

METHOD AND SYSTEM FOR COLLECTING FLUORESCENCE SPECTRAL DATA BY ULTRAVIOLET LIGHT, AND MEDIUM

TECHNICAL FIELD

5 The present invention relates to the field of optical data collection, and in particular to a method and system for collecting fluorescence spectral data by ultraviolet light, and a medium.

BACKGROUND

10 Optical testing technologies are classified into: absorption spectrometry, fluorescence spectroscopy, and the like. A fluorescence analysis method means a method for qualitative or quantitative analysis of fluorescence that can reflect characteristics of some substances and that is produced during a process in which the substances are in an excited state after being irradiated with ultraviolet light,
15 and molecules in an excited state undergo a de-excitation process of collision and emission. The fluorescence analysis method has advantages of high sensitivity, strong selectivity, a few samples demanded, simplicity, and the like, but different analytes excite different pieces of spectral data when irradiated by ultraviolet light. To obtain effective spectral data, peaks of spectral data are required to fall into a
20 recognizable range of an imaging spectral detection device. How to keep the peaks of the spectral data of different analytes within the recognizable range of the imaging spectral detection device is a problem to be solved urgently.

 A fluorescence radiation signal excited in the fluorescence method has a wide wavelength range, and a peak value of a spectral curve of the fluorescence radiation
25 signal may be beyond the recognizable range of the spectral detection apparatus, resulting in an inaccurate analysis result.

 A patent document CN118078277A discloses a non-invasive blood glucose

testing method based on hyperspectral data analysis, which implements non-invasive testing. The absorption spectroscopy is used in this document, and collected and analyzed spectral signals include not only a spectral signal of blood glucose, but also spectral signals of skin tissue and other components. Different
5 analytes excite different pieces of spectral data when irradiated by ultraviolet light. To obtain effective spectral data, peaks of spectral data are required to fall into a recognizable range of an imaging spectral detection device. The present invention does not address how the peaks of spectral data fall within the recognizable range of the imaging spectral detection device.

10 Patent document CN112842257B discloses a method and an apparatus for positioning blood vessels, including: obtaining a first infrared blood vessel image and a second infrared blood vessel image of at least two frames of continuously collected infrared blood vessel images, where the second infrared blood vessel image is collected at a moment later than a moment at which the first infrared blood
15 vessel image is collected; determining a target blood vessel in the first infrared blood vessel image; and based on the target blood vessel, using a pyramid optical flow tracing method to determine a position of the target blood vessel in the second infrared blood vessel image. The target blood vessel is determined from the first infrared blood vessel image, and then the position of the target blood vessel is
20 positioned from the second infrared image in the pyramid optical flow tracking method. Because the pyramid optical flow tracking method can adapt to large inter-frame variation, the position of the target blood vessel in the second infrared blood vessel image can be accurately determined in the pyramid optical flow tracking method based on the target blood vessel, thereby ensuring that the target
25 blood vessel is accurately positioned. However, different analytes excite different pieces of spectral data when irradiated by ultraviolet light. To obtain effective spectral data, peaks of spectral data are required to fall into a recognizable range of an imaging spectral detection device. Patent document CN117503123A discloses a multi-wavelength near-infrared non-invasive blood glucose testing system and

method, which uses a plurality of sensors and modules to simultaneously obtain a plurality of biological signals such as fingertip infrared information, facial infrared information, and forehead temperature information, and integrates a plurality of pieces of information for analysis and determining. This method has the following disadvantages: excessively many biological signals need to be collected at different positions, blood glucose concentrations in different parts of a body are also different, the costs are high, and it is impossible to achieve portable real-time testing effects, and excessively information causes a testing algorithm to be complicated.

Therefore, a new technical solution needs to be proposed to improve the above technical problems.

SUMMARY

To overcome defects in the prior art, the present invention is intended to provide a method and a system for collecting fluorescence spectral data by ultraviolet light, and a medium.

The method for collecting fluorescence spectral data by ultraviolet light according to the present invention includes:

ultraviolet light irradiation step: irradiating a first area by the ultraviolet light with a wavelength of 300–390 nm;

signal collection step: obtaining a fluorescence radiation signal that includes the fluorescence spectral data and that is emitted by the first area when excited, and performing imaging based on the fluorescence radiation signal, to obtain an image; and

spectral data collection step: selecting a data collection point from the image, substituting a grayscale value of the data collection point into a spectrum reconstruction algorithm, and obtaining the fluorescence spectral data by calculation, where a peak value of a spectral line of the fluorescence spectral data is

within 400–800 nm.

Preferably, the image is collected under irradiation of the ultraviolet light within a preset wavelength range; and

the image includes spectral data that indicate, in an imaging area, a reflection
5 signal or an excitation signal generated by the first area when irradiated by light.

Preferably, the signal collection includes:

based on grayscale values of pixels in the image, selecting one pixel whose
grayscale value meets preset requirements from a testing point candidate area as a
testing point or a combination of the one pixel and an adjacent pixel thereof as the
10 testing point, and selecting, from a reference point candidate area, another pixel
whose grayscale value is within a preset deviation range from the grayscale of the
selected testing point as a reference point or a combination of the another pixel and
a plurality of adjacent pixels thereof as the reference point, and calculating
fluorescence spectral data of the testing point and fluorescence spectral data of the
15 reference point.

Preferably, the preset wavelength range is beyond an effective response range
of an imaging spectrum detection apparatus; and

the light within the preset wavelength range can cause an analyte to excite a
fluorescence radiation signal, and a main peak of a fluorescence spectrum of the
20 fluorescence radiation signal is within the effective response range of the imaging
spectrum detection apparatus.

Preferably, the image includes spectral data that indicate, in the imaging area,
a fluorescence radiation signal excited by the first area when irradiated by
ultraviolet light; and

25 the spectral data collection step includes: based on grayscale distribution data,
dividing the imaging area into the testing point candidate area with a smaller
grayscale value and the reference point candidate area with a larger grayscale value,
selecting, based on the testing point candidate area, one pixel whose grayscale
value meets the preset requirements from the image as a testing point or a

combination of the one pixel and an adjacent pixel thereof as the testing point, and based on the reference point candidate area, selecting, from the image, another pixel whose grayscale value is within a preset deviation range from the grayscale value of the testing point as a reference point or a combination of the another pixel and a plurality of adjacent pixels thereof as the reference point; and

substituting a grayscale value of the testing point in the image into a spectrum reconstruction algorithm to obtain spectral data of the testing point, and substituting a grayscale value of the reference point in the image into the spectrum reconstruction algorithm to obtain spectral data of the reference point.

The present invention further provides method for testing an analyte, including the method for collecting fluorescence spectral data by ultraviolet light.

The present invention further provides a system for collecting fluorescence spectral data by ultraviolet light, and a medium, including:

an ultraviolet light irradiation module, configured to irradiate a first area by the ultraviolet light with a wavelength of 300–390 nm;

a signal collection module, configured to obtain a fluorescence radiation signal that includes the fluorescence spectral data and that is emitted by the first area when excited, and performing imaging based on the fluorescence radiation signal, to obtain an image; and

a spectral data collection module, configured to select a data collection point from the image, substitute a grayscale value of the data collection point into a spectrum reconstruction algorithm, and obtain the fluorescence spectral data by calculation, where a peak value of a spectral line of the fluorescence spectral data is within 400–800 nm.

Preferably, the image is collected under irradiation of the ultraviolet light within a preset wavelength range; and

the image includes spectral data that indicate, in an imaging area, a reflection signal or an excitation signal generated by the first area when irradiated by light.

Preferably, the collection module is configured to perform a step of:

based on grayscale values of pixels in the image, selecting one pixel whose grayscale value meets preset requirements from a testing point candidate area as a testing point or a combination of the one pixel and an adjacent pixel thereof as the testing point, and selecting, from a reference point candidate area, another pixel
5 whose grayscale value is within a preset deviation range from the grayscale of the selected testing point as a reference point or a combination of the another pixel and a plurality of adjacent pixels thereof as the reference point, and calculating fluorescence spectral data of the testing point and fluorescence spectral data of the reference point.

10 Preferably, the preset wavelength range is beyond an effective response range of an imaging spectrum detection apparatus; and

the light within the preset wavelength range can cause an analyte to excite a fluorescence radiation signal, and a main peak of a fluorescence spectrum of the fluorescence radiation signal is within the effective response range of the imaging
15 spectrum detection apparatus.

Preferably, the image includes spectral data that indicate, in the imaging area, a fluorescence radiation signal excited by the first area when irradiated by ultraviolet light; and

the spectral data collection module includes: based on grayscale distribution
20 data, dividing the imaging area into the testing point candidate area with a smaller grayscale value and the reference point candidate area with a larger grayscale value, selecting, based on the testing point candidate area, one pixel whose grayscale value meets the preset requirements from the image as a testing point or a combination of the one pixel and an adjacent pixel thereof as the testing point, and
25 based on the reference point candidate area, selecting, from the image, another pixel whose grayscale value is within a preset deviation range from the grayscale value of the testing point as a reference point or a combination of the another pixel and a plurality of adjacent pixels thereof as the reference point; and

substituting a grayscale value of the testing point in the image into a spectrum

reconstruction algorithm to obtain spectral data of the testing point, and substituting a grayscale value of the reference point in the image into the spectrum reconstruction algorithm to obtain spectral data of the reference point.

5 The present invention further provides system for testing an analyte, including the system for collecting fluorescence spectral data by ultraviolet light.

The present invention further provides a computer-readable storage medium that stores a computer program, and when the computer program is executed by a processor, the steps of the method for collecting fluorescence spectral data by ultraviolet light are implemented.

10 The present invention further provides an electronic device, including a memory, a processor, and a computer program stored in the memory and runnable on the processor, where when the computer program is executed by the processor, the steps of the method for collecting fluorescence spectral data by ultraviolet light are implemented.

15 Compared with the conventional technologies, the present invention has the following beneficial effects.

According to the technical solution in this application, peaks of spectral data of different analytes can be kept within a recognizable range of the imaging spectral detection device.

20 1. Electrochemical reaction with the analyte is not required in the technical solutions of this application, and a testing manner is more easily implemented, especially when an analyte in a living body is tested, so that non-invasive testing can be achieved.

25 2. In this application, spectral data of different areas can be obtained by utilizing uneven distribution of the analyte in the imaging area. Because other components in the imaging area except the analyte are distributed relatively uniformly, a difference in spectral data in different areas can directly demonstrate the information about the analyte (such as a concentration of the analyte) correlated to the spectral data after the influence of a non-analyte is excluded.

3. In this application, fluorescence spectroscopy is used for testing, preventing a traditional Raman method for measuring the analyte, thereby achieving low costs and miniaturization of the testing system and achieving real-time testing.

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BRIEF DESCRIPTION OF DRAWINGS

Other features, objects, and advantages of the present invention will become more apparent by reading detailed description of non-limiting embodiments with reference to the following drawings:

10 FIG. 1 is a flow chart in a first embodiment;

FIG. 2 is a schematic diagram of a first image collected in a second embodiment;

FIG. 3 is a schematic diagram of a second image collected in the second embodiment;

15 FIG. 4 is a schematic diagram of a testing model in the second embodiment;

FIG. 5 is a schematic diagram of testing point-reference point spectral data obtained in the second embodiment;

FIG. 6 is an experimental result of analysis result accuracy of an analysis model;

20 FIG. 7 is a schematic structural diagram of an analyte testing device provided in a fifth embodiment;

FIG. 8 is a schematic structural diagram of an electronic device provided in a sixth embodiment;

25 FIG. 9 is a schematic structural diagram of an analyte testing watch provided in the fifth embodiment;

FIG. 10 is a schematic diagram of a back of the analyte testing watch;

FIG. 11 is an exploded view of the analyte testing watch; and

FIG. 12 is a schematic diagram of a use state of the analyte testing watch.

In the accompanying drawings,
100: imaging area; 200: testing device;
201: light source; 202: imaging spectrum detection apparatus;
203: controller; 204: first bandpass filter;
5 205: lens; 206: second bandpass filter;
207: circuit board; 501: processor;
502: memory.

DETAILED DESCRIPTION OF EMBODIMENTS

10 The present invention is described below in detail with reference to specific embodiments. The following embodiments help those skilled in the art further understand the present invention, but do not limit the present invention in any form. It should be noted that those skilled in the art may make some improvements and transformation without departing from the idea of the present invention. The
15 improvements and transformation shall fall within the protection scope of the present invention.

First embodiment

FIG. 1 is a flow chart of this embodiment. This embodiment provides a method for testing an analyte, including the following steps.

20 Imaging: providing light within a preset wavelength range through a light source to irradiate a first area, and imaging the first area through an imaging spectrum detection apparatus, to obtain an image of an imaging area. The image can demonstrate, during light irradiation within a preset wavelength range, distribution data and spectral data in the imaging area of a reflection signal or an
25 excitation signal generated by the analyte when irradiated by light. The first area may be an area on a surface of a human skin. To prevent the influence of external light such as ambient light on testing, a collection window of the imaging spectrum detection apparatus needs to be tightly attached to the surface of the human skin in

the first area, and the imaging area means an area within a lens range of the imaging spectrum detection apparatus. In general, the imaging area may be a part of the first area, or may be the same area as the first area.

Because obtaining the distribution data and spectral data of the analyte
5 requires irradiation by light of different wavelength ranges, there are two
implementation manners: the light is light with a larger wavelength range provided
by one light source or light with smaller wavelength ranges provided by two light
sources respectively. When there is one light source, the wavelength range of the
light provided by the light source needs to cover both a wavelength range within
10 which analyte distribution data can be obtained and a wavelength range within
which analyte spectral data can be obtained. When there are two light sources, the
two light sources provide different pieces of light. A wavelength range of the
analyte distribution data can be obtained through a wavelength coverage of one
piece of light, and a wavelength range of the analyte spectral data can be obtained
15 through a wavelength coverage of the other piece of light. In addition, when there
is one light source, one image is formed; when there are two light sources, two
images are formed. For ease of processing, imaging areas of the two images are
required to be the same, that is, the collection window of the imaging spectrum
detection apparatus does not move on the surface of the human skin.

20 In this application, the analyte may be glucose, ketones, alcohol, lactate,
oxygen, hemoglobin A1C, acetylcholine, amylase, bilirubin, cholesterol, chorionic
gonadotropin, creatine kinase (such as CK-MB), creatine, creatinine, DNA,
fructosamine, glutamine, growth hormone, hormone, peroxide, prostate-specific
antigen, prothrombin, RNA, thyroid-stimulating hormone, or troponin in the blood
25 vessel, or may be a drug such as antibiotics (such as gentamicin, vancomycin, or
the like), digitoxin, digoxin, drugs of abuse, theophylline, or warfarin. In an
implementation in which two or more analytes are tested, the analytes may be
tested at the same or different time. In another embodiment, the analyte may
alternatively be another substance on the surface of the human body, and

non-invasive testing can be implemented according to the present invention.

Spectrum obtaining: obtaining, from the image by the imaging spectrum detection apparatus, spectral data that indicate uneven distribution in the imaging area of the reflection signal or the excitation signal generated by the analyte when
5 irradiated by light. Specifically, the imaging area can be partitioned based on different pieces of distribution data, so that a position of the spectral data can be selected from different partitions.

analyzing step: obtaining information about the analyte in the imaging area based on the obtained spectral data, where the information about the analyte
10 includes information about the analyte correlated to the spectral data. Due to a difference in distribution of the analyte in different areas, there is a difference in the reflection signal or excitation signal produced by the analytes when irradiated by light. For example, the human skin is divided into three parts: an epidermis tissue, a dermis tissue, and a subcutaneous tissue, and blood vessels such as veins
15 are located in the subcutaneous tissue. Corresponding spectral data can be obtained by irradiating a skin area with blood vessels and a skin area without blood vessels irradiated by ultraviolet light, or the corresponding spectral data can be obtained from a skin area with thicker blood vessels and a skin area with thinner blood vessels. Therefore, a difference between the two pieces of spectral data can
20 demonstrate the information about the analyte correlated to the spectral data in the blood vessels, such as intermediate information such as a degree of influence of the analyte on the spectral data for further analysis, or a concentration of the analyte and other information can be obtained directly through the analysis model.

Second embodiment

25 This embodiment is based on the first embodiment. For example, glucose testing in human blood vessels is used as an example, and a non-invasive glucose testing method is provided, including the following steps.

Imaging: irradiating, by infrared light, skin on the wrist and the back of the hand at which the veins are located, in a first wavelength range of 800–1,000 nm,

and collecting the first image of the imaging area, where the first wavelength range is preferably a near-infrared band; and irradiating, by ultraviolet light, a same position in a second wavelength range of 300-390 nm, to obtain the second image of the imaging area.

5 As shown in FIG. 2, an abscissa is a transverse coordinate of the first image, an ordinate is a longitudinal coordinate of the first image, white boxes represent selected testing point pixel blocks on venous blood vessels, and black boxes represent reference point pixel blocks on surrounding skin. In the first image, some of the infrared light penetrates the human skin and some is absorbed by the human
10 skin, and is also absorbed by the venous blood vessels in large quantities in an area at which the venous blood vessels are located, presenting a characteristic that a pixel grayscale value of the area at which the venous blood vessels are located is small, and a pixel grayscale value of the area at which non-venous blood vessels are located is large, thereby easily dividing the imaging area into the area at which
15 the venous blood vessels are located and the area at which the non-venous blood vessels are located.

As shown in FIG. 3, an abscissa is a transverse coordinate of the second image, an ordinate is a longitudinal coordinate of the second image, white boxes represent selected testing point pixel blocks on the venous blood vessels, and black
20 boxes represent reference point pixel blocks on surrounding skin. In the second image, because it is difficult to distinguish between the area at which the venous blood vessels are located and the area at which the non-venous blood vessels are located, the first image is required for distinguishing. Excitation light in the second wavelength range of 300–390 nm is used to obtain a high-quality effective
25 fluorescence spectral signal. A main response band of the imaging spectrum detection apparatus is 400–800 nm. When the wavelength of the excitation light is less than 300 nm, a main peak of a fluorescence spectrum of the excited fluorescence radiation signal is in a band <400 nm, and it is difficult for the imaging spectrum detection apparatus to obtain the high-quality effective

fluorescence spectral signal. When the wavelength of the excitation light used is >390 nm, the excitation light itself is visible light, the spectral signal of the excitation light and the fluorescence spectral signal are superimposed together, and it is difficult to eliminate the interference of the spectral signal of the excitation light and extract the effective fluorescence spectral signal. After absorbing ultraviolet light in the wavelength range of 300–390 nm, glucose in the venous blood vessels can emit a fluorescence radiation signal in a visible light band of 400–800 nm, and the band is in an effective response range of the imaging spectrum detection apparatus. A characteristic spectral intensity of the fluorescence radiation signal is positively correlated with the concentration of glucose, which has high fluorescence excitation efficiency.

Spectral obtaining: based on grayscale distribution of pixels in the first image, dividing the imaging area into the area at which the venous blood vessels are located and the area at which the non-venous blood vessels are located, selecting a testing point from a position corresponding to the second image in the area at which the venous blood vessels are located, selecting a reference point from a position corresponding to the second image in the area at which the non-venous blood vessels are located, and respectively obtaining spectral data of the testing point and spectral data of the reference point in the second image. Specifically, based on a grayscale value of the pixel in the second image, a pixel whose grayscale value meets preset requirements is selected from the area at which the venous blood vessels are located as the testing point, or a combination of the pixel and an adjacent pixel is selected as the testing point, and a pixel whose grayscale value has a deviation from the selected testing point within a preset deviation range is selected from the area at which the non-venous blood vessels are located as the reference point, or a combination of the pixel and a plurality of adjacent pixels is selected as the reference point, and the fluorescence spectral data of the testing point and the fluorescence spectral data of the reference point are calculated in the second image. The spectral data is taken from the testing point, a pixel of the

reference point, or a combined average of a plurality of pixels, which can be properly selected based on the width of the blood vessel. The combined average of the plurality of pixels can improve a signal-to-noise ratio, but is limited by the width of the blood vessel, preventing the spectral data from being taken from an extravascular area. Selection of a single pixel leads to high spatial resolution, which is proper for a thin blood vessel, but has a low signal-to-noise ratio. A preset requirement for the grayscale value may be using a point with the smallest grayscale value as the testing point, which is not limited in this application. A calculation result is shown in FIG. 5, an abscissa is the wavelength (in nm), an ordinate is a relative radiance (in W/nm), a solid line is the spectral data of the testing point, and a dashed line is the spectral data of the reference point. The grayscale value of the reference point and the grayscale value of the selected testing point are within the preset deviation range because the skin in the imaging area may be influenced by skin colors, pigmentation spots, cosmetics, and the like, which has a direct impact on the spectral data of the reference point, and the first image cannot distinguish the area including these influencing factors. These influencing factors can be effectively excluded by setting the preset deviation range of the grayscale value. In addition, the grayscale value and the grayscale value of the selected testing point are within the preset deviation range, which can ensure that the reference point is selected to be close to the testing point, such as an edge of the venous blood vessel. In this way, in addition to the blood vessels, the color, thickness, and other parameters of the epidermis tissue, the dermis tissue, and the subcutaneous tissues are approximately the same, so that a deviation between the spectral data of the testing point and the spectral data of the reference point can be excluded from the influence of a non-analyte as much as possible.

In addition to a spectrum reconstruction algorithm, the spectral data may alternatively be obtained by forming a radiometric calibration coefficient through previous radiometric calibration, and spectral lines are obtained by calculating the grayscale value * radiometric calibration coefficient.

When a combination of a plurality of pixels is selected at the testing point, the fluorescence spectral data at the testing point may be an average of fluorescence spectral data of these pixels. In addition, there may be one or more testing points and reference points. When there is a plurality of testing points and
5 reference points, the average of the fluorescence spectral data of all testing points and the average of the fluorescence spectral data of all reference points can be calculated respectively.

analyzing step: preprocessing the obtained spectral data of the testing points and reference points, inputting the preprocessed data into a trained testing model,
10 and outputting a concentration of glucose or an intermediate result of glucose correlated with the spectral data. When the testing model is trained, it is necessary to obtain spectral data and an accurate test result of a tested object, such as a blood collection test result, the spectral data is used as the input of the testing model, a blood collection test result is used as the output of the testing model, to train the
15 testing model.

The testing model may be a convolutional neural network model, including an input layer, at least two convolutional layers, at least two activation function layers, a Flatten layer, a fully connected layer, and an output layer in sequence, and the convolutional layer and the activation function layer are spaced apart. An
20 activation function used for the activation function layer is a Relu function.

In the convolutional neural network model, a size of each layer of convolutional kernels is 1, the number of convolutional kernels of a first convolutional layer is 32, and the number of convolutional kernels of a second layer is 64, which are both used to extract blood glucose features, and the output of
25 the convolutional layer is transformed nonlinearly by the activation function. The Flatten layer flattens the output of the convolutional layer into a one-dimensional vector, to connect the subsequent fully connected layers, resulting in a final output dimension of 1. The Adam optimizer is used for model training during model training, a mean square error is used as a loss function, and a mean absolute error is

calculated as a performance indicator of model evaluation.

When an output result of the testing model is glucose concentration, if an error between the output result and a measured standard glucose concentration value meets a preset condition, training is stopped, to obtain the testing model.

- 5 When the output result of the testing model is the intermediate result of glucose correlated to the spectral data, such as an intermediate neuron result, if the error between the output result and the intermediate neuron result meets the preset condition, training is stopped to obtain the testing model. The intermediate neuron result is further model-corrected to obtain the concentration of glucose.

- 10 As shown in FIG. 4, the input layer is a spectral data input layer, which is obtained after the original spectral data is preprocessed. A hidden layer is a middle hidden layer, and a finally predicted blood glucose concentration value is output in an output layer after feature combination through convolution operation deep learning. Alternatively, through convolution operation deep learning, a neuron
- 15 Output1 is output as an intermediate result value after feature combination, and model training is performed again on the intermediate result values Output1 and two infrared IR feature brightness values, to further correct a blood glucose prediction error, and the finally predicted blood glucose concentration value Output2 is output. Different parameters need to be set for a training degree of the
- 20 testing model according to the needs, and a plurality of extracted glucose eigenvalues are continuously learned based on setting of different parameters, until an error between an output result and a standard glucose value of the above label value meets the requirements, and then training is stopped, to obtain the testing model.

- 25 Through a plurality of iterations of training, neurons can learn corresponding changes between different glucose concentrations and glucose spectral characteristics of different sampled objects, thereby improving the universality of the testing model and implementing prediction of glucose concentrations for different users.

In the whole glucose testing process, there is no need to pierce the skin for collecting blood or pierce the skin for implantation, and spectral information of a person to be tested is obtained based on the fluorescence spectrum, and a glucose test result of the person to be tested is obtained based on the spectral information, preventing pain and discomfort, and improving testing comfort and convenience. In the method, a spectral signal for blood vessels and a spectral signal for the skin can be finely distinguished, which provides a possibility for subsequent accurate extraction of a glucose signal, and also enables the spectral signal to be strongly correlated with the glucose concentration, implementing accurate measurement of the glucose concentration, so that a test result is more accurate, and processing is more easily performed.

FIG. 6 shows a schematic diagram of an experimental effect of the trained testing model, where an abscissa is a reference blood glucose concentration (in mmol/L) collected by a blood glucose meter, and an ordinate is a blood glucose concentration (in mmol/L) predicted in the method in the patent. There are totally 2,037 samples of tested objects, including 1,537 samples from a training set and 500 samples from a prediction set. Distribution of test results for the testing model can be learned from the figure, a MARD value of the predicted samples is 11.32%, and most of the samples fall in areas A and B, among which 87.03% of the samples fall in area A and 12.77% of the samples fall in area B, indicating that testing accuracy of the testing model is high.

Third embodiment

This embodiment is based on the second embodiment. Infrared light is replaced with visible light, and another non-invasive method for glucose testing is provided, including the following steps.

Imaging: collecting the first image of the imaging area by irradiating visible light on the skin, such as the wrist and the back of the hand, at which the veins are located; and irradiating, by ultraviolet light, a same position in a second wavelength range of 300-390 nm, to obtain the second image of the imaging area.

In the first image, due to the difference between a color of the area at which the venous blood vessels are located and a color of the area at which the non-venous blood vessels are located, the imaging area can be easily divided into the area at which the venous blood vessels are located and the area at which the non-venous blood vessels are located.

In the second image, because it is difficult to distinguish between the area at which the venous blood vessels are located and the area at which the non-venous blood vessels are located, the first image is required for distinguishing. Excitation light in the second wavelength range of 300–390 nm is used to obtain a high-quality effective fluorescence spectral signal. A main response band of the imaging spectrum detection apparatus is 400–800 nm. When the wavelength of the excitation light is less than 300 nm, a main peak of a fluorescence spectrum of the excited fluorescence radiation signal is in a band <400 nm, and it is difficult for the imaging spectrum detection apparatus to obtain the high-quality effective fluorescence spectral signal. When the wavelength of the excitation light used is >390 nm, the excitation light itself is visible light, the spectral signal of the excitation light and the fluorescence spectral signal are superimposed together, and it is difficult to eliminate the interference of the spectral signal of the excitation light and extract the effective fluorescence spectral signal. After absorbing ultraviolet light in the wavelength range of 300–390 nm, glucose in the venous blood vessels can emit a fluorescence radiation signal in a visible light band of 400–800 nm, and the band is in an effective response range of the imaging spectrum detection apparatus. A characteristic spectral intensity of the fluorescence radiation signal is positively correlated with the concentration of glucose, which has high fluorescence excitation efficiency.

Spectral obtaining: based on grayscale distribution of pixels in the first image, dividing the imaging area into the area at which the venous blood vessels are located and the area at which the non-venous blood vessels are located, selecting a testing point from the area at which the venous blood vessels are located, selecting

a reference point from the area at which the non-venous blood vessels are located, and respectively obtaining spectral data of the testing point and spectral data of the reference point. Specifically, based on a grayscale value of the pixel in the second image, a pixel whose grayscale value meets preset requirements is selected from the area at which the venous blood vessels are located as the testing point, or a combination of the pixel and an adjacent pixel is selected as the testing point, and a pixel whose grayscale value has a deviation from the selected testing point within a preset deviation range is selected from the area at which the non-venous blood vessels are located as the reference point, or a combination of the pixel and a plurality of adjacent pixels is selected as the reference point, and the fluorescence spectral data of the testing point and the fluorescence spectral data of the reference point are calculated. The grayscale value of the reference point and the grayscale value of the selected testing point are within the preset deviation range because the skin in the imaging area may be influenced by skin colors, pigmentation spots, cosmetics, and the like, which has a direct impact on the spectral data of the reference point, and the first image cannot distinguish the area including all influencing factors. These influencing factors can be effectively excluded by setting the preset deviation range of the grayscale value.

When a combination of a plurality of pixels is selected at the testing point, the fluorescence spectral data at the testing point may be an average of fluorescence spectral data of these pixels. In addition, there may be one or more testing points and reference points. When there is a plurality of testing points and reference points, the average of the fluorescence spectral data of all testing points and the average of the fluorescence spectral data of all reference points can be calculated respectively.

analyzing step: preprocessing the obtained spectral data of the testing points and reference points, inputting the preprocessed data into a trained testing model, and outputting the concentration of glucose. When the testing model is trained, it is necessary to obtain spectral data and an accurate test result of a tested object, such

as a blood collection test result, the spectral data is used as the input of the testing model, a blood collection test result is used as the output of the testing model, to train the testing model.

5 The testing model may be a convolutional neural network model, including an input layer, at least two convolutional layers, at least two activation function layers, a Flatten layer, a fully connected layer, and an output layer in sequence, and the convolutional layer and the activation function layer are spaced apart. An activation function used for the activation function layer is a Relu function.

10 In the convolutional neural network model, a size of each layer of convolutional kernels is 1, the number of convolutional kernels of a first convolutional layer is 32, and the number of convolutional kernels of a second layer is 64, which are both used to extract blood glucose features, and the output of the convolutional layer is transformed nonlinearly by the activation function. The Flatten layer flattens the output of the convolutional layer into a one-dimensional
15 vector, to connect the subsequent fully connected layers, resulting in a final output dimension of 1. The Adam optimizer is used for model training during model training, a mean square error is used as a loss function, and a mean absolute error is calculated as a performance indicator of model evaluation.

20 If an error between the output result of the testing model and a standard glucose concentration meets a preset condition, training is stopped, to obtain the testing model.

Different parameters need to be set for a training degree of the testing model according to the needs, and a plurality of extracted glucose eigenvalues are continuously learned based on setting of different parameters, until an error
25 between an output result and a standard glucose value of the above label value meets the requirements, and then training is stopped, to obtain the testing model.

Through a plurality of iterations of training, neurons can learn corresponding changes between different glucose concentrations and glucose spectral characteristics of different sampled objects, thereby improving the universality of

the testing model and implementing prediction of glucose concentrations for different users.

In the whole glucose testing process, there is no need to collect blood or pierce the skin, and spectral information of a person to be tested is obtained based on the fluorescence spectrum, and a glucose test result of the person to be tested is obtained based on the spectral information, preventing pain and discomfort, and improving testing comfort and convenience. In the method, a spectral signal for blood vessels and a spectral signal for the skin can be finely distinguished, which provides a possibility for subsequent accurate extraction of a glucose signal, and also enables the spectral signal to be strongly correlated with the glucose concentration, implementing accurate measurement of the glucose concentration, so that a test result is more accurate, and processing is more easily performed.

Fourth embodiment

This embodiment provides a system for testing an analyte. The system for testing an analyte can be implemented by performing flow steps of the method for testing an analyte. In other words, those skilled in the art can understand the method for testing an analyte as a preferred implementation of the system for testing an analyte. The system for testing an analyte includes the following modules.

An imaging module, configured to provide light within a preset wavelength range through a light source to irradiate a first area, and image the first area through an imaging spectrum detection apparatus, to obtain an image of an imaging area. The image can demonstrate, during light irradiation within a preset wavelength range, distribution data and spectral data in the imaging area of a reflection signal or an excitation signal generated by the analyte when irradiated by light. Because different wavelength ranges are required to obtain distribution data and spectral data of the analyte, the light may be either light corresponding to two wavelength ranges or light with a larger wavelength range that covers the two desired wavelength ranges. When there are two pieces of light, there are two

obtained images. To facilitate processing, it is usually required that imaging areas of the two images be the same.

In this application, the analyte may be glucose, ketones, alcohol, lactate, oxygen, hemoglobin A1C, acetylcholine, amylase, bilirubin, cholesterol, chorionic gonadotropin, creatine kinase (such as CK-MB), creatine, creatinine, DNA, fructosamine, glutamine, growth hormone, hormone, peroxide, prostate-specific antigen, prothrombin, RNA, thyroid-stimulating hormone, or troponin in the blood vessel of an animal, or may be a drug such as antibiotics (such as gentamicin, vancomycin, or the like), digitoxin, digoxin, drugs of abuse, theophylline, and warfarin. In an implementation in which one or more analytes are tested, the analytes may be tested at the same or different time. In other embodiments, the analyte may alternatively be another substance in liquid.

A spectrum obtaining module, configured to obtain, from the image by the imaging spectrum detection apparatus, spectral data that indicate uneven distribution in the imaging area of the reflection signal or the excitation signal generated by the analyte when irradiated by light. Specifically, the imaging area can be partitioned based on different pieces of distribution data, so that a position of the spectral data can be selected from different partitions.

An analysis module, configured to obtain information about the analyte in the imaging area based on the obtained spectral data, where the information about the analyte includes information about the analyte correlated to the spectral data. Due to a difference in distribution of the analyte in different areas, there is a difference in the reflection signal or excitation signal produced by the analytes when irradiated by light. With this feature, the difference between the two pieces of spectral data can be obtained, to accurately demonstrate the information about the analyte correlated to the spectral data, such as the concentration of the analyte.

Those skilled in the art know that, in addition to implementing the system provided in the present invention and each apparatus, module, and unit thereof in a purely computer-readable program code mode, the system provided in the present

invention and each apparatus, module, and unit thereof can be enabled to implement same functions in the form of logic gates, switches, application-specific integrated circuits, programmable logic controllers, and embedded microcontrollers by performing logic programming of the method steps. Therefore, the system
5 provided in the present invention and each apparatus, module, and unit thereof may be regarded as hardware components, and each apparatus, module, and unit thereof for implementing various functions may also be regarded as structures within the hardware components; and each apparatus, module, and unit thereof for implementing various functions may alternatively be regarded as software modules
10 for implementing methods or structures within the hardware components.

Fifth embodiment

FIG. 7 shows an electronic device of this embodiment, which is specifically a testing device 200 of an analyte. The testing device 200 is a portable non-invasive testing device for a human body, which may be a single testing device or be
15 integrated into a watch or mobile phone, to implement easy and rapid testing of the analyte in the surface of the human body.

The testing device 200 includes: a light source 201, an imaging spectrum detection apparatus 202, a controller 203, a first bandpass filter 204, a second bandpass filter 206, and a lens 205. The controller 203 establishes an electric
20 connection or communication connection with the light source 201 and the imaging spectrum detection apparatus 202 separately.

The light source 201 can provide light within a preset wavelength range. Because obtaining the distribution data and spectral data of the analyte requires irradiation by light of different wavelength ranges, there are two implementation
25 manners: a light source that can provide light with a larger wavelength range or two light sources that can respectively provide light with smaller wavelength ranges. When there is one light source, the wavelength range of the light provided by the light source needs to cover both a wavelength range within which analyte distribution data can be obtained and a wavelength range within which analyte

spectral data can be obtained, such as a halogen lamp. When there are two light sources, the two light sources provide different pieces of light. A wavelength range of the analyte distribution data can be obtained through a wavelength coverage of one piece of light, and a wavelength range of the analyte spectral data can be obtained through a wavelength coverage of the other piece of light, such as an infrared lamp combined with an ultraviolet lamp, a visible light lamp combined with an ultraviolet lamp.

To enable the imaging area 100 to be irradiated evenly, a ring-shaped light source may be used. The light source has a plurality of light-emitting modules evenly distributed on a same circumference. When there are two light sources, the light-emitting modules of the two light sources are arranged alternately.

The imaging spectrum detection apparatus 202 can image the imaging area 100 based on an instruction to obtain a corresponding image, and can obtain corresponding spectral data based on the instruction. The imaging spectrum detection apparatus 202 includes a sensor and a periodic pixel-level light filtering structure arranged on a surface of the sensor. The periodic pixel-level light filtering structure is configured to perform spectral modulation on an incoming light signal, so that the sensor generates an image containing spectral information to be tested.

The periodic pixel-level light filtering structure includes a plurality of light filtering image element channels with pixel-level structures of different shapes. The plurality of light filtering image element channels have same specifications and sizes and are evenly arranged, and lengths and widths thereof are integer multiples of a pixel size in an image sensor. Light filtering image element channels of pixel-level light filtering structures of different shapes are corresponding to different spectral filter coefficients, and the pixel-level light filtering structures with different spectral filter coefficients are combined in a fixed order and then arranged periodically. The sensor modulates a received first testing light through the periodic pixel-level light filtering structure arranged on the surface of the sensor, to form a mosaic image containing spectral information, and then

reconstructs a grayscale image including the spectral information to be tested using an algorithm.

The controller 203 is configured to control the light source to provide light within a preset wavelength range to irradiate the first area, control the imaging spectrum detection apparatus to image the first area to obtain an image of an imaging area, and control the imaging spectrum detection apparatus to obtain, from the imaging area, spectral data that indicate uneven distribution in the imaging area of the reflection signal or excitation signal generated by the analyte when irradiated by light; and obtain information about the analyte in the imaging area based on the obtained spectral data, where the information about the analyte includes information about the analyte correlated to the spectral data. When there is one light source 201, one image is formed. When there are two light sources 201, two images are formed. When the first light source is turned on, the second light source is turned off. Similarly, when the second light source is turned on, the first light source is turned off. The two light sources do not interfere with each other.

The first bandpass filter 204 is located between the light source 201 and the imaging area 100. A function of the first bandpass filter 204 is to allow light within a preset wavelength range to pass through, while cutting off light outside the preset wavelength range, thereby reducing the impact of other external light on a test result.

The second bandpass filter 206 is located between the imaging spectrum detection apparatus 202 and the lens 205. A function of the second bandpass filter 206 is to allow light within a wavelength range in which the reflection signal or excitation signal generated by the analyte when irradiated by light is located to pass through, while cutting off light within another wavelength range, thereby reducing the influence of the reflection signal or excitation signal of a non-analyte on the test result.

The lens 205 can be configured to fix focus, to obtain a high-definition image. In another embodiment, the second bandpass filter 206 may alternatively be located

on a side of the lens 205 away from the imaging spectrum detection apparatus 202, which is not limited in the present invention.

Based on the above description, FIG. 9 shows an analyte testing watch provided in this embodiment. A front of the watch is a display, and as shown in FIG. 10, a back of the watch has a light-transmitting window with a built-in testing device 200. As shown in FIG. 11, the light source 201 and the first bandpass filter 204 are both annular structures. Light-emitting modules of the light source 201 are distributed in a ring shape, and emitted light is filtered by the first bandpass filter 204, and then light with a required wavelength is output, and is irradiated onto a human body through the light-transmitting window on the back of the watch. The reflection signal or excitation signal of the human body enters the light-transmitting window, passes through the light source 201 and a hollow part in the middle of the first bandpass filter 204, enters a second bandpass filter 206 through a lens 205, and enters an imaging spectrum detection apparatus 202 after being filtered by the second bandpass filter 206. The imaging spectrum detection apparatus 202 is mounted on a circuit board 207, and a controller 203 (not shown in the figure) of the testing device 200 is also mounted on the circuit board 207. As shown in FIG. 12, to more accurately identify a position of a venous blood vessel, the watch can be worn on an inner side of the wrist.

Sixth embodiment

FIG. 8 is a schematic structural diagram of an electronic device according to an embodiment of this application. As shown in FIG. 8, the electronic device includes at least one processor 501 and a memory 502 that is in communication connection with the at least one processor 501. The memory 502 stores instructions that can be executed by the at least one processor 501. The instructions are executed by the at least one processor 501, so that the at least one processor 501 can perform the above method for testing an analyte.

The memory 502 and the processor 501 are connected using a bus. The bus may include interconnected buses and bridges of any quantities. The bus connects

various circuits of one or more processors 501 and memories 502. The bus may further connect a peripheral device, a voltage regulator, and various other circuits such as a power management circuit, which are all well known in the art. A bus interface provides an interface between the bus and a transceiver. The transceiver
5 may be one component or a plurality of components, for example, a plurality of receivers and transmitters, to provide a unit that is configured to communicate with various other apparatuses on a transmission medium. Data processed by the processor 501 is transmitted on a wireless medium by using the antenna. Further, the antenna further receives data and transmits the data to the processor 501.

10 The processor 501 is responsible for managing the bus and general processing, and may further provide various functions, including timing, peripheral interfacing, voltage regulation, power management, and another control function. The memory 502 may be configured to store data used by the processor 501 when performing an operation.

15 The present invention also provides a computer-readable storage medium that stores a computer program, and when the computer program is executed by a processor, the above method for testing an analyte is implemented.

To be specific, those skilled in the art can understand that all or some of steps in the method of the above embodiments can be completed by instructing relevant
20 hardware through a program. The program is stored in a storage medium and includes several instructions to enable a device (which may be a single-chip microcontroller, a chip, or the like) or a processor (processor) to perform all or some of the steps of the method described in various embodiments of this application. The above storage medium includes any medium that can store
25 program code, such as a USB flash drive, a removable hard disk, a read-only memory (ROM, Read-Only Memory), a random access memory (RAM, Random Access Memory), a magnetic disk, or an optical disc.

Those skilled in the art can understand that the above implementations are specific embodiments for implementing the present invention, and in actual

application, various changes may be made thereto in form and detail without departing from the spirit and scope of the present invention.

Specific embodiments of the present invention are described above. It should be understood that the present invention is not limited to the foregoing specific
5 implementations. Those skilled in the art can make various variations or modifications within the scope of the claims, which does not affect the essence of the present invention. Embodiments in this application and the features in embodiments may be arbitrarily and mutually combined in the case of no conflict.

CLAIMS

1. A method for collecting fluorescence spectral data by ultraviolet light, comprising:

ultraviolet light irradiation step: irradiating a first area by the ultraviolet light with a
5 wavelength of 300–390 nm;

signal collection step: obtaining a fluorescence radiation signal that comprises the fluorescence spectral data and that is emitted by the first area when excited, and performing imaging based on the fluorescence radiation signal, to obtain an image; and

spectral data collection step: selecting a data collection point from the image,
10 substituting a grayscale value of the data collection point into a spectrum reconstruction algorithm, and obtaining the fluorescence spectral data by calculation, wherein a peak value of a spectral line of the fluorescence spectral data is within 400–800 nm.

2. The method for collecting fluorescence spectral data by ultraviolet light
15 according to claim 1, wherein the image is collected under irradiation of the ultraviolet light within a preset wavelength range; and

the image comprises spectral data that indicate, in an imaging area, a reflection signal or an excitation signal generated by the first area when irradiated by light.

20 3. The method for collecting fluorescence spectral data by ultraviolet light according to claim 2, wherein the signal collection comprises:

based on grayscale values of pixels in the image, selecting one pixel whose grayscale value meets preset requirements from a position that is in a testing point candidate area and that is corresponding to the image as a testing point or a combination

of the one pixel and an adjacent pixel thereof as the testing point, and selecting, from a reference point candidate area, another pixel whose grayscale value is within a preset deviation range from the grayscale of the selected testing point as a reference point or a combination of the another pixel and a plurality of adjacent pixels thereof as the
5 reference point, and calculating fluorescence spectral data of the testing point and fluorescence spectral data of the reference point.

4. The method for collecting fluorescence spectral data by ultraviolet light according to claim 3, wherein the preset wavelength range is beyond an effective
10 response range of an imaging spectrum detection apparatus; and

the light within the preset wavelength range enables to cause an analyte to excite a fluorescence radiation signal, and a main peak of a fluorescence spectrum of the fluorescence radiation signal is within the effective response range of the imaging spectrum detection apparatus.

15

5. The method for collecting fluorescence spectral data by ultraviolet light according to claim 4, wherein the image comprises spectral data that indicate, in the imaging area, the fluorescence radiation signal excited by the first area when irradiated by the ultraviolet light; and

20

the spectral data collection step comprises: based on grayscale distribution data, dividing the imaging area into the testing point candidate area with a smaller grayscale value and the reference point candidate area with a larger grayscale value, selecting, based on the testing point candidate area, one pixel whose grayscale value meets the preset requirements from the image as a testing point or a combination of the one pixel

and an adjacent pixel thereof as the testing point, and based on the reference point candidate area, selecting, from the image, another pixel whose grayscale value is within a preset deviation range from the grayscale value of the testing point as a reference point or a combination of the another pixel and a plurality of adjacent pixels thereof as the
5 reference point; and

substituting a grayscale value of the testing point in the image into a spectrum reconstruction algorithm to obtain spectral data of the testing point, and substituting a grayscale value of the reference point in the image into the spectrum reconstruction algorithm to obtain spectral data of the reference point.

10

6. A method for testing an analyte, comprising the method for collecting fluorescence spectral data by ultraviolet light according to any one of claims 1 to 5.

7. A system for collecting fluorescence spectral data by ultraviolet light,
15 comprising:

an ultraviolet light irradiation module, configured to irradiate a first area by the ultraviolet light with a wavelength of 300–390 nm;

a signal collection module, configured to obtain a fluorescence radiation signal that comprises the fluorescence spectral data and that is emitted by the first area when
20 excited, and performing imaging based on the fluorescence radiation signal, to obtain an image; and

a spectral data collection module, configured to select a data collection point from the image, substitute a grayscale value of the data collection point into a spectrum reconstruction algorithm, and obtain the fluorescence spectral data by calculation,

wherein a peak value of a spectral line of the fluorescence spectral data is within 400–800 nm.

8. The system for collecting fluorescence spectral data by ultraviolet light
5 according to claim 7, wherein the image is collected under irradiation of the ultraviolet light within a preset wavelength range; and

the image comprises spectral data that indicate, in an imaging area, a reflection signal or an excitation signal generated by the first area when irradiated by light.

10 9. The system for collecting fluorescence spectral data by ultraviolet light according to claim 8, wherein the signal collection module comprises:

based on grayscale values of pixels in the image, selecting one pixel whose grayscale value meets preset requirements from a position that is in a testing point candidate area and that is corresponding to the image as a testing point or a combination
15 of the one pixel and an adjacent pixel thereof as the testing point, and selecting, from a reference point candidate area, another pixel whose grayscale value is within a preset deviation range from the grayscale of the selected testing point as a reference point or a combination of the another pixel and a plurality of adjacent pixels thereof as the reference point, and calculating fluorescence spectral data of the testing point and
20 fluorescence spectral data of the reference point.

10. The system for collecting fluorescence spectral data by ultraviolet light according to claim 9, wherein the preset wavelength range is beyond an effective response range of an imaging spectrum detection apparatus; and

the light within the preset wavelength range enables to cause an analyte to excite a fluorescence radiation signal, and a main peak of a fluorescence spectrum of the fluorescence radiation signal is within the effective response range of the imaging spectrum detection apparatus.

5

11. The system for collecting fluorescence spectral data by ultraviolet light according to claim 10, wherein the image comprises spectral data that indicate, in the imaging area, the fluorescence radiation signal excited by the first area when irradiated by the ultraviolet light; and

10 the spectral data collection module comprises: based on grayscale distribution data, dividing the imaging area into the testing point candidate area with a smaller grayscale value and the reference point candidate area with a larger grayscale value, selecting, based on the testing point candidate area, one pixel whose grayscale value meets the preset requirements from the image as a testing point or a combination of the one pixel
15 and an adjacent pixel thereof as the testing point, and based on the reference point candidate area, selecting, from the image, another pixel whose grayscale value is within a preset deviation range from the grayscale value of the testing point as a reference point or a combination of the another pixel and a plurality of adjacent pixels thereof as the reference point; and

20 substituting a grayscale value of the testing point in the image into a spectrum reconstruction algorithm to obtain spectral data of the testing point, and substituting a grayscale value of the reference point in the image into the spectrum reconstruction algorithm to obtain spectral data of the reference point.

12. A system for testing an analyte, comprising the system for collecting fluorescence spectral data by ultraviolet light according to any one of claims 7 to 11.

13. A computer-readable storage medium that stores a computer program, wherein
5 when the computer program is executed by a processor, the steps of the method for collecting fluorescence spectral data by ultraviolet light according to any one of claims 1 to 5 are implemented.

14. An electronic device, comprising a memory, a processor, and a computer
10 program stored in the memory and runnable on the processor, wherein when the computer program is executed by the processor, the steps of the method for collecting fluorescence spectral data by ultraviolet light according to any one of claims 1 to 5 are implemented.