

Image reconstruction method for a two-layer tissue structure accounts for chest-wall effects in breast imaging

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Abstract. We develop a new tomographic imaging reconstruction algorithm for a two-layer tissue structure. Simulations and phantom experiments show more accurate reconstruction of target optical properties compared with those results obtained from a semi-infinite tissue model for layered structures. This improvement is mainly attributed to the more accurate estimation of background optical properties and more accurate estimation of weight matrix for imaging reconstruction by considering the light propagation effect in the second layer. Clinical results of breast lesions are also presented to demonstrate the utility of this new imaging algorithm. © 2008 Society of Photo-Optical Instrumentation Engineers. [DOI: 10.1117/1.3041497]

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1 Introduction

In the last two decades, many theoretical and numerical methods have been developed to model photon migration in layered tissue structures.¹⁻¹⁴ One well-established method is Monte Carlo (MC) simulation, which can be used to characterize the light propagation in complex tissue structures.^{7,10,11} However, the MC method is very computationally intensive. Analytical methods have been developed by many research groups to model light propagation in layered tissue structures. The publications in this regard have mostly focused on estimation of the dependence of the surface reflectance measurements on the optical properties of the two tissue layers.^{1,2,13} The widely referenced paper by Kienle et al.³ provided solutions of time- and frequency-domain diffusion equations for two-layer tissue structures. These solutions could be used to estimate tissue optical properties and in some cases the layer thickness by using nonlinear fitting methods to the measured reflectance data.

To the best of our knowledge, there has been little effort to develop tomographic reconstruction algorithms suitable for layered tissue structures. In breast imaging, a semi-infinite medium is an accurate approximation when the patient is scanned in a supine position and the breast thickness is 2 cm or more. However, the semi-infinite model is not accurate enough when breast tissue thickness is less than 1.5 to 2 cm with the chest wall underneath. Under this condition, the chest wall contributes to the total reflectance measurements at the skin surface. The optical properties of breast tissue and the chest wall can be estimated using the two-layer analytical solution with the layer thickness provided by coregistered ultrasound. The accuracy of reconstructed target optical properties can be improved by implementing a layered structure in

the imaging process as opposed to the semi-infinite tissue structure.¹⁵

In a recent publication, we discussed the development of the analytical solution for a two-layer tissue structure with a tilted interface to improve the estimation accuracy of the optical properties of breast tissue and the chest wall.¹² In this paper, we extend the analytical solution of a two-layer tissue structure developed by Kienle et al.³ to tomographic imaging. The Born approximation and the dual-zone mesh scheme we developed earlier are used in the image reconstruction.^{16,17} Simulations and phantom experiments are performed to validate the image reconstruction. Further, clinical examples are given to demonstrate the utility of this layer-structure-based tomography algorithm.

2 Theory

We briefly summarize the forward model of light propagation used in this paper in Sec. 2.1, and discuss our newly developed image reconstruction by combining the Born approximation with a dual-zone mesh scheme for the layered structure.

2.1 Forward Model

The diffusion equations were solved by Kienle et al.³ using the Fourier transform approach for a two-layer turbid medium with a semi-infinite second layer. We directly adopted this method for tomographic imaging and a brief description of the analytical solution is given here for the sake of completeness. A two-layer turbid medium is considered with upper layer thickness l and a semi-infinite second layer. An infinitely thin beam of frequency-modulated light is incident perpendicular on the top surface of a two-layer turbid medium. The beam is scattered isotropically at the depth of $z_0 = (\mu_{a1} + \mu'_{s1})^{-1}$, where the μ_{a1} and μ'_{s1} are the absorption and reduced scattering coefficients of the first layer, and z_0 is in the range of 1 to 2 mm for typical background breast-tissue ab-

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sorption and scattering properties. The diffusion equation for the photon density wave at two layer structure is given as

$$D_1 \nabla^2 \Phi_1(r, \omega) - \left(\mu_{a1} - j \frac{\omega}{v_m} \right) \Phi_1(r, \omega) = -\delta(x, y, z - z_0) \quad 0 < z < l, \quad (1)$$

$$D_2 \nabla^2 \Phi_2(r, \omega) - \left(\mu_{a2} - j \frac{\omega}{v_m} \right) \Phi_2(r, \omega) = 0 \quad l \leq z, \quad (2)$$

where $r = (x^2 + y^2 + z^2)^{1/2}$, $D_i = (1/3)(\mu_{ai} + \mu'_{si})^{-1}$ are the diffusion coefficients, μ_{ai} and μ'_{si} are the absorption and reduced scattering coefficients, and Φ_i are the fluence rates for the layers $i=1$ and 2 , respectively. The modulation frequency of the source is $\omega = 2\pi f$, v_i is the speed of light in the medium, and j is the imaginary unit.

The diffusion equations were first transformed to ordinary differential equations with the use of a 2-D Fourier transform

$$\phi_i(s_1, s_2, z, \omega) = \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} \Phi_i(x, y, z, \omega) \exp[j(s_1 x + s_2 y)] dx dy, \quad (3)$$

where s_1 and s_2 are spatial frequencies in the x and y directions. The derived equations are then solved by using the following boundary conditions,

$$\phi_1(z = -z_b, r) = 0, \quad (4)$$

$$\phi_1(r \rightarrow \infty) = 0, \quad (5)$$

$$\frac{\phi_1(l, r)}{\phi_2(l, r)} = \frac{n_1^2}{n_2^2}, \quad (6)$$

$$D_1 \frac{\partial \phi_1(z)}{\partial z} \Big|_{z=l} = D_2 \frac{\partial \phi_2(z)}{\partial z} \Big|_{z=l}. \quad (7)$$

The final results are then inverse Fourier transformed to obtain the solution of Eqs. (1) and (2). The results for $\phi_1(z, s, \omega)$ at different tissue layers are

$$\phi_1(z, s, \omega) = \frac{\sinh[\alpha_1(z_b + z_0)]}{D_1 \alpha_1} \frac{D_1 \alpha_1 \cosh[\alpha_1(l - z)] + (n_2/n_1)^2 D_2 \alpha_2 \sinh[\alpha_1(l - z)]}{D_1 \alpha_1 \cosh[\alpha_1(l + z_b)] + (n_2/n_1)^2 D_2 \alpha_2 \sinh[\alpha_1(l + z_b)]} - \frac{\sinh[\alpha_1(z_0 - z)]}{D_1 \alpha_1} \quad 0 \leq z < z_0, \quad (8)$$

$$\phi_1(z, s, \omega) = \frac{\sinh[\alpha_1(z_b + z_0)]}{D_1 \alpha_1} \frac{D_1 \alpha_1 \cosh[\alpha_1(l - z)] + (n_2/n_1)^2 D_2 \alpha_2 \sinh[\alpha_1(l - z)]}{D_1 \alpha_1 \cosh[\alpha_1(l + z_b)] + (n_2/n_1)^2 D_2 \alpha_2 \sinh[\alpha_1(l + z_b)]} \quad z_0 \leq z < l, \quad (9)$$

$$\phi_2(z, s, \omega) = \frac{\sinh[\alpha_1(z_b + z_0)] \exp[\alpha_2(l - z)]}{D_1 \alpha_1 \cosh[\alpha_1(l + z_b)] + (n_2/n_1)^2 D_2 \alpha_2 \sinh[\alpha_1(l + z_b)]} \quad z \geq l, \quad (10)$$

where $s^2 = s_1^2 + s_2^2$, $\alpha_i^2 = (D_i s^2 + \mu_{ai} - j\omega/v_i)/D_i$; z_b is the extrapolated boundary¹⁸ given by $z_b = 2D_1 \cdot (1 + R_{\text{eff}})/(1 - R_{\text{eff}})$; and R_{eff} depends on the refractive indices of the tissue layer and represents the fraction of photons internally diffused at the boundary.

The fluence rate in the spatial domain (x, y, z) can be obtained by spatial Fourier inversion:

$$\Phi_i(\rho, z, \omega) = \frac{1}{2\pi} \int_0^{\infty} \phi_i(s, z, \omega) s I_0(s\rho) ds, \quad (11)$$

where I_0 is the Bessel function of zero order, and $\rho = [(x - x_s)^2 + (y - y_s)^2]^{1/2}$, where x_s and y_s are spatial coordinates of the source position. Thus, Eq. (11) combined with either Eq. (8) (for depth $0 \leq z < z_0$), Eq. (9) (for depth $z_0 \leq z < l$), or Eq. (10) (for $z \geq l$) provides the incident fluence rate at any depth inside the two-layer medium.

2.2 Image Reconstruction Using Two-Layer Tissue Model

In the Born approximation, the photon density wave originating from a source at r_s and measured inside the tissue at r , is a linear superposition of its incident Φ_0 (homogeneous) and the scattered Φ_{sc} (heterogeneous) parts given as $\Phi(r_s, r, \omega) = \Phi_0(r_s, r, \omega) + \Phi_{\text{sc}}(r_s, r, \omega)$. By using the first-order solution to the heterogeneous equation under the approximation that $\Phi_{\text{sc}}(r_s, r, \omega) \ll \Phi_0(r_s, r, \omega)$, we obtain $\Phi_{\text{sc}}(r_d, r_s, \omega) = -(1/D_0) \int G(r, r_d, \omega) \Delta\mu_a(r) \Phi_0(r_s, r, \omega) d^3r$, where r_d represents detector position, $G(r, r_d, \omega)$ represents the scattered field at r_d due to the point scattered at r , $\Delta\mu_a(r)$ is the change of absorption coefficient at voxel at location r , and D_0 is the diffusion coefficient of the homogeneous medium. Figure 1 illustrates the source position r_s , field point r , detector position r_d , and ρ used for computing the incident field.

When using the two-layer model, $\Phi_0(r_s, r, \omega)$ is calculated based on the depth of the voxel to which the fluence is being computed. For voxels in the three regions the fluence rates are represented by Φ_{00} , Φ_{01} , and Φ_{02} , and D_0 is replaced by D_1

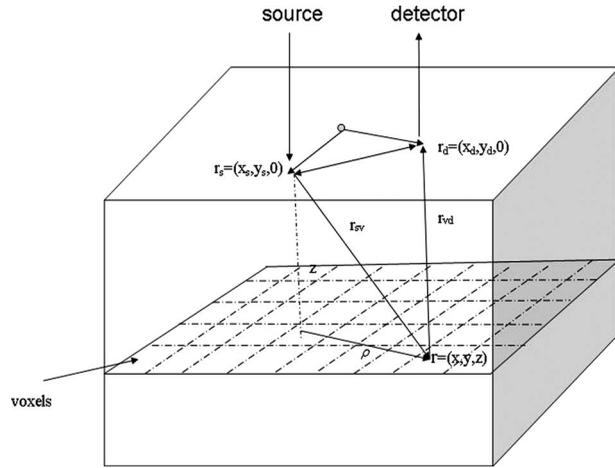


Fig. 1 Configuration for simulations and reconstructions.

or D_2 , depending on the voxel location in the layered structure. The Green function $G(r, r_d, \omega)$ of the homogeneous equation at the detector r_d is given as $G(r, r_d, \omega) = (D_i/S)\Phi_0(r, r_d, \omega) = \Phi_i[(x_d - x)^2 + (y_d - y)^2]^{1/2}, z, \omega$, which could be calculated again using spatial Fourier inversion from any one of Eq. (8) (if voxel depth $z < z_0$), Eq. (9) (if voxel depth $z_0 \leq z < l$), and Eq. (10) (if $z \geq l$), depending on the voxel depth from the surface.

The dual-zone mesh scheme introduced by us earlier¹⁶ was used to segment the imaging volume into lesion and background regions designated as L and B , respectively. The weight matrix is calculated using the two-layer model as $[\mathbf{U}_{sd}]_{M \times 1} = [\mathbf{W}_L, \mathbf{W}_B]_{M \times N} [\mathbf{M}_L, \mathbf{M}_B]^T$, where $\mathbf{W}_L = [-G(r_{vj}, r_{di})\Phi_0(r_{si}, r_{vj})/D_l]_{M \times N_L}$ and $\mathbf{W}_B = [-G(r_{vk}, r_{di})\Phi_0(r_{si}, r_{vk})/D_l]_{M \times N_B}$ are weight matrices for lesion and background volumes, respectively; $[\mathbf{M}_L] = [\int_{V_L} \Delta\mu_a(r')d^3r', \dots, \int_{N_L} \Delta\mu_a(r')d^3r']$ and $[\mathbf{M}_B] = [\int_{V_B} \Delta\mu_a(r')d^3r', \dots, \int_{N_B} \Delta\mu_a(r')d^3r']$ are the total absorption distributions of lesion and background volumes, respectively; and D_l equals D_1 or D_2 , depending on the voxel location in the layered structure. The dual-zone mesh scheme significantly reduces the total number of voxels with unknown optical properties and dramatically improves the convergence of inverse mapping of target optical properties.^{16,17}

3 Materials and Methods

3.1 Simulations

The imaging reconstruction algorithm developed in Sec. 2 was validated using forward data generated by the finite-element method (FEMLAB 6.2). The source-detector geometry is shown in Fig. 2. There are total of nine sources and 10 detectors. The middle slot was used for a commercial ultrasound transducer. The geometry used in simulation was the same as that used in phantom and clinical measurements.

Two sets of simulations were performed. In the first set of simulations, a spherical target of 1 cm diameter was embedded in the layered structure at a 1.5-cm depth from the surface. The target center was placed at 0.9 cm, and its absorption coefficient was varied from 0.07 to 0.30 cm^{-1} . Forward data was generated from two-layer structures whose optical

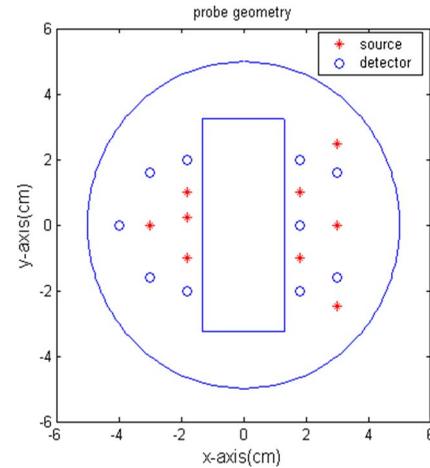


Fig. 2 Probe geometry. The red stars are sources, and the blue circles are the detectors. The middle slot is for a commercial ultrasound transducer. (Color online only.)

properties were in the range of typical breast tissue and chest wall. For each set of target data, a reference data set was also generated using the same layered structure with no target in the medium. Perturbation data were used for imaging reconstruction of the target. Two sets of imaging reconstructions were performed. One set of images was reconstructed from known optical properties of the two-layer structure, while another set was obtained by using fitted optical properties from the reflectance data. The reconstruction results for the first set provided the ideal condition when no fitting error for the two-layer optical properties was considered. The second reconstruction set is relevant to realistic clinical experiments, where the first- and second-layer optical properties must be estimated from the reflectance measurements. For this set of simulations, image reconstruction was also performed by using a semi-infinite model and a two-layer model with true background optical properties and fitted background properties.

In the second set of simulations, the absorption coefficient of the 1-cm-diam target was fixed at 0.2 cm^{-1} and the second-layer background optical absorption was varied from 0.02 to 0.15 cm^{-1} at 0.03- cm^{-1} intervals to simulate a range of chest-wall optical absorptions. For each set of background optical properties, forward data were generated with and without target. Image reconstruction was performed by using known background optical properties and fitted background optical properties. This set of simulations was intended to evaluate the effect of fitting error on reconstructed target absorption.

3.2 Phantom and Clinical Experiments

The experimental validation of the imaging algorithm was performed using an existing frequency-domain near-IR (NIR) system.¹⁹ It consisted of laser diodes of 780- and 830-nm wavelengths. The outputs of the laser diodes were amplitude modulated at 140 MHz and optically switched to nine positions on a hand-held probe. Ten optical fibers of 3 mm diameter were coupled to 10 parallel detection channels. Solid silicone phantoms were used to emulate the second layer over

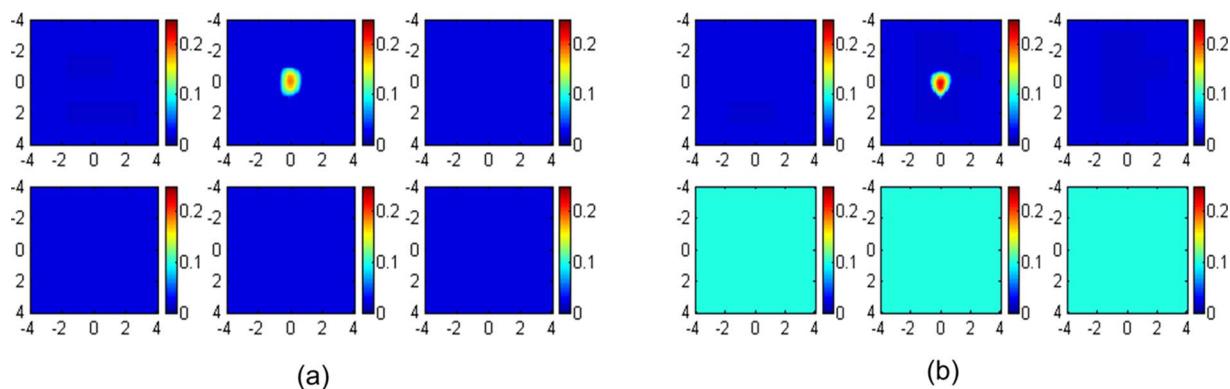


Fig. 3 Image reconstruction results using true background optical properties obtained using (a) a semi-infinite model for the two-layer structure and (b) a two-layer model for the layered structure. In each figure, the first slice is the spatial x - y image at a 0.5-cm depth from the surface and the last slice is at a 3.0-cm depth. The dimension of each slice is 8×8 cm and the spacing between the slices is 0.5 cm. The vertical scale is absorption coefficient in inverse centimeters.

which intralipid solution was used for the top layer. All the phantom experiments were performed using a flat interface between the layers. The measurements were first made on simple two-layer structures with various layer depths and used as the reference data. The measurements were repeated for the same structures with solid target phantoms immersed in the intralipid layer at known depths. Two types of target phantoms were used; one with low absorption to emulate a benign lesion and one with a high absorption to emulate a cancerous lesion in the breast.^{20,21} The layer thickness, target position, and target size were known in these phantom measurements as opposed to the clinical cases where we would estimate these parameters using a coregistered ultrasound probe.

The imaging algorithm was demonstrated with clinical examples. The study protocol was approved by a local Institutional Review Board (IRB) committee. Signed informed consent was obtained from all patients who agreed to participate in the study. In clinical experiments, NIR optical source fibers, detector fibers, and a commercial ultrasound transducer were integrated on a handheld probe.¹⁹ Patients were scanned in a supine position, while multiple sets of optical reflectance measurements were made with coregistered ultrasound images at lesion locations and symmetric locations of the contralateral normal breast. The measurements obtained at normal locations were used to estimate background optical properties, which were used to compute a weight matrix for imaging.

3.3 Estimation of Two-Layer Optical Properties and Image Reconstruction

Two-layer optical properties were estimated from fitting reflectance measurements made on the layered structure to analytical solution using the Nelder-Mead simplex algorithm as described in our previous paper.¹² This algorithm minimizes a scalar-valued nonlinear function of multiple independent parameters by using only the function values, and it is chosen due to its simplicity and good convergence in spite of the slow convergence rate. We chose various initial values and found that initial values are not critical as long as the function converges. For each case, the initial values were chosen as 0.05, 5.0, 0.05, and 5.0 cm^{-1} for μ_{a1} , μ'_{s1} , μ_{a2} , and μ'_{s2} , respec-

tively. In the phantom and clinical experiments, the thickness of the upper layer was estimated by using the coregistered ultrasound image. The nonlinear regression routine used here, *fminsearch*, is available in MATLAB.

In simulations, the voxel size was chosen as 0.1 cm in the x and y dimensions for fine mesh and 1.5 cm for coarse mesh. The corresponding voxel sizes were 0.25 and 1.5 cm in phantom studies, and 0.5 and 1.5 cm for clinical data. In all experiments, the voxel size in depth was 0.5 cm in the fine-mesh zone and 1.5 cm in the coarse-mesh zone. The choice of fine-mesh size is based on SNR and also spatial sampling considerations. The fine-mesh volume for all experiments was chosen at least 2.5 times larger than the target size and centered at the target position. Clinical experiments were performed by fitting the two-layer optical properties of the contralateral breast using the layer depth provided by coregistered ultrasound images. In all experiments, the dual-mesh algorithm was used for inversion with either known target location or size in simulation and phantom experiments or *a priori* information provided by coregistered ultrasound in clinical experiments.

4 Results

4.1 Simulation

In the first set of simulations, targets of various absorption properties were embedded in a layered structure of 1.5 cm upper layer thickness. The first and second layer known optical properties were $\mu_{a1}=0.02 \text{ cm}^{-1}$ and $\mu'_{s1}=7.0 \text{ cm}^{-1}$ and $\mu_{a2}=0.10 \text{ cm}^{-1}$ and $\mu'_{s2}=7.0 \text{ cm}^{-1}$, respectively. The target absorption coefficients were chosen as 0.07, 0.10, 0.15, 0.20, 0.25, and 0.30 cm^{-1} , and the reduced scattering coefficient was kept at 7.0 cm^{-1} . Reference data were also generated for the two-layer structure in the absence of the target. Figure 3 shows an example of reconstructed target absorption maps of an absorber ($\mu_a=0.30 \text{ cm}^{-1}$ and $\mu'_s=7.0 \text{ cm}^{-1}$) embedded in the layered-medium when the known optical properties were used in the reconstruction. Figure 3(a) shows the reconstruction result of using a semi-infinite model for the two-layer structure and known first-layer background values of $\mu_{a1}=0.02 \text{ cm}^{-1}$ and $\mu'_{s1}=7 \text{ cm}^{-1}$. Figure 3(b) shows the two-

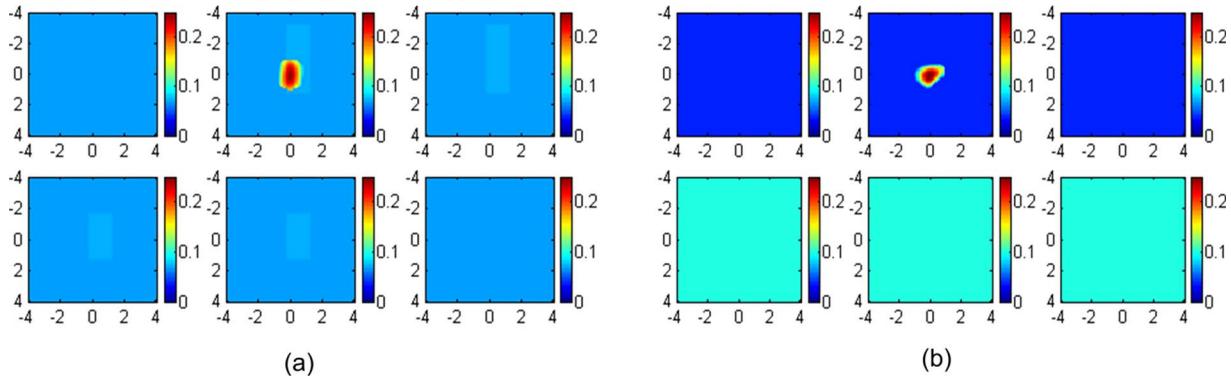


Fig. 4 Image reconstruction results using fitted background optical properties obtained using (a) the semi-infinite model with the fitted background optical properties and (b) the two-layer model with fitted background optical properties. In each figure, the dimension of each slice is 8×8 cm and the spacing between the slices is 0.5 cm. The vertical scale is absorption coefficient in inverse centimeters.

layer-model reconstruction using known two-layer background values of $\mu_{a1}=0.02 \text{ cm}^{-1}$, $\mu'_{s1}=7.0 \text{ cm}^{-1}$, $\mu_{a2}=0.10 \text{ cm}^{-1}$, and $\mu'_{s2}=7.0 \text{ cm}^{-1}$. The reconstructed maximum absorption values were 0.186 cm^{-1} (62%) and 0.220 cm^{-1} (73%), respectively. About 11% improvement in reconstruction accuracy was achieved using the layer-based reconstruction under this ideal scenario. When using a semi-infinite model to fit background optical properties of the two-layer structure, $\mu_a=0.069 \text{ cm}^{-1}$ and $\mu'_s=5.94 \text{ cm}^{-1}$ were obtained. Figure 4(a) shows the result of semi-infinite-model-based reconstruction and the reconstructed maximum absorption coefficient was 0.242 cm^{-1} [Fig. 4(a)]. When fitted two-layer optical properties of $\mu_{a1}=0.037 \text{ cm}^{-1}$, $\mu'_{s1}=7.24 \text{ cm}^{-1}$, $\mu_{a2}=0.10 \text{ cm}^{-1}$, and $\mu'_{s2}=5.09 \text{ cm}^{-1}$ were used in two-layer-model-based reconstruction, the reconstructed maximum absorption coefficient was 0.2613 cm^{-1} [Fig. 4(b)]. About 7% improvements in reconstruction accuracy were achieved even with fitting errors of the two-layer optical properties.

Figure 5 shows the reconstructed maximum absorption coefficient versus true target absorption value with known and fitted optical properties of two-layer model (black star and purple cross lines) and the semi-infinite approximation for the layered structure (green triangle and red square lines), respectively. It is interesting to note that the two-layer model improves the reconstruction accuracy only at higher contrast end when known optical properties of two-layers are used. This is due to the use of a more accurate weight matrix for the two-layer structure. Because both semi-infinite and two-layer models use the linear Born approximation, the reconstructed target absorption is reduced at higher contrast. When the fitted two-layer optical properties are used, the improvement is distributed at both high and low contrast. The improvement at low contrast is due to the use of more accurate first-layer background properties. Because reconstructed target $\mu_a = \mu_{a1} + \text{reconstructed } \Delta\mu_a$, a higher fitted background absorption using the semi-infinite model yields a higher target μ_a compared with using the two-layer model. This problem is less pronounced when the target to background ratio is higher.

In the second set of simulations, the first layer optical properties and the second layer reduced scattering coefficient were fixed at $\mu_{a1}=0.02 \text{ cm}^{-1}$, and $\mu'_{s1}=7 \text{ cm}^{-1}$, and $\mu'_{s2}=7 \text{ cm}^{-1}$. However, the second layer background μ_{a2} was

varied from 0.02 to 0.15 cm^{-1} to simulate a range of chest-wall absorption. Table 1 lists the true and fitted two-layer optical properties and the corresponding reconstructed maximum target absorption coefficient. Target reconstruction results using true and fitted background properties for each set of background values are within 13%. The small difference results from the fitting error of first-layer μ_{a1} .

4.2 Phantom Experiments

Phantom experiments were performed with various upper layer thicknesses and two 1-cm diameter spherical targets of high absorption (phantom 1) and low absorption (phantom 2) coefficients located at various depths. The calibrated absorption and reduced scattering coefficients obtained from the same phantom materials were $\mu_a=0.23 \text{ cm}^{-1}$ and $\mu'_s=5.45 \text{ cm}^{-1}$ for phantom 1 and $\mu_a=0.07 \text{ cm}^{-1}$ and μ'_s

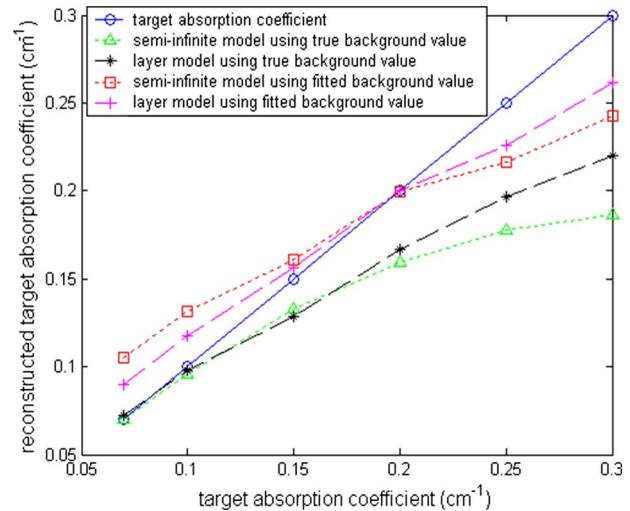


Fig. 5 Comparison of target absorption coefficients obtained with a semi-infinite-model-based reconstruction for the layered structure (green triangle line: using true background values; red squares: using fitted background values) and a two-layer-model based reconstruction for the layered structure (black stars: using true background values; purple crosses: using fitted background values). The blue circle line is the true target absorption coefficient. (Color online only.)

Table 1 Simulation results using the semi-infinite model and the two-layer model with fitted background values. Target center depth $C=0.9$ cm. Target radius $R=0.5$ cm. Second layer depth $D=1.5$ cm. Refractive index ($n_1=1.33$, $n_2=1.4$).

Two-layer Model with True Background Value		Two-layer Model with Fitted Background Value	
True Background Value [$\mu_{a1}\mu'_{s1}\mu_{a2}\mu'_{s2}$] (cm^{-1})	Reconstructed Maximum μ_a (cm^{-1})	Fitted Background Value [$\mu_{a1}\mu'_{s1}\mu_{a2}\mu'_{s2}$] (cm^{-1})	Reconstructed Maximum μ_a (cm^{-1})
[0.02 7 0.02 7]	0.1525 (76%)	[0.0335 6.9 0.0174 7.05]	0.1369 (68%)
[0.02 7 0.06 7]	0.1398 (70%)	[0.0339 6.92 0.053 6.88]	0.1331 (66%)
[0.02 7 0.09 7]	0.1486 (74%)	[0.0326 6.98 0.083 6.51]	0.1738 (87%)
[0.02 7 0.12 7]	0.1725 (86%)	[0.0353 7.01 0.1041 6.1]	0.1823 (91%)
[0.02 7 0.15 7]	0.1814 (91%)	[0.0355 7.04 0.1286 6.51]	0.1934 (97%)

Table 2 Reconstruction results for phantom 1 ($\mu_a=0.23 \text{ cm}^{-1}$ and $\mu'_s=5.45 \text{ cm}^{-1}$) using the semi-infinite model and the two-layer model with fitted background values.

	Target Center Depth (C)/ Second Layer Depth (D)	Semi-Infinite Model		Two-Layer Model	
		Fitted Background Value [$\mu_a\mu'_s$] (cm^{-1})	Reconstructed Maximum μ_a (cm^{-1})	Fitted Background Value [$\mu_{a1}\mu'_{s1}\mu_{a2}\mu'_{s2}$] (cm^{-1})	Reconstructed Maximum μ_a (cm^{-1})
Phantom 1 and silicon phantom 1	C 0.9 cm, D 1.5 cm	[0.048 5.33]	0.158 (68%)	[0.036 7.89 0.11 3.25]	0.201 (88%)
	C 1.5 cm, D 2.1 cm	[0.039 4.76]	0.169 (73%)	[0.037 5.92 0.079 5.17]	0.206 (90%)
	C 1.8 cm, D 2.4 cm	[0.037 5.40]	0.191 (83%)	[0.027 9.90 0.041 18.53]	0.198 (86%)
Phantom 1 and silicon phantom 2	C 0.9 cm, D 1.5 cm	[0.054 4.79]	0.165 (72%)	[0.020 5.62 0.083 20.21]	0.205 (89%)
	C 1.5 cm, D 2.1 cm	[0.037 5.42]	0.160 (70%)	[0.027 5.53 0.051 5.59]	0.173 (75%)
	C 1.8 cm, D 2.4 cm	[0.033 5.57]	0.186 (81%)	[0.018 7.71 0.051 8.07]	0.187 (81%)

Table 3 Reconstruction results for phantom 2 ($\mu_a=0.07 \text{ cm}^{-1}$ and $\mu'_s=5.50 \text{ cm}^{-1}$) using the semi-infinite model and the two-layer model with fitted background values.

	Target Center Depth (C)/ Second Layer Depth (D)	Semi-Infinite Model		Two-Layer Model	
		Fitted Background Value [$\mu_a\mu'_s$] (cm^{-1})	Reconstructed Maximum μ_a (cm^{-1})	Fitted Background Value [$\mu_{a1}\mu'_{s1}\mu_{a2}\mu'_{s2}$] (cm^{-1})	Reconstructed Maximum μ_a (cm^{-1})
Phantom 2 and silicon phantom 1	C 0.9 cm, D 1.5 cm	[0.065 4.82]	0.129 (184%)	[0.014 5.48 0.16 9.56]	0.0710 (101%)
	C 1.5 cm, D 2.1 cm	[0.045 5.02]	0.104 (148%)	[0.019 7.87 0.099 41.74]	0.0889 (127%)
	C 1.8 cm, D 2.4 cm	[0.042 4.88]	0.105 (150%)	[0.023 9.59 0.048 9.01]	0.0896 (128%)
Phantom 2 and silicon phantom 2	C 0.9 cm, D 1.5 cm	[0.054 5.04]	0.121 (173%)	[0.021 5.26 0.057 15.08]	0.0868 (124%)
	C 1.5 cm, D 2.1 cm	[0.035 5.11]	0.0944 (135%)	[0.023 5.83 0.053 5.557]	0.0899 (128%)
	C 1.8 cm, D 2.4 cm	[0.031 5.33]	0.103 (147%)	[0.031 5.93 0.058 6.26]	0.1000 (143%)

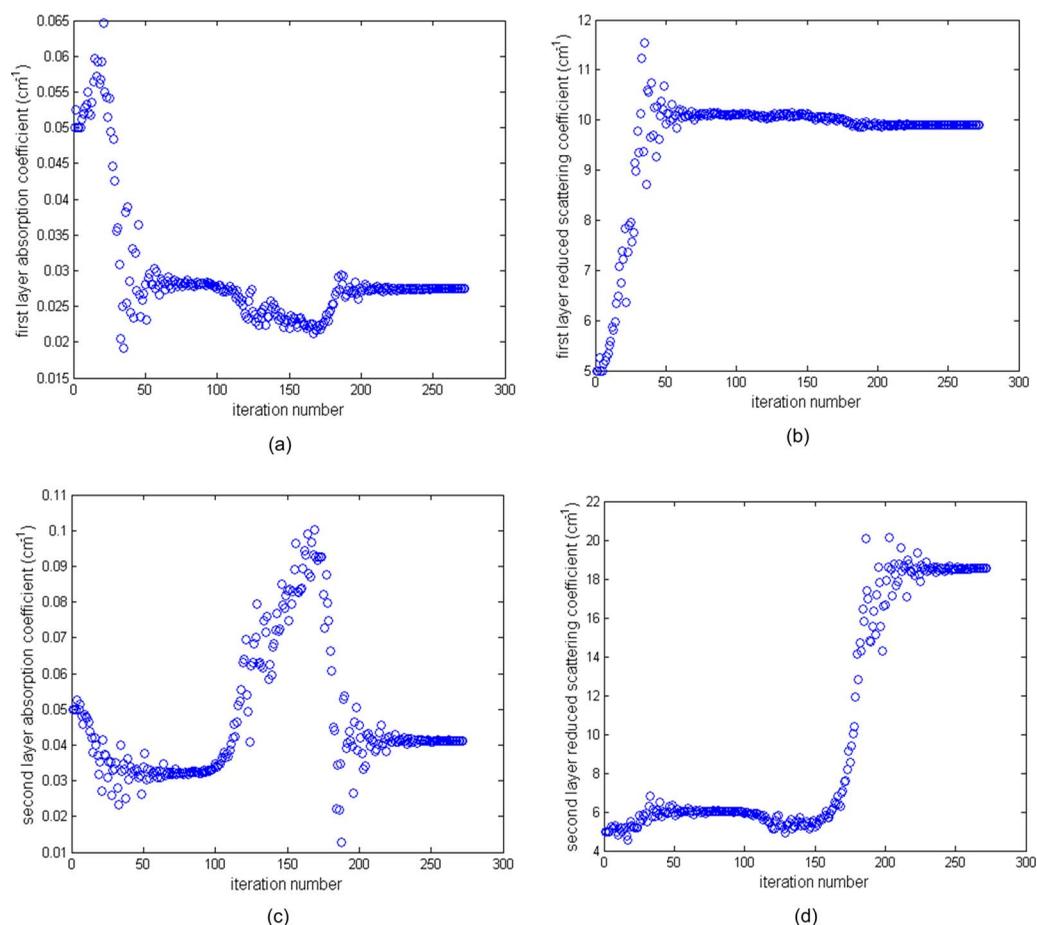


Fig. 6 Phantom experiments of fitted background optical properties versus iteration. Intralipid was used as the first layer and the silicon phantom 1 was used as the second layer. The depth of the intralipid layer was 2.4 cm. The measured optical properties of Intralipid were $\mu_a = 0.022 \text{ cm}^{-1}$ and $\mu'_s = 7.2 \text{ cm}^{-1}$, and those of the silicon phantom 1 were $\mu_a = 0.06 \text{ cm}^{-1}$ and $\mu'_s = 15 \text{ cm}^{-1}$. (a) and (c) Fitted layer absorption coefficient versus iteration for first and second layers, respectively, and (b) and (d) fitted reduced scattering coefficient versus iteration for first and second layers, respectively.

$= 5.50 \text{ cm}^{-1}$ for phantom 2 at 780 nm. The upper layer was intralipid solution of measured optical properties $\mu_a = 0.022 \text{ cm}^{-1}$ and $\mu'_s = 7.2 \text{ cm}^{-1}$, and the bottom layer comprised silicon phantoms of measured optical properties $\mu_a = 0.06 \text{ cm}^{-1}$ and $\mu'_s = 15 \text{ cm}^{-1}$ (silicon phantom 1) and $\mu_a = 0.12 \text{ cm}^{-1}$ and $\mu'_s = 15 \text{ cm}^{-1}$ (silicon phantom 2).

The reconstruction results were obtained by using fitted two-layer optical properties from the reflectance measurements. The results of six different experiments of phantoms 1 and 2 are tabulated in Tables 2 and 3. The reconstruction accuracy calculated from the calibrated target values using the semi-infinite and two-layer models are also given in the tables. For the low-contrast phantom 2, because the two-layer model provides more accurate background value, the reconstructed absorption coefficient is greater than 80% more accurate than that obtained from the semi-infinite model at a shallow depth, such as 1.5 cm. For the high-contrast phantom 1, the improvement of 10 to 20% is obtained when the two-layer analytic solution is used instead of the semi-infinite model. As expected, the difference in reconstructed values between the two models reduces as the layer depth increases. Because second-layer optical properties mainly affect distance

measurements acquired at longer source-detector pairs and these measurements have lower SNRs compared with shorter pairs, the fitted optical properties of the second layer are less accurate than those of the first layer. However, the reconstructed target contrast is not very sensitive to the second layer properties.

An example of imaging results of phantom 1 using silicon phantom 1 as the second layer is given in Figs. 6 and 7. Figure 6 shows the fitted background optical properties of μ_{a1} , μ'_{s1} , μ_{a2} , and μ'_{s2} versus iterations when the first layer depth is 2.4 cm (third line in Table 1 two-layer model). The fitting procedure converges and the fitted background values at iteration of 272 are $\mu_{a1} = 0.027 \text{ cm}^{-1}$, $\mu'_{s1} = 9.90 \text{ cm}^{-1}$, $\mu_{a2} = 0.041 \text{ cm}^{-1}$, and $\mu'_{s2} = 18.53 \text{ cm}^{-1}$.

Figure 7 shows the absorption maps of high-contrast phantom 1 placed in Intralipid layers of 0.9-, 1.5-, and 1.8-cm depth. The second layer (silicon phantom 1) was located at 1.5-, 2.1-, and 2.4-cm depths, respectively. Figure 7(a), 7(c), and 7(e) show reconstructed absorption maps obtained at 780 nm using a semi-infinite model for the layer structure. Reconstructed target maximum absorption coefficients were 0.158 (68%), 0.169 (73%), and 0.191 (83%) cm^{-1} , respec-

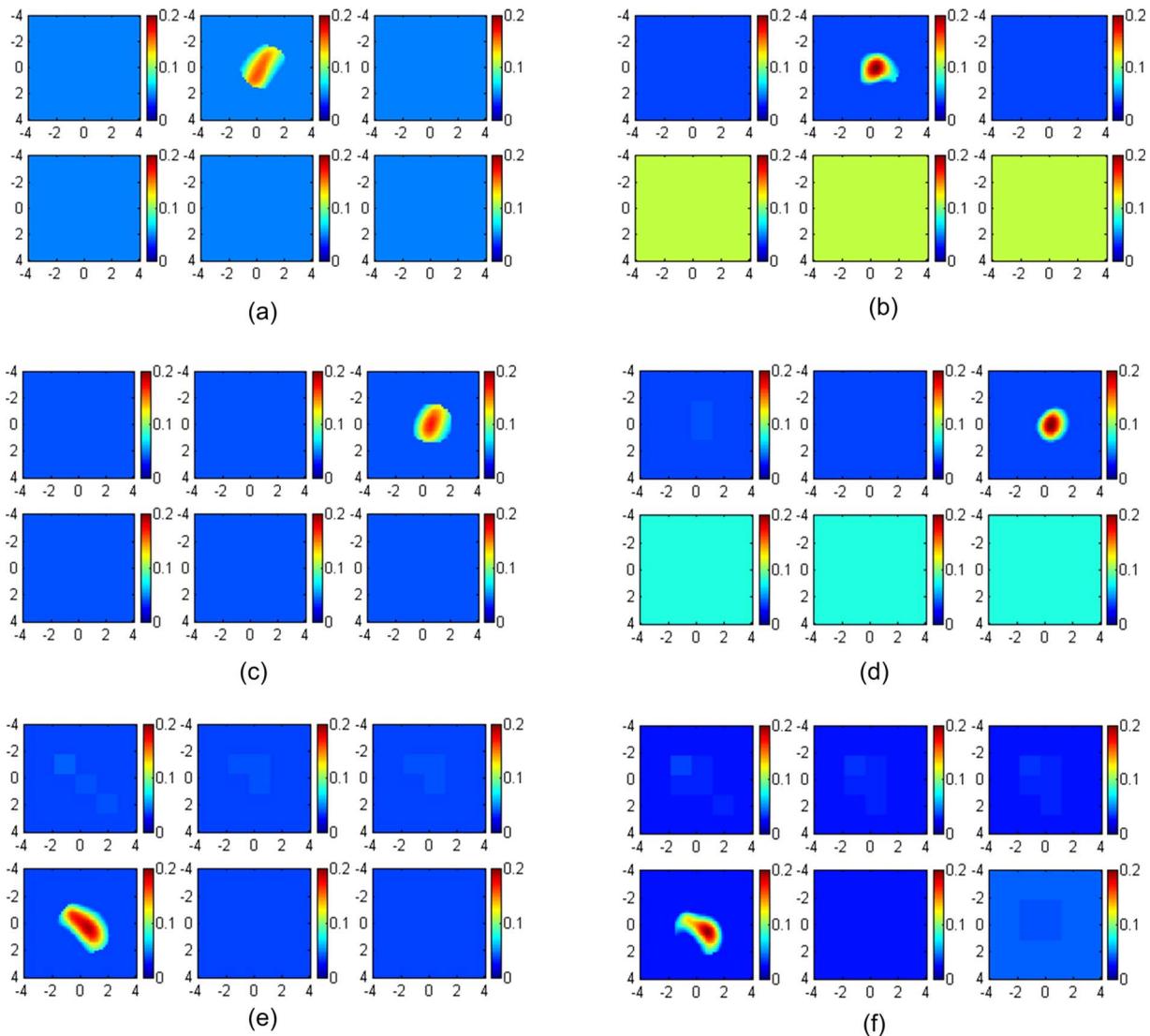


Fig. 7 Reconstructed absorption maps of phantom 1 placed in Intralipid layers of 0.9, 1.5, and 1.8 cm depth. The second layer silicon phantom was located at 1.5, 2.1, and 2.4 cm depth, respectively. (a) and (b), The target center was located at 0.9 cm depth. The first slice is the spatial x - y image of 8×8 cm obtained at a 0.4-cm depth, and the second and third slices at 0.9 and 1.4 cm, respectively. (c) and (d) The target center was located at a 1.5-cm depth. The first slice is the spatial x - y image of 8×8 cm obtained at a 0.5-cm depth, and the second and third cross sections at 1.0 and 1.5 cm, respectively. (e) and (f). The target center was located at a 1.8-cm depth. The first slice is the spatial x - y image of 8×8 cm obtained at a 0.3-cm depth, and the fourth cross-section slice is at a 1.8-cm depth. (a), (c), and (e) Reconstructed absorption maps obtained at 780 nm using fitted background values and a semi-infinite model for the layered structure. Reconstructed target maximum absorption coefficients were 0.158, 0.169, and 0.191 cm^{-1} , respectively. (b), (d), and (f) Reconstructed absorption maps obtained at the same wavelength using the two-layer model with fitted background values. The reconstructed target maximum absorption coefficients were 0.201, 0.206, 0.198 cm^{-1} , respectively.

tively. Figures 7(b), 7(d), and 7(f) show reconstructed absorption maps obtained at the same wavelength using the two-layer model. The reconstructed target maximum absorption coefficients were 0.201 (88%), 0.209 (91%), and 0.198 cm^{-1} (86%), respectively. About 3 to 20% improvement in reconstruction accuracy was obtained in this set of experiments. The reconstruction results obtained at 830 nm are very similar to the results at 780 nm. Therefore, images of 830 nm are not shown here.

4.3 Clinical Experiments

The two-layer-model-based reconstruction was demonstrated using clinical data. Two examples of benign and malignant

cases are given here to demonstrate the potential of this new reconstruction technique. Figure 8 shows the ultrasound image from the normal contralateral site [Fig. 8(a)] and lesion site [Fig. 8(b)] of a 54-year-old patient. The lesion was approximately 0.8 cm deep with approximately 0.3 cm radius in the z direction (depth) and the lateral radius or shape of the lesion was approximately 1.0 cm, as seen in the ultrasound image. We can see from Fig. 8(a) that the upper layer thickness is about 1.4 cm with an almost flat interface between the chest wall and the breast tissue. The fitted optical properties of the breast and chest-wall layer from the contralateral site with no lesion were $\mu_{a1}=0.041 \text{ cm}^{-1}$, $\mu'_{s1}=10.22 \text{ cm}^{-1}$, μ_{a2}

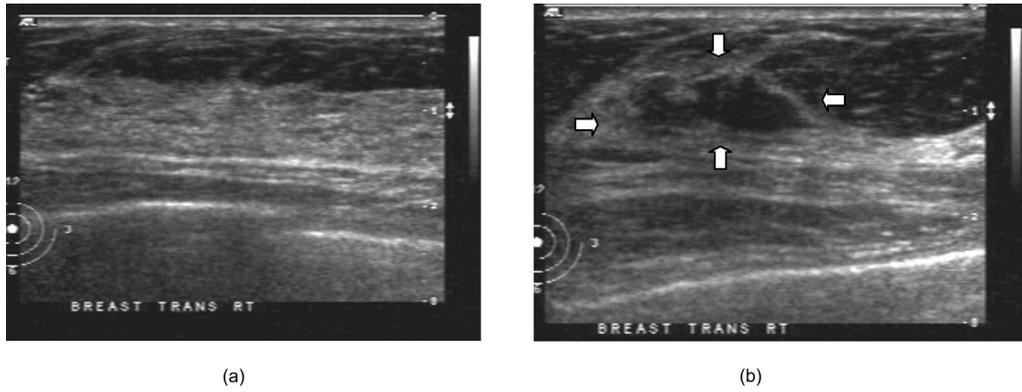


Fig. 8 Ultrasound image obtained from a 54-year-old patient with a suspicious mass: (a) contralateral normal site and (b) lesion site.

$=0.095 \text{ cm}^{-1}$, and $\mu'_{s2}=3.07 \text{ cm}^{-1}$ at 780 nm, and $\mu_{a1}=0.056 \text{ cm}^{-1}$, $\mu'_{s1}=9.05 \text{ cm}^{-1}$, $\mu_{a2}=0.073 \text{ cm}^{-1}$, and $\mu'_{s2}=3.4 \text{ cm}^{-1}$ at 830 nm. The fitted two-layer optical properties at 780 nm converge after 226 iterations, as shown in Figs. 9(a)–9(d). The fitted optical properties by using the semi-infinite method directly were $\mu_a=0.06 \text{ cm}^{-1}$, and $\mu'_s=6.77 \text{ cm}^{-1}$ at 780 nm and $\mu_a=0.061 \text{ cm}^{-1}$, and $\mu'_s=7.55 \text{ cm}^{-1}$ at 830 nm. Figures 10(a)–10(c) show the absorp-

tion maps at 780 and 830 nm and the computed total hemoglobin map, respectively, obtained by using a semi-infinite model. Figures 10(d)–10(f) show the corresponding absorption and total hemoglobin maps obtained using the two-layer model. Both images show low absorption and therefore lower total hemoglobin concentration with layer-based reconstruction providing slightly lower values. Ultrasound-guided core biopsy confirmed that the lesion was a benign fibroadenoma.

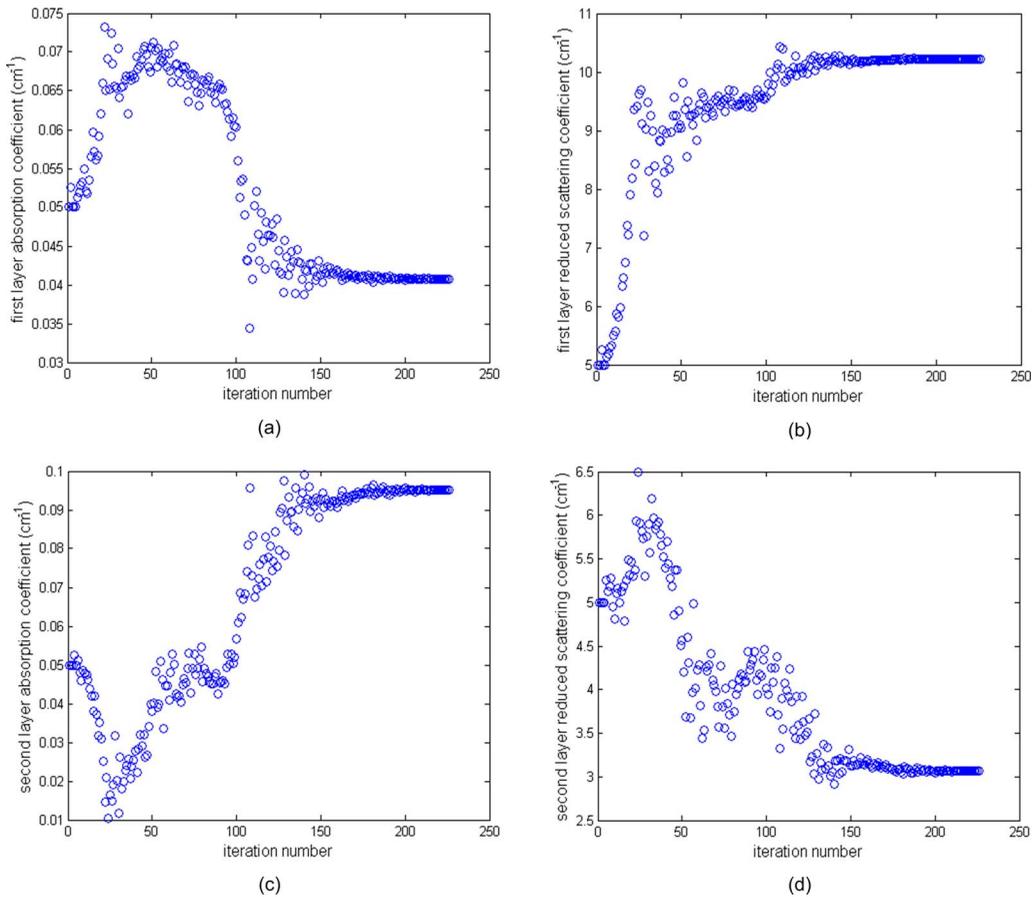


Fig. 9 Results of fitted background optical properties versus iteration numbers. (a) and (c) Fitted optical absorption coefficient versus iteration for first and second layers, respectively, and (b) and (d) fitted reduced scattering coefficient versus iteration for first and second layers, respectively. The fitting algorithm converges around iteration number 226.

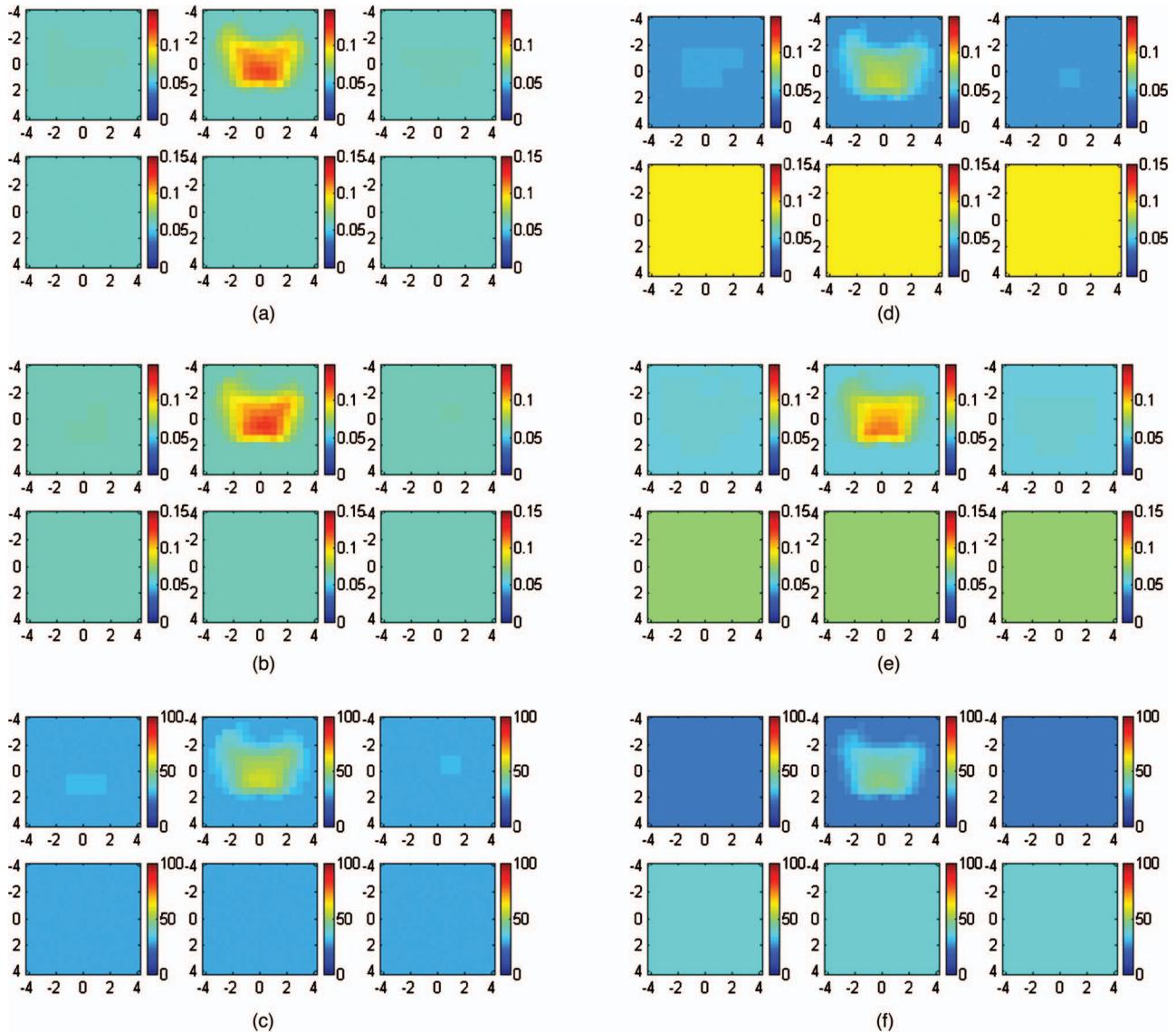


Fig. 10 (a) to (c) Reconstructed optical absorption maps at (a) 780 nm and (b) 830 nm and the computed total hemoglobin map (c) using semi-infinite model. Reconstructed absorption coefficients were 0.121 cm^{-1} (780 nm), 0.124 cm^{-1} (830 nm), and $57.69 \mu\text{mol/l}$ (total hemoglobin). (d) to (f) Reconstructed optical absorption maps at (d) 780 nm and (e) 830 nm and the computed total hemoglobin map (f) using the layer-based model. Reconstructed absorption coefficients were 0.0832 cm^{-1} (780 nm), 0.113 cm^{-1} (830 nm), and $47.09 \mu\text{mol/l}$ (total hemoglobin). In each figure, the first slice is the spatial x-y image 0.5 cm deep from the skin surface and the last slice is 3.0 cm toward the chest wall. The dimension of each slice is $9 \times 9 \text{ cm}$ and the spacing between the slices is 0.5 cm. The vertical scale is absorption coefficient in inverse centimeters or total hemoglobin concentration in micromoles per liter.

Figure 11 shows the ultrasound images obtained from the contralateral and lesion sites of a 61-year-old patient. It can be seen from Fig. 11(a) that the lesion is about 1.2 cm deep and the breast tissue and chest-wall layer interface is almost flat with a breast layer thickness of about 1.4 to 1.5 cm. The fitted background optical properties of the breast and chest-wall tissue obtained from the contralateral normal breast were $\mu_{a1}=0.079 \text{ cm}^{-1}$, $\mu'_{s1}=4.21 \text{ cm}^{-1}$, $\mu_{a2}=0.086 \text{ cm}^{-1}$, and $\mu'_{s2}=5.93 \text{ cm}^{-1}$ at 780 nm, and $\mu_{a1}=0.088 \text{ cm}^{-1}$, $\mu'_{s1}=4.38 \text{ cm}^{-1}$, $\mu_{a2}=0.11 \text{ cm}^{-1}$, and $\mu'_{s2}=6.76 \text{ cm}^{-1}$ at 830 nm. The fitted optical properties by using the semi-infinite method directly were $\mu_a=0.079 \text{ cm}^{-1}$, and $\mu'_s=4.29 \text{ cm}^{-1}$ at 780 nm and $\mu_a=0.091 \text{ cm}^{-1}$ and μ'_s

$=4.38 \text{ cm}^{-1}$ at 830 nm. The absorption maps and the computed total hemoglobin map obtained from with the semi-infinite approximation are shown in Figs. 12(a)–12(c) and those from the two-layer model are shown in Figs. 12(d)–12(f). Both images show high absorption and therefore high total hemoglobin concentration with layer-based reconstruction providing higher values. Ultrasound-guided core biopsy confirmed that the lesion was an early stage invasive carcinoma with ductal and lobular features. These examples demonstrate that the layer-based imaging reconstruction has a great potential to improve the contrast between malignant and benign lesions and therefore provide more accurate diagnosis.

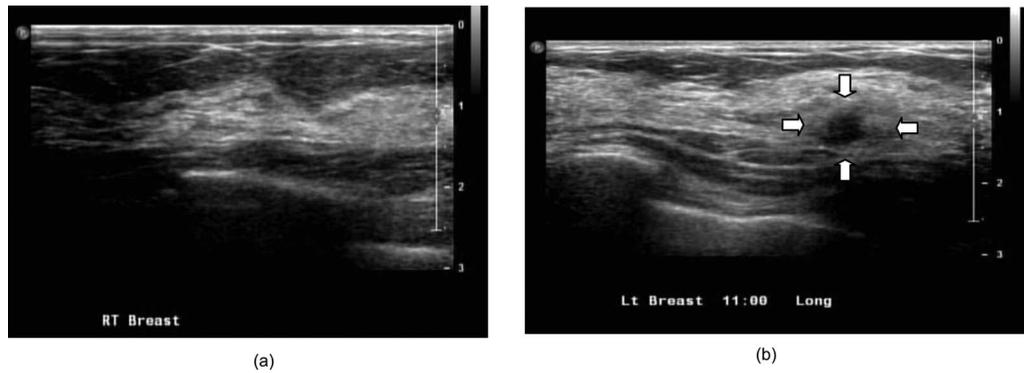


Fig. 11 Ultrasound images obtained from a 61-year-old patient who has a suspicious mass: (a) contralateral normal site and (b) lesion site.

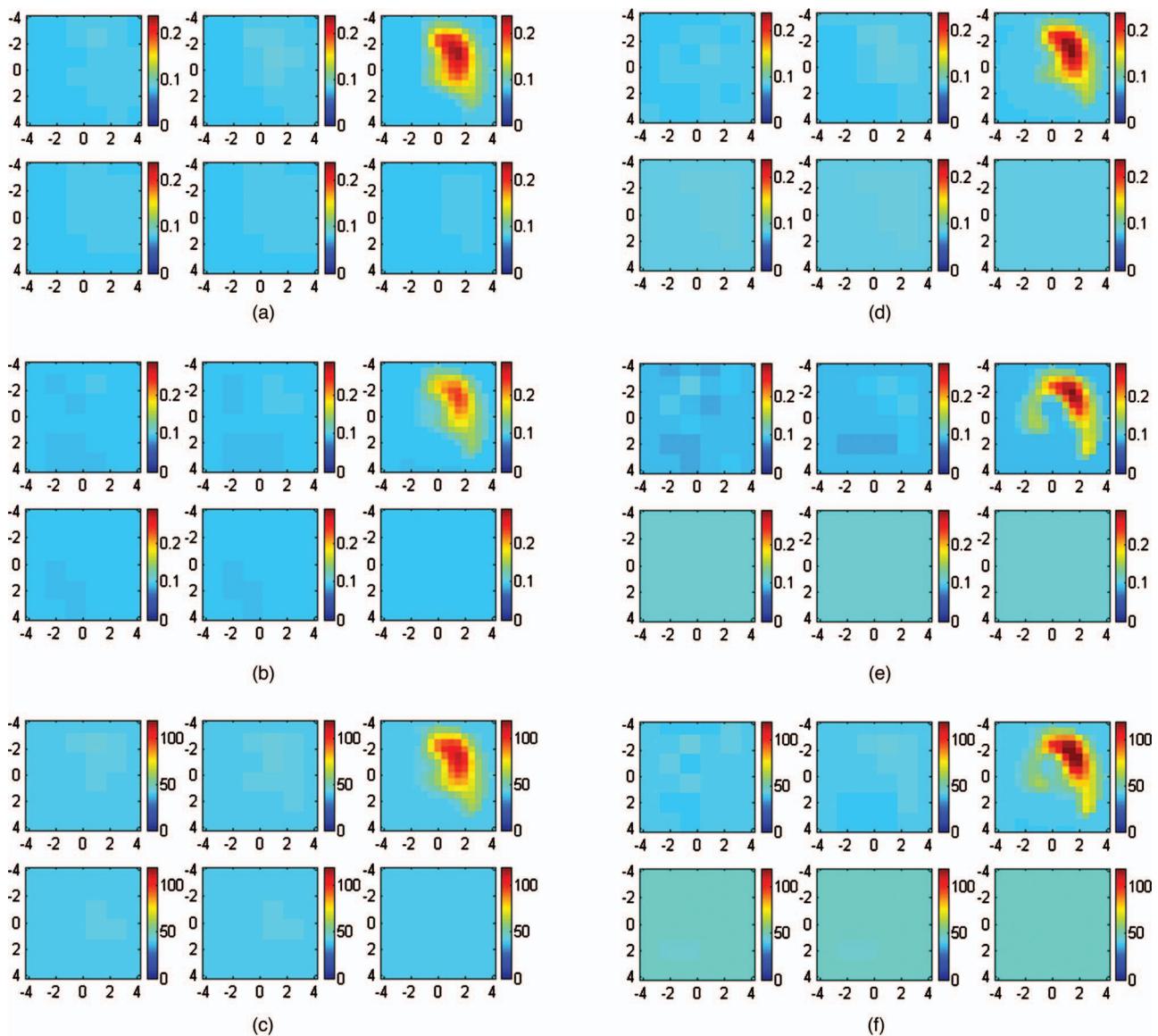


Fig. 12 (a) to (c) Reconstructed optical absorption maps at (a) 780 and (b) 830 nm and the computed total hemoglobin map (c) using the semi-infinite model. Reconstructed absorption coefficients were 0.232 cm^{-1} (780 nm), 0.238 cm^{-1} (830 nm), and $111.33 \mu\text{mol/liter}$ (total hemoglobin). (d) to (f) Reconstructed optical absorption maps at (d) 780 and (e) 830 nm and the computed total hemoglobin map (f) using the layer-based model. Reconstructed absorption coefficients were 0.247 cm^{-1} (780 nm), 0.293 cm^{-1} (830 nm), and $126.39 \mu\text{mol/liter}$ (total hemoglobin). In each figure, the first slice is the spatial x-y image of 0.2-cm depth from the skin surface and the last slice is 3.0 cm toward the chest wall. The dimensions of each slice are $9 \times 9 \text{ cm}$ and the spacing between the slices is 0.5 cm. The vertical scale is absorption coefficient in inverse centimeters or total hemoglobin in the unit of micromoles per liter.

5 Summary

A new optical tomography method based on a two-layer-model analytical solution was developed and compared with the semi-infinite model using simulated data and phantom experiments. Clinical examples were given to demonstrate the potential of this method. It was found that the reconstructed target optical properties were more accurate when the second layer was present in less than 2 cm depth and its effect was taken into account. This improvement is mainly attributed to the more accurate estimation of background optical properties and more accurate estimation of weight matrix for imaging reconstruction by considering the light propagation effect in the second layer. Our initial results indicate the potential of the layer-based imaging reconstruction method to improve discrimination between malignant and benign lesions.

The fitted first-layer optical properties were more accurate than those of the second layer. This is due to the lower SNR of measurements acquired at longer source-detector pairs. Simulations and phantom experiments showed that the reconstructed target absorption coefficient was not very sensitive to fitting errors of second-layer optical properties. Therefore, the fitting algorithm and the reported two-layer imaging reconstruction are more robust for use on clinical data where a range of optical properties and chest-wall depths may be present. The chest wall may potentially distort the distance measurements.

Optical properties of breast tissue and chest wall vary from patient to patient. From literature data and our own experience, the absorption coefficient μ_a is in the range of 0.01 to 0.08 cm^{-1} , and the reduced scattering coefficient μ'_s is in the range of 2.0 to 12.0 cm^{-1} . The optical properties of the chest wall are not available in the literature. The only available data are optical properties of muscles and bones. In Refs. 22–24, the muscle was reported to have a large absorption and medium scattering coefficients with μ_a in the range of 0.08 to 0.27 cm^{-1} and μ'_s in the range of 5 to 10 cm^{-1} . The bone is a highly absorbing and scattering medium. From the literature^{25,26}, the bone was reported to have large absorption and scattering coefficients with μ_a in the range of 0.05 to 0.15 cm^{-1} and μ'_s in the range of 20 to 30 cm^{-1} . We choose optical properties within this range for simulations and phantom experiments.

The two-layer analytical-solution-based imaging reconstruction is limited to a flat interface between first and second layers because the analytic solution is derived in the cylindrical coordinates with depth z as a parameter. The breast tissue and chest-wall interface may not be flat in many clinical cases. These factors may introduce errors in estimated two-layer background optical properties, and therefore reconstructed lesion optical properties. More clinical cases will be analyzed to evaluate the robustness of this layer-based tomography method. Finite element methods (FEMs) have been used by many research groups for optical tomography applications^{27,28} and are suitable for complex boundary conditions between the breast tissue and chest-wall layers. However, an FEM is computationally intensive, while the proposed analytical-solution-based reconstruction is computationally efficient.

In summary, a new tomographic method was introduced that has been shown to improve light quantification accuracy

of lesions imbedded in a two-layer tissue structure. The algorithm was demonstrated by using ultrasound prior to it; however, it can be used alone for optical imaging reconstruction. The robustness of this algorithm will be validated in a larger patient cohort of different breast tissue and chest wall optical properties.

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References

1. J. M. Schmitt, G. X. Zhuo, E. C. Walker, and R. T. Wall, "Multilayer model of photon diffusion in skin," *J. Opt. Soc. Am. A* **7**(11), 2141–2153 (1990).
2. I. Dayan, S. Havlin, and G. H. Weiss, "Photon migration in a two layer turbid medium: a diffusion analysis," *J. Mod. Opt.* **39**(7), 1567–1582 (1992).
3. A. Kienle, M. S. Patterson, N. Dögnitz, R. Bays, G. Wagnière, and H. Bergh, "Noninvasive determination of the optical properties of two-layered turbid media," *Appl. Opt.* **37**(4), 779–791 (1998).
4. Y. S. Fawzi, A. M. Youssef, M. H. El-Batany, and Y. M. Kadah, "Determination of the optical properties of a two-layer tissue model by detecting photons migrating at progressively increasing depth," *Appl. Opt.* **42**(31), 6398–6411 (2003).
5. T. J. Farrell, M. S. Patterson, and M. Essenpreis, "Influence of layered tissue architecture on estimates of tissue optical properties obtained from spatially resolved diffuse reflectometry," *Appl. Opt.* **37**(10), 1958–1972 (1998).
6. G. Alexandrakis, T. J. Farrell, and M. J. Patterson, "Accuracy of diffusion approximation in determining the optical properties of a two-layer turbid medium," *Appl. Opt.* **37**(31), 7401–7409 (1998).
7. G. Alexandrakis, T. J. Farrell, and M. S. Patterson, "Monte Carlo diffusion hybrid model for photon migration in a two-layer turbid medium in frequency domain," *Appl. Opt.* **39**(13), 2235–2244 (2000).
8. F. Martelli, S. D. Bianco, and G. Zaccanti, "Procedure for retrieving the optical properties of a two-layered medium from time resolved reflectance measurements," *Opt. Lett.* **28**(14), 1236–1238 (2003).
9. J. Tualle, J. Prat, E. Tinetti, and S. Avrillier, "Real-space Green's function calculation for the solution of the diffusion equation in stratified turbid media," *J. Opt. Soc. Am. A* **17**(11), 2046–2055 (2000).
10. L. Wang, S. L. Jacques, and L. Zheng, "MCML-Monte Carlo modeling of light transport in multi-layered tissues," *Comput. Methods Programs Biomed.* **47**, 131–146 (1995).
11. D. Boas, J. Culver, J. Stott, and A. Dunn, "Three dimensional Monte Carlo code for photon migration through complex heterogeneous media including the adult human head," *Opt. Express* **10**(3), 159–170 (2002).
12. M. Das, C. Xu, and Q. Zhu, "Analytical solution for light propagation in a two-layer tissue structure with a tilted interface for breast imaging," *Appl. Opt.* **45**(20), 5027–5036 (2006).
13. S. Tseng, C. Hayakawa, B. Tromberg, J. Spanier, and A. Durkin, "Quantitative spectroscopy of superficial turbid media," *Opt. Lett.* **30**(23), 3165–3167 (2005).
14. A. Vogel, V. Chernomordik, J. Riley, M. Hassan, F. Amyot, B. Dasgeb, S. Demos, R. Pursley, R. Little, R. Yarchoan, Y. Tao, and A. Gandjbakche, "Using noninvasive multispectral imaging to quantitatively assess tissue vasculature," *J. Biomed. Opt.* **12**(5), 051604:1–13 (2007).
15. M. Das and Q. Zhu, "Optical tomography for breast cancer imaging using a two-layer model to include chest-wall effects," *Proc. SPIE* **6511**, 65112L (2007).
16. Q. Zhu, N. Chen, and S. Kurtzman, "Imaging tumor angiogenesis using combined near infrared diffusive light and ultrasound," *Opt. Lett.* **28**(5), 337–339 (2003).
17. Q. Zhu, M. Huang, N. Chen, K. Zarfos, B. Jagjivan, M. Kane, P.

- Hegde, and S. Kurtzman, "Ultrasound-guided optical tomographic imaging of malignant and benign breast lesions: initial clinical results of 19 cases," *Neoplasia* **5**(5), 379–388 (2003).
18. R. C. Haskell, L. O. Svaasand, T. T. Tsay, T. C. Feng, M. Mc Adams, and B. J. Tromberg, "Boundary conditions for the diffusion equation in radiative transfer," *J. Opt. Soc. Am. A* **11**(10), 2727–2741 (1994).
 19. Q. Zhu, C. Xu, P. Guo, A. Aguirre, B. Yuan, F. Huang, D. Castillo, J. Gamelin, S. Tannenbaum, M. Kane, P. Hegde, and S. Kurtzman, "Optimal probing of optical contrast of breast lesions of different size located at different depths by US localization," *Technol. Cancer Res. Treat.* **5**(4), 365–380 (2006).
 20. B. Pogue and M. Patterson, "Review of tissue simulating phantoms for optical spectroscopy, imaging and dosimetry," *J. Biomed. Opt.* **11**(4), 041102 (2006).
 21. D. Grosenick, H. Wabnitz, K. T. Moesta, J. Mucke, P. M. Schlag, and H. Rinneberg, "Time-domain scanning optical mammography: II. Optical properties and tissue parameters of 87 carcinomas," *Phys. Med. Biol.* **50**(11), 2451–2468 (2005).
 22. W. Cheong, S. A. Prahl, and A. J. Welch, "A review of the optical property of biological tissues," *IEEE J. Quantum Electron.* **26**, 2166–2185 (1990).
 23. S. J. Matcher, M. Cope, and D. T. Delpy, "In vivo measurements of the wavelength dependence of tissue-scattering coefficients between 760 and 900 nm measured with time-resolved spectroscopy," *Appl. Opt.* **36**(1), 386–396 (1997).
 24. Y. Yang, O. Soyemi, M. Landry, and B. Soller, "Influence of a fat layer on the near infrared spectra of human muscle: quantitative analysis based on two-layered Monte Carlo simulations and phantom experiments," *Opt. Express* **13**(5), 1570–1579 (2005).
 25. P. Taroni, D. Comelli, A. Farina, A. Pifferi, and A. Kienle, "Time-resolved diffuse optical spectroscopy of small tissue samples," *Opt. Express* **15**(6), 3301–3311 (2007).
 26. Y. Xu, N. Ifimias, H. Jiang, L. L. Key, and M. B. Bolster, "Imaging of in vitro and in vivo bones and joints with continuous-wave diffuse optical tomography," *Opt. Express* **8**(7), 447–451 (2001).
 27. H. Jiang, N. Ifimias, Y. Xu, J. Eggert, L. Fajardo, and K. Klove, "Near-infrared optical imaging of the breast with model-based reconstruction," *Acad. Radiol.* **9**(2), 186–194 (2002).
 28. M. Schweiger and S. R. Arridge, "The finite-element method for the propagation of light in scattering media: frequency domain case," *Med. Phys.* **24**(6), 895–902 (1997).