Spatial variations in optical and physiological properties of healthy breast tissue

Natasha Shah  
Albert E. Cerussi  
Dorota Jakubowski  

University of California, Irvine  
Beckman Laser Institute  
1002 Health Sciences Road  
Irvine, California 92612

David Hsiang  
John Butler  

University of California Medical Center  
Department of Oncological Surgery  
101 The City Drive South  
Orange, California 92868

Bruce J. Tromberg  

University of California, Irvine  
Beckman Laser Institute  
Irvine, California 92612  
E-mail: tromberg@bli.uci.edu

Abstract. Near-infrared (NIR) diffuse optical spectroscopy (DOS) and diffuse optical imaging (DOI) show promise as noninvasive clinical techniques for breast cancer screening and diagnosis. Since NIR methods are based on optical contrast between healthy and diseased tissue, it is essential to characterize the sources of endogenous contrast in normal subjects. We report intra- and inter-subject variation and bilateral asymmetry of the optical and physiological parameters of 31 women using a seven-wavelength NIR frequency-domain photon migration (FDPM) instrument. Wavelength-dependent absorption and reduced scattering parameters (\(\mu_a\) and \(\mu_s^'\), respectively) were measured in four major quadrants and the areolar regions of left and right breasts. These values were used to determine tissue concentrations of oxy- (HbO\(_2\)) and deoxy- (Hb-R) hemoglobin, lipid content, water concentration, and tissue "scatter power." Mean total hemoglobin for premenopausal (PRE) women (20 to 30 \(\mu\)M) is approximately two-fold higher than for postmenopausal (POST) subjects at all positions. POST women have approximately 50% higher lipid content (50 to 60%) than PRE at all positions. Water concentration on average is 1.8-fold higher for PRE subjects (30 to 40%) than POST. These differences are most pronounced when comparing the areolar complex to the other regions of the breast. In premenopausal women, the areolar regions have 40 to 45% increased total hemoglobin concentration (THC), 20 to 25% lower lipid content, and 30 to 60% higher scatter power versus the quadrants. Small-scale (3 cm) changes in optical properties are negligible compared to large-scale variations over all quadrants, where the intrinsic spatial heterogeneity of healthy breast tissue is 20 to 40% for \(\mu_a\) and 5 to 12% for \(\mu_s^'\). Although no consistent right-left differences are observed in the study population, relative differences between symmetric positions ranged from 18 to 30% for THC, 10 to 40% for adipose, 10 to 25% for water, and 4 to 9% for scattering (674 nm) within an individual. © 2004 Society of Photo-Optical Instrumentation Engineers. [DOI: 10.1117/1.1695560]

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1 Introduction

A variety of near-infrared (NIR) optical methods for the detection, diagnosis, and clinical management of breast cancer are under development. In general, near-infrared (NIR) methods are based on photon migration models and technology, such as diffuse optical spectroscopy (DOS) and diffuse optical imaging (DOI). DOS andDOI methods provide unique quantitative physiological information that can be used in conjunction with conventional medical imaging techniques, such as magnetic resonance imaging and ultrasound. Recent applications of diffuse optical methods in breast cancer include monitoring chemotherapy, hormonal effects, characterizing tumors, and assessing disease risk.

In this work, we employ a DOS instrument based on frequency-domain photon migration (FDPM) technology. FDPM utilizes intensity-modulated light to separate and quantify intrinsic tissue absorption and scattering in vivo. Measurements of absorption coefficients (\(\mu_a\)) at multiple wavelengths are used to calculate concentrations of the principal NIR breast tissue chromophores: deoxy-hemoglobin, oxy-hemoglobin, lipids, and water. Measurements of the wavelength dependence of the reduced scattering parameter (\(\mu_s^'\)) is related to the size and distribution of biological scatterers. Hence, the spectral behavior of both \(\mu_a\) and \(\mu_s^'\) can be used for the noninvasive physiological characterization of soft tissues.

Several investigators have demonstrated the ability of DOS and DOI methods to distinguish malignant from benign breast
lesions in vivo. Preliminary studies have also described the effects of biological factors such as age, menopausal status, hormone use, and menstrual cycle fluctuations on optical properties in normal tissue. This study focuses on analyzing the intrinsic spatial variations of breast tissue optical properties in 31 healthy subjects.

Optical property variations across a small subsection of tissue (3 cm) are compared to intrasubject variations over the entire breast. The effects of age and menopausal status on the spatial distribution of optical and physiological parameters are also examined. In addition, since bilateral asymmetry in breasts is common in women, we present data on the typical differences in optical properties between left and right breasts. Our results provide insight concerning the glandular distribution, vascular patterns, and asymmetry within healthy breast tissue. These factors play an important role in characterizing normal breast tissue physiology and understanding the appearance of disease.

2 Materials and Methods

2.1 Clinical FDPM Measurements

A 1-GHz portable, multwavelength, high-bandwidth frequency-domain photon migration instrument has been designed and optimized for clinical optical property studies (Fig. 1). The FDPM instrument and theory has been described in detail elsewhere. Briefly, the network analyzer is used to produce modulation frequencies from 50 MHz to 1 GHz. A DC current source is mixed with RF power provided by the network analyzer in a bias network, which distributes power to one of seven laser diodes (674, 780, 803, 849, 894, 915, and 980 nm) to produce amplitude-modulated light. An optical switch delivers light serially from each diode to a 100-μm graded index optical fiber that delivers diode laser output to the tissue. A hand-held probe has been designed to house an avalanche photodiode (APD) that records the diffuse light signals after propagation through the tissue. The probe contains a plastic attachment on the casing to secure the source fiber at a fixed distance of 21 mm from the APD. The network analyzer measures the phase and amplitude of the electronic signal.

Amplitude-modulated light traverses through tissue as a photon density wave (PDW) with a distinct phase and amplitude. The phase shift and amplitude of the PDW after propagation through tissue was determined at each wavelength between 50 to 700 MHz at 1.375 MHz intervals. The range of modulation frequencies was swept repetitively so that each amplitude and phase value represents an average of four measurements. The measurement time over all seven wavelengths is less than a minute. The optical power launched into the subject ranged from 5 to 25 mW for each diode. The instrument response was calibrated before each measurement session on a tissue phantom of known optical properties.

FDPM data was collected from a total of 31 volunteers: 18 premenopausal women ages 18 to 48 years of age (PRE) and 13 postmenopausal women ages 52 to 64 (POST), six of whom were using hormone replacement therapy. Measurements on 28 of the subjects were performed at five distinct positions on each breast (Fig. 2): one position at each of the four quadrants midway between the nipple and the edge of the breast and the superior areolar border. The measurement locations were abbreviated as follows: R = right, L = left, U = upper, L = lower, O = outer, I = inner, and A = areolar, thus LDU denotes the left lower outer quadrant. Three repeat measurements were made by removing and replacing the probe in each location to ensure accuracy. Data from five normal premenopausal volunteers ages 27 to 47 were obtained to provide information on variations in a small region of tissue. A series of seven measurements were acquired in 5-mm increments over a 3-cm line in the upper outer quadrants of both breasts (Fig. 2, inset). Two repeat scans were performed. To ensure reproducible and accurate probe placement, all sites were measured and marked with a surgical pen. All data was collected with subjects resting in the supine position. Experiments were conducted in adherence to University of California (UC) Irvine IRB approved protocols 95-563 and 99-2183. After providing informed consent, volunteers filled out a brief questionnaire, which surveyed pertinent medical history.

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2.2 Theoretical Model

The $P_1$ approximation to the Boltzmann transport equation was used to extract tissue absorption ($\mu_a$) and reduced scattering ($\mu_s'$) coefficients from frequency-dependent phase and amplitude curves.\textsuperscript{21,22} The model employs an extrapolated boundary condition for a semi-infinite geometry.\textsuperscript{23} To determine the optical properties from a given set of frequency-dependent data, a Marquardt-Levenburg $\chi^2$ minimization algorithm was used to simultaneously fit the amplitude and phase by minimizing the difference between the measured values and those predicted by the $P_1$ approximation.

Physiological properties were calculated from the determined $\mu_a$ values measured at seven wavelengths by assuming breast tissue is composed of four principal NIR absorbers: deoxy-hemoglobin (Hb-R), oxy-hemoglobin (HbO$_2$), lipids, and water. Melanin was not included as an absorber, since the contribution of the epidermis to the total sampled volume of tissue is small.

Hemoglobin concentrations are measured in $\mu$M, lipid content is measured as a mass density percentage, and water is calculated relative to pure water, 55.6 M. Total hemoglobin concentration (THC) is \([Hb]+[HbO_2]\), and tissue hemoglobin oxygen saturation ($S_{O_2}$) is \([HbO_2]/THC\)*100%. The four chromophore concentrations are determined using a least-squares solution to $Ec = \mu_a$, where $E$ is a $7 \times 4$ matrix of the molar extinction coefficients and $c$ is the concentration of the chromophores.\textsuperscript{24–26} In matrix representation, the chromophore concentration is given by: $\tilde{c} = (\tilde{E}^T\tilde{E})^{-1}\tilde{E}^T\mu_a$, where $\tilde{E}^T$ and $\tilde{E}^{-1}$ denote the transpose and inverse of the matrix $\tilde{E}$, respectively.

The spatial heterogeneity in an individual was calculated from the normalized standard deviation (NSD) of the values determined at ten breast measurement locations (five positions on each breast). The NSD equals the standard deviation reflected as a percentage of the mean: $\left(\sigma/\bar{x}\right)\times 100\%$.

3 Results

To measure the variability in physiological and optical properties within a small subsection of tissue (3 cm), a series of seven measurements were made in 5-mm increments in the upper outer quadrants of five normal premenopausal subjects. The variability of each parameter was assessed using the normalized standard deviation. The results of the right breast are displayed in Table 1. The change over a 3-cm span is small for $\mu_s'$ (less than 10%), $S_{O_2}$ (less than 5%), and lipid content (less than 11%). Scatter power and water content are the most variable parameters (10 to 20%). Hemoglobin variability (oxy-, deoxy-, and total) for the five subjects ranges from 2 to 16%.

The spatial variations of healthy breast tissue over the entire breast and age-dependent trends in tissue heterogeneity are displayed in Figs. 3–6. Figure 3 illustrates the mean total

Table 1 Variation of optical and physiological properties in a small volume of tissue for five subjects. Values represent the normalized standard deviation of seven measurements made over a 3-cm span of tissue in the right upper outer quadrant.

<table>
<thead>
<tr>
<th>Optical/physiological parameter</th>
<th>Subject 1</th>
<th>Subject 2</th>
<th>Subject 3</th>
<th>Subject 4</th>
<th>Subject 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>27</td>
<td>35</td>
<td>35</td>
<td>40</td>
<td>47</td>
</tr>
<tr>
<td>$\mu_s'$</td>
<td>3</td>
<td>3</td>
<td>3</td>
<td>6</td>
<td>4</td>
</tr>
<tr>
<td>Scatter power</td>
<td>9</td>
<td>16</td>
<td>9</td>
<td>15</td>
<td>2</td>
</tr>
<tr>
<td>$S_{O_2}$</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>THC</td>
<td>9</td>
<td>16</td>
<td>9</td>
<td>14</td>
<td>2</td>
</tr>
<tr>
<td>Water</td>
<td>14</td>
<td>12</td>
<td>9</td>
<td>3</td>
<td>12</td>
</tr>
<tr>
<td>Lipid</td>
<td>3</td>
<td>5</td>
<td>5</td>
<td>5</td>
<td>4</td>
</tr>
</tbody>
</table>

Fig. 3 Mean total hemoglobin concentration at each measurement position for PRE ($n = 15$) and POST ($n = 13$) subject groups. Values are determined from wavelength-dependent absorption values for seven wavelengths ranging from 674 to 980 nm. Error bars represent the standard deviation of the mean.
hemoglobin concentration for PRE and POST subject groups at each quadrant position. Mean total hemoglobin for PRE women is approximately two-fold higher than for POST women at all positions. Areolar total hemoglobin levels are approximately 30 to 45% higher than average quadrant levels for PRE subjects, and 25 to 40% higher for POST subjects (89 and 94% confidence interval for left and right breast, respectively). The lower quadrants of PRE women have lower total hemoglobin concentrations relative to the upper quadrants (83% confidence interval).

Water concentration on average is 1.8-fold higher for PRE subjects than POST (Fig. 4). Areolar water concentration is 30 to 50% higher than water concentration for the quadrants for PRE and POST women (84 and 94% confidence interval for left and right breasts, respectively). For PRE women, however, the lower quadrants have more water than the upper quadrants (86% confidence interval); the same pattern is not evident for POST women.

Average lipid content for PRE and POST subject groups at each measurement position are shown in Fig. 5. POST women have approximately 50% higher lipid content than PRE women at all positions. Aerolar lipid content is 25 to 35% lower than average lipid content in the four quadrants for PRE and 10% lower in POST women (74% confidence interval). The lower outer quadrants have lower adipose content that the other three quadrants in PRE women (88% confidence interval); this pattern does not persist in POST women.

The mean \( S_{O_2} \) values calculated over the entire breast are 74±6 and 75±8% for PRE and POST subject groups, respectively. The left upper outer quadrant exhibits differences between subject groups (75.6±6% for PRE and 80±9% for POST), consistent with our previous reports, though this trend does not appear in the other measurement positions. Aerolar regions have significantly lower saturation values, (69.7±7% for PRE and 69.8±12% for POST, 97% confidence interval). There are no significant variations in the quadrants.

A plot of the reduced scattering coefficient at 674 nm for the two subject groups demonstrates that \( \mu_s' \) shows little positional variation (Fig. 6(a)). The inner quadrants have higher
scattering on average than outer quadrants (approximately 0.05 to 0.1 mm\(^2\)) for both subject groups (77 and 92\% confidence intervals for right and left breasts, respectively).

The wavelength dependence of scattering (i.e., scatter power) is influenced by the fat, collagen, and epithelial content. The scatter power (\(\mu_s' = A_k \cdot \text{scatter power}\)) is shown for each position in Fig. 6(b). PRE women have a scatter power that is 1.5-fold higher than POST women. The areolar scatter power is 30 to 60\% greater in PRE women, and 15 to 30\% greater in POST women (99 and 88\% confidence intervals for left and right breasts, respectively). The right areola has a larger scatter power than the left on average by about 15 to 25\% (97\% confidence interval).

The normalized standard deviation of the physiological parameters over the eight breast quadrants was calculated for each of the volunteers to quantify the variability typically found in healthy breast, and to investigate age-related changes in heterogeneity. Only lipid content showed significant changes in heterogeneity with age (Fig. 7). PRE and POST women typically (\(\geq 89\%\) of volunteers) have a NSD of 30\% or lower for adipose tissue, and lipid heterogeneity decreases with age (\(R=0.60\)). The NSD is typically (\(\geq 89\%\) of volunteers) less than 40\% for THC, less than 30\% for water, less than 12\% for \(\mu_s'\) (at 674 nm), and less than 25\% for the scatter power. The normalized standard deviation for \(S_{O2}\) is less than 12\% for all but one (97\%) of the 28 individuals.

### Table 2

<table>
<thead>
<tr>
<th>Quadrant</th>
<th>Water</th>
<th>THC</th>
<th>Scatter power</th>
<th>Lipid</th>
<th>(\mu_s')</th>
</tr>
</thead>
<tbody>
<tr>
<td>Upper outer</td>
<td>22</td>
<td>20</td>
<td>22</td>
<td>20</td>
<td>7</td>
</tr>
<tr>
<td>Other quadrants</td>
<td>15</td>
<td>10</td>
<td>20</td>
<td>15</td>
<td>7</td>
</tr>
<tr>
<td>Areolar</td>
<td>20</td>
<td>18</td>
<td>10</td>
<td>40</td>
<td>9</td>
</tr>
</tbody>
</table>

A small degree of asymmetry between breasts is common in women. To examine the differences between the right and left breasts of healthy individuals, the relative difference (right-left/average) *100\% for the four quadrants of each subject was measured (Table 2). Although differences between right and left breasts exist in individuals, the results found no consistent differences between the right and left breasts in the population for \(\mu_s'\), water, lipid, THC, or \(S_{O2}\). The right areolar region, however, has significantly higher mean scatter power values than the left, and equal or greater values were measured in 23/26 cases. The question of whether there is regular breast volume asymmetry remains unsolved. The relative difference between symmetric positions can range from 18 to 30\% for THC, 10 to 40\% for adipose, 10 to 25\% for water, and 4 to 9\% for scattering (674 nm), depending on position (Table 2).

### 4 Discussion

Optical techniques for breast cancer detection are based on the assumption that differences in tissue absorption and scattering can provide a basis for tumor characterization and clinical management. Thus, the accuracy and efficacy of optical mammography requires detailed study of the multiple factors affecting breast tissue heterogeneity and composition. Breast tissue is nonuniform; furthermore, physiological changes that occur over a woman’s lifetime can affect the distribution and composition of healthy breast tissue. With the onset of menopause, glandular tissue involutes and is replaced by adipose, which produces a new spatial distribution of chromophores. In addition, this process is nonuniform, which changes the inherent heterogeneity of the breast tissue. Reduced scattering (\(\mu_s'\) and tissue hemoglobin oxygen saturation (\(S_{O2}\)) showed \(\leq 12\%\) variation over the entire breast and \(\leq 6\%\) over a 3-cm region. Alterations in \(\mu_s'\) or \(S_{O2}\) may signify regions of breast disease. Previous studies have shown that increased spatial heterogeneity in scattering may be indicative of normal tissue or benign breast disease, while regions of more uniform scattering may suggest invasive cancer. Significantly higher \(\mu_s'\) values observed for the inner quadrants [Fig. 6(a)] may be due to contributions of the sternal border and pectoralis muscle to the optical signal. Changes in total hemoglobin over small regions of tissue (5 to 11\%) may be attributed to spatial variations in underlying blood vessels in the probed tissue volume.

In PRE women, the areolar region displayed 25 to 30\% lower lipid content, 30 to 50\% higher water content, 30 to 60\% higher scatter power, and 5\% lower \(S_{O2}\) values when compared to the quadrants (Figs. 3–6). These results are consistent with the unique physiology of the areola. Since lactiferous ducts open at the surface of the nipple, glandular tissue is concentrated and superficial at the areola. The ducts are surrounded by highly vascularized connective tissue; therefore, our measurements of elevated hemoglobin and reduced adipose content in this region of the breast are consistent with known breast structure. Lower \(S_{O2}\) at the areola may be due to the higher metabolic demands of glandular tissue. The areolar complex lacks fat and is mostly comprised of dense fibrous tissue and smooth muscle; the surrounding pigmented areola contains apocrine and eccrine glands. As a result, the areola has a greater scatter power, indicative of
smaller scattering particles. In POST women, the areolar region had 25 to 40% higher hemoglobin, 10% lower lipid content, and 15 to 30% greater scatter power. Since glandular tissue is replaced with fat after menopause, the optical contrast of the areolar region is less pronounced in POST subjects. 

Deeper pigmentation of the areolar region may be significant in an individual, although no consistent right-left differences are observed in a population. The intrinsic spatial heterogeneity of healthy breast tissue is 20 to 40% for \( \mu_s \) and 5 to 12% for \( \mu_s' \). These values are important because they establish the fundamental sensitivity limits for detecting breast disease based on optical contrast.

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