Optical scattering coefficient estimated by optical coherence tomography correlates with collagen content in ovarian tissue

Yi Yang
Tianheng Wang
Nrusingh C. Biswal
Xiaohong Wang
Melinda Sanders
Molly Brewer
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Yi Yang, a Tianheng Wang, a Nrusingh C. Biswal, a Xiaohong Wang, b Melinda Sanders, b Molly Brewer, a,c and Quing Zhu a

a University of Connecticut, Department of Electrical and Computer Engineering, Storrs, Connecticut 06269
b University of Connecticut Health Center, Division of Pathology, Farmington, Connecticut 06030
c University of Connecticut Health Center, Division of Gynecologic Oncology, Farmington, Connecticut 06030

Abstract. Optical scattering coefficient from ex vivo unfixed normal and malignant ovarian tissue was quantitatively extracted by fitting optical coherence tomography (OCT) A-line signals to a single scattering model. 1097 average A-line measurements at a wavelength of 1310 nm were performed at 108 sites obtained from 18 ovaries. The average scattering coefficient obtained from the normal tissue group consisted of 833 measurements from 88 sites was 2.41 mm−1 (±0.59), while the average coefficient obtained from the malignant tissue group consisted of 264 measurements from 20 sites was 1.55 mm−1 (±0.46). The malignant ovarian tissue showed significant lower scattering than the normal group (p < 0.001). The amount of collagen within OCT imaging depth was analyzed from the tissue histological section stained with Sirius Red. The average collagen area fraction (CAF) obtained from the normal tissue group was 48.4% (±12.3%), while the average CAF obtained from the malignant tissue group was 11.4% (±4.7%). A statistical significance of the collagen content was found between the two groups (p < 0.001). These results demonstrated that quantitative measurements of optical scattering coefficient from OCT images could be a potential powerful method for ovarian cancer detection.

1 Introduction

Ovarian cancer has the lowest survival rate of the gynecologic cancers because it is predominantly diagnosed in Stages III and IV due to the lack of distinctive early symptoms and efficacious screening and diagnostic techniques. Prophylactic oophorectomy could reduce ovarian cancer risk by more than 50% and has become accepted as the standard of care for high risk women. However, it has recently been found to increase the mortality of women undergoing oophorectomy prior to the age of 45 or even before the age of 55 to 60. These high risk women are not candidates for hormone replacement therapy because of their increased risks of breast cancer. As a result, there is an urgent need to develop more sensitive tools to effectively evaluate the ovary during minimally invasive surgery so that a surgeon can determine if an early-stage cancer is present, and thus avoid removal of normal ovaries.

Collagen fibers are the main scatterers in the stroma underlying the epithelium, and a study from fluorescence confocal microscopy have shown that collagen content and directivity in stroma may change as precancer or cancer develops. Optical coherence tomography (OCT), which measures backscattered light generated from an infrared light source directed to the tissue, could be a potential method for detecting ovarian cancer during minimally invasive surgery. In addition to offering high resolution morphological images, OCT is capable of quantitatively estimating total attenuation coefficient (μt) by fitting the A-line measurements. μt is the summation of absorption coefficient (μα) and scattering coefficient (μs). As μα is much smaller than μs, μs is almost equal to μt and is a good estimate of the local scattering properties. Therefore, the quantitative μt extracted from OCT A-lines could reflect the local collagen content. This paper, to the best of our knowledge, is the first to report μs and its correlation with the collagen content in ex vivo human ovaries. The results have shown that changes in collagen can be an indicator of malignancy, and quantitative analysis of OCT images has the potential to characterize ovarian tissue and detect ovarian cancer.

2 Methods and Materials

2.1 OCT Fitting Model

The technical details of the OCT system were described in our previous publication. Two models are mainly used for the description of the OCT signal: the single scattering model and the multiple scattering model. For weakly scattering media (μs < 6 mm−1), the single scattering model with dynamic focusing is valid to extract μs; for highly scattering media, the multiple scattering needs to be considered. In our case, it is suitable to use the single scattering model. The calculated numerical aperture 0.05 of the sample arm optics in our fixed focusing OCT system was very low, which ensured the superficial scanning depth within the focal zone. In order to obtain a more accurate fitting, a confocal point spread function (PSF) of fixed focusing geometry was taken into account. Therefore, the OCT signal as a function of depth z is described as the compound of confocal PSF and Beer’ law:

\[ i(z) \propto \sqrt{\exp[-2\mu_s z]/[1 + ((z - z_f)/z_R)^2]}, \]

(1)

where \( i(z) \) is the amplitude of the interference signal, \( z_f \) is the position of a focal plane, \( z_R \) is the “apparent” Rayleigh length (in our experiment, \( z_f = 0, z_R = 0.75 \) mm), and the factor 2 accounts for the round trip attenuation.
the OCT imaging plane, embedded in paraffin, and sectioned to 7-μm thickness. Once the slides that correspond to the imaged planes were identified, they were stained using Sirius Red (SR) which binds specifically to collagen. The digital histological images of ovarian tissue covering about 1-mm depth within the fitting range were acquired by a microscope. The amount of collagen was quantitatively analyzed using ImageJ (NIH). Collagen area fraction (CAF) was measured as “stained collagen area /tissue area.”

3 Results and Discussion

Figure 1 shows one set of examples from normal [Figs. 1(a)–1(c)] and malignant [Figs. 1(d)–1(f)] ovarian tissue. The μs extracted from the OCT fitting areas marked as the white dashed squares in Figs. 1(a) and 1(d) are 2.86 and 1.29 mm−1, respectively. The inset in Fig. 1(a) shows the average A-line profile. The stained red area in Figs. 1(c) and 1(f) represents the collagen bundles. Clearly, the collagen amount, structure, and arrangement are quite different between normal and malignant ovarian tissues. The normal ovary exhibits almost exclusively collagen with interspersed stromal cells and the collagen fibril is randomly oriented and wavy interlaced; the collagen fibers in the malignant tissue are unidirectionally organized into thicker bundles. A larger amount of collagen is found in normal tissue (CAF = 58.3%) than in malignant tissue (CAF = 8.4%).

A total of 1097 μs were extracted from 108 sites of 18 ovaries. Based on the pathology results, 833 measurements of 88 sites are from normal ovarian tissue of 15 ovaries including normal ovaries (n = 7), ovaries with large cysts (n = 1), calcifications (n = 2), focal lymphocytes (n = 1), large follicles (n = 2), corpus luteum (n = 1), and a benign dermoid tumor (n = 1). However, fitting was performed at the normal tissue area based on corresponding H&E slides. 264 measurements of 20 sites are from malignant ovarian tissue of 3 malignant ovaries. The normal tissue group shows higher scattering property at a wavelength of 1310 nm ranging from 0.50 to 4.16 mm−1 with a mean value of 2.41 mm−1 (± 0.59). The standard deviation per ovary of this group varies from 0.29 to 0.78 mm−1 (± 0.46). Inset table: specificity and sensitivity at different thresholds.

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2.2 Ovary and Collagen Quantification

A total of 18 ex vivo ovaries from 10 patients with age ranging from 32 to 79 (mean 56) were investigated. The detailed patient information can be found in a different study. The study protocol was approved by the Institutional Review Board of UCHC and signed informed consents were obtained from all patients. Ovaries were imaged immediately after they were excised. The imaged surfaces were positioned at the focal plane of the sample arm optics. The measurements were performed under rotational scanning geometry. To eliminate the effects of scattered photons outside of focal zone and tissue surface curvature, only a 300 μm region corresponding to 100 A-lines centrally around the perpendicular illumination was selected for averaging. μs was estimated by fitting the averaged depth profile to the model described by Eq. (1). The fitting started about 40 μm below the tissue surface to avoid the effect of the surface epithelium, which is typically composed of a single layer of cuboidal to columnar cells (typically 10 to 20-μm thick). After OCT imaging, the ovaries were fixed in formalin, cut in 5 mm blocks parallel to the OCT imaging plane, embedded in paraffin, and sectioned to 7-μm thickness. Once the slides that correspond to the imaged planes were identified, they were stained using Sirius Red (SR) which binds specifically to collagen. The digital histological images of ovarian tissue covering about 1-mm depth within the fitting range were acquired by a microscope. The amount of collagen was quantitatively analyzed using ImageJ (NIH). Collagen area fraction (CAF) was measured as “stained collagen area /tissue area.”

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Fig. 3 (a) Histograms of CAFs obtained from normal (n = 158) and malignant (n = 143) ovarian tissue groups, Gaussian distribution fits are shown. (b) Statistics of normal (mean ± std: 48.8 ± 12.3) and malignant groups (mean ± std: 11.4 ± 4.7). (c) ROC curve of CAF. Inset table: specificity and sensitivity at different thresholds.

The average CAF of the normal group was 48.4% (± 12.3%), while the average CAF of the malignant group was 11.4% (± 4.7%). Figure 3(b) shows statistical significance between these two groups (p < 0.001). To compare with the ROC curve obtained from μs, we also provide the ROC curve obtained from the CAF as shown in Fig. 3(c). The sensitivity and specificity based on the percentage of CAF are summarized in the inset in Fig. 2(c).

Figure 3(a) shows the histograms of CAFs obtained from normal (n = 158) and malignant (n = 143) ovarian tissue groups. The average CAF of normal group was 48.4% (± 12.3%), while the average CAF of the malignant group was 11.4% (± 4.7%). Figure 3(b) shows statistical significance between these two groups (p < 0.001). To compare with the ROC curve obtained from μs, we also provide the ROC curve obtained from the CAF as shown in Fig. 3(c). The sensitivity and specificity based on the percentage of CAF are summarized in the inset in Fig. 3(c). The AUC obtained from CAF is 0.89. Because SR stain binds specifically to collagen, the different collagen characteristics found in normal and malignant tissue groups could effectively explain the scattering properties estimated from OCT measurements obtained from these two groups. Note that elastin is another protein which may also perform a similar role in reducing optical scattering. However, by examining the SR and H&E stains, no elastic fiber is present in the ovarian stroma which was also reported by other researchers.11 There is probably elastin in the walls of vessels which could not be accounted as the major contributor to the μs obtained from malignant ovarian tissue. Regarding the effect of blood absorption on estimated μs, the ovaries were in saline water for a few minutes before imaging to remove the blood from the surface. For absorption of the ovarian tissue, our earlier study using diffused light showed that the ovarian tissue absorption coefficients were in the range of 0.006 to 0.018 mm⁻¹ which was less than 1% of the average scattering coefficients reported here.12 Thus the fitted μs is a good estimate of μs.

4 Summary

In this paper, optical scattering coefficients from normal and malignant ovarian tissue groups were quantitatively extracted by fitting the OCT signal to a single scattering model. CAFs were measured by analyzing microscopic images stained by SR. These results have shown that the malignant ovarian tissue has lower scattering coefficient and less collagen than that present in normal ovarian tissue. These initial findings suggest that quantitative analysis of ovarian tissue optical properties extracted from OCT images could be a powerful tool to reveal ovarian tissue neoplastic changes and to characterize ovarian cancers.

Acknowledgments

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References