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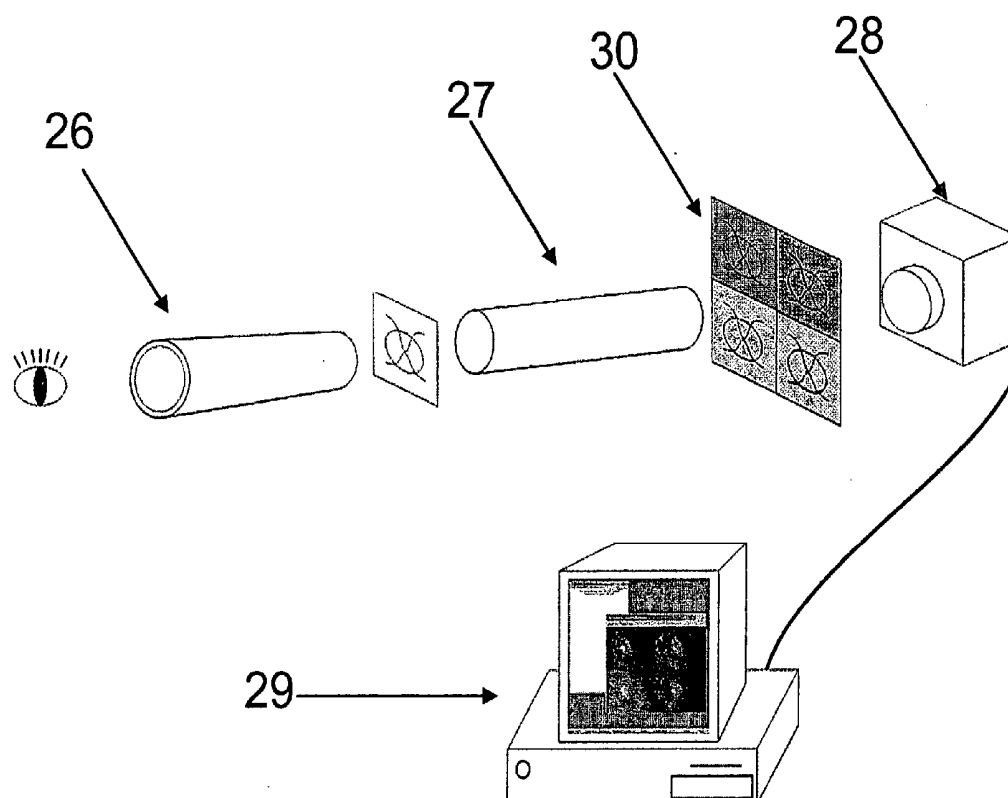
(19) **United States**(12) **Patent Application Publication** (10) **Pub. No.: US 2006/0276698 A1**
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Halldorsson et al.(54) **AUTOMATIC REGISTRATION OF IMAGES**(76) Inventors: **Gisli Hreinn Halldorsson**, Reykjavik (IS); **Einar Stefansson**, Reykjavik (IS); **Jon Atli Benediktsson**, Reykjavik (IS); **Robert Arnar Karlsson**, Reykjavik (IS); **Sveinn Hakon Hardarson**, Reykjavik (IS); **Thor Eysteinnsson**, Reykjavik (IS); **James M. Beach**, Slidell, LA (US)

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A61B 5/00 (2006.01)(52) **U.S. Cl.** **600/340**(57) **ABSTRACT**

The present invention relates to a system for automatically evaluating oxygen saturation of the optic nerve and retina, said system comprising: image capturing system further comprising: a fundus camera (26), a four wavelengths beam splitter (27), a digital image capturing device (28), a com-

puter system, image processing software performing in real-time the steps of: registering set of multi-spectral images (1) by, binarizing the multi-spectral image (7), find the all the border regions of each image by finding the region including the straight line that passes the most number of points in the region (8), use the orientation of the borders to evaluate the orientation of each spectral image (9), equalize the orientation of each spectral image by rotating the spectral image (10), edge detect each spectral image (11), estimate the translation between the spectral images based on the edges of adjacent images (12), transform the images to a stack of registered images (13), locating blood vessels (2) in each of the multi-spectral images by, retrieving registered spectral images (14), for each spectral image, remove defective pixels (15), enhance image contrast (16), perform top-hat transform using structuring element larger than the largest vessel diameter (17), binarize each image (18), apply filter for smoothing the image (19), combine all the spectral images using binary AND operator (20), skeletonize the resulting image (21), prune the skeleton image (22), re-grow the pruned image (23), locate junction points (24), evaluating the width of the blood vessels (3) by calculating the normal vector to the vessel direction to, estimate the direction and position of the vessel profile, evaluate positions of vessel walls based on the vessel profile obtained, evaluate vessel width based on the position of the vessel walls, selecting samples for calculating optical density, calculating the optical density ratio (4), evaluating oxygen saturation level (5), and presenting the results (6), wherein presenting the results may include presenting numerical and visual representation of the oxygen saturation.



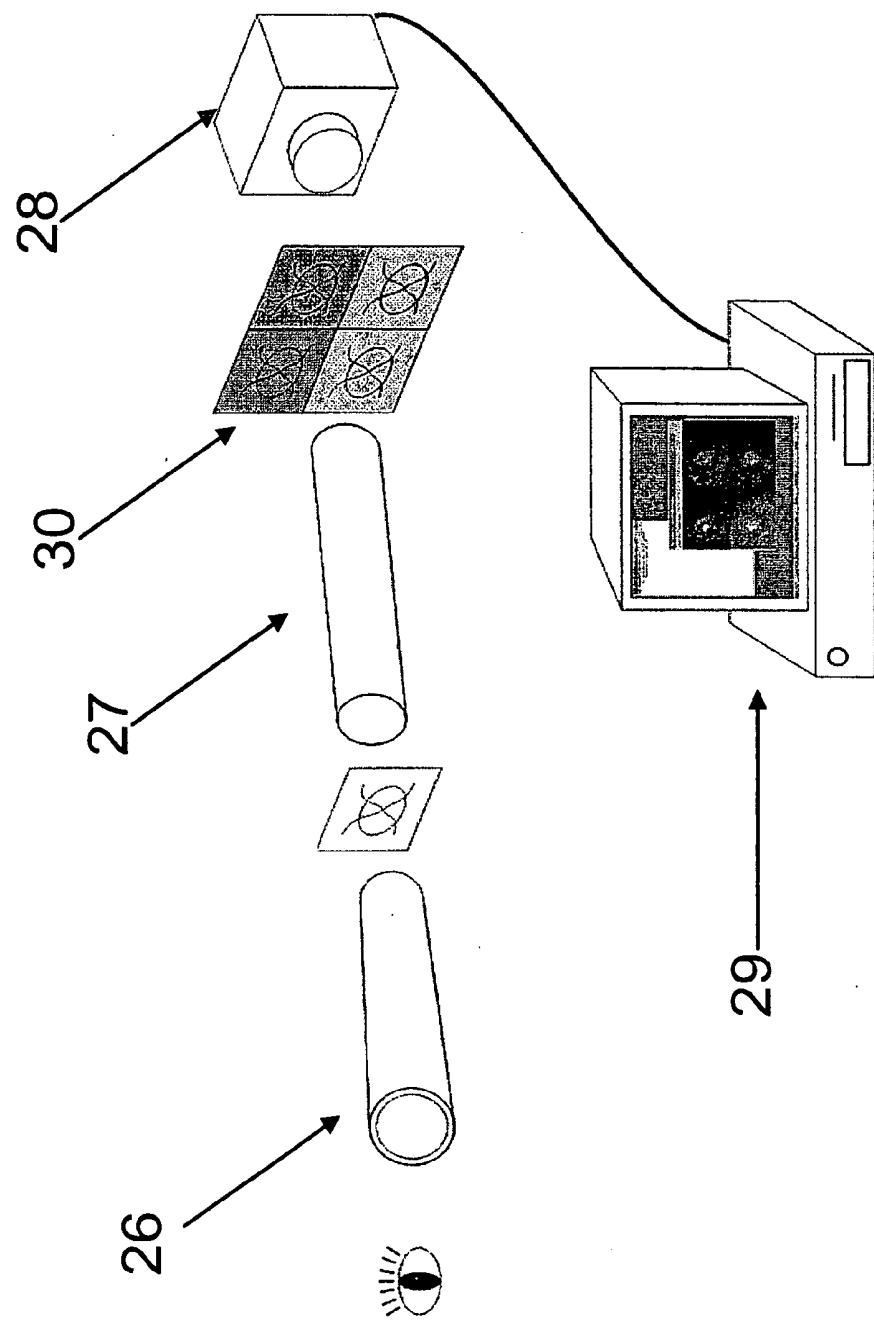


Figure 1

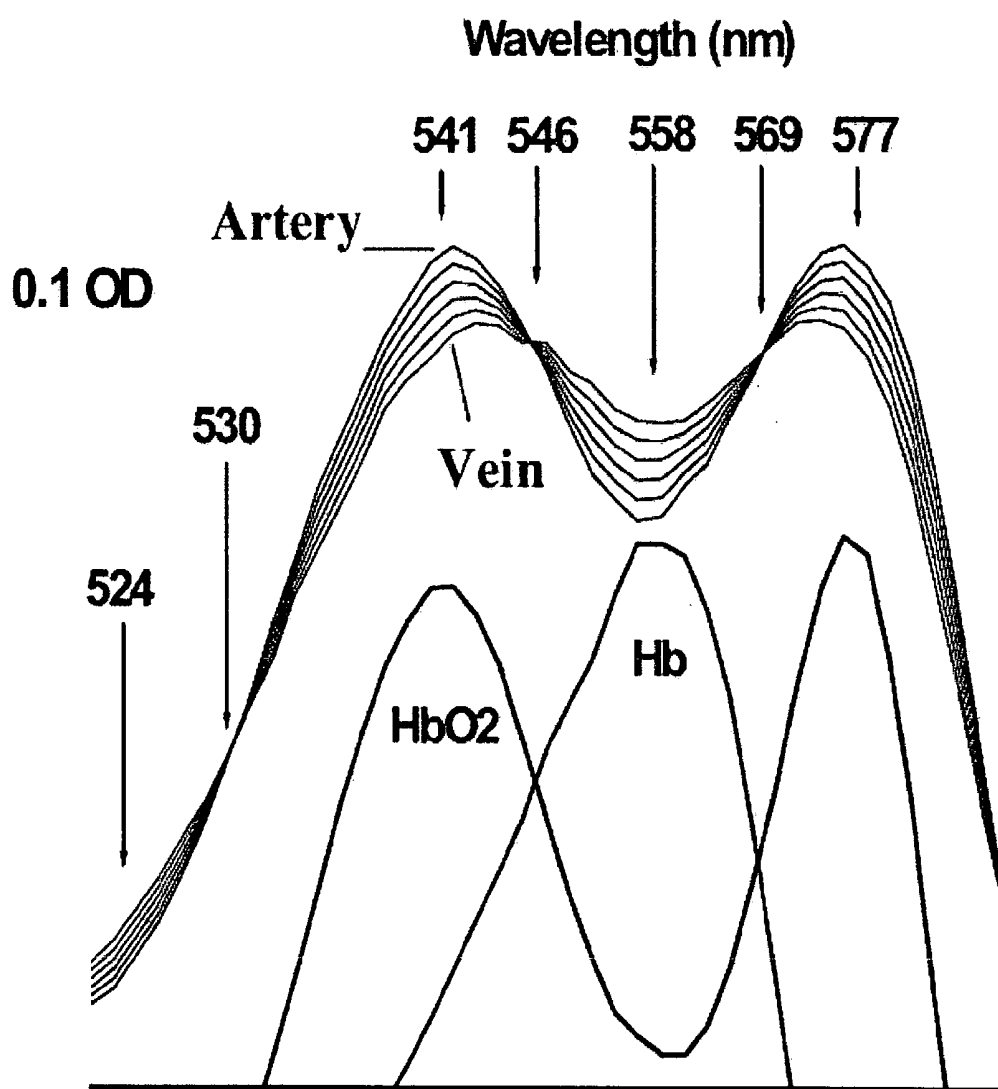
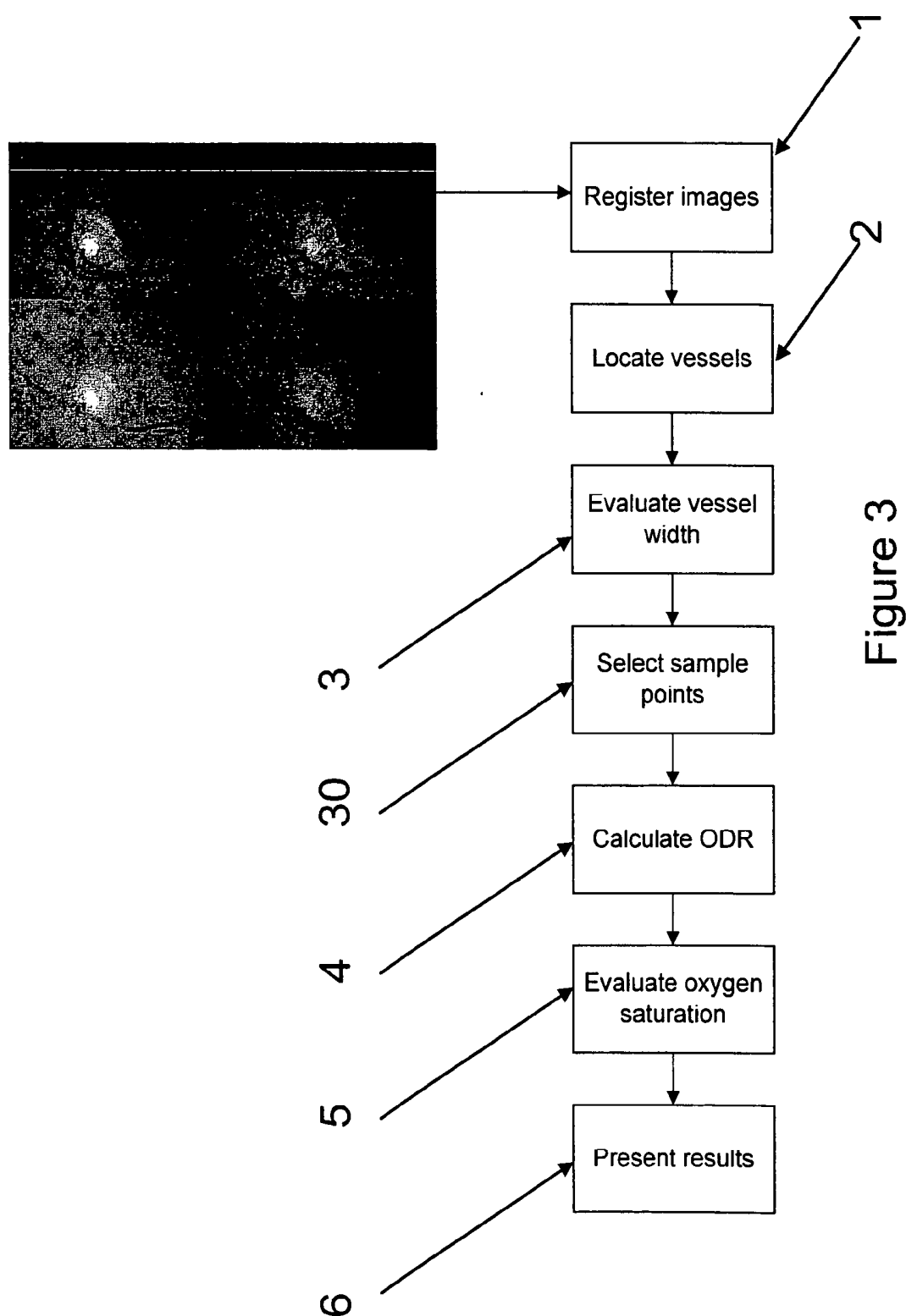
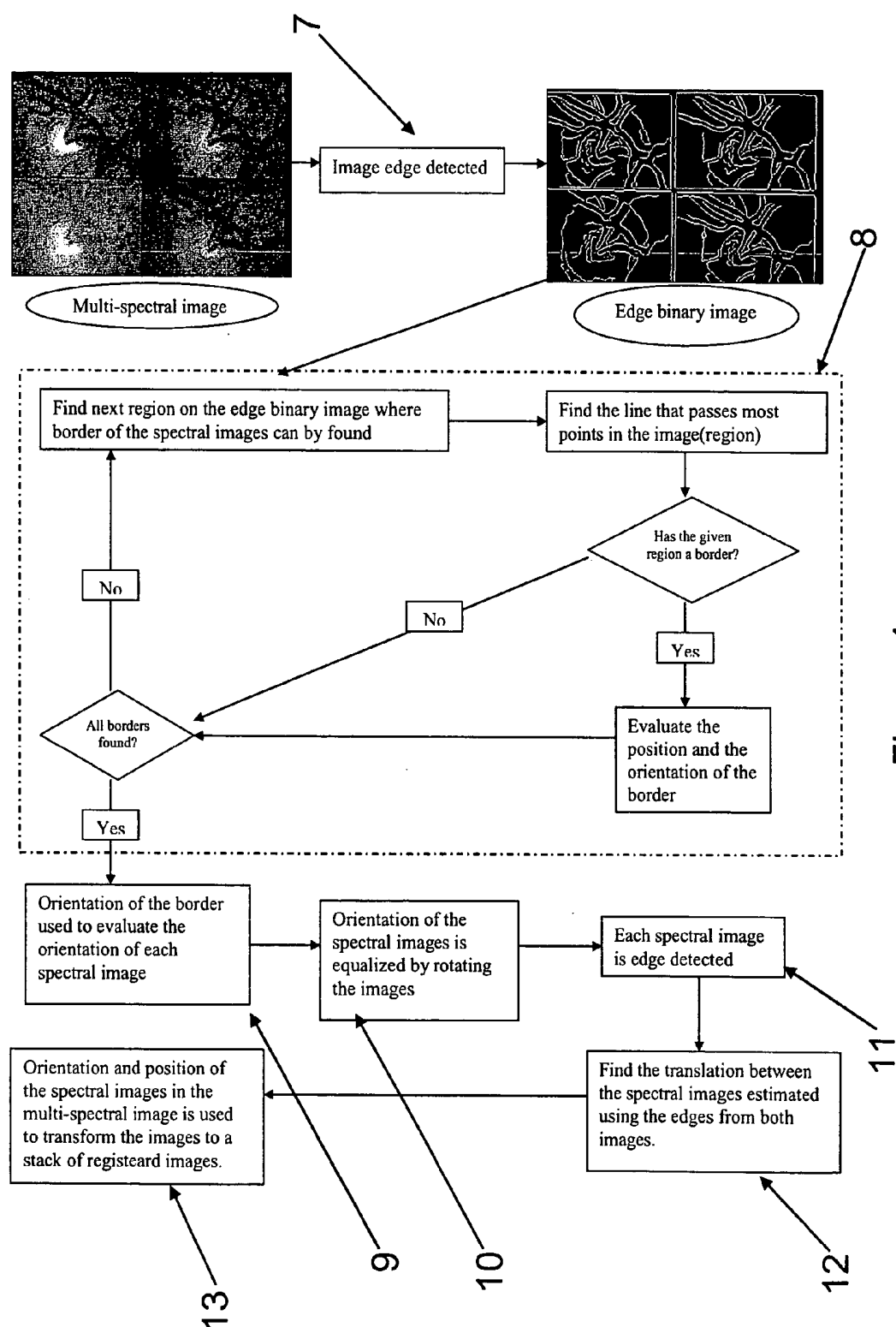


Figure 2





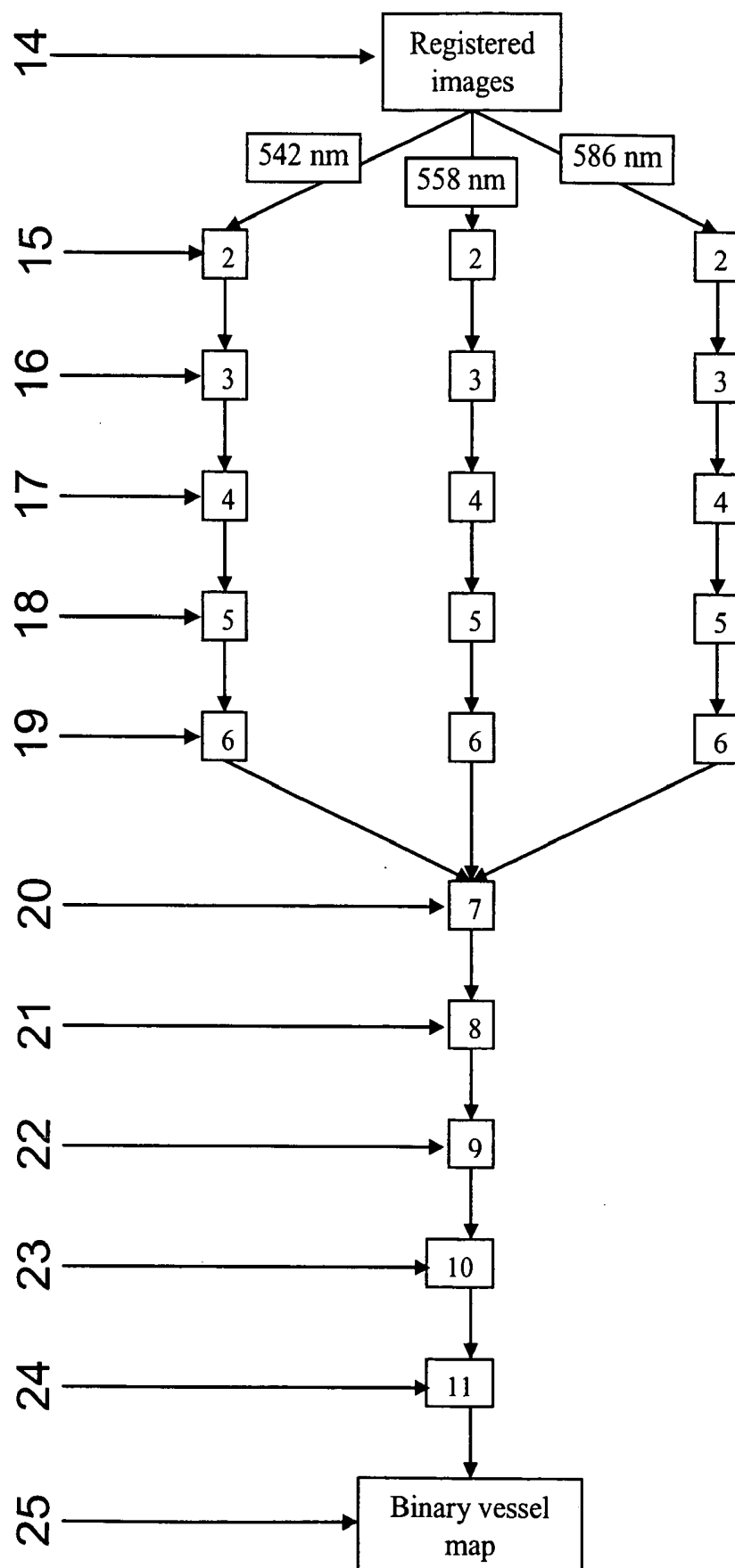


Figure 5

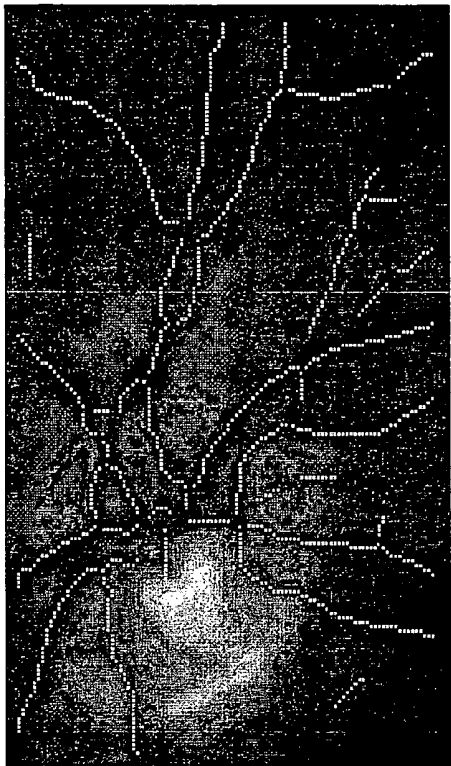


Figure 7

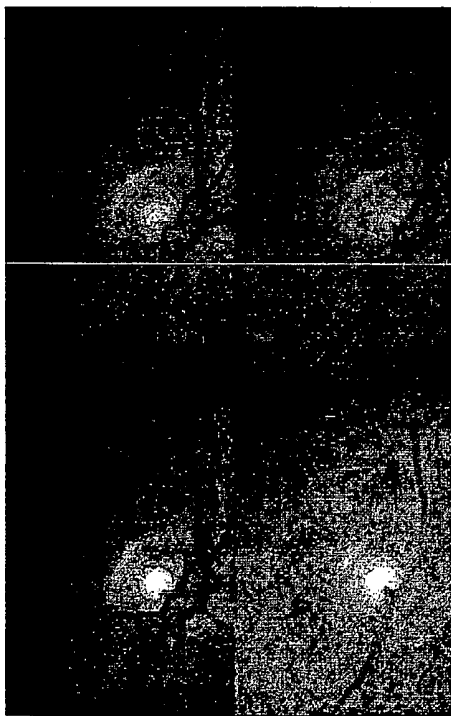


Figure 6

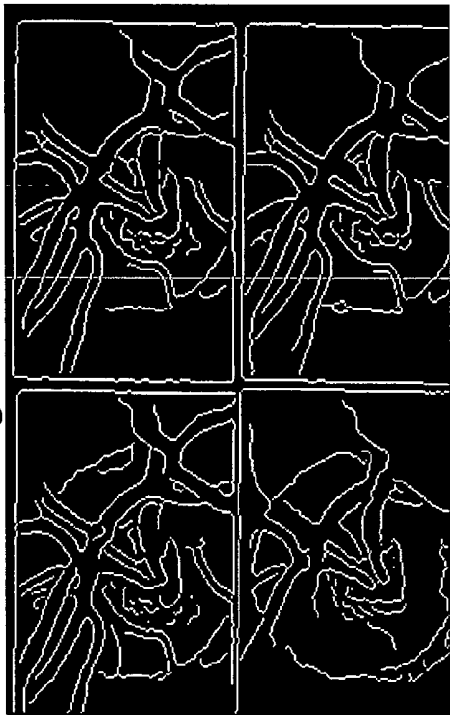


Figure 8

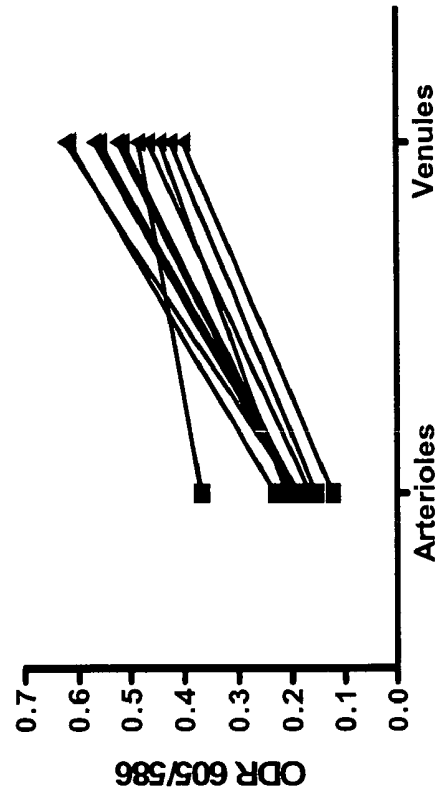


Figure 9

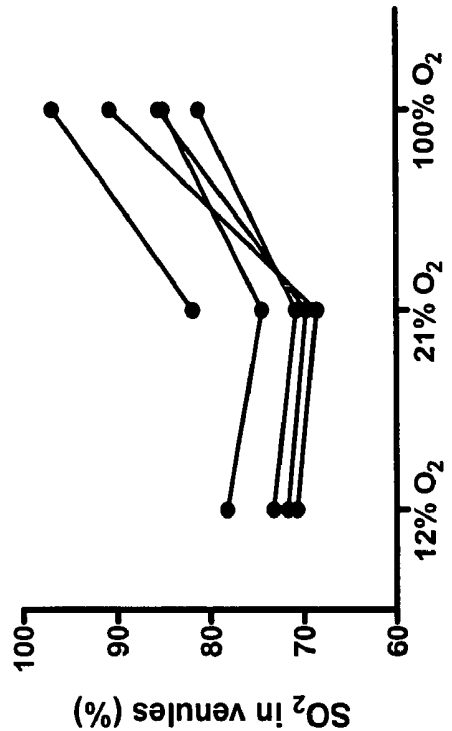


Figure 10

AUTOMATIC REGISTRATION OF IMAGES

TECHNICAL FIELD

[0001] The present invention relates to the field of image processing, more particularly to the automatic registration of multi-spectral images. The present invention, moreover, relates to the application of image processing in the field of ophthalmology for spectral, spatial and temporal analysis.

BACKGROUND ART

[0002] Some of the disease states that pose the greatest danger to vision are thought to involve abnormalities in the oxygen supply to the retina or the oxygen metabolism. These include diseases such as diabetic retinopathy, which is one of the most common causes of blindness in the world and retinopathy of prematurity, retinal vein and artery occlusions. It has also been suggested that ocular blood flow is compromised in glaucoma, including normal tension glaucoma, and that the loss of optic nerve fibers in glaucoma patients may be due to ischemia.

[0003] The principle of spectrophotometric hemoglobin oximetry is old and the method has been developed into clinically useful instrumentation in several specialties of medicine. The finger or earlobe oximeter, commonly used in anesthesiology and intensive care is an example of the successful application of oximetry in medicine.

[0004] Information on retinal and optic nerve oxygenation in health and disease is mostly derived from animal research. Non-invasive measurement of oxygenation in the human retina and optic nerve has proved difficult but considerable progress has been made, particularly in recent years as digital technology has evolved. The potential use of retinal oximetry covers a range of areas, including assessment of oxygen metabolism in disease, the efficacy of treatment of these diseases in restoring or improving the metabolic conditions, either by laser treatment, surgery, lowering of intraocular pressure or by medication. Retinal oximetry may also be of use in elucidating further the physiological processes involved in disease states, such as in glaucoma.

[0005] In the 1990s, an important version of retinal vessel oximetry was developed at the University of Virginia, by James Beach and James Tiedeman [1, 2]. Their version was based on a method proposed by Delori [3] with additional improvements. The main advantage of the Beach and Tiedeman method is the ability to obtain simultaneously two or more images of light reflectance at different wavelengths (called multi-spectral images) from the same fundus using a fundus camera. In this manner it is possible to record reflectance at both oxygen-sensitive and insensitive wavelengths from exactly the same area on the fundus, and at precisely the same time. This allows for precise quantification of the effects of oxygen binding on the light absorption spectra of hemoglobin. Oximetric measurements of this kind are achieved by a system whose main component consists of a modified fundus camera, although the internal xenon flash of the camera is still used as a light source, unmodified. Other components of the system are a beam splitter, a gray scale digital camera, possessing high quality linear performance, and a computer. The digital camera replaces the image acquisition mechanism of the fundus camera. Flashes from the fundus camera are synchronized with recordings by the digital camera electronically. The image thus obtained

with the digital camera is then fed into a computer for further analysis. Between the fundus camera and the digital camera there are image-splitting optics, which can project several separate images at the digital camera detector, as illustrated by FIG. 1.

[0006] In each of these optical pathways, which are separated by a beam splitter and dichroic mirrors, there is a narrow band pass interference filter. Each filter has a 5 nm half-bandwidth and different central (peak) wavelengths of transmission. The filters, or rather their peak wavelengths, are chosen on the basis of the absorption spectra of oxyhemoglobin (HbO_2) as before, and illustrated in FIG. 2. The filter bandwidth clearly affects measurements of the blood Optical Density (OD) and this can be evaluated by calculating the apparent light extinction coefficients of HbO_2 and hemoglobin with no oxygen (Hb) [2]. The extinction coefficients are nearly identical at the isosbestic wavelength of 586 nm, particularly at a narrow bandwidth, while considerably higher for Hb than HbO_2 at the oxygen-sensitive wavelength (600 or 605 nm). Thus, the apparent Optical Density Ratio (ODR) between the oxygen-sensitive and isosbestic wavelengths reflects the oxygen saturation in retinal vessels and tissue in an inverse manner. With higher oxygen saturation the OD at 605 nm is lower.

[0007] OD is a measure of absorption; higher OD means greater absorption [5]. One factor that affects the optical density measured in fundus oximetry is the extravascular pigmentation of the fundus, which can vary greatly between subjects. Pigmentation may cause underestimation of ODs in vessels, in particular at 586 nm, because it causes stronger absorption of light. The effect of retinal pigmentation can be estimated by measuring the ratio of extravascular light reflection near vessels at the oxygen-sensitive and insensitive wavelengths used [2]. The extravascular light reflection ratio (EVR) is calculated as:

$$EVR = \frac{I_{586}}{\eta I_{605}}$$

Where I_{586} is the intensity of 586 nm wave length light reflected, I_{605} is the intensity of 605 nm wave length light reflected, and η is a correction constant for light transmission through the instrument (evaluated separately), from the light source to the digital camera, via the beam splitter and optics. Experimentally it has been found that EVR is greater with higher fundus pigmentation [2]. In order to calculate vessel OD it is therefore necessary to correct for pigmentation by measuring intensity at both wavelengths in non-vascular areas.

[0008] One of the limitations of prior art retinal oximetry [13] includes difficulties in obtaining calibrated measures of actual oxygen saturation. An indirect calibration of the ODR, as it relates to actual oxygen saturation of vascular blood can be obtained by comparing the ODR and systemic O_2 saturation, the latter controlled by having subjects inhale air mixtures of different composition, with 100% oxygen and lower. The reliability of this approach can be further enhanced by measuring the systemic O_2 saturation with a finger pulse oximeter. It has been found experimentally in this manner that there is a fairly good inverse correlation between systemic O_2 saturation and the ODR (lower ODR

with higher systemic O_2) in retinal arterioles [2], and venules [4] in normal human volunteers.

[0009] Clearly, examination of images of the fundus is an important diagnostic tool in ophthalmology. Fundus images allow inspection of arteries and veins that are usually hidden beneath the skin. Consequently, fundus images may be used to diagnose many diseases that affect the vascular structure.

[0010] The problems with the prior art methods used in retinal oximetry is the need for an expert to perform the oximetry, the high variability of the results, and poor reproducibility. The prior art methods are time consuming, and manual registration of the spectral images is inaccurate because of the error prone manual selection of sample points.

DISCLOSURE OF THE INVENTION

[0011] The present invention concerns methods, software, and a system that enables in real time the automatic analysis of spectrophotometric measurements of oxygen saturation in the retinal vasculature. The invention enables any number of spectral images to be examined in the same coordinate system, which means that optical density ratios are more reliably and easily obtained. The method is suitable for measuring large part of the fundus at once. The current invention, moreover, automatically finds vessels in the spectral images and calculates the optical density ratios automatically. This has never been done before and results in more reliable measurements than when measurement points are chosen by hand, and can be done automatically and extended by a novice operator.

Terminology Used in the Present Context:

[0012] Registering, refers to the process of aligning the spectral images into one coordinate system.

[0013] Multi spectral image, refers to multiple images of the same scene acquired at different wavelengths.

[0014] Spectral image, is an image acquired at a specific wavelength.

[0015] Binarize, refers to the process of reducing the resolution of a gray scale image to black and white only.

[0016] Defective pixels, refers to pixels that contain invalid pixel values.

[0017] Edge detection, refers to the process of extracting edges or boundaries between two regions in an image.

[0018] Top-hat transform, refers to image morphological operator first presented by F. Meyer[12].

[0019] Structuring element, refers to a two dimensional matrix of pixel values 0 or 1 describing the neighborhood of an image operator.

[0020] Skeletonizing, refers to the process of generating a stick figure representation of the objects in the image being processed.

[0021] Prune, refers to the process of removing spurious lines in the image.

[0022] Re-Grow, refers to the process of expanding the skeletonized objects to original size.

[0023] Vessel map, refers to a map of the vessels in the image.

[0024] Vessel profile, refers to the side view of the vessels in the image.

[0025] Real-Time, refers to the processing of images on-line without noticeable delay to the human operator.

[0026] Isobestic, is by definition a wavelength which is absorbed equally by an oxygenated and deoxygenated hemoglobin solution.

[0027] In a one aspect the present invention relates to a method for evaluating oxygen saturation of the optic nerve and retina, said method comprising the steps of: registering set of multi-spectral images (1), locating blood vessels (2) in each of the multi-spectral images, evaluating the width of the blood vessels (3), selecting samples for calculating optical density, calculating the optical density ratio (4), where the optical density is the logarithm of the light reflected from background divided by light reflected from vessel and said optical density ratio is the ratio of said optical density at two different wavelengths, evaluating oxygen saturation level (5), and presenting the results (6), wherein presenting the results may include presenting numerical and visual representation of the oxygen saturation.

[0028] In another aspect the present invention relates to a method for automatic registration of multi-spectral images, said method comprises the steps of: binarize the multi-spectral image (7), find the all the border regions of each image, by, finding the region including the straight line that passes the most number of points in the region (8), use the orientation of the borders to evaluate the orientation of each spectral image (9), equalize the orientation of each spectral image by rotating the spectral image (10), edge detect each spectral image (11), estimate the translation between the spectral images based on the edges of adjacent images (12), transform the images to a stack of registered images (13), wherein the images are transformed based on orientation and position of each spectral image.

[0029] In another aspect the current invention relates to a method for detecting blood vessels in multi-spectral images, said method comprises the steps of: retrieving registered spectral images (14), for each spectral image, remove defective pixels (15), enhance image contrast (16), perform top-hat transform using structuring element larger than the largest vessel diameter (17), binarize each image (18), apply filter for smoothing the image (19), combine all the spectral images using binary AND operator (20), skeletonize the resulting image (21), prune the skeleton image (22), re-grow the pruned image (23), locate junction points (24), wherein the junction points are the intersection of two or more vessels.

[0030] In another aspect the current invention relates to a method for evaluating the width of the blood vessels, said method comprises the steps of: evaluating binary vessel map by calculating the normal vector to the vessel direction to, estimate the direction and position of the vessel profile, evaluate positions of vessel walls based on the vessel profile obtained, evaluate vessel width based on the position of the vessel walls, wherein the width is the distance between the vessel walls.

[0031] In another aspect the current invention relates to a method for selecting sample points for optical density calculation, said method comprising the steps of: obtaining a binary vessel map, obtaining vessel profile, locating the points of the vessel profile which have the lowest magnitude, wherein the selected points will provide samples with the least error contributed from the specular reflection of the vessel walls.

[0032] In another aspect the present invention relates to computer software realizing a method for evaluating oxygen saturation of the optic nerve and retina.

[0033] In another aspect the present invention relates to computer software realizing a method for automatic registration of multi-spectral images.

[0034] In another aspect the present invention relates to computer software realizing a method for detecting blood vessels in multi-spectral images.

[0035] In another aspect the present invention relates to computer software realizing a method for evaluating the width of blood vessels in multi-spectral images.

[0036] In another aspect the present invention relates to computer software realizing a method for selecting sample points for optical density calculation in multi-spectral images.

[0037] In another aspect the present invention relates to a system for automatically evaluating oxygen metabolism of the optic nerve and retina, said system comprising: image capturing system further comprising: a fundus camera (26), a four wavelengths beam splitter (27), a digital image capturing device (28), a computer system, image processing software performing in real-time the steps of: registering set of multi-spectral images (1) by, binarizing the multi-spectral image (7), find the all the border regions of each image by finding the region including the straight line that passes the most number of points in the region (8), use the orientation of the borders to evaluate the orientation of each spectral image (9), equalize the orientation of each spectral image by rotating the spectral image (10), edge detect each spectral image (11), estimate the translation between the spectral images based on the edges of adjacent images (12), transform the images to a stack of registered images (13), locating blood vessels (2) in each of the multi-spectral images by, retrieving registered spectral images (14), for each spectral image, remove defective pixels (15), enhance image contrast (16), perform top-hat transform using structuring element larger than the largest vessel diameter (17), binarize each image (18), apply filter for smoothing the image (19), combine all the spectral images using binary AND operator (20), skeletonize the resulting image (21), prune the skeleton image (22), re-grow the pruned image (23), locate junction points (24), evaluating the width of the blood vessels (3) by calculating the normal vector to the vessel direction to, estimate the direction and position of the vessel profile, evaluate positions of vessel walls based on the vessel profile obtained, evaluate vessel width based on the position of the vessel walls, selecting samples for calculating optical density, calculating the optical density ratio (4), evaluating oxygen saturation level (5), and presenting the results (6), wherein presenting the results may include presenting numerical and visual representation of the oxygen saturation.

BRIEF DESCRIPTION OF DRAWINGS

[0038] FIG. 1 shows multi-spectral image capturing system.

[0039] FIG. 2 shows the absorption spectra of the oxyhemoglobin.

[0040] FIG. 3 shows a functional flow graph of the retinal oxymetry process.

[0041] FIG. 4 shows a functional flow graph of the image registration process.

[0042] FIG. 5 shows a functional flow graph of the vessel location process.

[0043] FIG. 6 shows a typical multi-spectral image.

[0044] FIG. 7 shows results from the automatic vessel selection.

[0045] FIG. 8 shows binary edge detected image.

[0046] FIG. 9 shows results for arterioles and venules measured at 21% O₂. Each line represents one eye. Paired t-test, arterioles vs. venules: p<0.0001

[0047] FIG. 10 shows results for venules measured at different O₂ concentration in inhaled air. Paired t-test, 21% vs. 100%: p=0.0023.

BEST MODE FOR CARRYING OUT THE INVENTION

[0048] The retinal oxymetry process utilizes the fact that the amount of light absorbed by hemoglobin is dependent on the wavelength of the light and the oxygen saturation of the blood. To estimate the amount of light reflected, multi-spectral images containing sub-images at different wavelengths are acquired.

[0049] Therefore, the image from the digital camera is a multi-spectral image consisting of 2 or more sub-images. Each sub-image has been filtered with an optical narrow band-pass filter (with center wavelengths 542 nm, 558 nm, 586 nm, and 605 nm). The sub-images, here after referred to as spectral images, represent the same area of the retina. However, an accurate spectral image position and rotation on the multi-spectral image is not known because of the sensitivity of the beam splitter. For this reason, the spectral images may overlap and sometimes parts of the spectral images are outside of the multi-spectral image and therefore parts of the spectral image are missing.

[0050] To enable analysis of the retinal oximetry the spectral images have to go through a preprocessing stage were they are pair-wise registered together. The registration process is necessary in order to be able to compute the information extracted from the images in the same coordinate system. The registration algorithm is implemented in the software. The algorithm has been shown to be accurate (mean registration error between two images ≤ 1 pixel) and computationally inexpensive [7]

Software Specifications

[0051] FIG. 3 is a flow diagram that illustrates the sequence of events performed by the retinal oximetry software for processing the multi-spectral images.

[0052] The first task is preprocessing and registration of the spectral images and includes the following steps as shown in **FIG. 4**:

Image Edge Detected (7)

[0053] Edge detection is used to extract edges or boundaries between two regions in the image. Common edge detection methods such as Canny or Sobel are used. Mathematically the edges are the gray level gradient of the image and edge detection methods often involve defining the optimum gradient operators. Binary edge detected image, **FIG. 8**, is derived by thresholding the gradient of the image using for the Canny method.

Evaluate a Border Region (8)

[0054] The multi-spectral image, **FIG. 6**, contains four spectral images that are always ordered the same way and their position does not change dramatically. The mechanical sensitivity of the device to capture the images does only produce uncertainty in the image position (and rotation), which is no more than ± 20 pixels (concluded after examining large amount of data). Therefore, the area on the multi-spectral image, for finding the border of each spectral image, is evaluated as percent of the multi-spectral image size.

[0055] Find the line that passes most points in the image

[0056] The line that passes the most points is found by using the Hough transform. The line is located by transforming the region of the image, and finding the position of the maximum value in the Hough space.

[0057] Has the given region a border?

[0058] The only problem with the uncertainty in the position of the borders is that the spectral images can overlap and borders can be out side the multi-spectral image.

[0059] To evaluate if the region has a border a following methodology is used:

[0060] If the magnitude of the maximum value of the Hough transform is P % of the line length, the line is assumed to be a border.

[0061] To discriminate between two borders that lie close together on the binary edge image the edge gradient direction is used.

Find the Orientation of Each Spectral Image (9)

[0062] The orientation of the image can be estimated without using the similarity metric, since each spectral image has at least one visible border on the multi-spectral image. The Hough transformation is used to find how many points lie on lines that have orientation in a particular range. Therefore, the boarder orientation of the spectral images' is estimated by choosing the line that intersects the maximum number of points. Finally, the orientation of each spectral image is calculated using a weighted average of the border orientation, weighting each possible border with how many points the border line intersects.

Equalize the Orientation of the Spectral Images (10)

[0063] Position of each border, detected previously is used to cut large region around the spectral image out of the

multi-spectral image. Then, the orientation of the spectral images is equalized by rotating each image.

Find the Translation between the Spectral Images (12)

[0064] After the images have been rotated for orientation equalization, the translation between them is evaluated. The translation between the spectral images is estimated using the edges from both images. Then, by searching over all translation $((tx,ty)\pm 20)$, the most likely translation is the one which has the most edges with similar gradient direction and magnitude.

The Second Task is to Detect and Locate the Vessels:

[0065] First the registered spectral images are retrieved (14). The images are searched for defective pixels and these pixels are then removed from each of the spectral images (15). Next, each of the spectral images goes through a contrast enhancement process (16) to improve the contrast between the background and the features (vessels) of the images. Thereafter, the images are processed using the Black-Top-Hat transform using a structuring element in the shape of a disk with diameter larger than the diameter of the largest vessel (17). Next, each image is thresholded at for example 80% of the mean pixel value (18). After the thresholding procedure the images a median filter is applied to each of the images to smoothen them (19). After processing each image separately the resulting binarized images are combined by applying a binary AND operation on each pixel in the images (20). The resulting image is then skeletonised to find the centerline of the vessels (21). The skeletonized image is pruned (22), and the resulting image is then re-grown (23) to remove the spurs from the skeletonized image. Finally the junction points (24) of the vessels are located and binary vessel map is produced (25), **FIG. 8**.

[0066] The third task is to evaluate the vessel width and segment length. The binary vessel map is used to evaluate the direction and position of the vessel profile which is a perpendicular normal to the vessel direction. The profile is then used to evaluate accurate position of vessel walls (edges) and calculate the vessel width.

[0067] The fourth task is to select the sample points to use for the calculation of the optical density ratio. The points are selected both inside a vessel, to estimate the absorption of the blood and outside the vessels to estimate the amount of light entering the vessel. The points within a vessel are selected by locating the darkest spot within the vessel profile previously obtained and then average the value of the closest eight neighboring pixels within the vessel. The points outside the vessels are selected by moving away from the vessels in a perpendicular direction from the points selected within the vessel until a predetermined distance is reached for example 20 pixels. Again, the pixel values in its 8-connected neighborhood are averaged.

[0068] The fifth task is to evaluate the optical density. This is accomplished using following equation were the Light reflected by the background is the average pixel value from background and the Light reflected from vessel is the average pixel value within the selected vessel as described in the previous paragraph.

$$OD = \log \left(\frac{\text{Light reflected by background}}{\text{Light reflected from vessel}} \right)$$

[0069] OD is then calculated at each wavelength (OD_{542 nm}, OD_{558 nm}, OD_{586 nm} and OD_{605 nm}), and finally the optical density ratio at two different wavelengths is calculated

$$ODR = \frac{OD_X}{OD_Y}$$

[0070] The sixth task is to evaluate the oxygen saturation of the blood. As previously described, by selecting appropriate wavelengths when calculating this ratio (Y=586 nm and X={542,558,605}nm), a linear relationship exists between ODR and the oxygen saturation in the retinal vasculature, which in turn makes it possible to estimate the oxygen saturation.

[0071] The final step is to present the results. After the software has calculated OD values for each wavelength the optical density ratio (ODR) is calculated using one isobestic wavelength and one non-isobestic wavelength. The ODR information for any venule or arteriole is represented as numerical and visual information on the computer screen and stored in a data file for further investigation.

Vessel Classification

[0072] In order to measure automatically the difference between the oxygen content of the blood flowing to the retina of the eye and the blood going from it, the ability to classify the detected vessels as either arterioles or venules is quite important. In our software, a positive definite fuzzy classifier [9] is applied to the data collected by the vessel location method. Each pixel belonging to a vessel, as determined by the vessel classification algorithm, is classified as either a pixel belonging to a venule or an arteriole. In order to adjust for variability in illumination between images the input variables are scaled using the mean pixel value of its respective spectral band. The vessel itself is then classified according to which class the majority of the pixels within it are determined to belong to (see results in FIG. 4).

Experimental Results:

[0073] The software has been used to examine reproducibility and sensitivity of ODRs for different oxygen tensions. The results are given below.

Reproducibility:

[0074] Reproducibility was evaluated by measuring the same vessel on five consecutive images. Coefficient of variation (CV) was calculated as

$$CV = \frac{\text{Standard deviation of ODR (5 images of the same vessel)}}{\text{Mean ODR (5 images of the same vessel)}} \left(1 + \frac{1}{4n} \right)$$

[0075] where n=5 measurements (images). One arteriole and one venule were measured in images from 11 eyes. The

CV was calculated for measurements with both the automatic version of the software and with an older version of the software, which allows only manual placement of measurement points. The results are shown in Table 1.

TABLE 1

CV for automatic and manual software				
	Arterioles		Venules	
	Manual	Automatic	Manual	Automatic
CV (mean, range)	48% (15–139%)	16% (8–26%)	19% (5–81%)	8% (3–16%)

[0076] As seen in Table 1, the automatic approach to point selection clearly increases the reproducibility.

Sensitivity:

[0077] Sensitivity of the system was evaluated by comparing ODRs in arterioles and venules and by measuring changes in ODRs (or venular oxygen saturation) when healthy volunteers inhaled different concentration of oxygen.

[0078] FIG. 9 shows ODRs for arterioles and venules (subjects inhaling room air).

[0079] FIG. 9. Arterioles and venules measured at 21% O₂. Each line represents one eye. Paired t-test, arterioles vs. venules: p<0.0001.

[0080] FIG. 9 shows that the system is sensitive to the arterio-venous difference in ODRs (and therefore to the a.-v. difference in oxygenation). Furthermore, the size of the arterio-venous difference is similar between most individuals although the offset (arteriolar ODR) is variable.

[0081] Since oxygen tension in the retinal arterioles can be assumed to be similar to oxygen tension in arterial blood elsewhere in the body, calibration with a commercially available finger pulse oximeter is possible. This calibration would almost eliminate the intersubject variability in the arterioles in FIG. 9. Assuming a linear relationship between ODR and hemoglobin oxygen saturation (S_{O₂}) and using finger oximetry values for retinal arterioles we found that ΔS_{O₂}=−89ΔODR (preliminary result, 95% confidence interval for the slope is −125 to −53). This calibration was done with data from experiments where healthy volunteers inhaled either 12%, 21% or 100% oxygen to induce systemic hypoxia, normoxia and hyperoxia, respectively.

[0082] FIG. 10 shows the results from the hypoxia-normoxia-hyperoxia experiments for venules. One point from the hypoxia part of the experiment was omitted because the subject was not rendered hypoxic according to the finger pulse oximeter.

[0083] FIG. 10. Venules measured at different O₂ concentration in inhaled air. Paired t-test, 21% vs. 100%: p=0.0023.

[0084] The reason for the lack of difference between results for 12% and 21% inhaled oxygen concentration is most likely the well-known phenomenon of blood flow autoregulation [14]. When the oxygen concentration in arteriolar blood decreases the arterioles dilate, blood flow increases and similar amount of oxygen is delivered to the tissue. Therefore, there is little or no change in venular

oxygen tension. On the other hand, the system detects a significant difference in the venules when inhaled oxygen tension is changed from 21% to 100% (see FIG. 10). This is probably due to the effects of the choroid, which is a non-autoregulated vascular bed underneath the retina. ODRs are measured in the visible vasculature which nourishes the inner part of the retina. The choroid does not usually deliver oxygen to the inner part of the retina [14]. However, when 100% oxygen is inhaled, some of the oxygen will diffuse to the inner retina and this probably overwhelms the autoregulation of blood flow in the inner vasculature.

[0085] In summary, the results show that the system can detect expected differences in venular oxygenation when oxygen concentration in inhaled air is altered. Autoregulation of blood flow most likely explains part of the results.

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1. A method for evaluating oxygen saturation of the optic nerve and retina, said method comprising the steps of:
 - registering a set of multi-spectral images of a retina,
 - locating blood vessels in each of the multi-spectral images,
 - evaluating the width of the blood vessels,
 - selecting sample points for calculating optical density,
 - calculating the optical density ratio, where the optical density is the logarithm of the light reflected from background divided by light reflected from vessel and said optical density ratio is the ratio of said optical density at two different wavelengths,
 - evaluating oxygen saturation level, and
 - presenting results indicating oxygen saturation of the optic nerve and retina.
 2. The method of claim 1 wherein said presenting of results includes presenting numerical and/or visual representation of the oxygen saturation.
 3. The method of claim 1, wherein said registering a set of multi-spectral images of a retina comprises:
 - finding border regions of each image,
 - equalizing the orientation of each spectral image by rotating the spectral image,
 - edge-detecting each spectral image,
 - estimating the translation between the spectral images based on the edges of adjacent images,
 - transforming the images to a stack of registered images, wherein the images are transformed based on orientation and position of each spectral image.
 4. The method of claim 3, wherein said finding borders regions of each image comprises
 - binarizing the multi-spectral image,
 - finding the region including the straight line that passes the most number of bright points in the region, and
 - using the orientation of the borders to evaluate the orientation of each spectral image.
 5. The method of claim 1 wherein said step of locating blood vessels comprises
 - retrieving registered spectral images,
 - for each spectral image,
 - removing defective pixels,
 - enhancing image contrast,

performing top-hat transform using structuring element larger than the largest vessel diameter,

binarizing each image,

applying spatial filter for smoothing the image,

combining all the spectral images using binary AND operator,

skeletonizing the resulting image,

pruning the skeleton image,

re-growing the pruned image,

locating junction points,

wherein the junction points are the intersection of two or more vessels.

6. The method of claim 1 wherein said step of evaluating the width of the blood vessels comprises

evaluating binary vessel map by calculating the normal vector to the vessel direction to estimate the direction and position of the vessel profile,

evaluating positions of vessel walls based on the vessel profile obtained,

evaluating vessel width based on the position of the vessel walls,

wherein the width is the distance between the vessel walls.

7. The method of claim 1 wherein said step of selecting sample points for calculating optical density comprises selecting sample points within a vessel and outside a vessel, by:

locating a spot within a vessel profile previously obtained, with the lowest magnitude,

averaging the value of a predetermined number of symmetrically neighboring pixels,

moving away from a vessel in a predetermined distance and, with the provision that the location not being inside another vessel, averaging the value of a predetermined number of symmetrically neighboring pixels.

8. A method for automatic registration of multi-spectral images, said method comprising the steps of:

finding all border regions of each image by,

binarizing the multi-spectral image,

finding the region including the straight line that passes the most number of bright points in the region,

using the orientation of the borders to evaluate the orientation of each spectral image,

equalizing the orientation of each spectral image by rotating the spectral image,

edge detect each spectral image,

estimating the translation between the spectral images based on the edges of adjacent images,

transforming the images to a stack of registered images, wherein the images are transformed based on orientation and position of each spectral image.

9. A method for detecting blood vessels in multi-spectral images, said method comprising the steps of:

retrieving registered spectral images,

for each spectral image,

removing defective pixels,

enhancing image contrast,

performing top-hat transform using structuring element larger than the largest vessel diameter,

binarizing each image,

applying spatial filter for smoothing the image,

combining all the spectral images using binary AND operator,

skeletonizing the resulting image,

pruning the skeleton image,

re-growing the pruned image,

locating junction points,

wherein the junction points are the intersection of two or more vessels.

10. A method for evaluating the width of the blood vessels, said method comprising the steps of:

evaluating binary vessel map by calculating the normal vector to the vessel direction to estimating the direction and position of the vessel profile,

evaluating positions of vessel walls based on the vessel profile obtained,

evaluating vessel width based on the position of the vessel walls,

wherein the width is the distance between the vessel walls.

11. A method for selecting sample points for optical density calculation, said method comprising the steps of:

obtaining a binary vessel map,

obtaining vessel profile,

locating the points of the vessel profile which have the lowest magnitude,

wherein the selected points will provide samples with the least error contributed from the specular reflection of the vessel walls.

12. A computer software realizing the method of claim 1.

13. A computer software realizing the method of claim 8.

14. A computer software realizing the method of claim 9.

15. A computer software realizing the method of claim 10.

16. A computer software realizing the method of claim 11.

17. A system for automatically evaluating oxygen saturation of the optic nerve and retina, said system comprising:

an image capturing system further comprising:

a fundus camera,

a four wavelengths beam splitter,

a digital image capturing device,

a computer system,

image processing software performing in real-time the steps of:

- registering set of multi-spectral images,
- locating blood vessels in each of the multi-spectral images,
- evaluating the width of said blood vessels,
- selecting sample points for calculating optical density,
- calculating the optical density ratio,
- evaluating oxygen saturation level, and
- presenting results indicating oxygen saturation of the optic nerve and retina.

18. The system of claim 17 wherein said presenting results indicating oxygen saturation includes presenting numerical and/or visual representation of the oxygen saturation.

19. The system of claim 17 wherein said registering a set of multi-spectral images comprises:

- binarizing the multi-spectral image,
- finding border regions of each image by finding the region including the straight line that passes the most number of points in the region,
- using the orientation of the borders to evaluate the orientation of each spectral image,
- equalizing the orientation of each spectral image by rotating the spectral image,
- edge-detecting each spectral image,
- estimating the translation between the spectral images based on the edges of adjacent images,
- transforming the images to a stack of registered images, wherein the images are transformed based on orientation and position of each spectral image.

20. The system of claim 17, wherein said locating blood vessels in each of the multi-spectral images comprises

- retrieving registered spectral images,
- for each spectral image,
 - removing defective pixels,
 - enhancing image contrast,
 - performing top-hat transform using structuring element larger than the largest vessel diameter,
 - binarizing each image,
 - applying filter for smoothing the image,
- combining all the spectral images using binary AND operator,
- skeletonizing the resulting image,
- pruning the skeleton image,
- re-growing the pruned image,
- locating junction points,
- wherein the junction points are the intersection of two or more vessels.

21. The system of claim 17, wherein said evaluating the width of said blood vessels comprises:

- estimating the direction and position of the vessel profile,
- evaluating positions of vessel walls based on the vessel profile obtained,
- evaluating vessel width based on the position of the vessel walls,
- wherein the width is the distance between the vessel walls.

* * * * *

专利名称(译)	自动注册图像		
公开(公告)号	US20060276698A1	公开(公告)日	2006-12-07
申请号	US11/146278	申请日	2005-06-07
[标]申请(专利权)人(译)	HALLDORSSON 吉斯利 ^ h 斯蒂芬森埃纳尔 本尼迪克特森 JON 一个 卡尔松罗伯特 HARDARSON SVEINN ^ h EYSTEINSSON THOR 海滩 JAMES M		
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摘要(译)

本发明涉及一种自动评估视神经和视网膜氧饱和度的系统，所述系统包括：图像捕获系统，还包括：眼底照相机（26），四波长分束器（27），数字图像捕获装置（28），计算机系统，图像处理软件实时执行以下步骤：通过二值化多光谱图像（7）来记录多光谱图像（1）的集合，找到每个的所有边界区域通过找到包括通过区域（8）中最多点的直线的区域的图像，使用边界的方向来评估每个光谱图像（9）的方向，通过旋转来均衡每个光谱图像的方向光谱图像（10），边缘检测每个光谱图像（11），基于相邻图像（12）的边缘估计光谱图像之间的平移，将图像变换为一叠注册图像（13），定位血液ssels（2）在每个多光谱图像中，通过检索登记的光谱图像（14），对于每个光谱图像，去除缺陷像素（15），增强图像对比度（16），使用大于的结构元素执行顶帽变换最大血管直径（17），二值化每个图像（18），应用滤波器平滑图像（19），使用二进制AND运算符（20）组合所有光谱图像，

对结果图像进行骨架化（21），修剪骨架图像（22），重新生长修剪后的图像（23），定位连接点（24），通过计算血管方向的法向量来评估血管宽度（3），估计血管的方向和位置轮廓，根据获得的血管轮廓评估血管壁的位置，根据血管壁的位置评估血管宽度，选择用于计算光密度的样本，计算光密度比（4），评估氧饱和度（5），并展示结果（6），其中呈现结果可以包括呈现氧饱和度的数字和视觉表示。

