Use of kurtosis for locating deep blood vessels in raw speckle imaging using a homogeneity representation

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Abstract. Visualization of deep blood vessels in speckle images is an important task as it is used to analyze the dynamics of the blood flow and the health status of biological tissue. Laser speckle imaging is a wide-field optical technique to measure relative blood flow speed based on the local speckle contrast analysis. However, it has been reported that this technique is limited to certain deep blood vessels (about $\rho = 300 \mu m$) because of the high scattering of the sample; beyond this depth, the quality of the vessel’s image decreases. The use of a representation based on homogeneity values, computed from the co-occurrence matrix, is proposed as it provides an improved vessel definition and its corresponding diameter. Moreover, a methodology is proposed for automatic blood vessel location based on the kurtosis analysis. Results were obtained from the different skin phantoms, showing that it is possible to identify the vessel region for different morphologies, even up to $900 \mu m$ in depth.

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Keywords: blood vessel location; raw speckle images; kurtosis; co-occurrence matrix; image processing.

1 Introduction

The visualization of blood vessels is of fundamental importance for a wide variety of biological and biomedical applications, such as obstruction, stiffness, and response to an external stimulus. Laser speckle imaging (LSI) is a technique based on the spatial–temporal integration of the light scattered from a biological sample when illuminated with coherent light and imaged by an optical detector (i.e., CCD camera). Particle motion (i.e., blood cells) in the illuminated area causes a decrease in contrast, seen as a blurring effect in the image, which is related to the speed of the particles in the illuminated sample. Hence, LSI is used to measure the relative blood flow speed.

For deep vessels, LSI shows some limitations because of the strong scattering produced by static structures, such as the skull or the epidermis. Several approaches have been proposed to overcome these limitations. Pulsed photothermal radiometry (PPTR) photoinduces heating in blood vessels with a laser pulse to improve the visualization. Photothermal-LSI is a recently developed technique that combines the photoinduced heating of PPTR and the contrast of integrated intensity of LSI to improve the visualization of the deep blood vessels. Magnetomotive laser speckle imaging uses paramagnetic nanoparticles introduced in the vasculature. Under the influence of an external magnetic field, the mobility of those particles increases, allowing their visualization through the contrast of integrated speckle. Physicochemical tissue optical clearing (PCTOC) uses a topical substance that matches the refractive index of the skin. Although these techniques may increase the visualization of the deep blood vessels up to a few hundred micrometers, all of them require an external agent or stimulus. In a recent work, Postnov et al.

2 Materials and Methods

2.1 Texture and Co-occurrence Matrix

In computer vision, a texture is characterized by gray-level variations in an image that form small similar and repetitive regions. Thus, in order to describe a texture, it is necessary to analyze the distribution of its gray levels. While a histogram provides information about the distributions of gray levels, it...
does not take into account the relationship between a central pixel \( p \) and its neighbors.\(^{14} \) This information is important to provide a rich description of the texture. The gray level CM is one of the standard methods to analyze texture in digital image processing\(^{13} \) and can be used for describing the information of RSL, as we report in an initial approach.\(^{15} \) Through second-order statistics, this approach studies the way in which the gray levels are distributed in an image or region and their spatial relationships.\(^{16,17} \) A CM indicates the frequency at which a pair of pixels with gray values \( i \) and \( j \), located spatially at a distance of \( d \) pixels in the direction \( \theta \), occurs in an image \( I \). A distance value must be chosen carefully because a large distance may result in texture overlapping; oftentimes, \( d = 1 \) is used. The analysis direction takes values of \( \theta = \{0 \text{ deg, } 45 \text{ deg, } 90 \text{ deg, } 135 \text{ deg} \}; \) the analysis in all the directions is obtained by the addition of the four matrices.

Let \( p \) be a pixel in the image \( I^{H\times W} \) of \( H \) rows and \( W \) columns, where \( p \) takes gray values in the range \([0, G - 1] \), \( G \) being the range of possible values (256 for an 8-bit image). For instance, if \( I \) is an image with five possible gray values from 0 to 4 \((G = 5)\), as shown in Fig. 1(a), and the parameters \( d = 1 \) and \( \theta = 45 \text{ deg} \) are considered for its CM, then the procedure is as follows. A matrix \( CM^{G\times G} \) is generated and set to zeros; CM is always square and its dimensions depend on \( G \). To assign values to the CM, each pair of pixels in \( I \) is analyzed considering the parameters \( d \) and \( \theta \). Figure 1(b) shows that for the central pixel \( p \), the parameters indicate that \( q_1 \) is the neighbor pixel to analyze. Often, the neighborhood analysis is made symmetrically, hence, the coocurrence is also extended to \( q_2 \). Suppose we want to know the cooccurrences of a gray level \( i = 1 \) (reference level) with a gray level \( j \) (comparison level), denoted as CM\((i, j)\). For \( j = 0 \), the cooccurrences with \( i \) happen when the coordinates of \( p \) are \( \{I(1,0), I(1,1)\} \), therefore CM\((1,0) = 2 \). For \( j = 1 \), the cooccurrences with \( i \) happen in coordinates \( \{I(2,0), I(2,1)\} \) by considering a neighborhood in \( q_1 \). Also, in this case, the cooccurrences happen in a symmetrical way, when \( p \) has coordinates \( \{I(1,1), I(1,2)\} \) by considering a neighborhood in \( q_2 \); therefore, CM\((1,1) = 4 \). The final CM is obtained by analyzing the cooccurrences between all the possible values of \( i \) and \( j \), as shown in Fig. 1(c) for this example.

From the CM calculus, two important things should be noted: (a) a new representation of the gray levels in \( I \) is provided by the CM, which is considered a symmetric two-dimensional histogram, but it does not characterize a texture by itself;\(^{13,18} \) (b) although the amount of information in \( I \) has been reduced in the CM, a large amount of data remain for texture characterization, i.e., \( G \times G \) characteristics.

### 2.2 Feature Extraction

In order to use the information provided by the CM, some features must be calculated. Haralick et al.\(^{17} \) proposed 14 textural features that can be calculated from the CM to identify difference among textures. One of the most used features is the inverse difference moment, a measure of the local homogeneity, calculated as

\[
H_g = \frac{\sum_{i=1}^{G} \sum_{j=1}^{G} P(i, j|d, \theta)}{1 + |i - j|},
\]

where \( P(i, j|d, \theta) \) is the probability density function obtained by dividing the CM matrix by the total pixels of image \( I (H \times W) \); \( i \) and \( j \) are the reference and comparison levels of the CM matrix described in Sec. 2.1.\(^{16,20} \) This feature provides information regarding the differences between gray levels in the image, taking values in the range \([0, 1]\), where 1 indicates a limited range of gray levels (high homogeneity) and 0 indicates a wide diversity of gray levels (high roughness). For instance, texture T1 in Fig. 2(a) contains coarse structures (bricks) identified by having close gray values and smooth transitions among them. Thus, high occurrences are expected near the CM diagonal of this image with \( H_g = 0.3631 \). In the opposite case, when an image contains fine granulated structures (T2), the scattered occurrences are expected throughout the CM, as in the case of Fig. 2(b), and consequently, a lower homogeneity \( H_g = 0.1169 \). Thus, it is possible to find differences among textures by calculating features, such as homogeneity based on the CM values.

### 2.3 Skin Phantoms

These skin phantoms consist of two parts: dermis and epidermis with specific characteristics, as suggested by Saager et al.\(^{21} \) For the dermis phantom [Fig. 3(a)], we used a transparent resin with the necessary amount of titanium dioxide \((\text{TiO}_2 \ 1.45 \text{ mg/mL})\) to simulate the scattering coefficient of the skin \( (\mu_s = 2 \text{ mm}^{-1}) \). The epidermis phantom [Fig. 3(b)] was made of silicone (polydimethylsiloxane) mixed with \( \text{TiO}_2 \) powder (2 mg/mL) to simulate the corresponding scattering properties \( (\mu_s = 3.0 \text{ mm}^{-1}) \) and dried coffee (10 mg/mL) to simulate the absorption
 coefficient ($\mu_a = 0.2 \text{ mm}^{-1}$). Different thicknesses of phantom epidermis layers, namely 200, 400, 500, 600, 700, and 900 $\mu$m, were placed on top of the phantom dermis to simulate different depths of the vessels [Fig. 3(c)].

2.3.1 **In vitro vessels**

For this work, two types of in vitro vessels were used: straight and bifurcated. For the straight vessel, a glass capillary with an
inner diameter of 700 μm was used, whereas for the bifurcated vessel, a slide with 300-μm-inner diameter microchannels (thinXXS Microtechnology AG, Germany) was used. Both in vitro vessels were placed above silicon layers containing TiO2 powder to mimic the scattering properties of the soft biological tissues. The phantoms described in Sec. 2.3 were placed on top of the slide to simulate the depth of the vessels [Fig. 3(e)].

2.4 Experimental Setup

The LSI setup (Fig. 4) consists of laser He–Ne (632.8 nm) that impinges on a static engineered diffuser (Model ED1-C20, Thorlabs Inc.), which in turn illuminates homogeneously a circle with a 1-cm diameter on the phantom’s surface. If the phantom is not homogeneously illuminated, the intensity distribution can influence the spatial contrast. A lens coupled to a CCD camera (Model Retiga 2000R, QImaging, Canada) images the surface of the phantom on the CCD. The magnification of the optical system is $M = 0.285$ and the corresponding speckle size is $≈7.9 \mu m$. A linear polarizer is placed in front of the lens with its transmission axis crossed with the He–Ne polarization to minimize the specular reflection. The CCD camera is connected to a computer to save the captured speckle images. The camera exposure time is set to $T = 10$ ms because this value is quite common for in vivo LSI applications. This exposure time is approximately three-orders of magnitude larger than the correlation time reported for a similar LSI experiment at a flow speed of 8 mm/s. An infusion pump (Model NE-500, New Era Pump System Inc.) was used to inject the intralipid at 3% in water as blood substitute, whose pump (Model NE-500, New Era Pump System Inc.) was used to inject the intralipid at 3% in water as blood substitute, whose distribution can influence the spatial contrast.

3.2 Homogeneity Representation

It must be taken into account that the vessel region usually occupies a small area in the image; therefore, the homogeneity value must be computed for the small regions. Hence, the image $P_{xy}$ is divided into subimages or subregions through an analysis window $P_{xy}$ [Fig. 5(a)], and a CM is calculated for this area, just as the contrast image is calculated. With the aim of increasing the precision in the location of the vessel, a certain level of information redundancy is necessary. There must be an overlap between each pair of subregions, i.e., $J_m+1$ must have a displacement $\Delta$ with regard to $J_m$ with $1 \leq \Delta \leq (h-1)/2$ where $h$ is the size of the analysis window. In this way, we generate a new representation of the RSI through the homogeneity image, shown in Figs. 5(d) and 5(e), in which it is possible to improve the discrimination between dynamic and static regions. As shown, high and low values of homogeneity will be associated to dynamic and static regions, respectively. Homogeneity representation enables smoother transitions among consecutive regions and defines the vessel better.

Often, the analysis window has a small size, such as $5 \times 5$ or $7 \times 7$, but if a large area is analyzed, it is possible to find statistics that better describe the characteristics of that area. This is particularly interesting because the speckle pattern is generated randomly. If the analysis is made using a small window, a higher dispersion is found [as shown in Figs. 5(b)–5(c) and Figs. 5(d)–5(e)]; but if a larger window is used, that dispersion is attenuated because the occurrence of gray levels becomes higher, i.e., some of them are no longer seen as isolated occurrences and the relationship among pixels is higher, although they still reflect local values. As the use of a larger window improves the homogeneity representation by avoiding strong variations, the use of a larger window with $h = 31$ and $\Delta = 5$ will be explored. As shown in Figs. 5(f) and 5(g), the use of a larger window improves the homogeneity representation by avoiding strong variations.

The transversal view of $I_{hh}$ ($P_{hh}$, hereinafter named as homogeneity profile) shows with more detail that low homogeneity values are related to static region, whereas high homogeneity values are related to dynamic regions, i.e., regions of high
contrast. We can also observe that the homogeneity rapidly increases in the regions closest to the vessel until it reaches its maximum level at the center of the vessel. In this new representation, it is clearer to distinguish the static and the dynamic regions, thereby improving the vessel location. Although this homogeneity profile corresponds to an RSI at $\rho = 0\, \mu m$ in depth (which we call the reference case), this analysis can be extended to deeper vessels.

### 3.3 Homogeneity Profile Characterization

In LSI, it is known that the vessel region is associated with a blurring effect due to the blood flow (dynamic region); however, as the depth increases, this effect is reduced, making it difficult to identify the vessel region. Although a certain level of blurring remains in the image, the dynamic region becomes very similar to the static region. Therefore, we need to find other ways to identify the vessel region from an image that appears to be completely noisy. The homogeneity representation will be used for this purpose.

For the proposed methodology, an initial set of RSI (S1) was tested. The set contains a package of images obtained using the experimental setup described in Secs. 2.3 and 2.4. Table 1 describes the characteristics of this package, consisting of 30 RSI images at $\rho = 0\, \mu m$ in depth. This is done in order to study whether there is a characteristic pattern in the homogeneity behavior. The starting point (reference case) is $\rho = 0\, \mu m$. All the homogeneity profiles were found to present a distribution with a tall narrow peak and long tails due to the extreme deviation from the central tendency of the values, this feature suggest the use of kurtosis. This deviation was caused by the rapid increase of homogeneity values in the dynamic region [similar to Figs. 5(e)–5(g)].

The kurtosis is a measure that describes the shape of a distribution based on the ratio between the central tendency and the deviation of the data. When values are greatly deviated from the central tendency, the kurtosis takes values $>3$, and the distribution is called leptokurtic. Given that a superficial vessel at $\rho = 0\, \mu m$ has a characteristic leptokurtic homogeneity distribution, kurtosis can be used as a parameter of reference for vessel location. In order to prove this statement, the kurtosis value ($k$) was calculated for the 30 profiles, according to

$$
\kappa = \frac{n \sum_{i=1}^{n} (\bar{x} - \mu)^4}{\sigma^4},
$$

where $\bar{x}$ is the mean, $\mu$ is the central tendency, and $\sigma$ is the standard deviation.

#### Table 1 Characteristics of the reference set (S1) of RSI.

<table>
<thead>
<tr>
<th>Image dimensions (pixels)</th>
<th>Vessel diameter ($\mu m$)</th>
<th>$\rho$ ($\mu m$)</th>
<th>Speed (mm/s)</th>
<th>$\kappa$</th>
<th>PSNR</th>
</tr>
</thead>
<tbody>
<tr>
<td>329 × 200</td>
<td>700</td>
<td>0</td>
<td>20</td>
<td>4.42 ± 0.02</td>
<td>32.70</td>
</tr>
</tbody>
</table>

Fig. 5 (a) An RSI of a superficial vessel (i.e., $\rho = 0\, \mu m$) and (b, c) its contrast image computed with an analysis window of $5 \times 5$. The homogeneity computed images using analysis windows of (d, e) $5 \times 5$ and (f, g) $31 \times 31$, respectively. Left and right columns show the transversal and frontal views of the images, respectively.
where \( \mu^4 \) is the fourth moment about the mean, i.e., the sum of the difference between the mean \( \mu \) and each \( z \) value raised to the fourth power in a given set of \( n \) values; \( \sigma \) is the standard deviation. The obtained mean kurtosis \( \bar{k} \) of the reference case \( \rho = 0 \mu m \) is 4.42 \( \pm 0.02 \) confirming its leptokurtic distribution (Table 1). In order to determine the quality of the set S1, the peak signal-to-noise ratio (PSNR) for all RSI images of S1 was estimated using

\[
\text{PSNR} = 10 \log_{10} \left( \frac{G - 1}{\sum_{a=0}^{N-1} \sum_{b=0}^{W-1} (I(a, b) - \bar{I})^2} \right),
\]

where \( \bar{I} \) is the mean value of the image \( I \).28 The images of S1 were found to have a high PSNR; therefore, we use its mean kurtosis value \( k = 4.42 \) as a reference for the proposed analysis.

Once the reference homogeneity profile has been established, it is necessary to locate the vessel and later to estimate its diameter, as shown in Sec. 4.1. A transversal cut of the homogeneity profile [Fig. 6(a)] shows that it is highly correlated with the actual dimension of the blood vessel [Fig. 6(b)]. Also, it notes that the dotted blue lines correspond to the actual vessel location, whereas the region comprised between a continuous red line and a dotted blue line is associated to the area around the vessel due to the migration of photons from the vessel to the surrounding regions [Figs. 6(a) and 6(b)]. The actual vessel region in the homogeneity representation was found to be best defined in the interval comprised between the maximum value (vessel center) and a standard deviation of the homogeneity profile, i.e., \( \max(P_{H_g}) \pm \sigma \) (dotted blue lines).

Nonetheless, the supposed location depends on a distribution with high kurtosis since the standard deviation may comprise a wide or narrow region, depending on the peak width of the homogeneity distribution. As observed in Fig. 7(a), the deeper the vessel, the wider the homogeneity profile (and therefore, the lower the kurtosis). Figure 7(b) shows a close up around the peak and the vertical dotted lines indicate the actual vessel location for reference purposes only. In the same figure, the circular markers correspond to \( \max(P_{H_g}) = \sigma \) for the different depths of the vessels. Note that the criteria for vessel location established for the reference case is no longer valid for deeper vessels. Hence, the one standard deviation is not a good parameter for locating deep vessel.

We propose a weighted standard deviation \( (\sigma_w) \) parameter [Eq. (5)] that compensates the flattening of the distribution caused by the increasing depth. The weight \( w \) depends on the ratio between the kurtosis of the homogeneity profile at a given depth \( d \left( k_d \right) \) and the kurtosis of reference \( k_0 = 4.42 \). The deeper the vessel, the lower the \( \sigma_w \). Thus, \( w \) controls the proportion that must be considered from the standard deviation at a given depth \( (\sigma_d) \); therefore, it restricts the peak width associated to the vessel region. Figure 7(c) shows the vessel location obtained with \( \sigma_w \), as we observe it fits best with the actual vessel location,

\[
\sigma_w = \frac{k_d}{k_0} \cdot \sigma_d.
\]

The proposed methodology is summarized in Fig. 8, where the input is an RSI image processed by an analysis window, just as it is in a traditional LSI, in order to calculate its corresponding CM and homogeneity values. Once the homogeneity has been calculated for the whole image, the kurtosis analysis is made by sections and the blood vessel is located according to Eq. (5). Finally, a segmented image that shows only the vessel region is obtained.

4 Results

4.1 Linear Vessel

In order to assess the proposed methodology under different conditions, we tested it using another set of RSI obtained at different depths as described in Table 2.

For the set S2, the vessel was manually located among the rows [141, 168] for \( \rho = 0 \mu m \), and this location is used as a reference for future analysis, whereas for other depths, the location of the straight vessels was done automatically by searching for the peak of the homogeneity profiles and performing the kurtosis analysis. The proposed methodology was first tested in straight vessels by processing homogeneity profiles for each column and computing the kurtosis analysis. Figure 9 shows
the vessel extracted from S2 at different depths and with a velocity of 10 mm/s. As one can see, the located vessel is better defined at lower depths. At higher depths, the image is noisier, and therefore, the homogeneity values in static and dynamic regions become similar and harder to define.

The goal of performing in vitro tests at different depths is to demonstrate that $\sigma_w$ allows the estimated vessel location to match with the real vessel location, and therefore, it is possible to measure the vessel diameter. For our skin phantoms, it is a known value (700 $\mu$m). Thus, based on the obtained location at each depth, we were able to estimate the vessel diameter and its deviation from its actual value. Figure 10 shows the plot of this estimation for set S2 by comparing the results at two velocities. We can observe that the speed does not seem to affect the estimation significantly of the vessel diameter as the majority of the values lies within 10% of the actual vessel diameter (dashed lines). The plot shows how close the automatic location is to the reference diameter (solid line).

### 4.2 Bifurcated Vessel

In in vivo situations, the vessels are not straight but takes complicated forms and even bifurcations. In order to apply our proposed kurtosis analysis, a bifurcated vessel with a diameter of 300 $\mu$m and a depth of $\rho = 400 \mu$m was tested [Fig. 11(b)].

<table>
<thead>
<tr>
<th>Set</th>
<th>Image dimensions (pixels)</th>
<th>Vessel diameter ($\mu$m)</th>
<th>$\rho$ ($\mu$m)</th>
<th>Speed (mm/s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>S2</td>
<td>298 × 313</td>
<td>700</td>
<td>0, 200, 400, 500, 600,700, 900</td>
<td>10,20</td>
</tr>
</tbody>
</table>

Table 2 Characteristics of the set S2 of RSI used for testing the proposed methodology.
In this vessel morphology, the analysis of distributions is performed along each column and each row. Figure 11(a) shows the unidimensional profile corresponding to column $c=10$, where the two main peaks indicate a high homogeneity area related to the left branches of the vessel [Fig. 11(b)]. The same analysis is performed but along the row $r=165$ [Fig. 11(c)]. It is important to remark that only the most prominent peaks are considered, and each one is analyzed as an individual distribution. Evidently, these peaks match with the center of the vessel branches but they cover a wider area than the reference as expected. For instance, the upper peak in Fig. 11(a) comprises rows [210, 270]. However, using the proposed parameter $\sigma_w$, a better location of the vessel is achieved once again now between rows [233, 251]. The final location of the vessel is given by adding both results, and it is used as a marker. When this marker overlaps with the homogeneity representation, it generates the segmented image of the vessel. Figures 11(d) and 11(e) show the result of automatic vessel location performed in the contrast and homogeneity representations. Although the vessel is well located in both representations, it presents a poorer definition in the contrast representation unlike the homogeneity representation, which retains better the vessel shape. The white lines in Fig. 11(b) indicate the ideal definition and location of the vessel.

4.3 Discussion

The full width half maximum (FWHM) is a common measure often used to estimate the width of a function or signal. In this way, the FWHM and $\sigma_w$ accomplish a similar function. Here, we compare the performance for vessel location of the FWHM and $\sigma_w$ measurements for each depth (set S2). Figure 12 shows the differences in the vessel diameter when the vessel location is estimated with $\sigma_w$ and with FWHM. We can observe that the FWHM has a marked deviation from the actual diameter, mainly for higher depths since the homogeneity profiles tend toward a normal distribution. As shown in Fig. 10, the deviation of the automatic location proposed here is smaller than the 10% (dashed lines) of the diameter of the reference (black continuous line).
With the aim of measuring the error between $\sigma_w$ and FWHM, the coefficient of the variation of the root-mean-square error \( \text{CV}(\text{RMSE}) \) was calculated by Eq. (6), where $r_v$ is the actual vessel diameter, $v_e$ is the diameter estimated for each depth, and $\bar{v}$ is the mean value of the estimations. Error estimation for S2 (Fig. 12) indicates values of $\text{CV}(\text{RMSE})_{\sigma_w} = 0.089$ and $\text{CV}(\text{RMSE})_{\text{FWHM}} = 0.561$. Although the main objective of this work is to locate the vessel, we were able to provide an accurate estimation of its diameter based on $\sigma_w$. Since the homogeneity provides a representation where the transitions between the static and dynamic regions are smoother, the kurtosis analysis allows locating the blood vessels accurately. Although the kurtosis analysis can also be applied to the traditional contrast representation, it provides less accurate results because of the high variations in its representation [Fig. 5(b)] as shown in Fig. 11.

$$\text{CV}(\text{RMSE}) = \sqrt{\frac{\sum_{i=1}^{n} (v_i - \bar{v})^2}{n}}.$$ (6)

Fig. 11 (a) Homogeneity profile for $c = 10$, (b) bifurcated vessel at $\rho = 400 \mu m$ in the RSI, (c) homogeneity profile for $r = 165$, and the segmentation of the bifurcated vessel according to the estimated location using kurtosis analysis over (d) a contrast and (e) a homogeneity representation.

Fig. 12 Comparison of vessel diameter estimation using $\sigma_w$ and FWHM in set S2.
Table 3: Reported works for visualizing deep blood vessels.

<table>
<thead>
<tr>
<th>Method</th>
<th>Diameter vessel</th>
<th>Speed or flow</th>
<th>Depth</th>
<th>Brief description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Photothermal LSI</td>
<td>320 μm</td>
<td>4 mm/s</td>
<td>400 μm epidermal phantom</td>
<td>Improve the visualization of blood vessels by increasing local temperature with a 595-nm laser pulse. It could not be applied on temperature-sensitive areas. The diameter vessel is not estimated.</td>
</tr>
<tr>
<td>PCTOC</td>
<td>Not specified</td>
<td>0 and 5 mL/s</td>
<td>1.45 mm ex vivo porcine skin sample</td>
<td>PCTOC method was employed to enhance the contrast of LSI in ex vivo and in vivo experiments. The diameter vessel is not estimated.</td>
</tr>
<tr>
<td>LSI-SFDI</td>
<td>2 mm</td>
<td>6 mm/s</td>
<td>2 and 4 mm/s epidermal phantom</td>
<td>Depth-sensitive speckle contrast is shown in skin phantoms by separating a shallow vessel (2 mm) from a deep vessel (4 mm) using a high spatial frequency of illumination and LSI analysis. The vessel diameter is not estimated.</td>
</tr>
<tr>
<td>Gradient analysis and LSI</td>
<td>200 μm</td>
<td>Not specified</td>
<td>Exposed blood vessels, in vivo</td>
<td>Use an image-processing algorithm based on second-order gradient analysis of LSI images to estimate the vessel diameter and blood flow from rat mesenterium exposed and mouse cortex.</td>
</tr>
<tr>
<td>This work</td>
<td>700 μm</td>
<td>10 and 20 mm/s</td>
<td>0 to 900 μm epidermal phantom</td>
<td>External agent is not required neither invasive procedures. Vessel location is estimated only from the RSI. Vessel diameter also can be estimated.</td>
</tr>
</tbody>
</table>

However, both representations can complement each other. On one hand, contrast is a well-known representation and provides information to calculate the speckle flow index (SFI), which is proportional to the blood flow velocity. On the other hand, the homogeneity representation and its kurtosis analysis provide a better blood vessel location even at higher depths. Therefore, the result of the latter can be used as a marker to indicate the region where the former should be calculated and still obtain the SFI.

Table 3 shows a comparison among different methods based on LSI analysis to improve the visualization of deep (hundreds of microns) blood vessel. Some of them use an external agent, for example, photothermal-LSI uses a 595-nm laser pulse to thermally excite the blood flow, resulting in a local increase of temperature and thus, a transient decrease in speckle contrast; this technique should not be applied on temperature-sensitive areas, such as brain. PCTOC uses glycerol to reduce the scattering of in vivo and ex vivo skin samples; 60 min after the application of PCTOC, a sequence of RSI is acquired and processed with the LSI algorithm. LSI-SFDI uses structured illumination and LSI to separate a 2-mm-diameter shallow vessel (2 mm depth) from a deeper one (4 mm depth) but the vessel diameter is not estimated. Postnov et al. proposed an image-processing algorithm based on second-order gradient analysis of LSI images to estimate the vessel diameter and blood flow from an exposed rat mesenterium and a mouse cortex. Our results compare quite well with the previous results but, in addition, can improve the visualization of deep vessels and estimate its diameter without the use of external agents and in a noninvasive way.

Although in all image-processing methods there is a loss of information, due to the required processes for generating the homogeneity representation and the kurtosis analysis, there is also a trade-off in the location of the blood vessel that has demonstrated to be accurate. Moreover, the homogeneity representation visually improves the blood vessel definition, and whereas the loss of information is inevitable, new information is also gained (vessel location). It is important to remark that several versions of the information are generated as the information is being processed and then, at the end of our process, the segmented image can be used as a marker. The resulting marker in our proposal can be used just for indicating the vessel location, as the white lines in Fig. 11(b), over the homogeneity representation or over the raw speckle. In this way, original and new information can be complemented. Even more, if a contrast representation is also computed, the marker can be used over it.

5 Conclusions

In this work, it has been demonstrated that by exploring a representation based on the homogeneity values and using second-order statistics, it is possible to improve the visualization and definition of deep blood vessel regions in RSI obtained from skin phantoms. Moreover, a relationship between the vessel region and high kurtosis levels has been demonstrated. The kurtosis analysis tests prove that it is possible to obtain well-defined vessels and provide a good accurate estimation of their corresponding diameter. This work provides a methodology that works automatically and can locate vessels with different morphologies and depths as large as 900 μm in depth, three times the depth previously reported by other authors. Nevertheless, it is necessary to apply the proposed methodology in in vivo experiments. This research topic will be addressed in future work.

Disclosures

The authors have no conflicts of interest to declare in this manuscript.

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