Digital Breast Tomosynthesis: Three-Dimensional Measurement of Breast-Absorbed Dose Distribution

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Abstract

Background: The dose exposure associated with digital breast tomosynthesis (DBT) is evaluated by multiplying the mean glandular dose (MGD) of mammography (MMG) with the correction coefficient of the angle of X-ray incidence. However, it has been pointed out that there are various problems when using the MGD as a standard for risk assessment in breast cancer screening. Therefore, the aim of this study was to assess the breast-absorbed dose for different breast sizes for dose assessment. Furthermore, in this study, by measuring the dose distribution three-dimensionally, we aimed to examine the MGD correction method using the breast size as a factor. **Methods:** A simulated breast phantom with a diameter of 40–160 mm and a total thickness of 40 mm, made with polymethyl methacrylate, was created by simulating the phantom shape used in the simulation calculation for calculating the MGD. It was made with polymethyl methacrylate. Radiochromic films were placed at different depths, which measured the breast-absorbed dose distribution three-dimensionally. The MGD was calculated from the breast-absorbed dose distribution obtained. **Results:** The three-dimensional dose distribution revealed that there was a difference in the distribution of MMG and DBT with increasing depth. In addition, a lower X-ray energy and a smaller breast size resulted in a greater difference in the absorbed dose between DBT and MMG. **Conclusion:** Incorporating the DBT correction according to the breast size into the MGD improves the accuracy of dose evaluation by the MGD. Additionally, a corrected MGD provides useful information for risk assessment when DBT is used for breast cancer screening.

Keywords: Digital breast tomosynthesis- breast dose distribution- MGD- radiochromic film

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Introduction

Currently, mammography (MMG) has been scientifically proven to be an effective technique for breast cancer screening. The greatest advantage of breast cancer screening is the reduction of mortality via the early detection of breast cancer. In contrast, its disadvantages include radiation exposure, false-positive or negative diagnosis, over diagnosis, physical burden, and psychological burden. The effectiveness of the screening depends on a comprehensive judgment that considers the balance between the advantages and disadvantages (Sylvia et al., 2011).

In a dense breast, it is difficult to diagnose the presence of a tumor using MMG as MMG imaging overlaps with the mammary glands and tumors (Thomas et al., 2002). Therefore, MMG may be less effective for screening individuals with dense breasts, and other methods are needed to improve the sensitivity of breast cancer screening (Suzuki et al., 2021).

Digital breast tomosynthesis (DBT) is attracting attention as an additional function of the mammography

system (Alakhras et al., 2013). In DBT, approximately 10 X-ray irradiations are performed using the step-andshot method, and the projected images at each angle are collected (Uematsu, 2013). The use of DBT minimizes or eliminates the limitation of MMG associated with the mammary glands and mass overlap (Helvie, 2010; Sechopoulos, 2013). Therefore, this technique reduces the rate of false positives and false negatives. The use of DBT images has been reported to potentially improve the diagnosis and examination accuracy in individuals with dense breasts, reduce the requirement for re-examination, improve the conviction of interpretation, and identify the location of lesions (Helvie, 2010; Houssami and Skaane, 2013; Skaane et al., 2013; Gilbert et al., 2015; Chong et al., 2019; Kulkami et al., 2021; Mackenzie et al., 2021).

Irrespective of the effectiveness, radiation exposure is unavoidable for every individual undergoing breast cancer screening using X-rays. Therefore, it is necessary to consider the risk of radiation exposure (Ali et al., 2015).

Dance et al., (2011) proposed a method for estimating the mean glandular dose (MGD) of DBT using Eq. (1). This is a method of assessing the average absorbed dose

Department of Radiological Sciences, Ibaraki Prefectural University of Health Sciences, Ibaraki Prefecture, Japan. *For Correspondence: nakajimae@ipu.ac.jp by the mammary glands in a compressed breast. MGD is simulated using a polymethyl methacrylate (PMMA) phantom with a diameter (symbol: φ) of 160 mm and a thickness of 40 mm as a substance simulating a standard breast (compression breast thickness, 45 mm; mammary gland/fat ratio, 41/59). The MGD is a product of air kerma (*K*), on the surface of the breast with coefficient factor *g*, corresponding to 50% of the mammary gland mass coefficient factor *c*, for correcting the difference in breast composition from 50% glandularity, and coefficient factor *s* for the target/filter combination. Additionally, it is calculated using a coefficient factor *T* that considers the effect of obliquely incident X-rays.

$$MGD = K \times g \times c \times s \times T \tag{1}$$

The MGD estimates the average of the breast-absorbed dose that changes depending on the subject's breast thickness and mammary gland concentration (Dance et al., 2011). In the case of MMG, the MGD evaluated using the PMMA phantom does not match the actual breastabsorbed dose (Dance et al., 2005; Sechopoulos et al., 2012; Hernandez et al., 2015; Wang et al., 2017; Sarno et al., 2018; Chang et al., 2020; Huda and Antar, 2020). In DBT cases, it has been reported that when the shape of the phantom is made closer to that of the actual breast, the peripheral part of the breast absorbs a higher dose than the central part at a depth of 20 mm from the surface (Flores et al., 2019). In addition, as X-rays are obliquely incident, there is an effect of scattered radiation from the compression plate and breast support. In particular, it has been reported that the contribution becomes more pronounced as the projection angle increases (Diaz et al., 2019). Therefore, the dose distribution in the breast is presumed to be complicated. Furthermore, the composition of the human breast has been elucidated using breast-dedicated CT, and the coefficient factor T increases when the dose is evaluated considering the breast model (Sarno et al., 2018).

In a previous study, to evaluate the validity of the MGD of DBT by Monte Carlo simulation, the two-dimensional dose distribution in the phantom was actually measured by inserting a radiochromic film (RCF) into the phantom for each depth (Sarno et al., 2017). However, the analyzed area included only three positions on the chest wall side and one position on the nipple side, but not the entire breast. In another study, the dose distribution by depth in the phantom was evaluated considering the shape of the breast. However the setting of the region of interest was limited only to the chest wall side and nipple side (Flores et al., 2019).

In accordance with the direction of X-ray incidence in DBT, the difference in breast size is considered to be related to the exposure dose. Currently, the evaluation of MGD is performed by simulating the breast size of European/US populations. The breast size is simulated using a PMMA phantom with a diameter (ϕ) of 160 mm and a thickness of 45 mm. However, the average breast diameter (ϕ) in Japanese women was reported to be 89.4 mm (National Institute of Bioscience and Human-Technology, 1996). To date, only a few studies have examined the exposure dose owing to the difference in breast size.

To evaluate the dose absorbed by the breast for dose evaluation and to examine the risks associated with the use of DBT for breast cancer screening, this study focused on breast size and changes in breast dose distribution associated with DBT. Furthermore, our aim was to examine the MGD correction method using breast size and X-ray energy as factors by measuring dose distribution three-dimensionally.

Materials and Methods

Study design

The PMMA phantom sizes used in this study were φ160 mm (Dance, 1990), which is the standard breast phantom size; $\varphi 80$ mm, which is close to the average breast size of Japanese women; and $\varphi 40$ mm, which is close to the clinically smallest size. The direction of and the direction perpendicular to the chest wall-nipple are described as directions Y and X, and the X-ray tube-breast support direction is described as direction Z. The origins of directions X and Y were the center of the incident X-ray and the edge of the chest wall, respectively; the origin of direction Z was 40 mm above the breast support. The size of the phantom in direction Y was half of that in direction X. The total thickness was 40 mm by combining 5-mm and 10-mm phantoms. A schematic diagram of the prepared phantom is shown in Figure 1. The RCFs were sized 170 mm \times 85 mm, 90 mm \times 45 mm, and 50 mm \times 25 mm, for the phantom sizes of φ 160 mm, 80 mm, and 40 mm, respectively. At depths of 0-10 mm and 20-40 mm from the surface of the phantom, RCFs were loaded at 5 mm and 10 mm intervals, respectively. There were six layers in total.

Dose evaluation using RCF

The type of RCF was XR-SP2 (lot#01212101; ASHLAND, Covington, KY, USA). The RCF used the background correction method to convert the *Apixel* values into air kerma K (Saur and Frengen, 2008). For irradiation, we used Senographe Pristina (GE, Fairfield, CT, USA) equipment. The irradiation conditions were target/filter combination of Mo/Mo and tube voltage of 24–32 kV. Additionally, to simulate the actual human body, a 40-mm thick PMMA plate that simulates the pectoralis major muscle was placed by the phantom. Figure 2 shows the measurement layout figure and photo when the K in the phantom was measured using RCF. The phantom was placed at the center of the X-ray machine in direction X and fixed with a compression plate in direction Y such that it was in line with the edge of the chest wall. Furthermore, the average energy (E_{ave}) of incident X-ray at each depth was calculated using the simulation calculation software MoXS-5 (Molybdenum target X-ray Spectrum, ver.5). Using E_{max} , we referred to the data of the National Institute of Standards and Technology to obtain the mass energy absorption coefficient ratio of air and phantom $\left(\frac{\mu_{en}}{\rho}\right)_{PMMA,air}$ (data by National Institute of Standards and Technology). The absorbed dose in the phantom for each depth (D_d) was calculated from Eq. (2).

$$D_{\rm d} = K \times \left(\frac{\mu_{\rm en}}{\rho}\right)_{\rm PMMA, air}$$
(2)

Examination of correction coefficient of average absorbed dose

The average dose absorbed by the phantom (\overline{D}_{all}) measured using RCF was calculated from Eq. (3).

$$\overline{D}_{\text{all}} = \frac{1}{n} \sum_{1}^{n} \overline{D}_{\text{d}}$$
(3)

The ratio R_T of \overline{D}_{all} with respect to MMG and DBT was calculated from Eq. (4).

$$R_{\rm T} = \frac{\overline{D}_{\rm all, DBT}}{\overline{D}_{\rm all, MMG}} \tag{4}$$

Here, $\overline{D}_{all,DBT}$ indicated the average dose absorbed by the phantom subjected to DBT, and $\overline{D}_{all,MMG}$ indicated the average dose absorbed by the phantom subjected to MMG. Furthermore, the ratio of \overline{D}_{all} because of the difference in the phantom size was calculated. The ratios f_{80} and f_{40} for phantoms of sizes φ 80 mm and φ 40 mm, respectively, were calculated from Eq. (5) and Eq. (6) with φ 160 mm as the reference.

$$f_{80} = \frac{D_{\text{all},\varphi 80}}{\overline{D}_{\text{all},\varphi 160}} \tag{5}$$

$$f_{40} = \frac{\overline{D}_{\text{all},\phi40}}{\overline{D}_{\text{all},\phi160}} \tag{6}$$

In Eq. (5), $\overline{D}_{all,\varphi80}$ and $\overline{D}_{all,\varphi160}$ indicated the average doses absorbed by the $\varphi80$ -mm and $\varphi160$ mm phantoms. In Eq. (6), $\overline{D}_{all,\varphi40}$ indicates the average dose absorbed by the $\varphi40$ mm phantom.

Results

Dose evaluation using RCF

The ratio of the mass energy absorption coefficient of air calculated from the X-ray spectrum to the mass energy absorption coefficient of PMMA ($\left(\frac{\mu en}{\rho}\right)_{PMMA,air}$) was approximately 0.62–0.63 regardless of the depth, Z. The absorbed dose, D_{ap} in the phantom in the XY direction

Table 1. Average Absorbed Dose by All Phantoms (\overline{D}_{all}) .

Tube Voltage [kV]	φ [mm]	$\overline{D}_{ m all} \left[\mu { m Gy/mAs} ight]$	
		MMG	DBT
24	160	8.97	8.83
	80	8.22	7.9
	40	8.15	9.5
28	160	16.9	17
	80	16.1	16.6
	40	15.3	16.8
32	160	26.2	25.2
	80	22.9	23.4
	40	23.4	24

DBT, digital breast tomosynthesis; MMG, mammography

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of MMG and DBT with a tube voltage of 24 kV and φ 40 mm is shown in the color map by depth (Figure 3a, b). In addition, regions that were 7 mm away from the edge of the chest wall were out of the irradiation field. In both cases, the dose decreased with an increase in the depth, Z. Moreover, with an increase in the depth, the dose at the phantom edge was higher for DBT than for MMG.

Figure 4 shows the color map of the dose absorbed D_d by the phantom on the XZ plane when the tube voltage was 24 kV and the phantom sizes were φ 160 mm, 80 mm, and 40 mm. The smaller the phantom size, the greater the difference in the absorbed dose distribution with respect to MMG and DBT. Especially, at a deep position in direction Z, the DBT-associated dose at the edge was higher than the MMG-associated dose. This phenomenon became more remarkable with a decrease in tube voltage.

Table 1 shows the results of \overline{D}_{all} obtained from Eq. (3). \overline{D}_{all} was between 8.15 μ Gy/mAs and 26.2 μ Gy/mAs in MMG. In DBT, it was between 7.90 and 25.2 μ Gy/mAs. In both MMG and DBT, the higher the energy regardless of the phantom size, the higher the \overline{D}_{all} in the phantom. With MMG, \overline{D}_{all} was smaller for all tube voltages as the phantom size was smaller.

Examination of correction coefficient of average absorbed dose

Using Eq. (4), the dose ratio $R_{\rm T}$ of $\overline{D}_{\rm all}$ of DBT and MMG was calculated. The results are presented in Table 2. Concerning the phantom size, when the phantom size was small, the $R_{\rm T}$ increased. The largest $R_{\rm T}$ was 1.166 when the tube voltage was 24 kV and phantom size was $\varphi 40$ mm. From Table 2, $R_{\rm T}$ values at the phantom size of $\varphi 160$ mm were 0.985, 1.008, and 0.963 at tube voltages of 24, 28, and 32 kV, respectively, with an average of 0.985. Furthermore, the averages for the phantom sizes of $\varphi 80$ mm and $\varphi 40$ mm were 1.006 and 1.098, respectively.

Figure 5 shows f_{80} and f_{40} of \overline{D}_{all} due to the difference in phantom size based on $\varphi 160$ mm phantoms. It was 0.87–0.95 for both f_{80} and f_{40} regardless of the tube voltage for MMG. With respect to DBT, f_{80} was lower than 1 regardless of tube voltage, as in the case of MMG, and f_{40} was lower than 1 at tube voltages of 28 kV and 32 kV. Especially, \overline{D}_{all} values at $\varphi 80$ mm and $\varphi 40$ mm were lower than that at $\varphi 160$ mm. However, the f_{40} at a tube voltage of 24 kV was 1.08. The relational expression between f_{40} and tube voltage in DBT is presented in Eq. (7).

$$f_{40} = 0.0016x^2 - 0.1038x + 2.67 \tag{7}$$

In this equation, x indicates the tube voltage.

Table 2. Ratio $R_{\rm T}$ of $\overline{D}_{\rm all}$ between DBT and MMG.

φ [mm]	Tub	Tube Voltage[kV]			
	24	28	32		
160	0.985	1.008	0.963	0.985	
80	0.962	1.032	1.023	1.006	
40	1.166	1.102	1.026	1.098	

DBT, digital breast tomosynthesis; MMG, mammography



Figure 1. Semi-Cylindrical PMMA Phantom Diagram. PMMA, polymethyl methacrylate

Discussion

In this study, the dose distribution in the phantom of MMG and DBT was measured three-dimensionally using RCF. With the metal-oxide-semiconductor field effect transistor and thermoluminescent dosimeter used to measure the dose distribution in the phantom, the measurement location is limited to the representative points (Fedon et al., 2018). However, using the RCF, the dose distribution in the phantom was obtained with high spatial resolution. By using RCF, the dose distribution at the periphery of the phantom could be measured in detail. Figure 3 shows that a deeper depth indicated the use of lower doses of MMG and DBT. The difference between MMG and DBT was noticeable at the edges. Especially when compared at a depth Z = 40 mm, it was visually confirmed that the dose at the margin was higher in DBT than in MMG. As shown in Figure 4, as the phantom size decreased, the absorbed dose D_d at the edge of the twodimensional dose distribution of DBT tended to increase with respect to MMG. In particular, when comparing the two-dimensional distributions of absorbed dose D_d with a $\varphi 40$ mm phantom, the dose distributions of MMG and DBT differed at the phantom edge, and the dose of DBT



(a) figure Figure 2. Layout for the Measurement of K in the Phantom.

(b) photo



(b) DBT

Figure 3. Color Maps of Absorbed dose D_d in Phantom in XY Direction (tube voltage 24 kV, φ 40 mm). DBT, digital breast tomosynthesis; MMG, mean glandular dose

at the phantom edge was higher than that at a deep position in the phantom. This is because the oblique incidence of X-rays causes the primary X-rays to be incident on the sides of the phantom and the increased contribution of scattered radiation from the breast support increases the dose of the phantom. Previous studies have shown that DBT is associated with a higher edge dose than MMG, and our results corroborate this finding (Diaz et al., 2019).

Table 1 shows that \overline{D}_{all} increased with an increase in the energy of both MMG and DBT regardless of the phantom size. This result is consistent with the previous ones because the coefficients factors g and c used for calculating MGD also increase with an increase in the energy (Dance et al., 2009).

Regarding the phantom size, \overline{D}_{all} of MMG decreased with a decrease in the phantom size. This can be explained as follows: when the phantom size is small, the decrease in the scattered dose in the phantom contributes more than the increase in the scattered radiation from the breast support. In contrast, \overline{D}_{all} of DBT was larger in the φ 40-mm phantom at a tube voltage of 24 kV than in the φ 160-mm phantom. This can be explained by the fact that the rate of increase in the scattered radiation from the breast support is larger than the rate of decrease in the scattered dose in the phantom because of the oblique incidence of X-rays.

The coefficient factor *T* for converting MGD of MMG to MGD of DBT reported by Dance et al., (2011) is 0.990 at a PMMA phantom thickness of 40 mm. As shown in Table 2, the $R_{\rm T}$ values measured using RCF were 0.985, 1.008, and 0.963 at tube voltages of 24, 28, and 32 kV, respectively. The average value was 0.985. These values are the same as those reported by Dance et al., (2011). Furthermore, the averages for φ 80 mm and φ 40 mm were 1.005 and 1.098, respectively. A smaller phantom size resulted in a larger $R_{\rm T}$. This can be explained as follows: when the phantom size decreases, X-rays are obliquely incident on DBT and, therefore, the primary X-rays are incident on the side surface of the phantom, and the dose in



(b) DBT

Figure 4. Color Map of D_d by the Depth to Differences in Phantom Size in the XZ Direction. (tube voltage, 24 kV). It presents at Y=10 mm position.

the phantom increases due to the contribution of scattered rays from the breast support. These are consistent with previous studies reporting the use of higher doses at the edge of the phantom when the angle increased (Flores et al., 2019). In addition, there is no contradiction with a previous report, which stated that the dose distribution at the depth Z of 10 mm was higher in the peripheral area owing to the effect of oblique incidence even in the absence of a compression plate (Fedon et al., 2018).

As shown in Figure 5, f_{40} and f_{80} values of MMG and DBT were <1 at most tube voltages and phantom sizes. Especially, the φ 160-mm phantoms currently used for dose evaluation overestimate these values for subjects with small breast sizes. However, f_{40} at a tube voltage of 24 kV is 1.08, which is underestimated at the phantom size of φ 160 mm. The risk assessment for breast cancer screening is assessed by stochastic effects. The cancer mortality rate due to radiation exposure is estimated by multiplying the effective dose by the lifetime risk factor according to the age at the time of exposure. If the MGD is 2 mGy/exposure with MMG, the lifetime fatal carcinogenic risk is 0.0006% by ICRP (ICRP, 2007). Considering that this study found f_{40} to be 1.08, if the individual's breast size is φ 40 mm, the lifetime fatal carcinogenic risk would be 0.000648%. Previous dose assessments revealed that six people/million would develop breast cancer and die



Figure 5. Ratio of D_{all} Difference by Phantom Size Based on the φ 160 mm Phantom with Respect to DBT and MMG. DBT, digital breast tomosynthesis; MMG, mammography

because of DBT-based breast cancer screening; however, after taking the phantom size into account, 6.48 people/ million were expected to die if the individual's breast size is $\varphi 40$ mm. In the efficacy evaluation of DBT-based cancer screening, the risk of death owing to other factors is individually determined, the total value of these factors is compared with the life-saving effect of the screening, and a profit analysis is performed. Therefore, dose evaluation is one of the important factors that helps determine the effectiveness of breast cancer screening (Miglioretti et al., 2016).

Breast thickness was only 40 mm in this study. Also, the shape was evaluated only with a semi-cylindrical shape. In the future, it is necessary to change the phantom thickness, phantom size and phamtom shape in detail, in consideration of clinical conditions, in order to examine the correction method of MGD according to the phantom size more accurately. In the future, by applying this evaluation method to other DBT equipment and calculating correction values for each equipment, the geometric conditions of the breast can be incorporated as MGD correction coefficients for each equipment.

Author Contribution Statement

Erika Nakajima and Hitoshi Sato substantially contributed to the study conceptualization and to data analysis and interpretation. All authors critically reviewed and revised the manuscript draft and approved the final version for submission.

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This article does not contain any studies with human participants.

Conflict of interest No conflict of interest.

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