at about 12 mGy/100 mAs (11.6 ± 0.2 mGy/100 mAs for thoracic examinations and 11.8 ± 0.4 mGy/100 mAs for abdominal examinations). The correlation between nSSDE and D_{eff} for thoracic and abdominal examinations are shown in Figure 4. The nSSDE decreases exponentially with increasing D_{eff} . The average nSSDE values were 18.9 mGy/100 mAs (± 1.7 mGy/100 mAs) and 18.7 mGy/100 mAs (± 1.7 mGy/100 mAs). The nSSDE was found to overestimate $nCTDI_{vol}$ by about 63% and 59% for thoracic and abdominal examinations respectively.

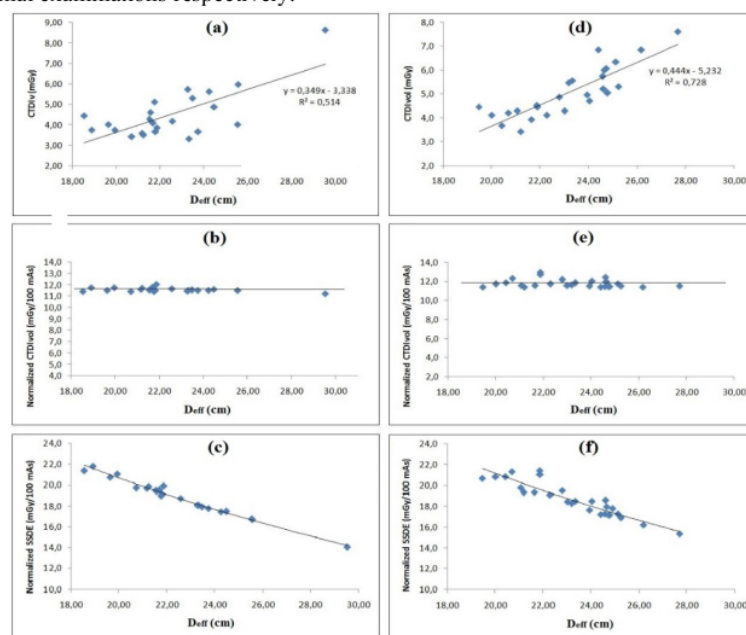


Figure 4. The correlations between $CTDI_{vol}$ and D_{eff} , $nCTDI_{vol}$ and D_{eff} , and nSSDE and D_{eff} for thoracic (a-c) and abdominal examinations (d-f).

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3.2. Effective Dose

The normalized standard effective dose and normalized size-specific effective doses for thoracic and abdominal are shown in Figure 5. The standard normalized effective dose calculated using ImPACT is independent of D_{eff} . It was constant at about 6 mSv/100 mAs (6.3 ± 0.1 mSv/100 mAs and 5.6 ± 0.2 mSv/100 mAs, for thoracic and abdominal examinations respectively). However the normalized size-specific effective dose decreases exponentially with increasing D_{eff} . For the largest patients in the thoracic examinations, the normalized size-specific effective dose was found to be about 56% higher than for smallest patients; and in the abdominal examinations, the normalized size-specific effective dose was found to be 39% higher than that for smallest patients. The average of the normalized size-specific effective doses were $10.4 (\pm 1.0)$ mSv/100 mAs and $8.8 (\pm 0.8)$ mSv/100 mAs for thoracic and abdominal examinations respectively.

The same pattern was obtained by using the Sahbaee formula [21]. The differences between our approach and the Sahbaee formula were $4.5 \pm 2.5\%$ for the thoracic examinations and $19.5 \pm 3.4\%$ for the abdominal examinations. As comparison, the differences between the normalized size-specific effective dose using the Sahbaee formula and the standard normalized standard effective dose using ImPACT (which neglects patient size) were $35.7 \pm 6.4\%$ and $23.7 \pm 7.6\%$ for thoracic and abdominal examinations respectively.

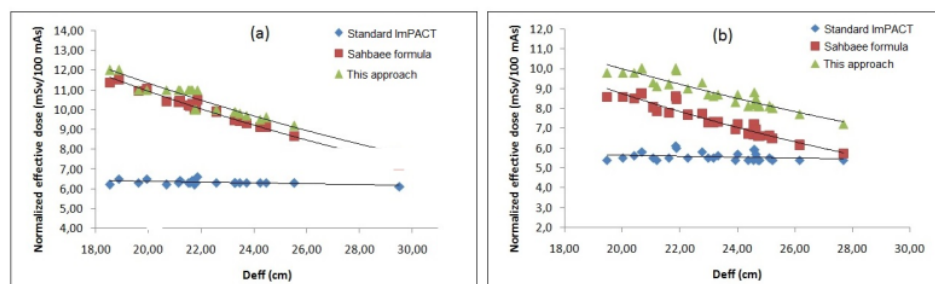


Figure 5. The normalized standard effective dose and normalized size-specific effective doses for (a) thoracic examinations and (b) abdominal examinations.

4. Discussion

Effective dose is a descriptor that reflects the cancer risk of CT examinations. Several previous studies reported that the effective dose can be estimated from the DLP by applying a k -factor [11,19]. The k -factor values have been expanded to take into account age and gender. However, it is still limited to a phantom which may differ from the actual size of patient. By ignoring the size of the patient, effective dose can vary by 100% or more [11]. The k -factor has now been reported for various sizes and for some specific protocols [21], based on validated Monte Carlo simulations on mathematical model phantoms. In addition to using the DLP and k -factor, the simplest method for calculating the effective dose is to use a specific calculator such as ImPACT. However, ImPACT uses a standard phantom representing the standard patient size. This study has attempted to develop a technique for calculating size-specific effective dose using ImPACT, without having to change the ImPACT software itself.

The results showed that the normalized size-specific effective dose decreases exponentially with increasing D_{eff} . The adult patients evaluated in this study with D_{eff} values from 18.5 cm to 29.5 cm, the differences between the largest and smallest normalized size-specific effective dose were about 55% and 40% for thoracic and abdominal examinations respectively. The largest of the normalized size-specific effective dose compared to the normalized standard effective dose (not taking into account D_{eff}) were 90% and 80% for thoracic and abdominal examinations respectively. For pediatric patients with D_{eff} under 15 cm, the differences may be more 100%. Therefore in order to achieve accurate estimates of effective dose, it is critical to take into account the size of the patient.

The results of this approach were consistent with Sahbaee formula [21]. The differences between both were quite small, which are about 5% and 20% for thoracic and abdominal examinations. The discrepancies are possibly due to: First, the values of scan length used in this approach was constant (i.e., 26 cm and 24 cm for thoracic and abdominal examinations), while Sahbaee formula was obtained using many mathematical phantoms with varying size and length. Second, the Sahbaee formula was obtained from GE Light-Speed VCT system and this approach was used to calculate the effective dose from Siemens Somatom-6 CT system. However, the relatively small differences indicated that the size-specific effective dose can be accurately estimated by this approach using the ImPACT software. One advantage of using this software rather than using k -factor and DLP, is that the size-specific effective dose can be estimated for different protocols and scan ranges.

5. Conclusions

The normalized size-specific effective dose has been calculated using ImPACT software, by replacing $nCTDI_{vol}$ with $nSSDE$. The results show that the normalized size-specific effective dose decreases exponentially with increasing patient size. The results of normalized size-specific effective dose were consistent to within 20% with a Sahbaee formula.

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