



US 20200113453A1

(19) **United States**(12) **Patent Application Publication**
PARK et al.(10) **Pub. No.: US 2020/0113453 A1**(43) **Pub. Date: Apr. 16, 2020**(54) **APPARATUS AND METHOD FOR
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Oct. 10, 2018 (KR) 10-2018-0120640

Publication Classification(51) **Int. Cl.****A61B 5/021** (2006.01)**A61B 5/00** (2006.01)(52) **U.S. Cl.**CPC **A61B 5/02108** (2013.01); **A61B 5/02255**
(2013.01); **A61B 5/7242** (2013.01)

(57)

ABSTRACT

A blood pressure estimating apparatus is provided. The blood pressure estimating apparatus may include: a sensor configured to measure a bio-signal waveform; and a processor configured to obtain a first feature based on a first area under the bio-signal waveform in a first time interval, obtain a second feature based on a second area under the bio-signal waveform in a second time interval which is different from the first time interval, and estimate blood pressure based on the first feature and the second feature.

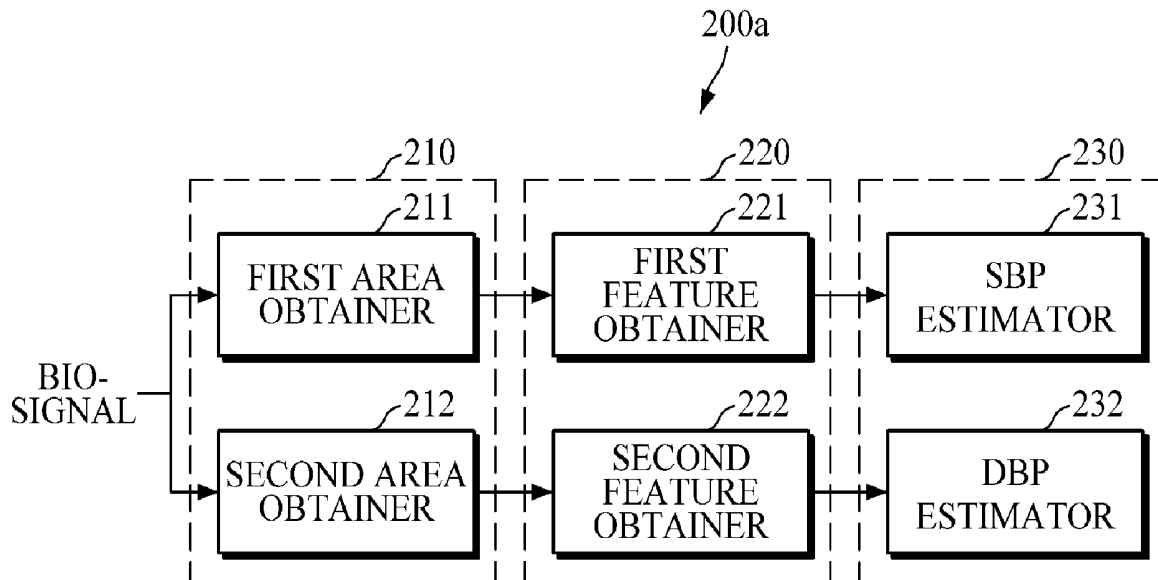


FIG. 1A

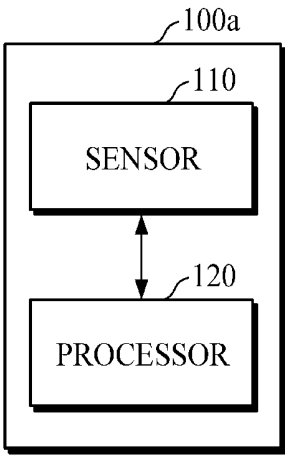


FIG. 1B

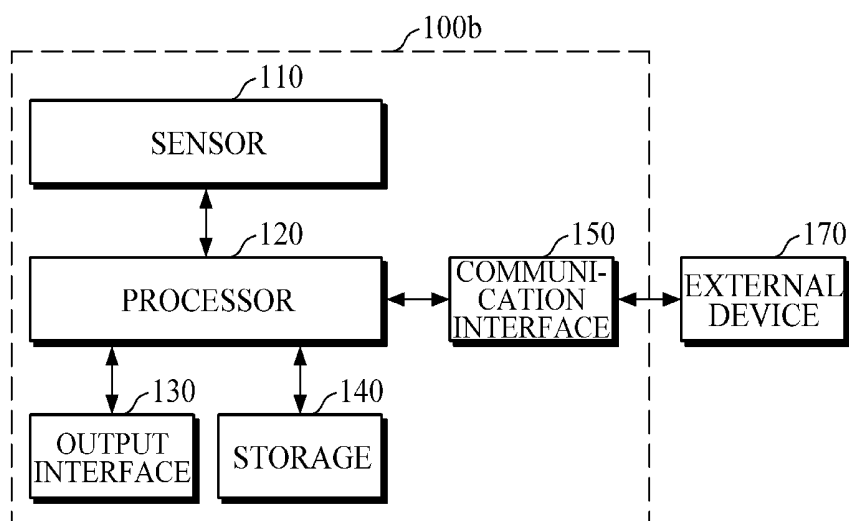


FIG. 2A

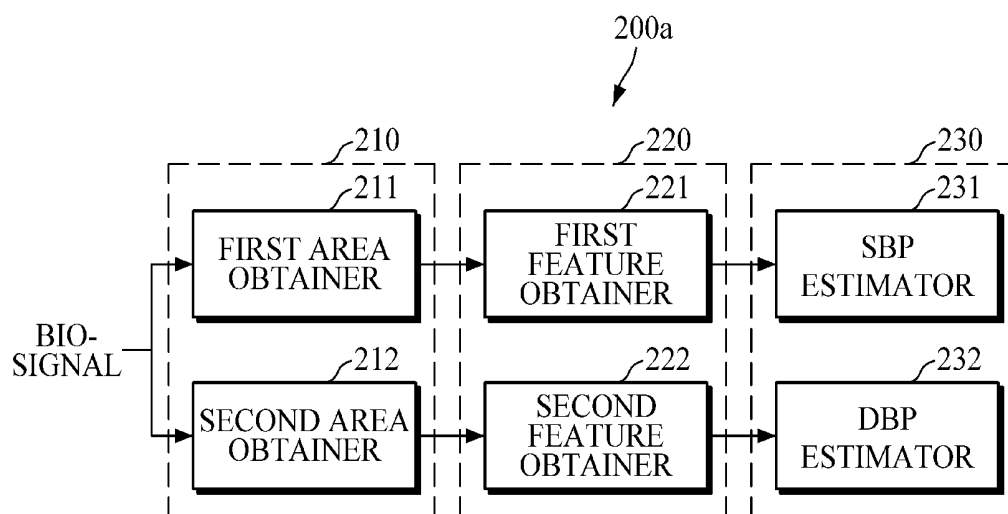


FIG. 2B

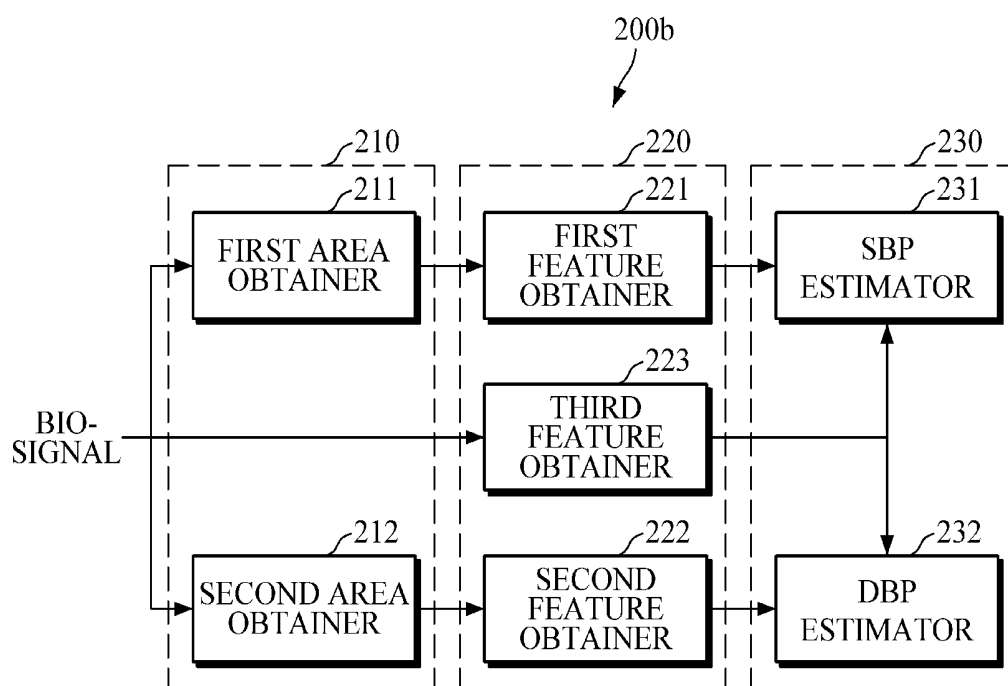


FIG. 3A

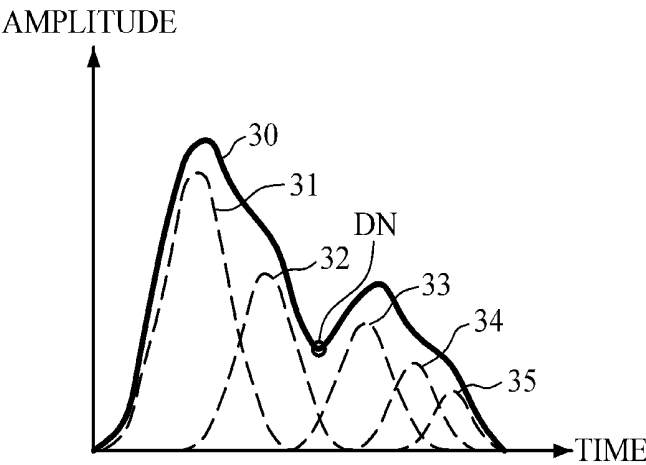


FIG. 3B

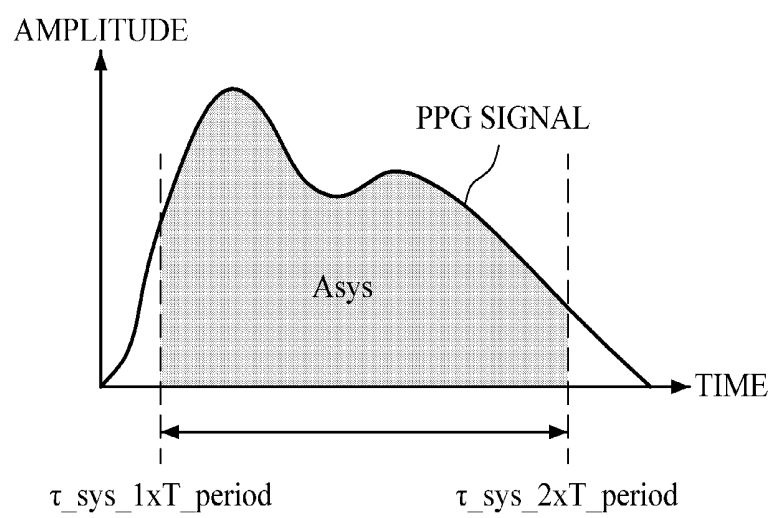


FIG. 3C

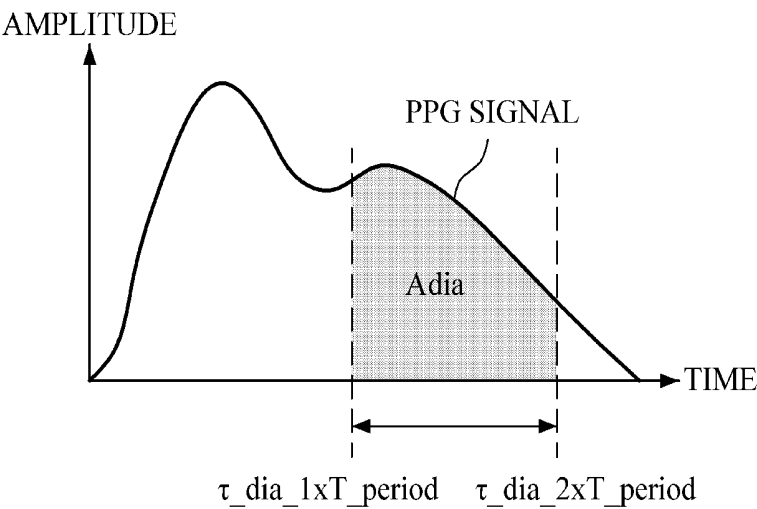


FIG. 3D

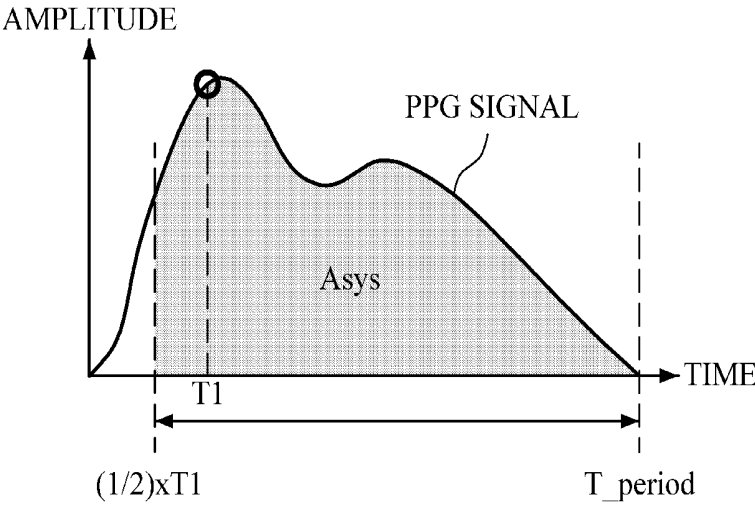


FIG. 3E

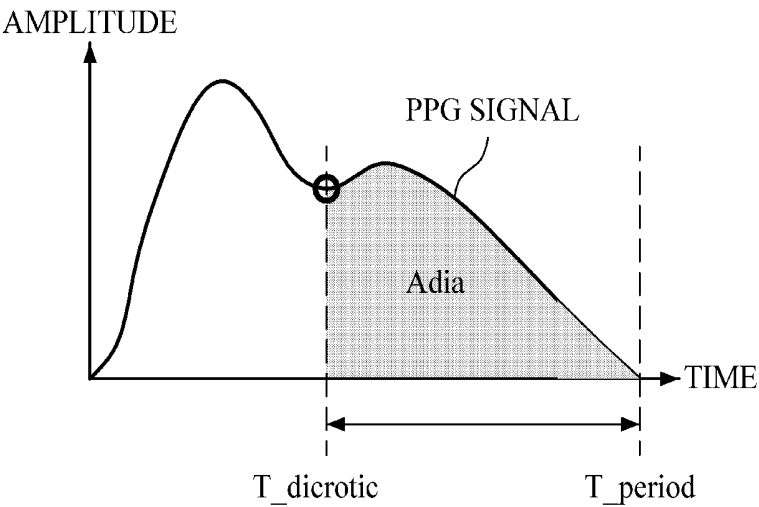


FIG. 4

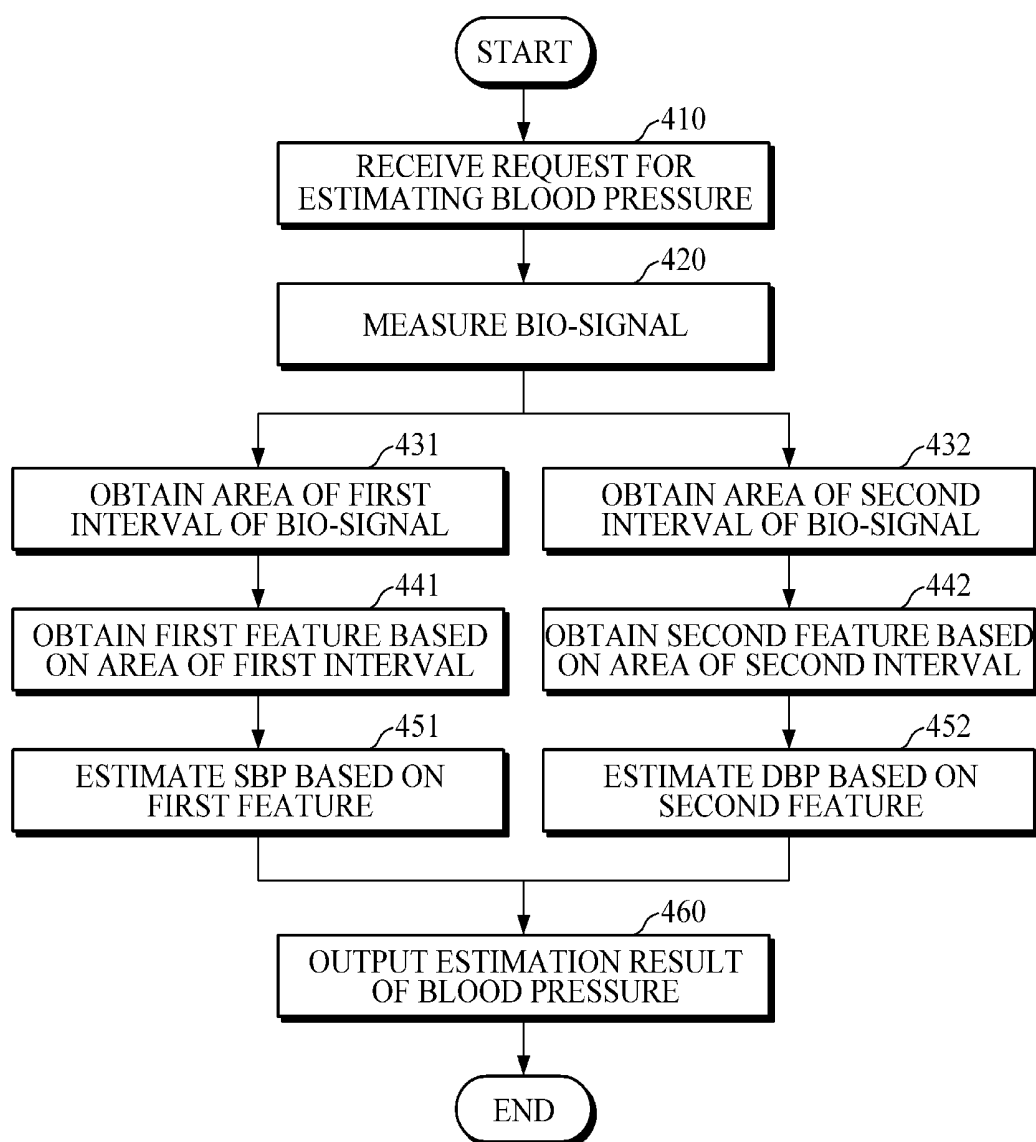


FIG. 5

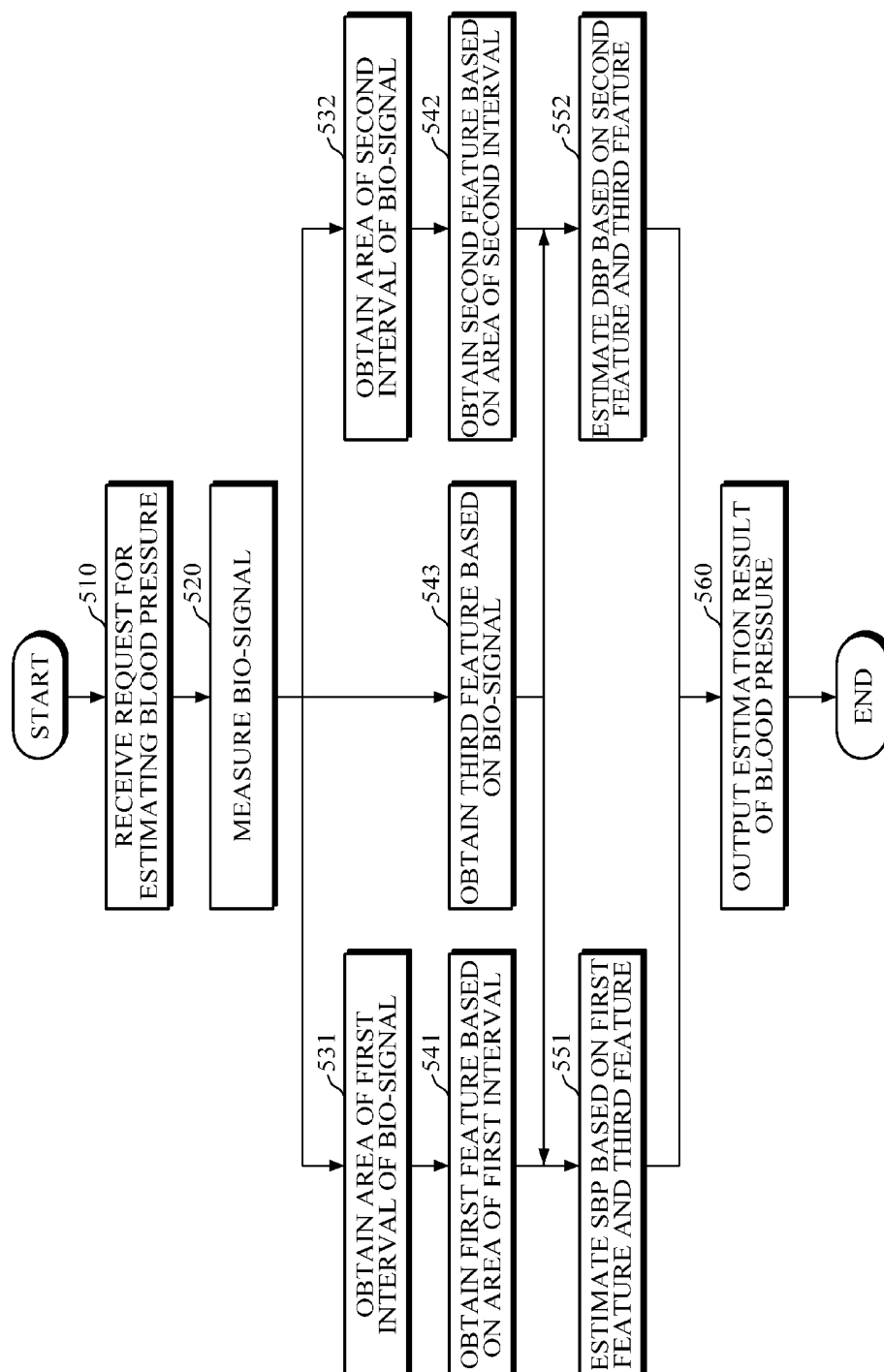


FIG. 6A

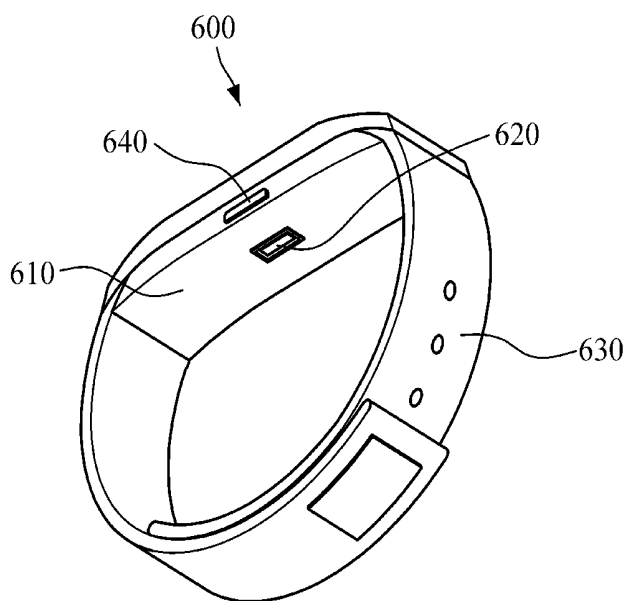


FIG. 6B

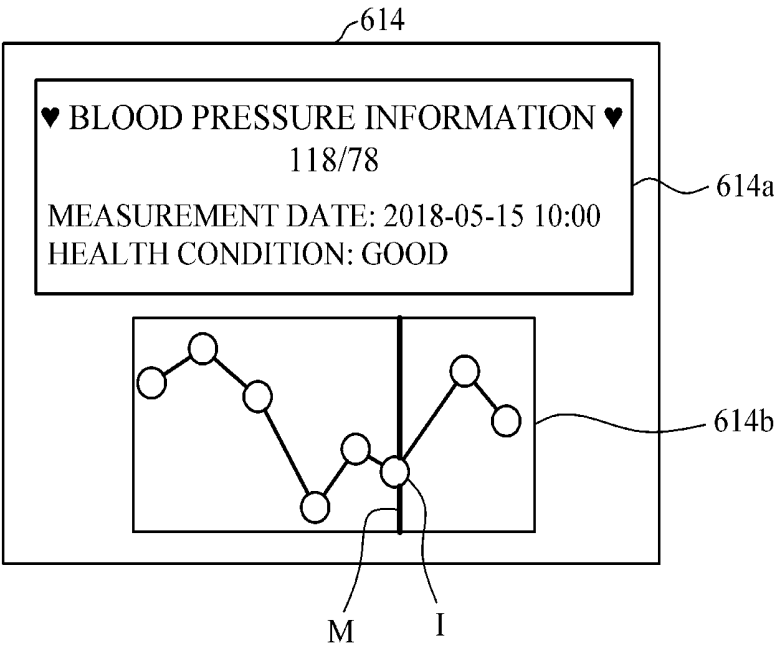
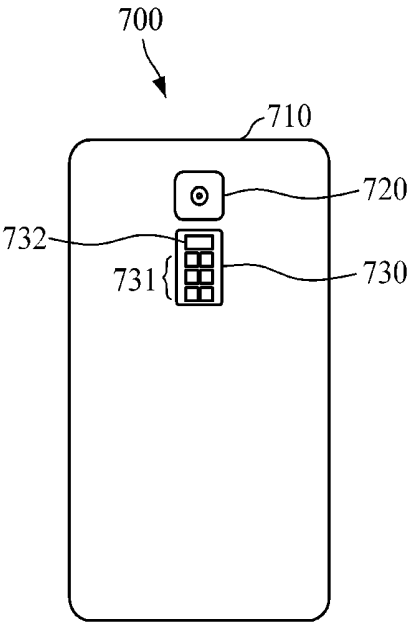


FIG. 7



APPARATUS AND METHOD FOR ESTIMATING BLOOD PRESSURE

CROSS-REFERENCE TO RELATED APPLICATION(S)

[0001] This application claims priority from Korean Patent Application No. 10-2018-0120640, filed on Oct. 10, 2018 in the Korean Intellectual Property Office, the disclosure of which is incorporated herein by reference in its entirety.

BACKGROUND

1. Field

[0002] Apparatuses and methods consistent with example embodiments relate to cufflessly estimating blood pressure.

2. Description of the Related Art

[0003] Recently, with the aging population, soaring medical costs, and a lack of medical personnel for specialized medical services, research is being actively conducted on tech convergence for medical devices. Particularly, monitoring of patients' health condition is not limited to places such as hospitals, but is expanding to mobile healthcare fields that may allow the patients to monitor their health condition anywhere and anytime in daily life including their home or office. Typical examples of bio-signals, which indicate the health condition of individuals, include an electrocardiography (ECG) signal, a photoplethysmogram (PPG) signal, an electromyography (EMG) signal, and the like, and various bio-signal sensors are being developed to measure these signals in daily life. Particularly, a PPG sensor may estimate blood pressure of a human body by analyzing a pulse waveform which reflects a condition of the cardiovascular system of the human body.

[0004] According to studies on the PPG signal, the entire PPG signal is a superposition of propagation waves starting from the heart toward the distal end portions of the body and reflection waves returning back from the distal end portions. Further, it has been known that information for estimating blood pressure may be obtained by extracting various features related to the propagation waves or the reflection waves.

SUMMARY

[0005] According to an aspect of an example embodiment, there is provided a blood pressure estimating apparatus including: a sensor configured to measure a bio-signal waveform; and a processor configured to obtain a first feature based on a first area under the bio-signal waveform in a first time interval, obtain a second feature based on a second area under the bio-signal waveform in a second time which is different from the first interval, and estimate blood pressure based on the first feature and the second feature.

[0006] The processor may estimate systolic blood pressure (SBP) and diastolic blood pressure (DBP) by applying a blood pressure estimation model to each of the first feature and the second feature.

[0007] The processor may determine a start time and an end time of each of the first time interval and the second time interval based on a bio-signal period of the bio-signal

waveform, and a reference starting value and a reference ending value of each of the first time interval and the second time interval.

[0008] In this case, the reference starting value and the reference ending value of each of the first time interval and the second time interval may include at least one of a constant value and a variable value which is adaptively changed according to a shape of the bio-signal waveform.

[0009] The processor may obtain the first area of the first time interval and the second area of the second time interval by integrating differences between an amplitude value of the bio-signal waveform and an amplitude value of a baseline of the bio-signal waveform, for the first time interval and the second time interval, respectively.

[0010] In this case, the amplitude value of the baseline may include at least one of an amplitude value of the bio-signal waveform at a measurement start time of the bio-signal waveform, a minimum amplitude value of the bio-signal waveform, and an amplitude value of a straight line formed by connecting the start time and the end time of each of the first time interval and the second time interval.

[0011] The processor may determine a start time and an end time of each of the first time interval and the second time interval based on at least one of a start time and an end time of measurement of the bio-signal waveform, a first time of a position of a pulse waveform component present in the bio-signal waveform, a second time of a diastolic notch position of the bio-signal waveform, times calculated by multiplying the first and the second times by a predetermined coefficient, and times of an internally dividing point obtained by applying a weighted value to the first and the second times.

[0012] The processor may perform differentiation on the bio-signal waveform to obtain a differential signal, and may determine a local minimum point of the differential signal to be the position of the pulse waveform component.

[0013] The weighted value may be determined based on at least one of an amplitude of the differential signal and an amplitude of the bio-signal waveform corresponding to the first and the second times.

[0014] The processor may obtain the first feature and the second feature by normalizing the first area and the second area based on a first reference value and a second reference value, respectively.

[0015] The processor may normalize the first area and the second area by dividing the first area and the second area by the first reference value and the second reference value respectively, or by dividing the first reference value and the second reference value by the first area and the second area, respectively.

[0016] The processor may set the first reference value and the second reference value based on at least one of a predetermined value, an amplitude value of the bio-signal waveform in a systolic phase or a diastolic phase of the bio-signal waveform, an amplitude value of an internally dividing point between two pulse waveform components of the bio-signal waveform, and a maximum amplitude value of the bio-signal waveform in the systolic phase or the diastolic phase.

[0017] The processor may further obtain a third feature from the bio-signal waveform, and may estimate systolic blood pressure based on the first feature and the third feature and diastolic blood pressure based on the second feature and the third feature.

[0018] The processor may obtain the third feature from the bio-signal based on at least one of a shape of the bio-signal waveform, a time value and an amplitude value of a maximum point of the bio-signal waveform, a time value and an amplitude value of a minimum point of the bio-signal waveform, a time value and an amplitude value of a position of a pulse waveform component present in the bio-signal waveform, and an area under the bio-signal waveform in the third interval.

[0019] The sensor may include: a light source configured to emit light onto an object; and a detector configured to detect light scattered from the object.

[0020] In addition, the blood pressure estimating apparatus may further include an output interface configured to output a processing result of the processor.

[0021] According to an aspect of another example embodiment, there is provided a blood pressure estimating method including: measuring a bio-signal waveform; obtaining a first area under the bio-signal waveform in a first time interval and a second area under the bio-signal waveform in a second time interval which is different from the first time interval; obtaining a first feature and a second feature based on the first area and the second area, respectively; and estimating blood pressure based on the first feature and the second feature.

[0022] The estimating of the blood pressure may include independently estimating systolic blood pressure (SBP) and diastolic blood pressure (DBP) by applying a blood pressure estimation model to each of the first feature and the second feature.

[0023] The obtaining of the first and the second areas may include determining a start time and an end time of each of the first time interval and the second time interval based on a bio-signal period of the bio-signal waveform, and a reference starting value and a reference ending value of each of the first time interval and the second time interval.

[0024] The reference starting value and the reference ending value of each of the first time interval and the second time interval may include at least one of a constant value and a variable value which is adaptively changed according to a shape of the bio-signal waveform.

[0025] The obtaining of the first and the second areas may include obtaining the first area and the second area by integrating differences between an amplitude value of the bio-signal waveform and an amplitude value of a baseline of the bio-signal waveform, for the first time interval and the second time interval, respectively.

[0026] The amplitude value of the baseline may include at least one of an amplitude value of the bio-signal waveform at a measurement start time of the bio-signal waveform, a minimum amplitude value of the bio-signal waveform, and an amplitude value of a straight line formed by connecting the start time and the end time of each of the first time interval and the second time interval.

[0027] The obtaining of the first and the second areas may include determining a start time and an end time of each of the first time interval and the second time interval based on at least one of a start time and an end time of measurement of the bio-signal waveform, a first time of a position of a pulse waveform component present in the bio-signal waveform, a second time of a diastolic notch position of the bio-signal waveform, times calculated by multiplying the first and the second times by a predetermined coefficient,

and times of an internally dividing point obtained by applying a weighted value to the first and the second times.

[0028] The obtaining of the first feature and the second feature may include normalizing the first area and the second area based on a first reference value and a second reference value, respectively.

[0029] The obtaining of the first feature and the second feature may include setting the first reference value and the second reference value based on at least one of a predetermined value, an amplitude value of the bio-signal waveform in a systolic phase or a diastolic phase of the bio-signal waveform, an amplitude value of an internally dividing point between two pulse waveform components of the bio-signal waveform, and a maximum amplitude value of the bio-signal waveform in the systolic phase or the diastolic phase.

[0030] In addition, the blood pressure estimating method may further include obtaining a third feature from the bio-signal waveform, wherein the estimating of the blood pressure may include estimating systolic blood pressure based on the first feature and the third feature and diastolic blood pressure based on the second feature and the third feature.

[0031] The obtaining of the third feature may include obtaining the third feature from the bio-signal waveform based on at least one or two or more of a shape of the bio-signal waveform, a time value and an amplitude value of a maximum point of the bio-signal waveform, a time value and an amplitude value of a minimum point of the bio-signal waveform, a time value and an amplitude value of a position of a pulse waveform component present in the bio-signal waveform, and an area under the bio-signal waveform in a third time interval.

[0032] Moreover, the blood pressure estimating method may further include outputting an estimation result of blood pressure.

[0033] According to an aspect of another example embodiment, there is provided a non-transitory computer readable storage medium storing a program that is executable by a computer to perform a method including: obtaining a bio-signal waveform; setting a systolic time interval and a diastolic time interval of the bio-signal waveform to be different from each other; calculating a first area under the bio-signal waveform in the systolic time interval, and a second area under the bio-signal waveform in the diastolic time interval; and estimating a blood pressure based on the first area in the systolic time interval and the second area in the diastolic time interval.

BRIEF DESCRIPTION OF THE DRAWINGS

[0034] The above and/or other aspects will be more apparent by describing certain example embodiments, with reference to the accompanying drawings, in which:

[0035] FIGS. 1A and 1B are block diagrams illustrating a blood pressure estimating apparatus according to example embodiments;

[0036] FIGS. 2A and 2B are block diagrams illustrating a processor of FIGS. 1A and 1B;

[0037] FIGS. 3A, 3B, 3C, 3D, and 3E are diagrams explaining extracting features for estimating systolic blood pressure (SBP) and diastolic blood pressure (DBP) according to example embodiments;

[0038] FIG. 4 is a flowchart illustrating a blood pressure estimating method according to an example embodiment;

[0039] FIG. 5 is a flowchart illustrating a blood pressure estimating method according to another example embodiment;

[0040] FIGS. 6A and 6B are diagrams illustrating a wearable device according to an example embodiment; and

[0041] FIG. 7 is a diagram illustrating a smart device for estimating blood pressure according to an example embodiment.

DETAILED DESCRIPTION

[0042] Example embodiments are described in greater detail below with reference to the accompanying drawings.

[0043] In the following description, like drawing reference numerals are used for like elements, even in different drawings. The matters defined in the description, such as detailed construction and elements, are provided to assist in a comprehensive understanding of the example embodiments. However, it is apparent that the example embodiments can be practiced without those specifically defined matters. Also, well-known functions or constructions are not described in detail since they would obscure the description with unnecessary detail.

[0044] It will be understood that, although the terms first, second, etc. may be used herein to describe various elements, these elements should not be limited by these terms. These terms are only used to distinguish one element from another. Any references to singular may include plural unless expressly stated otherwise. In addition, unless explicitly described to the contrary, an expression such as “comprising” or “including” will be understood to imply the inclusion of stated elements but not the exclusion of any other elements. Also, the terms, such as or ‘module’, etc should be understood as a unit that performs at least one function or operation and that may be embodied as hardware, software, or a combination thereof.

[0045] Expressions such as “at least one of,” when preceding a list of elements, modify the entire list of elements and do not modify the individual elements of the list. For example, the expression, “at least one of a, b, and c,” should be understood as including only a, only b, only c, both a and b, both a and c, both b and c, all of a, b, and c, or any variations of the aforementioned examples.

[0046] FIGS. 1A and 1B are block diagrams illustrating a blood pressure estimating apparatus according to example embodiments. The blood pressure estimating apparatuses 100a and 100b may be embedded in an electronic device, such as a smartphone, a tablet PC, a desktop computer, a laptop computer, and the like, or in a wearable device that may be worn on an object. In particular, examples of the wearable device may include a wristwatch-type wearable device, a bracelet-type wearable device, a wristband-type wearable device, a ring-type wearable device, a glasses-type wearable device, a hairband-type wearable device, or the like, but the wearable device is not limited thereto, and may be embedded in a medical device manufactured for use in medical institutions to measure and analyze bio-information.

[0047] Referring to FIGS. 1A and 1B, each of the blood pressure estimating apparatuses 100a and 100b includes a sensor 110 and a processor 120.

[0048] The sensor 110 may measure a bio-signal from an object. In particular, the bio-signal may be a pulse wave signal including a photoplethysmogram (PPG) signal. However, the bio-signal is not limited thereto, and may include various bio-signals, such as an electrocardiography (ECG)

signal, an electromyography (EMG) signal, and the like, which may be modeled by adding a plurality of waveform components. The object may be a body part which comes into contact with or is adjacent to the sensor 110, and may be a body part where pulse waves may be easily measured. For example, the object may be an area of skin on the wrist that is adjacent to the radial artery or a human skin area through which veins or capillaries pass. However, the object is not limited thereto, and may be peripheral body portions, such as fingers, toes, and the like, which have a high density of blood vessels.

[0049] The sensor 110 may include a light source and a detector. The light source may emit light onto the object, and the detector may detect light scattered or reflected from the object. The light source may include a light emitting diode (LED), a laser diode (LD), a fluorescent body, and the like, and may be arranged in one or two or more arrays. The detector may include one or more pixels, each of which includes a photo diode, a photo transistor (PTr), an image sensor, and the like, which detects light and converts the detected light into an electric signal. The sensor 110 may be implemented as a spectrometer.

[0050] The processor 120 may be electrically connected to the sensor 110. The processor 120 may control the sensor 110 in response to a request for estimating blood pressure, and may receive a bio-signal from the sensor 110. The request for estimating blood pressure may be input by a user, or may be generated at predetermined estimation intervals. Upon receiving an electrical bio-signal from the sensor 110, the processor 120 may perform preprocessing such as filtering for removing noise, amplification of the bio-signal, conversion of the bio-signal into a digital signal, and the like.

[0051] The processor 120 may estimate blood pressure based on the bio-signal received from the sensor 110. In particular, the processor 120 may independently estimate systolic blood pressure (SBP) and diastolic blood pressure (DBP) by analyzing a waveform of a bio-signal. For example, based on the area under a waveform of a bio-signal, the processor 120 may obtain a first feature for estimating SBP and a second feature for estimating DBP, and may estimate SBP by using the first feature and DBP by using the second feature.

[0052] Referring to FIG. 1B, the blood pressure estimating apparatus 100b may further include an output interface 130, a storage 140, and a communication interface 150.

[0053] The output interface 130 may output results processed by the sensor 110 and the processor 120. For example, the output interface 130 may visually output an estimated blood pressure value by using a display screen, or may output the estimated blood pressure value in a non-visual manner through voice, vibration, tactile sensation, and the like, by using a speaker, a haptic motor, and the like. The output interface 130 may divide a display area into two or more areas according to a setting, and may output a bio-signal graph used for estimating blood pressure, a blood pressure estimation result, and the like in a first area, and a blood pressure estimation history in the form of a graph and the like in a second area. In this case, if an estimated blood pressure value falls outside a normal range, the output interface 130 may output warning information in various manners, such as highlighting an abnormal value in red and the like, displaying the abnormal value along with a normal

range, outputting a voice warning message, adjusting a vibration intensity, and the like.

[0054] The storage **140** may store results processed by the sensor **110** and the processor **120**. Further, the storage **140** may store various criteria required for estimating blood pressure. For example, the criteria may include user feature information such as a user's age, gender, health condition, and the like. In addition, the criteria may also include information such as a reference blood pressure value, a blood pressure estimation model, a blood pressure estimation period, and the like. Further, the criteria may include a method of setting an interval for calculating an area, a start reference value and an end reference value of each interval, a baseline for calculating an area, a first reference value and a second reference value for normalization, and the like, but are not limited thereto.

[0055] In particular, the storage **140** may include at least one storage medium of a flash memory type memory, a hard disk type memory, a multimedia card micro type memory, a card type memory (e.g., an SD memory, an XD memory, etc.), a Random Access Memory (RAM), a Static Random Access Memory (SRAM), a Read Only Memory (ROM), an Electrically Erasable Programmable Read Only Memory (EEPROM), a Programmable Read Only Memory (PROM), a magnetic memory, a magnetic disk, and an optical disk, and the like, but the storage medium is not limited thereto.

[0056] The communication interface **150** may communicate with an external device **170** using wired or wireless communication techniques under the control of the processor **120**, and may transmit and receive various data to and from the external device **170**. For example, the communication interface **150** may transmit a blood pressure estimation result to the external device **170**, and may receive various criteria required for estimating blood pressure from the external device **170**. Examples of the external device **170** may include a cuff-type blood pressure measuring device, and an information processing device such as a smartphone, a tablet PC, a desktop computer, a laptop computer, and the like.

[0057] In particular, examples of the communication techniques may include Bluetooth communication, Bluetooth Low Energy (BLE) communication, Near Field Communication (NFC), WLAN communication, Zigbee communication, Infrared Data Association (IrDA) communication, Wi-Fi Direct (WFD) communication, Ultra-Wideband (UWB) communication, Ant+ communication, WIFI communication, Radio Frequency Identification (RFID) communication, 3G communication, 4G communication, 5G communication, and the like. However, this is merely an example and is not intended to be limiting.

[0058] FIGS. 2A and 2B are block diagrams illustrating a processor according to example embodiments of FIGS. 1A and 1B. FIGS. 3A to 3E are diagrams explaining extracting features for estimating systolic blood pressure (SBP) and diastolic blood pressure (DBP) according to example embodiments.

[0059] FIG. 3A illustrates a measured pulse wave signal **30** formed by superposition of five pulse waveforms **31**, **32**, **33**, **34**, and **35** according to an example embodiment. The pulse wave signal is a superposition of propagation waves of a pumping pulse and reflection waves, wherein the pumping pulse represents blood flow that starts from the heart toward the distal end portions of the body, and the reflection waves represent blood flow that returns back from the distal end

portions. Based on the combination of time and amplitude values of the pulse waveforms **31**, **32**, **33**, **34**, and **35**, features having a high correlation with blood pressure may be extracted. For example, the first, second and third pulse waveforms **31-33** may be used to estimate blood pressure. Pulse waveforms after the third pulse waveform **33** may not be observed depending on individuals in some cases and are difficult to be found due to noise, or have a low correlation with estimation of blood pressure.

[0060] Among the pulse waveforms **31**, **32**, **33**, **34**, and **35** of the pulse wave signal **30**, the first pulse waveform **31** and the second pulse waveform **32** may be present in a systolic phase, and the pulse waveform **34** and **35** following the third pulse waveform **33** may be present in a diastolic phase. Here, the systolic and the diastolic phases may be divided based on a position of a downwardly concave dicrotic notch (DN) which is located between the second pulse waveform **32** and the third pulse waveform **33**. For example, the systolic phase may correspond to the bio-signal between a start time T_{start} of the bio-signal and a time T_{DN} when the DN occurs, and the diastolic phase may correspond to the bio-signal between the time T_{DN} and an end time T_{end} of the bio-signal. In defining the systolic phase and the diastolic phase, an offset time T_{offset} may be added to the start time T_{start} and may be subtracted from the end time T_{end} .

[0061] Referring to FIG. 2A, a processor **200a** includes an area obtainer **210**, a feature obtainer **220**, and a blood pressure estimator **230**.

[0062] The area obtainer **210** may include a first area obtainer **211** and a second area obtainer **212**. The first area obtainer **211** may determine a first area under a bio-signal waveform in a first interval to estimate SBP. The area under the bio-signal waveform may be referred to Area Under the Curve (AUC). For example, the area obtainer **210** may obtain the first area by using the following Equation 1.

$$Asys = \sum_{t=T_{sys_1}}^{t=T_{sys_2}} (P(t) - P_{base}) \quad \text{[Equation 1]}$$

[0063] Herein, $Asys$ denotes the first area of the first interval which is to be used to estimate SBP; T_{sys_1} and T_{sys_2} denote a start time and an end time of the first interval; $P(t)$ denotes an amplitude value at a time (t); and P_{base} denotes a baseline amplitude value. In this case, the amplitude value of the baseline may be an amplitude value at the initial start time of the bio-signal or an amplitude value in the entire interval of the bio-signal; or may be an amplitude value of a straight line formed by connecting the start time and the end time, e.g., an amplitude value corresponding to the time (t) on the straight line with respect to the time (t), but the amplitude value of the baseline is not limited thereto.

[0064] The first area obtainer **211** may analyze the bio-signal to determine the start time and the end time of the first interval. For example, the first area obtainer **211** may obtain the start time of the first interval by multiplying a bio-signal period by a reference starting value, and may obtain the end time of the first interval by multiplying the bio-signal period by a reference ending value, as represented by the following Equation 2. FIG. 3B illustrates an example of obtaining the start time and the end time of the first interval of the PPG

signal, and obtaining an area $Asys$ between the start time to the end time, as represented by the following Equation 2

$$T_{sys_1} = \tau_{sys_1} \times T_{period}$$

$$T_{sys_2} = \tau_{sys_2} \times T_{period} \quad [\text{Equation 2}]$$

[0065] Herein, τ_{sys_1} and τ_{sys_2} denote the reference starting value and the reference ending value of the first interval respectively; and T_{period} denotes the period of the bio-signal.

[0066] The reference starting value and the reference ending value may be values predetermined through preprocessing. For example, the reference starting value may be selected from a range of 0 to 0.3 and the reference ending value may be selected from a range of 0.7 to 1.0, so that the first interval of the bio-signal may include most of the entire interval thereof. The values are merely examples for convenience of explanation, and are not limited thereto. In this case, the reference starting value and the reference ending value may be predetermined for each user, or for groups classified according to various criteria such as age, gender, health condition, and the like.

[0067] Alternatively, the reference starting value and the reference ending value may be values which are adaptively changed according to a shape of a bio-signal. That is, once a bio-signal is measured, the first area obtainer **211** may analyze the measured bio-signal to adjust the reference starting value and the reference ending value. For example, in the case where a heart rate, extracted from the bio-signal, increases higher or decreases lower than a threshold, or in the case where it is determined that a rate of change in the bio-signal over time is greater than a reference rate, the first area obtainer **211** may increase or decrease the reference starting value or the reference ending value according to a predetermined adjustment rate. In this case, the adjustment rate may be set based on comparison between an increase/decrease in the heart rate and the threshold, and/or comparison between the rate of change in the bio-signal over time and the reference rate.

[0068] In another example, the first area obtainer **211** may obtain the start time and the end time of the first interval by using time values associated with the pulse waveform components present in the bio-signal. In particular, the first area obtainer **211** may perform secondary differentiation on the bio-signal, and may detect a local minimum point of the differential signal, to determine the position of the local minimum point as a position of the pulse waveform component.

[0069] For example, the time values associated with the pulse waveform components may include a time of a position of the pulse waveform component, a time of a position of the DN, a time calculated by multiplying the times by a predetermined coefficient, a time of an internally dividing point obtained by applying a weighted value to the times, a start time and an end time of a bio-signal period, a time calculated by linearly combining at least some of the times, and the like. In this case, the weighted value to be applied to the times may be determined by using an amplitude of a differential signal of positions corresponding to the times, and/or an amplitude of the bio-signal. For example, the amplitude of the differential signal and/or the amplitude of the bio-signal, corresponding to each of the times, may be set as a weighted value to be applied to each time, so that a higher weighted value is applied to time values corresponding to a higher amplitude of a differential signal and/or a

higher amplitude of the bio-signal, and an internally dividing point may be obtained by integrating the time values, to which weighted values are applied, and by dividing the added value by the sum of the weighed values. However, the weighted value is not limited thereto.

[0070] FIG. 3D illustrates an example of determining, as a start time of a first interval, a value $\frac{1}{2} \times T1$ calculated by multiplying a time $T1$, corresponding to a position of a first pulse waveform component of a PPG signal, by a predetermined coefficient ($\frac{1}{2}$); determining, as an end time of the first interval, a time T_{period} corresponding to an end time of a period of the PPG signal; and obtaining an area $Asys$ of the first interval.

[0071] In order to estimate DBP, the second area obtainer **211** may obtain a second area under a bio-signal waveform in a second interval. In this case, the second interval may be an interval which is different from the first interval on a time axis of the bio-signal. For example, the second area obtainer **211** may obtain the second area of the second interval by using the following Equations 3 and 4. FIG. 3C illustrates an example of obtaining a start time and an end time of the second interval of the PPG signal using the following Equation 4, and obtaining an area $Adia$ between the start time and the end time.

$$Adia = \sum_{t=T_{dia_1}}^{t=T_{dia_2}} (P(t) - P_{base}) \quad [\text{Equation 3}]$$

$$T_{dia_1} = \tau_{dia_1} \times T_{period}$$

$$T_{dia_2} = \tau_{dia_2} \times T_{period} \quad [\text{Equation 4}]$$

[0072] Herein, $Adia$ denotes the second area of the second interval for estimating DBP; T_{dia_1} and T_{dia_2} denote a start time and an end time of the second interval; $P(t)$ denotes an amplitude value at a time (t); and P_{base} denotes an amplitude value of a baseline. In this case, the amplitude value of the baseline may be an amplitude value of the initial start time or a minimum amplitude value in the entire interval of the bio-signal; or may be an amplitude value of a straight line formed by connecting the start time and the end time, e.g., an amplitude value corresponding to the time (t) on the straight line with respect to the time (t), but the amplitude value of the baseline is not limited thereto. Further, τ_{dia_1} and τ_{dia_2} denote a reference starting value and a reference ending value of the second interval respectively; and T_{period} denotes the period of the bio-signal.

[0073] As in the first area obtainer **211** described above, the second area obtainer **212** may set predetermined fixed values as the reference starting value and the reference ending value. For example, the second area obtainer **212** may set an interval, which is 0.4 to 0.9 times the period of the bio-signal, as the second interval, so that the diastolic interval may be mainly used for estimating DBP. For example, the second area obtainer **212** may set 0.5 as a reference starting value, and 0.8 as a reference ending value. However, the values are merely examples for convenience of understanding, and are not limited thereto. In another example, the second area obtainer **212** may analyze the waveform of the bio-signal to adaptively adjust the reference starting value and the reference ending value, which is

described above with reference to the first area obtainer **211**, such that detailed description thereof will be omitted.

[0074] Further, instead of obtaining the start time and the end time by using the above Equation 4, the second area obtainer **212** may obtain the start time and/or the end time of the second interval by using time values associated with the pulse waveform components present in the bio-signal, which is described above with reference to the first area obtainer **211**, such that detailed description thereof will be omitted. FIG. 3E illustrates an example of determining a time T_{dirotic} , corresponding to the position of the DN of the PPG signal, as the start time of the second interval, and a time T_{period} corresponding to the end of the period of the bio-signal as the end time of the second interval, and obtaining an area A_{dia} of the second interval.

[0075] The feature obtainer **220** may extract features for estimating blood pressure by using the area obtained by the area obtainer **210**. The feature obtainer **220** may include a first feature obtainer **221** and a second feature obtainer **222** as illustrated therein.

[0076] The first feature obtainer **221** may obtain a first feature for estimating SBP based on the area of the first interval obtained by the first area obtainer **211**. For example, the first feature obtainer **221** may obtain the first area of the first interval as it is as the first feature for estimating SBP. In another example, the first feature obtainer **221** may normalize the area of the first interval by using a predetermined first reference value, and may obtain the normalized value as the first feature. For example, the first feature obtainer **221** may normalize the first area of the first interval by dividing the first area of the first interval by the first reference value, or conversely, by dividing the first reference value by the first area of the first interval. In this case, the first reference value may include a bio-signal amplitude value of the position of the first pulse waveform component in the systolic phase, a bio-signal amplitude value of the position of the second pulse waveform component in the systolic phase, a bio-signal amplitude value of an internally dividing point between the position of the first pulse waveform component and the position of the second pulse waveform component, a maximum amplitude value in the systolic phase, and the like. However, the first reference value is not limited thereto.

[0077] The second feature obtainer **222** may obtain a second feature for estimating DBP based on an second area of the second interval obtained by the second area obtainer **212**. As in the first feature obtainer **221**, the second feature obtainer **222** may obtain the second area of the second interval as it is as the second feature, or may obtain the second feature by normalizing the second area of the second interval by using a second reference value. In this case, the second reference value may be the same as or different from the first reference value, and may include a pulse waveform component in the diastolic interval, i.e., an amplitude value of the position of the third pulse waveform component, an amplitude value of the position of the fourth pulse waveform component, an amplitude value of an internally dividing point between the position of the third pulse waveform component and the position of the fourth pulse waveform component, a maximum amplitude value in the diastolic interval, and the like. However, the second reference value is not limited thereto.

[0078] The blood pressure estimator **230** may estimate blood pressure by using the first feature and the second feature obtained by the feature obtainer **220**. The blood

pressure estimator **230** may independently estimate SBP and DBP. To this end, the blood pressure estimator **230** may include an SBP estimator **231** and a DBP estimator **232**.

[0079] The SBP estimator **231** may estimate SBP based on the first feature. For example, the SBP estimator **231** may estimate SBP by applying an SBP estimation model as represented by the following Equation 5

$$SBP = a_{sys} \times f_1 + b_{sys} \quad [\text{Equation 5}]$$

[0080] Herein, the SBP denotes systolic blood pressure; a_{sys} and b_{sys} denote predetermined coefficients for estimating SBP; and f_1 denotes the first feature.

[0081] The DBP estimator **232** may estimate DBP based on the second feature. For example, the DBP estimator **232** may estimate DBP by applying a DBP estimation model as represented by the following Equation 6.

$$DBP = a_{dia} \times f_2 + b_{dia} \quad [\text{Equation 6}]$$

[0082] Herein, the DBP denotes diastolic blood pressure; a_{dia} and b_{dia} denote predetermined coefficients for estimating DBP; and f_2 denotes the second feature. In this case, the coefficients for estimating SBP may be the same as or different from the coefficients for estimating DBP.

[0083] Referring to FIG. 2B, a processor **200b** includes an area obtainer **210**, a feature obtainer **220**, and a blood pressure estimator **230**. In the embodiment, the feature obtainer **220** may further include a third feature obtainer **223** as illustrated in FIG. 2B. Detailed description of components described above with reference to FIG. 2A will be omitted.

[0084] The third feature obtainer **223** may obtain an additional third feature, which may be applied collectively for estimating both SBP and DBP, from a bio-signal. For example, the third feature obtainer **223** may obtain a third feature from the bio-signal by combining one or two or more of the following: a shape of a bio-signal waveform, a time value and an amplitude value of a maximum point, a time value and an amplitude value of a minimum point, a time value and an amplitude value of a position of the pulse waveform component present in the bio-signal, and an third area of a third interval of the bio-signal. However, the third feature is not limited thereto, and the third feature obtainer **223** may obtain the third feature by combining the first feature and the second feature by, for example, multiplying the first feature by the second feature. In this case, the first feature and the second feature may be combined in various manners, such as addition, subtraction, division, multiplication, logarithmic value, regression equation, and a combination thereof, with no specific limitation.

[0085] The following Equation 7 represents an example of obtaining the third feature by combining bio-signal amplitude values P_1 , P_2 , and P_3 corresponding to positions of the first, the second, and the third pulse waveform components.

$$f_3 = \frac{P_1 + P_2}{P_3} \quad [\text{Equation 7}]$$

[0086] The SBP estimator **231** may combine the first feature, which is obtained by the first feature obtainer **221**, and the third feature which is obtained by the third feature obtainer **223**, and may estimate SBP by applying a blood pressure estimation model represented by the following Equation 8.

$$SBP = a_{sys} \times (w_1 f_1 + w_3 f_3) + b_{sys} \quad [\text{Equation 8}]$$

[0087] Herein, the SBP denotes systolic blood pressure; a_{sys} and b_{sys} denote predetermined coefficients for estimating SBP; f_1 and f_3 denote the first and the third features respectively; and w_1 and w_3 denote weighted values applied to the first and the third features respectively. The weighted values may not be applied as needed, and may be applied to any one of the features.

[0088] The DBP estimator 232 may combine the second feature, which is obtained by the second feature obtainer 222, and the third feature which is obtained by the third feature obtainer 223, and may estimate SBP by applying a blood pressure estimation model such as the following Equation 9.

$$DBP = a_{dia} \times (w_2 f_2 + w_3 f_3) + b_{dia} \quad [\text{Equation 9}]$$

[0089] Herein, the DBP denotes diastolic blood pressure; a_{dia} and b_{dia} denote predetermined coefficients for estimating DBP; f_2 and f_3 denote the first and the third features respectively; and w_2 and w_3 denote weighted values applied to the second and the third features respectively. The weighted values may not be applied as needed, and may be applied to any one of the features.

[0090] FIG. 4 is a flowchart illustrating a blood pressure estimating method according to an example embodiment. The embodiment of FIG. 4 may be an example of a blood pressure estimating method performed by the blood pressure estimating apparatuses 100a and 100b.

[0091] The blood pressure estimating apparatus may receive a request for estimating blood pressure in operation 410. The blood pressure estimating apparatus may provide an interface for interaction with a user. The user may request estimation of blood pressure through the interface provided by the blood pressure estimating apparatus. Alternatively, the blood pressure estimating apparatus may receive the request for estimating blood pressure from an external device. In particular, the request for estimating blood pressure may include a request for providing an estimation result of blood pressure. In the case where the external device includes a blood pressure estimation algorithm, the request for estimating blood pressure may include a request for providing obtained features. Examples of the external device may include a smartphone, a tablet PC, a laptop computer, a wearable device, and the like.

[0092] Then, the blood pressure estimating apparatus may measure a bio-signal in operation 420. The blood pressure estimating apparatus may control a sensor to measure a bio-signal, including a pulse wave signal, from an object.

[0093] Subsequently, the blood pressure estimating apparatus may analyze the bio-signal to obtain a first area of a first interval and a second area of a second interval of the bio-signal in operations 431 and 432. In this case, the first interval for estimating SBP and the second interval for estimating DBP may be intervals in different ranges on a time axis of a bio-signal. For example, the blood pressure estimating apparatus may obtain the first area and the second area by integrating differences between an amplitude value of each time, ranging from a start time to an end time of each interval, and an amplitude value of a baseline. In this case, the start time and the end time of each interval may be determined by using a reference starting value and a reference ending value, which are predefined for each interval, a bio-signal period, and the like. Further, the reference starting value and the reference ending value may be adaptively adjusted by analyzing the shape of a bio-signal. Altern-

tively, the start time and the end time of each interval may also be determined by using time values associated with pulse waveform components present in the bio-signal.

[0094] Then, the blood pressure estimating apparatus may obtain a first feature based on the first area of the first interval in operation 441, and may obtain the second feature based on the second area of the second interval in operation 442. For example, the blood pressure estimating apparatus may obtain the first feature by normalizing the first area of the first interval using a first reference value, and may obtain the second feature by normalizing the second area of the second interval using a second reference value. In this case, the first reference value and the second reference value may be different from each other, and may be determined by combining amplitude values of the positions of the pulse waveform components, an amplitude value of the bio-signal at the internally dividing point of the positions of the pulse waveform components, and the like.

[0095] Subsequently, the blood pressure estimating apparatus may estimate SBP based on the first feature in operation 451, and may estimate DBP based on the second feature in operation 452. As described above, the blood pressure estimating apparatus may estimate blood pressure by inputting each feature into a blood pressure estimation model. In this case, the blood pressure estimation model may be defined differently for SBP and DBP.

[0096] Next, the blood pressure estimating apparatus may output an estimation result of blood pressure in operation 460. For example, the blood pressure estimating apparatus may use various visual methods to output the estimation result of blood pressure through a visual output device such as a display and the like. Alternatively, the blood pressure estimating apparatus may output the estimation result of blood pressure in a non-visual manner through voice, vibration, tactile sensation, and the like, by using a speaker and/or a haptic motor. Further, the blood pressure estimating apparatus may determine a user's health condition based on estimated bio-information, and may provide guide information, such as a warning, a response action, and the like, based on the determination.

[0097] FIG. 5 is a flowchart illustrating a blood pressure estimating method according to another example embodiment. The embodiment of FIG. 5 may be an example of a blood pressure estimating method performed by the blood pressure estimating apparatuses 100a and 100b of FIGS. 1A and 1B.

[0098] Upon receiving a request for estimating blood pressure in operation 510, the blood pressure estimating apparatus may measure a bio-signal in operation 520.

[0099] Then, the blood pressure estimating apparatus may analyze the bio-signal to obtain a first area of a first interval for estimating SBP and a second area of a second interval for estimating DBP in operations 531 and 532.

[0100] Subsequently, the blood pressure estimating apparatus may obtain a first feature based on the first area of the first interval in operation 541 and may obtain a second feature based on the second area of the second interval in operation 542. Further, the blood pressure estimating apparatus may analyze the bio-signal, and may obtain a third feature which may be applied collectively for estimating both SBP and DBP in operation 543. In this case, the blood pressure estimating apparatus may obtain the third feature by combining the first feature and the second feature, and by

combining time or amplitude values of positions of pulse waveform components present in the bio-signal.

[0101] Next, the blood pressure estimating apparatus may estimate SBP based on the first feature and the third feature in operation 551, and may estimate DBP based on the second feature and the third feature in operation 552. In this case, a predefined blood pressure estimation model may be applied, which may be defined differently for SBP and DBP.

[0102] Then, the blood pressure estimating apparatus may output an estimation result of blood pressure to a user by using various visual/non-visual devices and methods in operation 560.

[0103] FIGS. 6A and 6B are diagrams illustrating a wearable device according to an example embodiment. Various embodiments of the above-described blood pressure estimating apparatuses 100a and 100b may be embedded in a smart watch worn on the wrist or a smart band-type wearable device. However, this is merely examples for convenience of explanation, and the blood pressure estimating apparatuses 100a and 100b may be applied to an information processing terminal such as a smartphone, a tablet PC, a laptop computer, a desktop computer, and the like.

[0104] Referring to FIGS. 6A and 6B, the wearable device 600 includes a main body 610 and a strap 630.

[0105] The main body 610 may be formed to have various shapes, and may include modules which are mounted inside or outside of the main body 610 to perform the aforementioned function of estimating blood pressure as well as various other functions. A battery may be embedded in the main body 610 or the strap 630 to supply power to various modules of the wearable device 600.

[0106] The strap 630 may be connected to the main body 610. The strap 630 may be flexible, so as to be bent around a user's wrist. The strap 630 may be bent in a manner that allows the strap 630 to be detached from the user's wrist or may be formed as a band that is not detachable. Air may be injected into the strap 630 or an airbag may be included in the strap 630, so that the strap 630 may have elasticity according to a change in pressure applied to the wrist, and the change in pressure of the wrist may be transmitted to the main body 610.

[0107] The main body 610 may include a sensor 620 for measuring a bio-signal. The sensor 620 may be mounted on a rear surface of the main body 610, which comes into contact with a user's wrist, and may include a light source for emitting light onto the skin of the wrist and a detector for detecting light scattered or reflected from the object. The sensor 620 may further include a contact pressure sensor for measuring contact pressure of the object.

[0108] A processor may be mounted in the main body 610. The processor may be electrically connected to various modules, mounted in the wearable device 600, to control operations thereof. In addition, the processor may estimate blood pressure by using the bio-signal measured by the sensor 620. As described above, the processor may obtain an area for estimating SBP and an area for estimating DBP from the bio-signal, and may independently estimate SBP and DBP based on each of the areas.

[0109] In the case where the sensor 620 includes a contact pressure sensor, the processor may monitor a contact state of the object based on contact pressure between the wrist and the sensor 620, and may provide a user with a guide to a contact position and/or a contact state through a display.

[0110] Further, the main body 610 may include a storage which stores a processing result of the processor and various types of information. In this case, various types of information may include criteria for estimating blood pressure as well as information associated with functions of the wearable device 600.

[0111] In addition, the main body 610 may also include a manipulator 640 which receives a control command of a user and transmits the received control command to the processor. The manipulator 640 may include a power button to input a command to turn on/off the wearable device 600.

[0112] With reference to FIGS. 6A and 6B, a display 614 may be mounted on a front surface of the main body 610, and may include a touch panel for touch input. The display 614 may receive a touch input from a user, may transmit the received touch input to the processor, and may display a processing result of the processor.

[0113] For example, the display 614 may display the estimated blood pressure information. In this case, the display 614 may also display additional information such as a blood pressure estimation date, a health condition, and the like. When a user requests detailed information by operating the manipulator 640 or by touching the display 614 for touch input, the display 614 may output detailed information in various manners.

[0114] Referring to FIG. 6B, the display 614 may output detailed information in a first region 614a of the display 614 and may output a blood pressure history graph in a second region 614b of the display 614. In this case, the blood pressure history graph may include an object (e.g., a figure such as a circle, a square, etc.) which indicates a blood pressure estimation time. Further, an identification mark M, indicating an object I currently selected by a user, may be displayed on the blood pressure history graph. The identification mark M is shown as a vertical line, but is not limited thereto, and may have various shapes, such as a polygonal shape including a circular shape, a square shape, and the like, an arrow indicating a position, and the like. Once the blood pressure history graph is displayed in the second region 614b, detailed information may be output in the first region 614a when a user touches the object I of a specific time, or moves the graph to the right and left sides to place the object I of the specific time on the identification mark M. In this case, information, such as an estimated blood pressure value, a measurement date, a health condition at the time point, and the like, may be output in the first region 614a. However, the information output in the first region 614a is not limited thereto.

[0115] Moreover, a communication interface, provided for communication with an external device, such as a mobile terminal of a user, may be mounted in the main body 610. The communication interface may transmit an estimation result of bio-information to an external device, e.g., a user's smartphone, to display the result to a user. However, this is merely an example, and the communication interface may transmit and receive various necessary information.

[0116] FIG. 7 is a diagram illustrating a smart device including a blood pressure estimating apparatus according to an example embodiment. Examples of the smart device may include a smartphone, a tablet PC, and the like.

[0117] Referring to FIG. 7, the smart device 700 includes a sensor part 730 mounted on one surface of a main body 710. The sensor part 730 may include a pulse wave sensor which includes one or more light sources 731 and a detector

732. As illustrated in FIG. 7, the sensor part **730** may be mounted on a rear surface of the main body **710**, but is not limited thereto. Further, the sensor part **730** may be configured in combination with a fingerprint sensor or a touch panel mounted on a front surface.

[0118] In addition, a display may be mounted on a front surface of the main body **710**. The display may visually display an estimation result of bio-information and the like. The display may include a touch panel, and may receive various types of information input through the touch panel and transmit the received information to the processor.

[0119] Moreover, an image sensor **720** may be mounted in the main body **710**. The image sensor **720** may embodied as a camera and/or a fingerprint scanner. When a user's finger approaches the sensor part **730** to measure a pulse wave signal, the image sensor **720** may capture an image of the finger and may transmit the captured image to the processor. In particular, based on the image of the finger, the processor may identify a relative position of the finger with respect to an actual position of the sensor part **730**, and may provide the relative position of the finger to the user through the display, so that pulse wave signals may be measured with improved accuracy.

[0120] The processor may independently estimate SBP and DBP by using the bio-signal measured by the sensor part **730**. As described above in detail, the processor may obtain features, related to areas for estimating each of SBP and DBP, from the bio-signal, and may estimate SBP and DBP by using the obtained features related to the areas. Various modules for performing various other functions may be mounted in the smart device **700**, and detailed description thereof will be omitted.

[0121] While not restricted thereto, an example embodiment can be embodied as computer-readable code on a computer-readable recording medium. The computer-readable recording medium is any data storage device that can store data that can be thereafter read by a computer system. Examples of the computer-readable recording medium include read-only memory (ROM), random-access memory (RAM), CD-ROMs, magnetic tapes, floppy disks, and optical data storage devices. The computer-readable recording medium can also be distributed over network-coupled computer systems so that the computer-readable code is stored and executed in a distributed fashion. Also, an example embodiment may be written as a computer program transmitted over a computer-readable transmission medium, such as a carrier wave, and received and implemented in general-use or special-purpose digital computers that execute the programs. Moreover, it is understood that in example embodiments, one or more units of the above-described apparatuses and devices can include circuitry, a processor, a microprocessor, etc., and may execute a computer program stored in a computer-readable medium.

[0122] The foregoing example embodiments are not to be construed as limiting. The present teaching can be readily applied to other types of apparatuses. Also, the description of the example embodiments is intended to be illustrative, and not to limit the scope of the claims, and many alternatives, modifications, and variations will be apparent to those skilled in the art.

What is claimed is:

1. A blood pressure estimating apparatus comprising:
a sensor configured to measure a bio-signal waveform;
and

a processor configured to obtain a first feature based on a first area under the bio-signal waveform in a first time interval, obtain a second feature based on a second area under the bio-signal waveform in a second time interval which is different from the first time interval, and estimate a blood pressure based on the first feature and the second feature.

2. The blood pressure estimating apparatus of claim 1, wherein the processor is further configured to estimate a systolic blood pressure (SBP) and a diastolic blood pressure (DBP) by applying a blood pressure estimation model to each of the first feature and the second feature.

3. The blood pressure estimating apparatus of claim 1, wherein the processor is further configured to determine a start time and an end time of each of the first time interval and the second time interval based on a bio-signal period of the bio-signal waveform, and a reference starting value and a reference ending value of each of the first time interval and the second time interval.

4. The blood pressure estimating apparatus of claim 3, wherein the reference starting value and the reference ending value of each of the first time interval and the second time interval comprise at least one of a constant value and a variable value which is adaptively changed according to a shape of the bio-signal waveform.

5. The blood pressure estimating apparatus of claim 3, wherein the processor is further configured to obtain the first area of the first time interval and the second area of the second time interval by integrating differences between an amplitude value of the bio-signal waveform and an amplitude value of a baseline of the bio-signal waveform, for the first time interval and the second time interval, respectively.

6. The blood pressure estimating apparatus of claim 5, wherein the amplitude value of the baseline comprises at least one of an amplitude value of the bio-signal waveform at a measurement start time of the bio-signal waveform, a minimum amplitude value of the bio-signal waveform, and an amplitude value of a straight line formed by connecting the start time and the end time of each of the first time interval and the second time interval.

7. The blood pressure estimating apparatus of claim 1, wherein the processor is further configured to determine a start time and an end time of each of the first time interval and the second time interval based on at least one of a start time and an end time of measurement of the bio-signal waveform, a first time of a position of a pulse waveform component present in the bio-signal waveform, a second time of a dicrotic notch position of the bio-signal waveform, times calculated by multiplying the first and the second times by a predetermined coefficient, and times of an internally dividing point obtained by applying a weighted value to the first and the second times.

8. The blood pressure estimating apparatus of claim 7, wherein the processor is further configured to perform differentiation on the bio-signal waveform to obtain a differential signal, and determine a local minimum point of the differential signal to be the position of the pulse waveform component.

9. The blood pressure estimating apparatus of claim 8, wherein the weighted value is determined based on at least one of an amplitude of the differential signal and an amplitude of the bio-signal waveform corresponding to the first and the second times.

10. The blood pressure estimating apparatus of claim **1**, wherein the processor is further to obtain the first feature and the second feature by normalizing the first area and the second area based on a first reference value and a second reference value, respectively.

11. The blood pressure estimating apparatus of claim **10**, wherein the processor is further configured to normalize the first area and the second area by dividing the first area and the second area by the first reference value and the second reference value, respectively, or by dividing the first reference value and the second reference value by the first area and the second area, respectively.

12. The blood pressure estimating apparatus of claim **10**, wherein the processor is further configured to set the first reference value and the second reference value based on an amplitude value of the bio-signal waveform in a systolic phase or a diastolic phase of the bio-signal waveform, an amplitude value of an internally dividing point between two pulse waveform components of the bio-signal waveform, and a maximum amplitude value of the bio-signal waveform in the systolic phase or the diastolic phase.

13. The blood pressure estimating apparatus of claim **1**, wherein the processor is further configured to:

obtain a third feature from the bio-signal waveform based on at least one of a shape of the bio-signal waveform, a time value and an amplitude value of a maximum point of the bio-signal waveform, a time value and an amplitude value of a minimum point of the bio-signal waveform, a time value and an amplitude value of a position of a pulse waveform component present in the bio-signal waveform, and an area under the bio-signal waveform in a third time interval;

estimate a systolic blood pressure based on the first feature and the third feature; and

estimate a diastolic blood pressure based on the second feature and the third feature.

14. A blood pressure estimating method comprising:
measuring a bio-signal waveform;

obtaining a first area under the bio-signal waveform in a first time interval and a second area under the bio-signal waveform in a second time interval which is different from the first time interval;

obtaining a first feature and a second feature based on the first area and the second area, respectively; and

estimating a blood pressure based on the first feature and the second feature.

15. The blood pressure estimating method of claim **14**, wherein the estimating the blood pressure comprises estimating a systolic blood pressure (SBP) and a diastolic blood pressure (DBP) by applying a blood pressure estimation model to each of the first feature and the second feature.

16. The blood pressure estimating method of claim **14**, wherein the obtaining the first area and the second area comprises determining a start time and an end time of each of the first time interval and the second time interval based on a bio-signal period of the bio-signal waveform, and a reference starting value and a reference ending value of each of the first time interval and the second time interval.

17. The blood pressure estimating method of claim **16**, wherein the reference starting value and the reference ending value of each of the first time interval and the second time interval comprise at least one of a constant value and a variable value which is adaptively changed according to a shape of the bio-signal waveform.

18. The blood pressure estimating method of claim **16**, wherein the obtaining the first area and the second area comprises obtaining the first area and the second area by integrating differences between an amplitude value of the bio-signal waveform and an amplitude value of a baseline of the bio-signal waveform, for the first time interval and the second time interval, respectively.

19. The blood pressure estimating method of claim **18**, wherein the amplitude value of the baseline comprises at least one of an amplitude value of the bio-signal waveform at a measurement start time of the bio-signal waveform, a minimum amplitude value of the bio-signal waveform, and an amplitude value of a straight line formed by connecting the start time and the end time of each of the first time interval and the second time interval.

20. The blood pressure estimating method of claim **14**, wherein the obtaining the first area and the second area comprises determining a start time and an end time of each of the first time interval and the second time interval based on at least one of a start time and an end time of measurement of the bio-signal waveform, a first time of a position of a pulse waveform component present in the bio-signal waveform, a second time of a dicrotic notch position of the bio-signal waveform, times calculated by multiplying the first and the second times by a predetermined coefficient, and times of an internally dividing point obtained by applying a weighted value to the first and the second times.

21. The blood pressure estimating method of claim **14**, wherein the obtaining the first feature and the second feature comprises normalizing the first area and the second area based on a first reference value and a second reference value, respectively.

22. The blood pressure estimating method of claim **21**, wherein the obtaining the first feature and the second feature comprises setting the first reference value and the second reference value based on at least one of an amplitude value of the bio-signal waveform in a systolic phase or a diastolic phase of the bio-signal waveform, an amplitude value of an internally dividing point between two pulse waveform components of the bio-signal waveform, and a maximum amplitude value of the bio-signal waveform in the systolic phase or the diastolic phase.

23. The blood pressure estimating method of claim **14**, further comprising obtaining a third feature from the bio-signal waveform based on at least on one or two or more of a shape of the bio-signal waveform, a time value and an amplitude value of a maximum point of the bio-signal waveform, a time value and an amplitude value of a minimum point of the bio-signal waveform, a time value and an amplitude value of a position of a pulse waveform component present in the bio-signal waveform, and an area under the bio-signal waveform in a third time interval,

wherein the estimating the blood pressure comprises estimating a systolic blood pressure based on the first feature and the third feature, and estimating a diastolic blood pressure based on the second feature and the third feature.

24. A non-transitory computer readable storage medium storing a program that is executable by a computer to perform a method comprising:

obtaining a bio-signal waveform;

setting a systolic time interval and a diastolic time interval of the bio-signal waveform to be different from each other;

calculating a first area under the bio-signal waveform in the systolic time interval, and a second area under the bio-signal waveform in the diastolic time interval; and estimating a blood pressure based on the first area in the systolic time interval and the second area in the diastolic time interval.

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专利名称(译)	估计血压的装置和方法		
公开(公告)号	US20200113453A1	公开(公告)日	2020-04-16
申请号	US16/374327	申请日	2019-04-03
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IPC分类号	A61B5/021 A61B5/00		
CPC分类号	A61B5/7242 A61B5/0002 A61B5/02108 A61B5/02255 A61B5/681 A61B5/02141 A61B5/742 A61B5/02116 A61B5/02416 A61B5/6898 A61B5/7235		
优先权	1020180120640 2018-10-10 KR		
外部链接	Espacenet USPTO		

摘要(译)

提供一种血压估计装置。该血压估计装置可以包括：传感器，被配置为测量生物信号波形；以及处理器，用于在第一时间间隔内根据所述生物信号波形下的第一面积获得第一特征，在第二时间间隔内根据所述生物信号波形下的第二面积获得第二特征。第一时间间隔，并根据第一特征和第二特征估算血压。

