Analysis of percent density estimates from
digital breast tomosynthesis projection images

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ABSTRACT

Women with dense breasts have an increased risk of breast cancer. Breast density is typically measured as the percent
density (PD), the percentage of non-fatty (i.e., dense) tissue in breast images. Mammographic PD estimates vary, in
part, due to the projective nature of mammograms. Digital breast tomosynthesis (DBT) is a novel radiographic method
in which 3D images of the breast are reconstructed from a small number of projection (source) images, acquired at
different positions of the x-ray focus. DBT provides superior visualization of breast tissue and has improved sensitivity
and specificity as compared to mammography. Our long-term goal is to test the hypothesis that PD obtained from DBT
is superior in estimating cancer risk compared with other modalities. As a first step, we have analyzed the PD estimates
from DBT source projections since the results would be independent of the reconstruction method. We estimated PD
from MLO mammograms (PD_M) and from individual DBT projections (PD_T). We observed good agreement between
PD_M and PD_T from the central projection images of 40 women. This suggests that variations in breast positioning, dose,
and scatter between mammography and DBT do not negatively affect PD estimation. The PD_T estimated from
individual DBT projections of nine women varied with the angle between the projections. This variation is caused by
the 3D arrangement of the breast dense tissue and the acquisition geometry.

Keywords: Methods: Quantitative image analysis, Effect of physical imaging parameters; Modalities:
Mammography, Digital breast tomosynthesis; Diagnostic Task: Risk assessment.

1. INTRODUCTION

Women with dense breasts have an increased risk of breast cancer.1-11 The ability to estimate breast cancer risk is of
great importance since it may allow customization of breast cancer detection and treatment, especially for patients at
high risk of breast cancer. It is hypothesized that by estimating breast density one can estimate cancer risk. The most
commonly used description of breast density is the American College of Radiology standardized four-point classification
scheme, Breast Imaging Reporting and Data System.12 While the BI-RADS classification represents a qualitative
description, a typically used quantitative measure of breast density is the percent density (PD), the percentage of non-
fatty (i.e., dense) tissue in breast images.

Mammographic PD estimates vary, in part, due to the projective nature of mammograms. Digital breast tomosynthesis
(DBT) is a radiographic method in which images of parallel slices of the breast are reconstructed from a small number of
projection (source) images, acquired at different positions of the x-ray focus.13, 14 DBT provides superior tissue
visualization and has improved sensitivity and specificity compared to mammography.15, 16

Our long-term goal is to test the hypothesis that PD obtained from DBT is superior in estimating cancer risk compared
with mammography. As a first step, we have analyzed the PD estimates from DBT source projections. Analysis of
projection images has value as it is independent of DBT reconstruction algorithms. Our focus was to evaluate the effects
of differences in acquisition parameters between mammography and DBT on PD estimation. DBT projections are
acquired using breast positioning similar to an MLO mammographic view; however, the two modalities differ in the
amount of breast compression, amount of scatter, and radiation dose. In addition, in DBT, multiple projections of the
breast are acquired at different angles. To evaluate the effects of the differences in acquisition parameters, we performed

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two studies. In the first study, we compared PD estimates from the MLO mammograms and central DBT projections. In the second study we tested the effect of varying the x-ray tube angle by comparing the PD estimates from individual DBT projections.

2. MATERIALS AND METHODS

2.1 Acquisition of clinical mammograms and DBT images
To date we have acquired DBT images of 52 women as a part of a clinical evaluation of multimodality breast imaging at the University of Pennsylvania. Each women in the study also had a digital mammogram. The DBT projections have been acquired using a Senographe 2000D (General Electric Medical Systems, Milwaukee, WI) digital mammography machine, modified to allow the x-ray focus to be positioned at nine angles, each 6.25° apart; covering a total angular range of [-25°,25°]. The nine DBT projections were taken using the same total dose as a standard 2-view mammographic exam and a slightly lower amount of compression. Figure 1 shows an example of clinical images used in this study.

This multimodality breast imaging study included volunteers with a high risk of breast cancer as computed by the Gail and Claus models, women referred to biopsy, and follow-up cancer cases. The clinical DBT images were acquired from the latter two groups. We have selected digital mammograms and DBT images from 40 women (mean age 51.4 years; range 31-80). We estimated PD from the breast contralateral to any finding, since the presence of a lesion could erroneously increase the PD estimate. Twelve women were excluded from our analysis. We excluded women with bilateral lesions and those for whom a unilateral study was performed. Women with breast implants were also excluded. The clinical images were acquired in the period between September 2004 and April 2005. The average lifetime Gail risk value for the population of 40 patients was 10.78% (range 1.8-30.3%).

2.2 Estimation of breast PD from clinical breast images
We have estimated PD from DBT projections using Cumulus, a software package developed at the University of Toronto. This software package provides the user with the ability to exclude a region of the breast from PD
estimation (e.g., the pectoral muscle region). The PD estimate is then computed as the ratio between the area within the manually thresholded region representing breast dense tissue and the area within the manually thresholded breast outline. The Cumulus package has been widely validated. Alternative methods for estimating PD from clinical mammograms have also been reported.

2.3 Correlation between PD estimated from MLO mammograms and central DBT projections

We have compared mammographic PD estimates \((PDM)\) and DBT PD estimates \((PDT)\) from the central DBT projections of the same breast. The \(PDM\) and central projection \(PDT\) values were compared by calculating their Pearson correlation coefficient and by calculating the slope and intercept of their linear regression.

Linear regression of the \(PDM\) and \(PDT\) values is formulated as: 
\[
(PDT)_{LR} = S \cdot PDM + I, 
\]
where \(S\) and \(I\) represent the linear regression slope and intercept, respectively, and are computed using the following equations:
\[
S = \frac{\sum(PDM \cdot PDT) - (\sum PDM)(\sum PDT)/N}{\sum (PDM)^2 - (\sum PDM)^2/N}, \quad \text{and} \quad I = E(PDT) - b \cdot E(PDM),
\]
where average values \(E(PDM)\) and \(E(PDT)\) have been computed over \(N\) selected cases.

The Pearson correlation coefficient \(r\) is defined as:
\[
r = \frac{\sum(PDM \cdot PDT)}{\sqrt{\sum PDM^2 \sum PDT^2}}. 
\]

The summation in Equations (1) and (2) is computed over \(N\) selected cases.

2.4 Correlation between PD estimates from individual DBT projections

Availability of clinical DBT projection images acquired at different angles of the x-ray tube, offers a unique opportunity to analyze the dependence of PD estimation on acquisition angle. We tested the angular dependence by computing the average values and the standard deviation (SD) of \(PDT\) estimates for individual DBT projections taken with a given tube angle. We also computed the average Pearson correlation coefficient between individual \(PDT\) estimates of DBT projections acquired at a given angular separation and between \(PDT\) of the central DBT projection and \(PDT\) of DBT projections acquired at a given angular distance from the central projection.

3. RESULTS

3.1 Comparison of \(PDM\) and central projection \(PDT\)

Figure 2 shows a scatter plot and the linear regression of PD computed from the central DBT projections and the corresponding MLO mammograms of 40 women. The Pearson coefficient of correlation between the \(PDM\) and the central projection \(PDT\) values is equal to 0.90, and the slope and intercept values of the linear regression are equal to 1.06 and 1.65%, respectively.

In order to test the reproducibility of the observers, we calculated the intra-observer variations of the PD estimates obtained using Cumulus for a subset of 9 women. Figure 3 compares the results of the reproducibility studies for \(PDM\) values (left) and the projection \(PDT\) values (right). The intra-observer Pearson correlation coefficient is equal to 0.92 for \(PDM\) and 0.94 for the central projection \(PDT\). The slope values of the linear regression are equal to 0.92 for \(PDM\) and 0.94 for \(PDT\), and the intercept values are equal to 6.61% for \(PDM\) and 2.05% for \(PDT\).
Figure 2: Correlation between PD estimates from mammograms and central DBT projections. Scatter plot of PD estimates from clinical mammograms (PD_M) and from central DBT projections (PD_T), obtained from contralateral breasts of 40 women. The central DBT projection is acquired with the same position of the x-ray focus as the corresponding MLO mammogram. Plotted is also the linear regression of the central projection PD_T values as a function of the PD_M values. The corresponding Pearson correlation coefficient is equal to 0.90.

Figure 3: Intra-observer variation. Scatter plot of mammographic PD_M estimates (left) and central DBT projection PD_T estimates (right) from two repeated studies of a subset of clinical images from 9 women. The DBT data and the corresponding mammograms were obtained from the contralateral breasts of 9 women. Plotted are also the intra-observer linear regressions of the PD_M and PD_T values. The Pearson correlation coefficients are equal to 0.92 for PD_M and 0.94 for PD_T.
3.2 Comparison of PD\textsubscript{T} from individual DBT projections
We have evaluated the dependence of PD on acquisition angle by comparing the PD\textsubscript{T} values calculated for each of the nine DBT projection angles; specifically, we computed the mean and SD of the PD at each DBT angle, and the correlation of PD estimates between different DBT angles. Figure 4 shows the mean and SD of the PD\textsubscript{T} estimates for nine women as a function of DBT projection angle.

![Figure 4: PD\textsubscript{T} estimates from individual DBT projections. Plotted are the mean and standard deviation (SD) of the PD\textsubscript{T} values for nine women (grey line) as a function of angle. Also, shown is the mean and SD of PD\textsubscript{M} for these women (black line). Error bars show ± one SD.](image)

Figure 5: Correlation between PD estimates from mammograms and individual DBT projections. Plotted are the mean and SD of the Pearson correlation coefficients between PD estimates from individual DBT projections acquired at a given angular separation (black line). Also, shown are the average values of the Pearson correlation coefficients between PD\textsubscript{T} of the central DBT projection and PD\textsubscript{T} at a given angular separation (grey line). Error bars, when shown, are equal to ± one SD.
Figure 5 shows the average values of the Pearson correlation coefficients between $PD_T$ for DBT projections which are separated by the same angular distance (black line). The average correlations were computed as follows. We first computed the Pearson correlation between any two vectors of $PD_T$ values, corresponding to DBT projections separated by the same angle. (The vector elements are the $PD_T$ values for different women.) For example, the average correlation value for an angular distance of 6.25° was obtained by averaging the Pearson correlations corresponding to eight pairs of DBT projections, each pair separated by 6.25° (i.e., $T_1$ and $T_2$, $T_2$ and $T_3$, ..., $T_8$ and $T_9$; see Figure 1). Similarly, the average correlation value for the angular distance of 12.5° was obtained by averaging the Pearson correlation coefficients corresponding to seven pairs of projections each 12.5° apart (i.e., $T_1$ and $T_3$, $T_2$ and $T_4$, ..., $T_7$ and $T_9$). This procedure was repeated for all angular separations.

We have also computed the corresponding SD values for each angular separation. Error bars in Figure 5 (black line) show ± one SD. Note that the average correlation value for an angular separation of 50° was computed using the Pearson correlation for a single pair of projections $T_1$ and $T_9$ and that the average correlation for the zero separation was set to 1 by default. Thus, the SD values for angular separations of 0° and 50° are not shown.

Figure 5 also shows the average values of the Pearson correlations between the central projection $PD_T$ and $PD_T$’s from individual DBT projections acquired at a given angular distance from the central projection (grey line). Error bars show ± one SD.

4. DISCUSSION

4.1 Effects of differences in acquisition parameters between mammography and DBT on PD estimation

It can be seen from Figure 2 that there is good agreement between $PD_M$ and the central projection $PD_T$. These results suggest that the differences in acquisition parameters (i.e., differences in the amount of compression, radiation dose, and scatter, and variation in positioning between mammography and DBT) do not significantly influence PD estimation. These results are also supported by studies published in the literature of PD estimates obtained using very low radiation dose and with variation in acquisition parameters.

4.2 Angular dependence of $PD_T$ estimates

Figure 4 shows that the $PD_T$ does not vary significantly with acquisition angle. We also observed that the SD of $PD_T$, computed over all DBT angles, for each individual woman is very low (range 1%-7%). Although the observed variations were small, we were concerned that the changes in collimation as a function of angle would lead to a bias in our results. Note that a portion of each projection image is occluded by the collimator (see Figure 1). This occurs because the fulcrum of the x-ray tube rotation does not coincide with the position of the breast (see Figure 6).

We performed two additional studies to elucidate the effects of the collimation. First, we masked the central projection image to match the collimation for each DBT projection. We then calculated the difference in $PD_T$ estimate for the image acquired at that angle and the central projection image masked according to the collimation at that angle.

Figure 7 (upper graph) shows the mean and SD of the difference between $PD_T$ and the PD of the central projection image masked to match the collimation at a given angle. Error bars show ± one SD. For comparison, we show the SD of the $PD_T$ of the central projection on the same graph. It can be seen that the SD of the difference between the projection $PD_T$ and the $PD_T$ from the masked central projection are 2-6 times smaller than the SD of the individual projection $PD_T$. This suggest that $PD_T$ values from the masked central projection are highly correlated with the $PD_T$ values from individual projection images.
Second, we applied the same mask to each projection image and calculated the PD_T. We used a mask formed by the union of the collimated regions of the breast. As a result, only the same central portion of each projection image was used to estimate PD_T.

It can be seen from Figure 7 (lower graph) that there are differences between the PD_T estimated at a given angle. A possible cause of these differences is that, due to the projection acquisition geometry, there is motion of the fibroglandular (i.e., dense) tissue region in one DBT projection relative to the other (see Figure 6). In combination with the applied masking, this motion results in different amounts of dense tissue being visible in the central image portion and hence used to estimate PD_T. This effect is illustrated in Figure 8. Without collimation, the whole area of the dense tissue will be visualized in each projection, thus reducing the effects of the projective geometry on PD estimation.

Our department has recently installed a new breast imaging system, Senographe DS (General Electric Medical Systems, Milwaukee, WI), which is optimized for DBT acquisition. The collimation effects, described in this section, have been eliminated in the new system.
Figure 7: Analysis of the effects of non-uniform collimation on PD estimation from DBT projections. (Upper graph) Plotted are the mean and SD of the differences between PD_T of individual projections and PD_T of the central projection masked to match the collimation for each DBT projection (black line). Solid error bars show ± one SD. Also, plotted is an error bar corresponding to ± one SD for the central projection PD_T (grey line). (Lower graph) Plotted are the mean and SD of PD_T of individual DBT projection (black line) and the PD_T estimated when the same mask is applied to each projection image (grey line). The mask was formed by the union of the collimated regions in all projections. Error bars show ± one standard error.
5. CONCLUSIONS

We have estimated the PD from DBT projections (PD_T) and mammograms (PD_M) for 40 women. We have computed the Pearson correlation coefficients and the linear regression between PDM and PDT. There is little variation in PDM and the PDT of the central DBT projection, suggesting that the differences in acquisition parameters between mammography and DBT do not affect PD estimation. In addition, we analyzed the angular dependence of PD_T for 9 women. The obtained results do not show a significant variation with acquisition angle. We did observe small variations due to the projective geometry and the variation in collimation as a function of angle. To the best of our knowledge, this study represents the first analysis of the variation in 2D estimates of the breast percent density as a function of the projection angle.

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