Evaluation of Freshness of Soft Tissue Samples with Optical Coherence Tomography Assisted by Low Frequency Electric Field

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We present an optical coherence tomography based methodology to determine freshness of soft tissue samples by evaluation of their interaction with low frequency electric field. Various biological tissues samples of different stages of freshness were exposed by low frequency electric current. The influence of the low frequency electric field on tissues was observed and quantified by the double correlation optical coherence tomography (dcOCT) approach developed in house. The quantitative evaluation of electric field — tissue interaction by dcOCT shows an increase of its relative magnitude with the tissue freshness from fresh to the non-fresh ones. The presented approach has a strong potential in food sciences for evaluation of meat quality.

Key words: optical coherence tomography; low frequency electric field; soft biological tissue samples.

Optical coherence tomography (OCT) is a non-invasive, high resolution optical imaging technique that can perform in vivo high resolution, cross-sectional microstructure imaging of biological tissues and other scattering materials [1, 2]. First successful application of OCT was demonstrated in ophthalmology, in particular, to visualize an anterior/posterior eye segment [3] and to measure a blood flow in the retina vessels [4, 5]. The field of OCT applications continues growing very intensively. The superior capability of imaging of the scattering media and inner flows with high spatial and time resolution allows one to conduct detailed investigations of the structure and the functioning of the blood circulatory system of humans and animals [6, 7], imaging of embryonal development [8], as well as the study of cell motility dynamics [9].

The effects of electromagnetic fields interaction with biological tissues have been studied widely in the past [10]. A number of studies have been performed to evaluate the effects of different electrical stimulation procedures to obtain tenderer and high quality meat [11]. Recently it was demonstrated that pulsed electric fields affect the moisture content of meat [12].

Recently, the double correlation OCT (dcOCT) approach was introduced and applied to observe molecular diffusion in skin tissues [13, 14], visualization vascular skin bed in vivo [15], investigate electro-kinetic response of biological tissues influenced by electric field [16, 17] and its changes with optical clearing [18].

In current paper, the dcOCT approach, a functional extension of conventional OCT, has been applied to study the electro-kinetic properties of tissues when a low frequency electric field is applied to tissue samples ex vivo.

Soft biological tissue samples are often used in biomedical research for validation of diagnostic methods and in vitro studies. Obviously, the freshness of tissue samples is an important issue that potentially can effect on the validity of experimental laboratory studies. Surprisingly there are no standards of freshness until now. The researchers rely on the very subjective parameters such as colour and smell of the samples. In current study we consider the application of dcOCT-based methodology to determine and/or quantify the freshness of soft tissue samples ex vivo stored at different conditions.

Materials and Methods. Recently it has been demonstrated that dcOCT is sensitive to visualize the degree of influence of the applied electric field in biological tissues [16–18]. In order to determine freshness of tissue samples, a standard swept source OCT (OCM1300SS, Thorlabs, Inc., USA) with central wavelength of 1325 nm and spectral bandwidth of 100 nm operated in polarization sensitive mode without phase retardation has been used to acquire OCT images of tissue samples ex vivo. The
samples are taken from fresh chicken breast pectoralis muscles and aged under different conditions: 0°C during 6 days, 4°C during 8 days, and 15°C during 8 days.

During the experiment, two electrodes (Ø=0.3 mm) made of stainless steel were inserted into the tissue sample towards each other at the depth of about 1 mm parallel to the direction of the collagen fibres. The distance between the tips of the electrodes inside the sample was of 1.5 cm. AC electric current/voltage from function generator (GFC 2100, Iso-Tech, UK) was applied to the electrodes. The amplitude of the signal was in the range of 1–20 V, the frequency varies within 0.1–5.0 Hz. OCT B-scans were taken from the area between the electrodes perpendicular to the direction of current flow. To evaluate the degree of similarity of OCT images obtained after and before the exposure of the electric field the dcOCT is applied. The dcOCT approach can be summarized in 3 steps, as shown in Figure 1. Step 1 involves the acquisition of the image before and after the exposure of the electric field. Step 2 consists of the Wiener filtering, while step 3 consists of the calculation of the cross correlation image and the relative magnitude of influence of the electric field. Detailed description of dcOCT approach is given in [15].

**Image processing techniques.** Background noise from the experimental setup and/or sudden movements of OCT probe result imaging artefacts (such as blur, physical offsets of tissue boundaries) and, therefore, makes it quite challenging to obtain images with high spatial resolution by using the image-quality demanding correlation procedure. In the cross-correlation analysis the size of the grid should also be carefully chosen as it is a trade-off between the processing time and the final quality of the outcome [19]. If a large grid (e.g. 40×40 pixels) is used, ‘blurring’ and a loss of structural signal occurs [19]. For a small grid (e.g. 7×7 pixels), the background noise of OCT has a significant impact on the structural signal which results in decorrelation [19]. Experimental evidence suggests that the optimal size of the grid for the biological tissues lies within the range 5–25 pixels [19].

Thus, the background noise should be suppressed. The adaptive Wiener filtering procedure removes the unwanted background noise from the images which allows better cross-correlation outcomes to be obtained for smaller grid sizes. It should be pointed out that the other techniques to eliminate the weak correlation effects could also be used (e.g. structural mask) [19].

The original implementation of the filtering procedure suggested by Wiener was applied [20]. Commonly used in image processing, Wiener filtering utilizes a sliding window (or grid) of size of M by N pixels centred on each pixel of the input image (Figure 2).
In the frequency-domain the optimal Wiener filter $W(l)$ is defined as [20, 21]:

$$W(l) = \frac{P_S(l)}{P_S(l) + P_N(l)},$$

(1)

where $P_S(l)$ and $P_N(l)$ are the power spectrums of signal and noise, respectively. $P_S(l)$ is obtained by taking the Fourier transform of that grid’s autocorrelation according to the Wiener–Khinchin theorem [20], whereas the power spectrum of noise is estimated by calculating the local variance for each grid [21]. Thus, the desired OCT signal is obtained by subtracting an estimate of the noise from the original signal. The procedure is repeated for all grids and resulting OCT images are obtained. Figure 3 shows an example of noise removal images by the Wiener filtering.

There are a number of other approaches to create Wiener filter coefficients, including Bayesian implementation of the Wiener filter, hidden Markov models, additive noise reduction, etc. [20].

Double correlation OCT imaging works by comparing two successive OCT images that correspond to the same structural information so that one is able to map out the regions of interest. The technique performs some extra image processing steps designed to enhance the experimental data. In the correlation mapping OCT approach [19] a grid $I_A$ from image $A(x, z)$ is compared to the same grid $I_B$ from image $B(x, z)$ defined over $X$–$Z$ plane. The $M$ by $N$ grid is shifted pixel-by-pixel across the images, forming a correlation plane $C(x, z)$. Normalized cross-correlation function can be written as:

$$C(x, z) = \frac{\sum_{p=0}^{M-1} \sum_{q=0}^{N-1} I_A(p, q)I_B(p, q)}{\sqrt{\sum_{p=0}^{M-1} \sum_{q=0}^{N-1} |I_A(p, q)|^2 \sum_{p=0}^{M-1} \sum_{q=0}^{N-1} |I_B(p, q)|^2}}.$$  

(2)

The output has values in the range $\pm 1.0$, indicating low correlation to both high positive and high negative correlation. Direct computation of normalized cross-correlation is very computationally expensive, time consuming and can be optimized by various techniques such as sum-tables [21]. However, more efficient approach to calculate the cross-correlation is based on the fast Fourier transform (FFT). The Fourier transform is widely used in a large class of signal processing algorithms for fast and efficient function decomposition and analysis [20]. Our approach utilizes the cross-correlation theorem, stating that the cross-correlation is equal to the inverse Fourier transform of the product between individual Fourier transforms of two functions, presuming one of them is represented by a complex conjugate [20, 21]. Thus, to find the cross-correlation between two grids the following equation is used [20, 21]:

$$C(x, z) = \text{FFT}^{-1}(\text{FFT}(I_A) \times \text{FFT}(I_B)).$$

(3)

**GPU acceleration of image processing.** The analysis of the images has been performed on GPUs utilizing recently developed CUDA framework in frequency-domain (Figure 4). CUDA framework gives access to the CUDA fast Fourier transform library [22]. The algorithm allows carrying out FFT computations in parallel utilizing hundreds of GPU cores providing a significant speed up compared to the standard CPU approach (up to 1000× faster) [23].

It should be pointed out that if the images do not have the same dimensions, the wraparound error could occur [20, 21]. To cancel the effects of spatial aliasing, the zero-padding procedure is typically employed: to prevent the periodic replicas from overlapping, the size of the images in the GPU memory is increased and the empty spaces are filled with zeros [22]. It is also suggested that before obtaining the Fourier transform both images are zero-mapped [20–22]. The procedure ensures that the sum of all pixels is zero and the maximum of the correlation function corresponds to the right position in the image instead of the brightest pixel [20, 21].

Moreover, the software development kit provided by the device manufacturer (Thorlabs, Inc., USA), allows
bypassing the step of manual acquisition of the OCT images and provides the opportunity for direct export of the OCT hardware FFT-encoded image data to the processing software. Incorporating this ability in the dcOCT method significantly improves the quality of imaging outcomes and speeds up the image processing (up to 30–40 times, i.e. makes it possible to proceed in real time).

Results and Discussion. The soft biological tissues at different stages of freshness and corresponding conventional OCT images are presented in Figure 5. As one can see no changes in colour and in structural integrity correlating with the sample freshness can be observed and/or distinguished.

To distinguish the samples of biological tissues in order of their freshness we apply dcOCT approach presented above. Figure 6 (a) shows an example of typical OCT image of chicken sample aged at 15°C during 8 days, while Figure 6 (b) corresponds to cross correlation image calculated using the dcOCT approach. At both figures the arc-shaped pattern corresponds to the reflection from the surface of electrode.

The relative magnitude of influence of the low-frequency external electric field (See Figure 6 (b)) is counted based on the dcOCT image as [17]:

$$\Psi = 1 - \frac{1}{M \times N} \sum_{x=1}^{M-1} \sum_{z=1}^{N-1} C(x, z). \quad (4)$$

Further, we investigated the changes of this relative magnitude vs the tissue samples freshness. Figure 7 shows the results of the calculation of the relative magnitude of influence of the electric field in chicken meat samples. As one can see fresh tissue has the lower relative magnitude and it follows a trend in which the non-fresh tissue stored at 15°C during 8 days has the highest relative magnitude. This suggests that the evaluation of the relative magnitude of influence of the electric field can be used to assess relative freshness of soft biological tissues (here, we present only the results for chicken breast tissues for the purpose of brevity, but similar results were also observed for other soft biological tissues).

We suggest that the observed behaviour is attributed
to the tissue tendering process. The tenderness increases due to natural enzymatic changes that take place in tissue samples. Less tender tissue results in high correlation which itself reduces the amount of affection of the electric field.

**Conclusions.** The obtained results show that freshness of soft biological tissues affects the propagation of ac current in the samples that can be observed with dcOCT. It has been shown that the relative magnitude of influence of the low-frequency external electric field on the biological tissue ($\Psi$) increases from fresh to non-fresh samples as an inverse exponential function. This potentially can be used to build a phenomenological model to assess quantitatively the freshness of soft biological tissues. The possibility of real-time data processing essential for the real-life application was also demonstrated. We foresee the further development of the approach in applications in modern biomedicine as well as in the food industry.

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