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(54) **BLOOD PRESSURE MEASUREMENT
DEVICE HAVING FUNCTION OF
DETERMINING REST CONDITION OF
PATIENT**

USPC 600/300, 301, 481, 483–485, 500, 501,
600/509, 513
See application file for complete search history.

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(57) **ABSTRACT**

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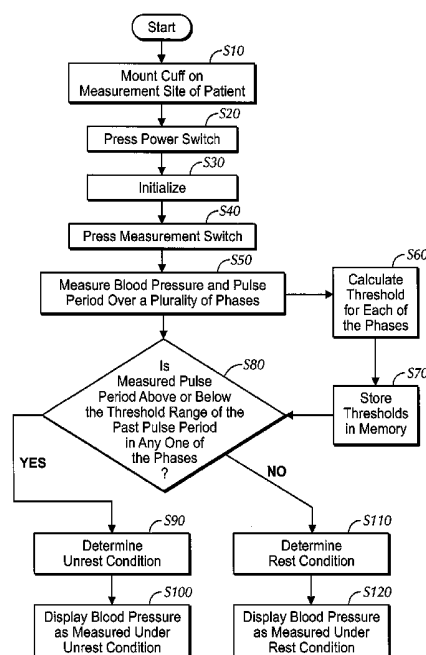
(51) **Int. Cl.**
A61B 5/09 (2006.01)
A61B 5/024 (2006.01)
A61B 5/022 (2006.01)
A61B 5/00 (2006.01)

A method includes pressurizing a cuff of a blood pressure measurement device that is wrapped around a measurement site of a patient, measuring the patient's blood pressure and pulse period during and before depressurization of the cuff, calculating a threshold value range of the pulse period based on data of a past measured pulse period of the patient, determining whether the measured pulse period is within the threshold value range of the past measured pulse period, if the measured pulse period is determined to be within the threshold value range of the past measured pulse period both during and before depressurization, displaying that the patient was in the rest condition, and if the measured pulse period is determined not to be within the threshold value range of the past measured pulse period either during or before depressurization, displaying that the patient was in the unrest condition.

(52) **U.S. Cl.**
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(2013.01); **A61B 5/024** (2013.01); **A61B 5/742**
(2013.01)

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CPC A61B 5/022; A61B 5/024; A61B 5/742;
A61B 5/7282; A61B 18/22; A61B 18/2277;
G02B 6/262; G02B 6/0283; G02B 6/06327

10 Claims, 7 Drawing Sheets



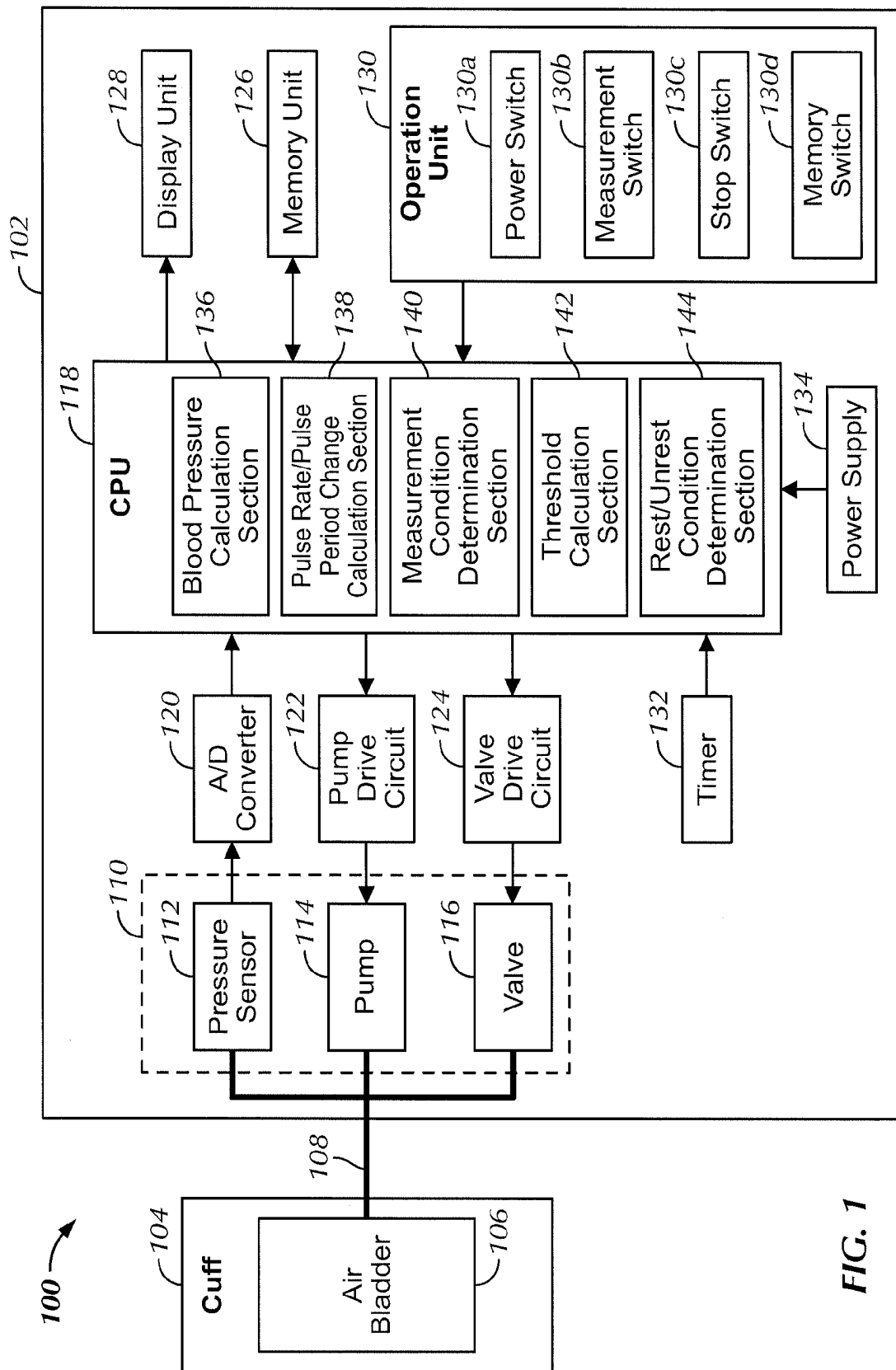


FIG. 1

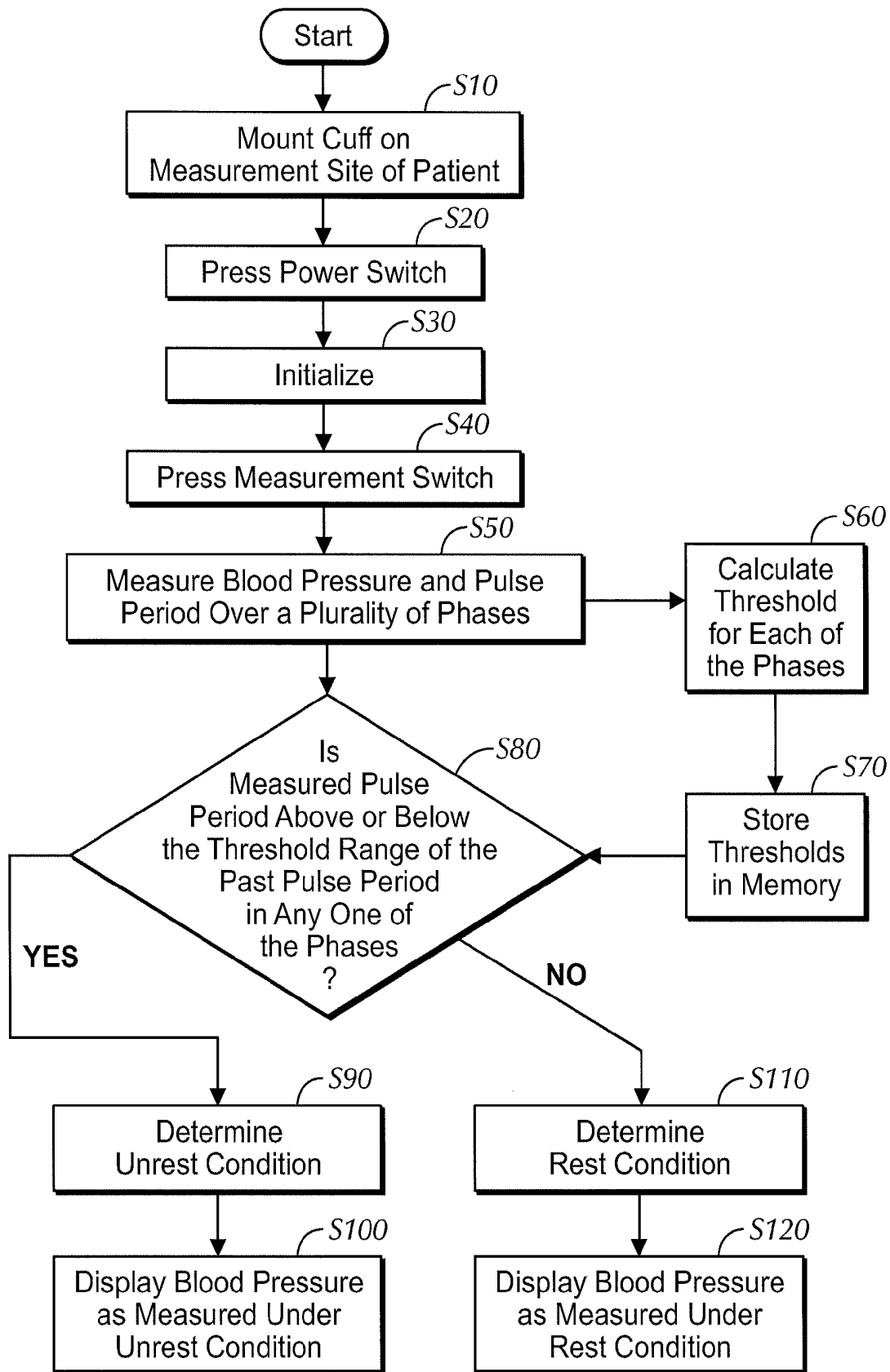


FIG. 2

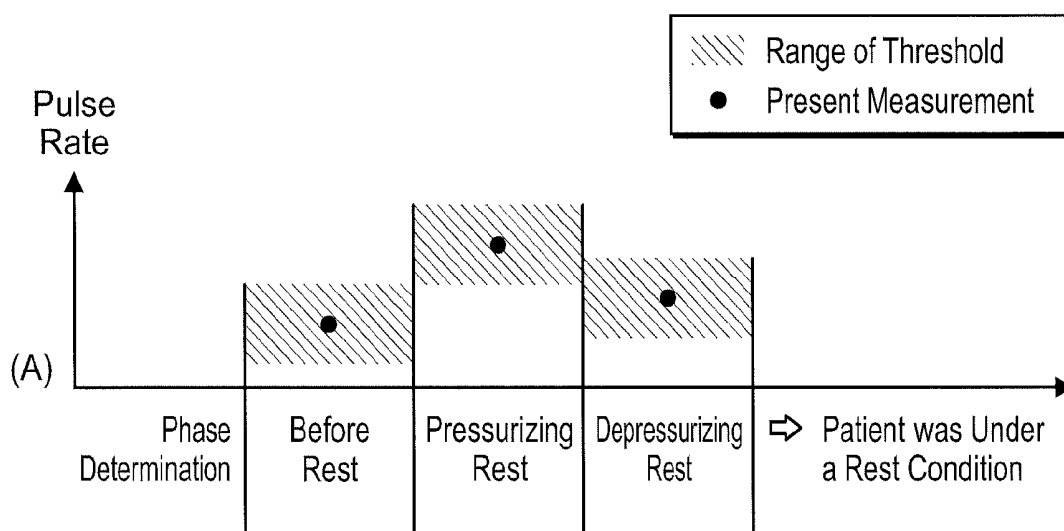


FIG. 3A

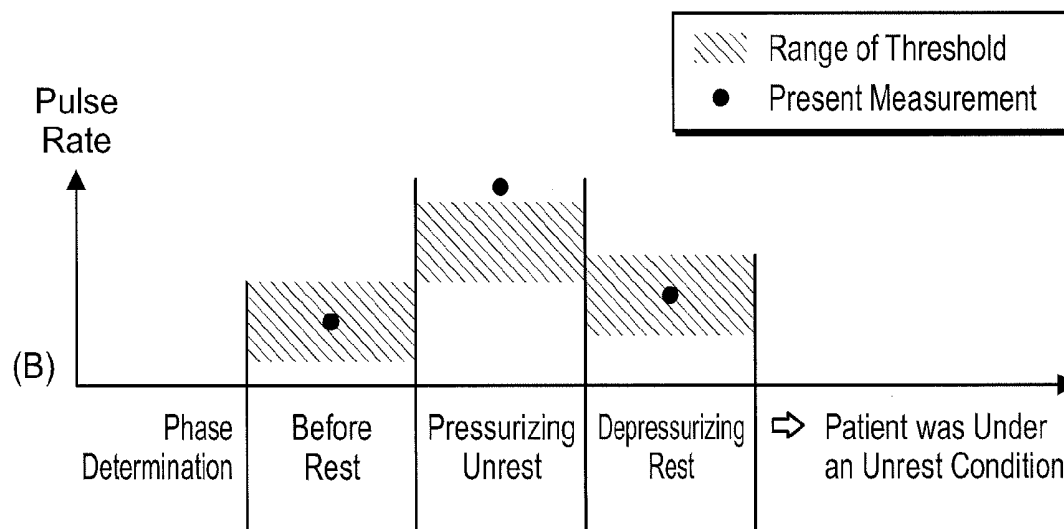


FIG. 3B

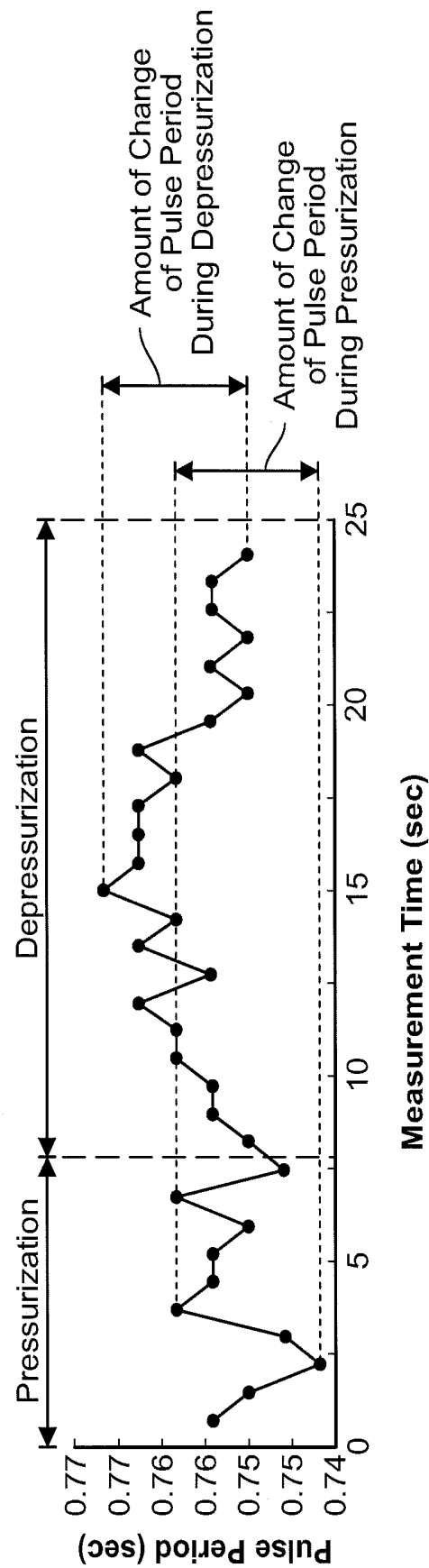
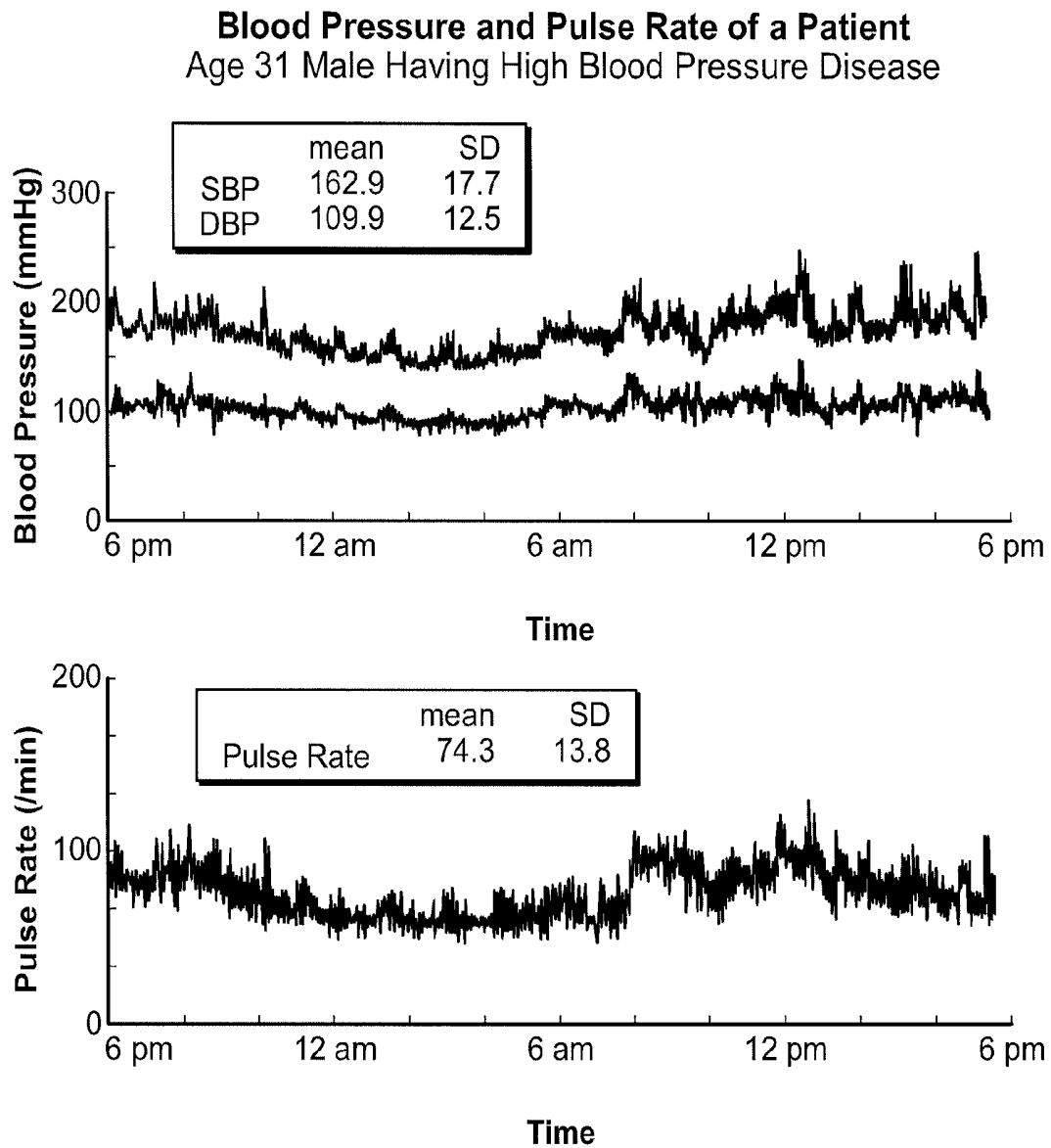
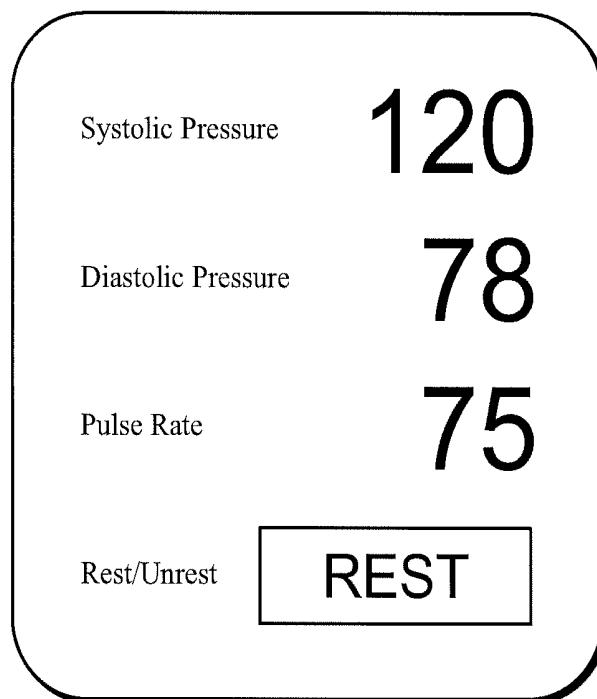


FIG. 4A

	Time Lapse	Pulse Period
Pressurization Phase	0.725	0.754167
	1.479167	0.75
	2.229167	0.741667
	2.970833	0.745833
	3.716667	0.754167
	4.475	0.754167
	5.229167	0.754167
	5.983333	0.75
	6.733333	0.758333
	7.491667	0.745833
Depressurization Phase	8.2375	0.75
	8.9875	0.754167
	9.741667	0.754167
	10.49583	0.758333
	11.25417	0.758333
	12.0125	0.7625
	12.775	0.754167
	13.52917	0.7625
	14.29167	0.758333
	15.05	0.766667
	15.81667	0.7625
	16.57917	0.7625
	17.34167	0.7625
	18.10417	0.758333
	18.8625	0.7625
	19.625	0.754167
	20.37917	0.75
	21.12917	0.754167
	21.88333	0.75
	22.63333	0.754167
	23.3875	0.754167
	24.14167	0.75
	24.89167	0

FIG. 4B

**FIG. 5**

*FIG. 6*

BLOOD PRESSURE MEASUREMENT DEVICE HAVING FUNCTION OF DETERMINING REST CONDITION OF PATIENT

BACKGROUND

(1) Field of the Invention

The present invention relates to a blood pressure measurement device and methods for measuring blood pressure. More specifically, the present invention relates to apparatus and methods for determining whether a patient was in a rest condition during blood pressure measurement.

(2) Description of Related Art

A rest condition of a patient is important when measuring blood pressure. If the patient is not in a rest condition, the measured blood pressure may differ from the value measured in the rest condition. Thus, it is necessary to know whether the patient was in the rest condition when the blood pressure was being measured.

As a measure to determine whether a patient was in a rest condition, measuring pulse rate and pulse wave period is known. Generally, the patient's pulse wave period becomes short or unstable after hard exercise or mental distress. Therefore, it is possible to determine whether the patient was in the rest condition by examining whether the patient's pulse wave period was stable during a certain period of time.

Rossmax Hemodynamic Stability Determination (HSD) discloses that when a patient is not in a rest condition due to mental or physical load, his or her pulse period may become short and/or unstable, which may cause blood pressure measurement error.

JPH2012-120206 discloses the relationship between an increase of pulse rate and pulse period change at the time when a patient received stress.

However, because most blood pressure devices require use of a cuff to be mounted on an arm of the patient, some patients may feel stress during blood pressure measurement. In that event, the pulse period of the patient may be significantly affected by the stress that was caused by the cuff. Therefore, even if the patient was a under rest condition before measurement, the pulse period measured by using the blood pressure measurement device may become different from the pulse period of the patient in the rest condition.

SUMMARY OF INVENTION

Therefore, a blood pressure measurement device or method according to one or more embodiments of the present invention accurately determines whether a patient was in a rest condition during blood pressure measurement while taking into account (or regardless of) a change of the pulse period that may be caused by use of a cuff of the blood pressure measurement device.

According to one or more embodiments of the present invention, during measurement of blood pressure, the pulse period of the patient is measured over a plurality of phases: before, during and after pressurization of the cuff. The pulse periods measured during those phases are each compared with the pulse period of past measurement(s) of corresponding phases.

According to one or more embodiments of the present invention, a comparison of the pulse period may be made in terms of: a time period between pulse waves (pulse rate), a change of time period between the pulse waves (pulse rate variability), or both.

According to one or more embodiments of the present invention, pulse rate in a phase may be calculated by the time period of the phase divided by the number of pulses during the phase. According to one or more embodiments of the present invention, a change of pulse period in a phase may be calculated by the difference between a maximum time period of a pulse and a minimum time period of a pulse during the phase.

According to one or more embodiments of the present invention, a threshold value range for comparing a present measurement with past measurement(s) may be calculated by using a pulse period (time period of a pulse wave and/or change of pulse period) measured in the past measurement(s) of the same patient during the phases.

According to one or more embodiments of the present invention, if the pulse period (time period of a pulse wave and/or change of pulse period) measured in any one of those phases is beyond or below the threshold value range, the patient is determined to have been under an unrest condition at the time of measurement.

According to one or more embodiments of the present invention, a method of determining a rest or an unrest condition of a patient during measurement of blood pressure of the patient includes: providing a cuff of a blood pressure measurement device that is configured to be wrapped around a measurement site of the patient, and pressurizing the cuff; measuring the blood pressure and a pulse period of the patient during and before depressurization of the cuff; calculating a threshold value range of the pulse period measured during and before depressurization of the cuff based on data of a pulse period measured in a past measurement of the patient and stored in a memory of the blood pressure measurement device; determining whether the pulse period of the patient measured during and before depressurization of the cuff is within the threshold value range of the pulse period stored in the memory; if the pulse period measured during and before depressurization is determined to be within the threshold value range of the pulse period stored in the memory both during and before depressurization, displaying on a display of the blood pressure measurement device that the blood pressure was measured when the patient was in the rest condition, and if the pulse period measured during and before depressurization is determined not to be within the threshold value range of the pulse period stored in the memory either during or before depressurization, displaying on the display of the blood pressure measurement device that the blood pressure was measured when the patient was in the unrest condition.

According to one or more embodiments of the present invention, a blood pressure measurement device that determines a rest or an unrest condition of a patient includes: a cuff for mounting on a measurement site of the patient, wherein the cuff is configured to be pressurized and depressurized to measure a blood pressure and a pulse period of the patient; means for measuring the blood pressure and the pulse period of the patient during and before depressurization of the cuff; a memory that stores data of a pulse period measured in a past measurement of the patient by the measuring means; a threshold calculation section that calculates a threshold value of the pulse period measured during and before depressurization of the cuff based on data of the pulse period measured in the past measurement of the patient; a rest/unrest condition determination section that determines whether the patient was in a rest condition or an unrest condition during blood pressure measurement by comparing the pulse period measured during and before depressurization of the cuff with the threshold value based on data of the pulse period measured in the past measurement of the patient; a display unit that displays the

measured blood pressure and whether the blood pressure was measured during the rest condition or the unrest condition of the patient.

According to one or more embodiments of the present invention, a blood pressure measurement device that determines a rest or an unrest condition of a patient includes: a cuff for mounting on a measurement site of the patient, wherein the cuff includes an air bladder and is configured to be pressurized and depressurized to measure a blood pressure and a pulse period of the patient; a measurement air system comprising a pressure sensor, a pump, and a valve; an air tube that connects the air bladder of the cuff to the measurement air system; a blood pressure calculation section that calculates a blood pressure of the patient based on an internal pressure of the air bladder as detected by the pressure sensor; a heart rate/pulse period change calculation section that calculates a pulse rate of the patient during and before depressurization of the cuff and a pulse period change of the patient during and before depressurization of the cuff based on the measured pulse period of the patient; a memory that stores data of a pulse period measured in a past measurement of the patient; a threshold calculation section that calculates a threshold value of at least one of the pulse rate of the patient calculated during and before depressurization of the cuff and the pulse period change of the patient calculated during and before depressurization of the cuff based on data of the pulse period measured in the past measurement of the patient; a rest/unrest condition determination section that determines whether the patient was in a rest condition or an unrest condition during blood pressure measurement by comparing at least one of the pulse rate of the patient calculated during and before depressurization of the cuff and the pulse period change of the patient calculated during and before depressurization of the cuff with the threshold value based on data of the pulse period measured in the past measurement of the patient; and a display unit that displays the measured blood pressure and whether the blood pressure was measured during the rest condition or the unrest condition of the patient.

BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 is a functional block diagram showing a configuration of a blood pressure measurement device (sphygmomanometer) according to one or more embodiments of the present invention.

FIG. 2 is a flow chart of processing related to the determination of whether a patient was in a rest condition or an unrest condition during blood pressure measurement according to one or more embodiments of the present invention.

FIGS. 3(A) and 3(B) show a comparison of measured pulse periods with past pulse periods in three phases according to one or more embodiments of the present invention.

FIGS. 4(A) and 4(B) show measurement data of pulse periods during pressurization and depressurization phases of the cuff of the blood pressure measurement device (sphygmomanometer) according to one or more embodiments of the present invention.

FIG. 5 shows an example of how blood pressure and pulse rate fluctuate depending on the time zone of measurement.

FIG. 6 is an example display of data on a display unit of a blood pressure measurement device (sphygmomanometer) according to one or more embodiments of the present invention.

DETAILED DESCRIPTION OF INVENTION

Hereinafter, embodiments of a blood pressure measurement device according to one or more embodiments of the

present invention will be described with reference to the drawings. In the following description, the same reference numerals are given to the same components and constituent elements. The names and functions thereof are also the same.

FIG. 1 is a functional block diagram showing a configuration of a blood pressure measurement device (sphygmomanometer) according to one or more embodiments of the present invention.

Referring to FIG. 1, the blood pressure measurement device (sphygmomanometer) 100 is provided with a device main body 102 and a cuff 104. When blood pressure measurement is performed by the sphygmomanometer 100, the cuff 104, which has a belt-like outer shape, is wrapped around a measurement site of the patient. The cuff 104 houses an air bladder 106 serving as a fluid bag for compressing the measurement site.

The cuff 104 and the device main body 102 are connected by an air tube 108 serving as a connecting tube. According to one or more embodiments of the present invention, the air tube 108 consists of a flexible tube. One end of the air tube 108 is connected to a measurement air system 110 that is provided in the device main body 102. The other end of the air tube 108 is connected to the air bladder 106 of the cuff 104.

The measurement air system 110 supplies air to or discharges air from the air bladder 106 contained in the cuff 104 via air tube 108. The measurement air system 110 includes a pressure sensor 112 that detects the pressure inside the air bladder 106, and a pump 114 and a valve 116 for expanding and contracting the air bladder 106. Supplying air to the air bladder 106 pressurizes the cuff 104, while discharging air from the air bladder 106 depressurizes the cuff 104. The blood pressure measurement device (sphygmomanometer) 100 also includes a central processing unit (CPU) 118, an A/D converter 120, a pump drive circuit 122, and a valve drive circuit 124. The CPU 118 controls the entirety of the blood pressure measurement device (sphygmomanometer) 100. The A/D converter 120, the pump drive circuit 122, and the valve drive circuit 124 are connected to the measurement air system 110.

The pressure sensor 112 detects the internal pressure of the air bladder 106 and inputs a detection signal to the A/D converter 120. The input detection signal is converted to a digital signal by the A/D converter 120, and input to the CPU 118. The CPU 118 executes predetermined processing based on the internal pressure of the air bladder 106 obtained from the pressure sensor 112 and outputs control signals to the pump drive circuit 122 and the valve drive circuit 124 in accordance with the result of the predetermined processing.

The device main body 102 of the blood pressure measurement device (sphygmomanometer) includes a memory unit 126, which stores programs executed by the CPU 118 and results of measurement. That is, the memory unit 126 is constituted by a storage medium. Memory unit 126 may be realized by a single storage medium or more than one storage media. Exemplary storage media include media for storing programs in a non-volatile manner such as CD-ROM (Compact Disc-Read Only Memory), DVD-ROM (Digital Versatile Disk-Read Only Memory), USB (Universal Serial Bus) memory, memory card FD (Flexible Disk), hard disk, magnetic tape, cassette tape, MO (Magnetic Optical Disc), MD (MiniDisc), IC (Integrated Circuit) card (excluding memory card), optical card, mask ROM, EPROM, and EEPROM (Electrically Erasable Programmable Read-Only Memory).

The device main body 102 of the blood pressure measurement device (sphygmomanometer) 100 also includes a display unit 128 and an operation unit 130. The display unit 128 displays the results of measuring blood pressure values, pulse

rates, states of rest or unrest, etc. using numerical values, labels, indicia, graphs, and the like in a manner that allows visible confirmation. A liquid crystal panel or the like, for example, may be used as the display unit **128**. According to one or more embodiments of the present invention, the operation unit **130** includes a power switch **130a**, a measurement switch **130b**, a stop switch **130c**, and a memory switch **130d**.

The device main body **102** of the blood pressure measurement device (sphygmomanometer) **100** also includes a timer **132** and a power supply **134**. The timer **132** has a clock function, and the power supply **134** supplies power to the CPU **118**. According to one or more embodiments of the present invention, the power supply **134** supplies power to the CPU **118** from an external power supply. According to other embodiments of the present invention, the power supply **134** may be a battery or similar element that supplies power to the CPU **118** without receiving power from an external power supply.

As previously described, the operation unit **130** includes the power switch **130a**, the measurement switch **130b**, the stop switch **130c**, and the memory switch **130d** according to one or more embodiments of the present invention. The power switch **130a** receives input of an instruction for turning the power supply on or off. The measurement switch **130b** receives a measurement start instruction. The stop switch **130c** receives a measurement stop instruction. Finally, the memory switch **130d** receives an instruction to read out information such as blood pressure recorded in memory unit **126**.

The CPU **118** includes a blood pressure calculation section **136** that calculates blood pressure values (a systolic blood pressure value, a diastolic blood pressure value, and/or an average blood pressure value) based on the internal pressure of the air bladder **106** obtained from the pressure sensor **112**. The CPU **118** outputs the blood pressure values calculated by the blood pressure calculation section **136** to the display unit **128** to display them as a result of measurement.

The CPU **118** also includes a pulse rate/pulse period change calculation section **138** that calculates the pulse rate and pulse period change of a patient based on a measured pulse period of the patient for a particular phase lasting for a period of time. According to one or more embodiments of the present invention, the measured pulse period may be based on a time period of a detected pulse waveform and/or a change of pulse period. According to one or more embodiments of the present invention, the pulse rate/pulse period change calculation section **138** calculates the pulse rate of the patient in a phase based on the time period of the phase divided by the number of pulses during the phase. According to one or more embodiments of the present invention, the pulse rate/pulse period change calculation section **138** calculates the pulse period change of the patient in a phase based on the difference between the maximum time period of a pulse and the minimum time period of a pulse during the phase. The CPU **118** outputs the pulse rate calculated by the pulse rate/pulse period change calculation section **138** to the display unit **128** to display it as a result of measurement.

The CPU **118** also includes a measurement condition determination section **140** that determines which of a plurality of measurement conditions applies during blood pressure measurement. According to one or more embodiments of the present invention, the plurality of measurement conditions are a plurality of phases, which may include before pressurization of the cuff **104**, during pressurization of the cuff **104**, and after pressurization of the cuff **104**.

The CPU **118** also includes a threshold calculation section **142** that calculates a threshold range for the pulse rate and pulse period change calculated by the pulse rate/pulse period

change calculation section **138**, with respect to each of the plurality of measurement conditions (i.e., each of the plurality of phases) determined by the measurement condition determination section **140**. The CPU **118** outputs the threshold ranges calculated by the threshold calculation section **142** to the memory unit **126** for storage.

The CPU **144** also includes a rest/unrest condition determination section **144** that determines whether the patient is in a rest condition or an unrest condition during blood pressure measurement by comparing present pulse rate and/or pulse period change measurements with the threshold range of past pulse rate and/or pulse period change measurements. If it is determined that the present pulse rate and/or pulse period changes are outside (that is, above or below) the threshold range of past pulse rate and/or pulse period change measurements with respect to any of the plurality of phases, then the rest/unrest condition determination section **144** determines that the patient was in an unrest condition during blood pressure measurement. On the other hand, if it is determined that the present pulse rate and/or pulse period changes are within the threshold range of past pulse rate and/or pulse period change measurements with respect to any of the plurality of phases, then the rest/unrest condition determination section **144** determines that the patient was in a rest condition during blood pressure measurement. The CPU **118** outputs the determination of whether the patient was in an unrest or a rest condition during blood pressure measurement as determined by the rest/unrest condition determination section **144** to the display unit **128** to display it as a result of measurement.

Displaying the blood pressure measurement of the patient on the display unit **128** together with the determination of whether the patient was in an unrest or a rest condition during the blood pressure measurement allows the patient to decide whether the measured blood pressure is reliable.

FIG. 2 is a flow chart of processing related to the determination of whether a patient was in a rest condition or an unrest condition during blood pressure measurement according to one or more embodiments of the present invention. In the blood pressure measurement device (sphygmomanometer) **100**, this processing is realized by the CPU **118** executing a program stored in the memory unit **126** (or in a recording medium detachable from the device main body **102**).

Referring to FIG. 2, in the blood pressure measurement processing, at step S10, the cuff **104** of the blood pressure measurement device (sphygmomanometer) **100** is mounted on a measurement site of the patient.

At step S20, the CPU **118** stands by until the power switch **130a** is operated, and advances the processing to step S30 when it is judged that the power switch **130a** has been operated.

At step S30, the CPU **118** initializes the blood pressure measurement device (sphygmomanometer) **100**. The internal pressure of the air bladder **106** of the cuff **104** is thereby initialized. After initialization, the CPU **118** advances the processing to step S40.

At step S40, the CPU **118** stands by until the measurement switch **130b** is operated. When it is judged that the measurement switch **130b** has been operated, the CPU **118** advances the processing to step S50.

At step S50, the CPU **118** performs the processing to measure the blood pressure and pulse period of the patient over a plurality of phases. According to one or more embodiments of the present invention, the blood pressure may be a systolic blood pressure value, a diastolic blood pressure value, and/or an average blood pressure value. According to one or more embodiments of the present invention, the measured pulse period may be used to calculate a time period of

a detected pulse waveform, that is, the pulse rate, and/or a change of pulse period. According to one or more embodiments of the present invention, the plurality of phases may include before pressurization of the cuff 104, during pressurization of the cuff 104, and after pressurization of the cuff 104. The CPU 118 then advances the processing to step S60.

At step S60, the CPU 118 uses the blood pressure and pulse period of the patient over the plurality of phases as measured in step S50 to perform the processing to calculate a threshold value of the pulse rate and pulse period change of the patient with respect to each of the plurality of phases. According to one or more embodiments of the present invention, the threshold value may be a range of threshold values. Examples of threshold value ranges of the pulse rate of a patient with respect to each of a plurality of phases are shown in FIGS. 3A and 3B, which are further described below. The threshold value ranges are shown with shading.

According to one or more embodiments of the present invention, at step S60, the CPU 118 performs the processing to calculate a threshold value range of the pulse rate of the patient with respect to a plurality of phases as further described below. As previously described, the CPU 118 performs the processing to measure the pulse period of the patient over a plurality of phases in step S50. At step S60, the CPU 118 uses the measured pulse period of the patient with respect to a particular phase from step S50 to calculate the pulse rate of the patient. According to one or more embodiments of the present invention, the CPU 118 calculates the pulse rate of the patient with respect to a particular phase by dividing the time length of the phase by the number of pulses that occurred during the phase. As previously described, the plurality of phases may include before pressurization of the cuff 104, during pressurization of the cuff 104, and after pressurization of the cuff 104, according to one or more embodiments of the present invention. As such, at step S60, the CPU 118 calculates the pulse rate of the patient with respect to each of the following phases: a phase that occurs before pressurization of the cuff 104, a phase that occurs during pressurization of the cuff 104, and a phase that occurs after pressurization of the cuff 104, according to one or more embodiments of the present invention.

According to one or more embodiments of the present invention, at step S60, the CPU 118 uses the calculated pulse rates of the patient with respect to each of the plurality of phases to calculate a pulse rate threshold range with respect to each of the plurality of phases. The pulse rate threshold range is delimited by an upper end threshold value and a lower end threshold value. According to one or more embodiments of the present invention, the upper end threshold value and the lower end threshold value for the pulse rate threshold range with respect to each of the plurality of phases may be calculated using Equation 1 and Equation 2, as shown below.

$$\text{Upper end of threshold value} = \text{pulse rate during a phase} \times (1 + \alpha) \quad \text{Equation 1.}$$

$$\text{Lower end of threshold value} = \text{pulse rate during a phase} \times (1 - \alpha) \quad \text{Equation 2.}$$

According to one or more embodiments of the present invention, “ α ,” as used in Equation 1 and Equation 2, may be set discretionarily in view of a tendency of the blood pressure value and the pulse rate of the patient. Referring now to FIG. 5, an example of how blood pressure and pulse rate fluctuate depending on the time zone of measurement is shown for a particular patient. As shown in FIG. 5, the patient’s fluctuation of pulse rate tends to be larger during the daytime zone, as compared with the night time zone. Therefore, if the blood

pressure measurement was conducted during the daytime time zone, α of Equation 1 and Equation 2 may be set to be 0.15. If the blood pressure measurement was conducted during the night time zone, α of Equation 1 and Equation 2 may be set to be 0.10.

Referring back to FIG. 2, according to one or more embodiments of the present invention, at step S60, the CPU 118 performs the processing to calculate a threshold value range of the pulse period change of the patient with respect to a plurality of phases as further described below. As previously described, the CPU 118 performs the processing to measure the pulse period of the patient over a plurality of phases in step S50. At step S60, the CPU 118 uses the measured pulse period of the patient with respect to a particular phase from step S50 to calculate the pulse period change of the patient. According to one or more embodiments of the present invention, the CPU 118 calculates the pulse period change of the patient with respect to a particular phase based on a difference between a maximum pulse period and a minimum pulse period that occurred during the phase. As previously described, the plurality of phases may include before pressurization of the cuff 104, during pressurization of the cuff 104, and after pressurization of the cuff 104, according to one or more embodiments of the present invention. As such, at step S60, the CPU 118 calculates the pulse period of the patient with respect to each of the following phases: a phase that occurs before pressurization of the cuff 104, a phase that occurs during pressurization of the cuff 104, and a phase that occurs after pressurization of the cuff 104, according to one or more embodiments of the present invention.

According to one or more embodiments of the present invention, at step S60, the CPU 118 uses the calculated pulse period changes of the patient with respect to each of the plurality of phases to calculate a pulse period change threshold range with respect to each of the plurality of phases. The pulse period change threshold range is delimited by an upper end threshold value and a lower end threshold value. According to one or more embodiments of the present invention, the upper end threshold value and the lower end threshold value for the pulse period change threshold range with respect to each of the plurality of phases may be calculated using Equation 3 and Equation 4, as shown below.

$$\text{Upper end of threshold value} = \text{pulse period change during a phase} \times (1 + \alpha) \quad \text{Equation 3.}$$

$$\text{Lower end of threshold value} = \text{pulse period change during a phase} \times (1 - \alpha) \quad \text{Equation 4.}$$

According to one or more embodiments of the present invention, “ α ,” as used in Equation 3 and Equation 4, may be set discretionarily in view of a tendency of the blood pressure value and the pulse rate of the patient. Referring now to FIG. 5, an example of how blood pressure and pulse rate fluctuate depending on the time zone of measurement is shown for a particular patient. As shown in FIG. 5, the patient’s fluctuation of pulse rate tends to be larger during the daytime zone, as compared with night time zone. Therefore, if the blood pressure measurement was conducted during the daytime time zone, α of Equation 3 and Equation 4 may be set to be 0.15. If the blood pressure measurement was conducted during the night time zone, α of Equation 3 and Equation 4 may be set to be 0.10.

According to one or more embodiments of the present invention, the pulse period may be measured a plurality of times at step S50 so that, at step S60, an average of the plurality of pulse period measurements may be used to calculate the threshold value ranges with respect to each of the

plurality of phases. Using an average of the plurality of pulse period measurements in accordance with one or more embodiments of the present invention may result in more reliable threshold value ranges. After the CPU 118 performs the processing to calculate the pulse rate and/or pulse period change threshold value ranges with respect to each of the plurality of phases, the CPU 118 advances the processing to step S70.

At step S70, the CPU 118 stores in the memory unit 126 the threshold value ranges with respect to each of the plurality of phases, as calculated at step S60. After step S70, the CPU 118 advances the processing to step S80.

At step S80, the CPU 118 performs the processing to determine whether the presently measured pulse rate or the presently measured pulse period change as based on the presently measured pulse period is within or beyond the threshold value range of the past-measured pulse rate or the past-measured pulse period change as based on the past-measured pulse period (or an average of past-measured pulse periods) in at least one of the corresponding plurality of phases. If the presently measured pulse rate or the presently measured pulse period change is beyond the threshold value range of the past-measured pulse rate or the past-measured pulse period change in at least one of the corresponding plurality of phases, then the CPU 118 advances the processing to step S90 and determines that the patient was in an unrest condition during blood pressure measurement. After step S90, the CPU 118 advances the processing to step S100 to display on the display unit 128 the presently measured blood pressure was measured when the patient was in an unrest condition. On the other hand, if the presently measured pulse rate or the presently measured pulse period change is within the threshold value range of the past-measured pulse rate or the past-measured pulse period change in at least one of the corresponding plurality of phases, then the CPU 118 advances the processing to step S110 and determines that the patient was in a rest condition during blood pressure measurement. After step S110, the CPU 118 advances the processing to step S120 to display on the display unit 128 the presently measured blood pressure was measured when the patient was in a rest condition.

FIG. 3 shows an example of a comparison of measured pulse periods with past pulse periods in three phases according to one or more embodiments of the present invention. As shown in FIG. 3(A), if the presently measured pulse period (which as shown here, was used to calculate the pulse rate) is within the threshold value range in all of the plurality of phases (e.g., the phase that occurs before pressurization of the cuff 104, the phase that occurs during pressurization of the cuff 104, and the phase that occurs after pressurization of the cuff 104), then it is determined that the patient was under a rest condition during blood pressure measurement, and the blood pressure is displayed as measured under the rest condition. On the other hand, if the presently measured pulse period is beyond or below the threshold value range in any one of the plurality of phases as shown in FIG. 3 (B), then it is determined that the patient was under an unrest condition during blood pressure measurement, and the blood pressure is displayed as measured under the unrest condition.

FIGS. 4(A) and 4(B) show measurement data of pulse periods during pressurization and depressurization phases of the cuff of the blood pressure measurement device (sphygmomanometer) according to one or more embodiments of the present invention.

As shown in FIGS. 4(A) and 4(B), the total time lapsed during the pressurization phase of blood pressure measurement is 7.491667 seconds. As also shown in FIGS. 4(A) and

4(B), a pulse was measured 10 times during the pressurization phase of blood pressure measurement. As such, in this example, the pulse period in the pressurization phase of blood pressure measurement is 7.491667 seconds/10 times=0.7491667 sec. Therefore, the pulse rate during the pressurization phase is calculated by: 60 sec/0.7491667 sec=80.01.

As further shown in FIGS. 4(A) and 4(B) the total time lapsed during the depressurization phase of blood pressure measurement is 24.16167 seconds-7.491667 seconds=16.67 seconds. As also shown in FIGS. 4(A) and 4(B), a pulse was measured 22 times during the depressurization phase of blood pressure measurement. As such, in this example, the pulse period in the depressurization phase of blood pressure measurement is 16.67 seconds/22 times=0.757 sec. Therefore, the pulse rate during the depressurization phase is calculated by: 60 sec/0.757 sec=79.26.

In view of the above, the threshold range of the pulse rate calculated from past measurements during the pressurization phase as shown in FIGS. 4(A) and 4(B) may be calculated as follows, based on Equation 1 and Equation 2:

Upper end of threshold value of pulse rate during pressurization phase is $80.01 \times (1 + 0.15) = 92.01$.

Lower end of threshold value of pulse rate during pressurization phase is $80.01 \times (1 - 0.15) = 68.01$.

As shown in this example, "α" of Equation 1 and Equation 2 has been set to 0.15 to account for the tendency of the blood pressure value and the pulse rate of the patient. With respect to this example, if the presently measured pulse rate of the patient during the pressurization phase is measured to be beyond 92.01 or below 68.01, then it is determined that the presently measured pulse rate was conducted when the patient was in an unrest condition. On the other hand, if the presently measured pulse rate of the patient during the pressurization phase is measured to be within the threshold range of 68.01 to 92.01, then it is determined that the presently measured pulse rate was conducted when the patient was in a rest condition.

Further, the threshold range of the pulse rate calculated from past measurements during the depressurization phase as shown in FIGS. 4(A) and 4(B) may be calculated as follows, based on Equation 1 and Equation 2:

Upper end of threshold value of pulse rate during depressurization phase is $79.26 \times (1 + 0.15) = 91.15$.

Lower end of threshold value of pulse rate during depressurization phase is $79.26 \times (1 - 0.15) = 67.37$.

As shown in this example, "α" of Equation 1 and Equation 2 has been set to 0.15 to account for the tendency of the blood pressure value and the pulse rate of the patient. With respect to this example, if the presently measured pulse rate of the patient during the depressurization phase is measured to be beyond 91.15 or below 67.37, then it is determined that the presently measured pulse rate was conducted when the patient was in an unrest condition. On the other hand, if the presently measured pulse rate of the patient during the depressurization phase is measured to be within the threshold range of 67.37 to 91.15, then it is determined that the presently measured pulse rate was conducted when the patient was in a rest condition.

Still referring to FIGS. 4(A) and 4(B), the maximum pulse period during the pressurization phase is 0.758333 sec. The minimum pulse period during the pressurization phase is 0.741667 sec. As such, in this example, the pulse period

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change in the pressurization phase is 0.758333 sec. (max)–0.741667 sec. (min)=0.01666 sec.

On the other hand, the maximum pulse period during the depressurization phase is 0.766667 sec. The minimum pulse period during the depressurization phase is 0.75 sec. As such, in this example, the pulse period change in the depressurization phase is 0.766667 sec. (max)–0.75 sec. (min)=0.01667 sec.

In view of the above, the threshold range of the pulse period change calculated from past measurements during the pressurization phase as shown in FIGS. 4(A) and 4(B) may be calculated as follows, based on Equation 3 and Equation 4:

Upper end of threshold value of pulse period change during pressurization phase is $0.01666 \text{ sec.} \times (1 + 0.15) = 0.019159 \text{ sec.}$

Lower end of threshold value of pulse period change during pressurization phase is $0.01607 \text{ sec.} \times (1 - 0.15) = 0.014169 \text{ sec.}$

As shown in this example, “ α ” of Equation 3 and Equation 4 has been set to 0.15 to account for the tendency of the blood pressure value and the pulse rate of the patient. With respect to this example, if the presently measured pulse change period of the patient during the pressurization phase is measured to be beyond 0.019159 sec. or below 0.014169 sec., then it is determined that the presently measured pulse change period was conducted when the patient was in an unrest condition. On the other hand, if the presently measured pulse change period of the patient during the pressurization phase is measured to be within the threshold range of 0.014169 sec. to 0.019159 sec., then it is determined that the presently measured pulse change period was conducted when the patient was in a rest condition.

Further, the threshold range of the pulse period change calculated from past measurements during the depressurization phase as shown in FIGS. 4(A) and 4(B) may be calculated as follows, based on Equation 3 and Equation 4:

Upper end of threshold value of pulse period change during depressurization phase is $0.01667 \text{ sec.} \times (1 + 0.15) = 0.0191705 \text{ sec.}$

Lower end of threshold value of pulse period change during depressurization phase is $0.01667 \text{ sec.} \times (1 - 0.15) = 0.0141695 \text{ sec.}$

As shown in this example, “ α ” of Equation 3 and Equation 4 has been set to 0.15 to account for the tendency of the blood pressure value and the pulse rate of the patient. With respect to this example, if the presently measured pulse change period of the patient during the depressurization phase is measured to be beyond 0.0191705 sec. or below 0.0141695 sec., then it is determined that the presently measured pulse change period was conducted when the patient was in an unrest condition. On the other hand, if the presently measured pulse change period of the patient during the depressurization phase is measured to be within the threshold range of 0.0141695 sec. to 0.0191705 sec., then it is determined that the presently measured pulse change period was conducted when the patient was in a rest condition.

Therefore, the blood pressure measurement device (sphygmomanometer) 100 according to one or more embodiments of the present invention accurately determines whether a patient was in a rest condition during blood pressure measurement while taking into account (or regardless of) a change of the pulse period that may be caused by use of a cuff of the blood pressure measurement device.

FIG. 6 is an example display of data on a display unit of a blood pressure measurement device (sphygmomanometer)

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according to one or more embodiments of the present invention. As shown in FIG. 6, the example display may show the blood pressure (systolic blood pressure and diastolic blood pressure) and the pulse rate of a patient together with a determination of whether the patient was in a rest or an unrest condition during blood pressure measurement, according to one or more embodiments of the present invention.

While the invention has been described with respect to a limited number of embodiments, those skilled in the art, having benefit of this disclosure, will appreciate that other embodiments can be devised which do not depart from the scope of the invention as disclosed herein. Accordingly, the scope of the invention should be limited only by the attached claims.

The invention claimed is:

1. A method of determining a rest or an unrest condition of a patient during measurement of blood pressure of the patient, the method comprising:

providing a cuff of a blood pressure measurement device that is configured to be wrapped around a measurement site of the patient, and pressurizing the cuff;

measuring the blood pressure and a pulse period of the patient during and before depressurization of the cuff;

calculating a threshold value range of the pulse period measured during and before depressurization of the cuff based on data of a pulse period measured in a past measurement of the patient and stored in a memory of the blood pressure measurement device,

wherein an upper end of the threshold value range comprises a product of: the pulse period measured either during or before depressurization of the cuff; and a discretionary constant plus one, and

wherein a lower end of the threshold value range comprises a product of: the pulse period measured either during or before depressurization of the cuff; and one minus the discretionary constant;

determining whether the pulse period of the patient measured during and before depressurization of the cuff is within the threshold value range of the pulse period stored in the memory;

if the pulse period measured during and before depressurization is determined to be within the threshold value range of the pulse period stored in the memory both during and before depressurization, displaying on a display of the blood pressure measurement device that the blood pressure was measured when the patient was in the rest condition, and

if the pulse period measured during and before depressurization is determined not to be within the threshold value range of the pulse period stored in the memory either during or before depressurization, displaying on the display of the blood pressure measurement device that the blood pressure was measured when the patient was in the unrest condition.

2. The method of measuring the blood pressure of the patient according to claim 1, further comprising adjusting the threshold value range of the pulse period stored in the memory depending on a condition of at least one of the blood pressure measurement and the pulse period measurement.

3. The method of measuring the blood pressure of the patient according to claim 1, further comprising determining whether the patient was in the rest condition or the unrest condition during blood pressure measurement by examining both a pulse rate and a pulse period change of the patient, and if at least one of the pulse rate and the pulse period change is found to be different from a pulse rate and a pulse period change of past measurement of the patient by more than a

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threshold value, determining that the blood pressure was measured when the patient was in the unrest condition.

4. The method of measuring the blood pressure of the patient according to claim 3,

wherein the pulse rate and the pulse period change of the patient is based on the measured pulse period of the patient, and

wherein the pulse rate and the pulse period change of past measurement of the patient is based on the pulse period measured in the past measurement of the patient.

5. A blood pressure measurement device that determines a rest or an unrest condition of a patient, the device comprising:

a cuff for mounting on a measurement site of the patient, wherein the cuff is configured to be pressurized and depressurized to measure a blood pressure and a pulse period of the patient;

means for measuring the blood pressure and the pulse period of the patient during and before depressurization of the cuff;

a memory that stores data of a pulse period measured in a past measurement of the patient by the measuring means;

a threshold calculation section that calculates a threshold value of the pulse period measured during and before depressurization of the cuff based on data of the pulse period measured in the past measurement of the patient, wherein the threshold value is a threshold value range, wherein if the pulse period was measured during depressurization of the cuff,

an upper end of the threshold value range comprises a product of: the pulse period measured during depressurization of the cuff; and a discretionary constant plus one, and

a lower end of the threshold value range comprises a product of: the pulse period measured during depressurization of the cuff; and one minus the discretionary constant, and

wherein if the pulse period was measured before depressurization of the cuff,

an upper end of the threshold value range comprises a product of: the pulse period measured before depressurization of the cuff; and the discretionary constant plus one, and

a lower end of the threshold value range comprises a product of: the pulse period measured during depressurization of the cuff; and one minus the discretionary constant;

a rest/unrest condition determination section that determines whether the patient was in a rest condition or an unrest condition during blood pressure measurement by comparing the pulse period measured during and before depressurization of the cuff with the threshold value based on data of the pulse period measured in the past measurement of the patient; and

a display unit that displays the measured blood pressure and whether the blood pressure was measured during the rest condition or the unrest condition of the patient.

6. The blood pressure measurement device according to claim 5, wherein the threshold calculation section adjusts the threshold value depending on a measurement condition of the patient.

7. The blood pressure measurement device according to claim 5,

wherein the rest/unrest condition determination section determines whether the patient was in the rest condition or the unrest condition during blood pressure measurement by examining both a pulse rate of the patient mea-

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sured during and before depressurization of the cuff and a pulse period change of the patient measured during and before depressurization of the cuff, and if at least one of the pulse rate of the patient measured during and before depressurization of the cuff and the pulse period change of the patient measured during and before depressurization of the cuff is found to be different from a pulse rate of past measurement of the patient and a pulse period change of past measurement of the patient by more than a corresponding threshold value, the patient is determined to have been in the unrest condition during blood pressure measurement,

wherein the pulse rate of the patient measured during and before depressurization of the cuff and the pulse period change of the patient measured during and before depressurization of the cuff are based on the pulse period of the patient measured during and before depressurization of the cuff, and

wherein the pulse rate of past measurement of the patient and the pulse period change of past measurement of the patient are based on the pulse period measured in the past measurement of the patient.

8. A blood pressure measurement device that determines a rest or an unrest condition of a patient, the device comprising:

a cuff for mounting on a measurement site of the patient, wherein the cuff comprises an air bladder and is configured to be pressurized and depressurized to measure a blood pressure and a pulse period of the patient;

a measurement air system comprising a pressure sensor, a pump, and a valve;

an air tube that connects the air bladder of the cuff to the measurement air system;

a blood pressure calculation section that calculates a blood pressure of the patient based on an internal pressure of the air bladder as detected by the pressure sensor;

a heart rate/pulse period change calculation section that calculates a pulse rate of the patient during and before depressurization of the cuff and a pulse period change of the patient during and before depressurization of the cuff based on the measured pulse period of the patient;

a memory that stores data of a pulse period measured in a past measurement of the patient;

a threshold calculation section that calculates a threshold value of at least one of the pulse rate of the patient calculated during and before depressurization of the cuff and the pulse period change of the patient calculated during and before depressurization of the cuff based on data of the pulse period measured in the past measurement of the patient,

wherein the threshold value is a threshold value range, wherein, with respect to the threshold value of the pulse rate or the pulse period change of the patient calculated during depressurization of the cuff,

an upper end of the threshold value range comprises a product of: the pulse rate or the pulse period change calculated during depressurization of the cuff; and a discretionary constant plus one, and

a lower end of the threshold value range comprises a product of: the pulse rate or the pulse period change calculated during depressurization of the cuff; and one minus the discretionary constant,

wherein, with respect to the threshold value of the pulse rate or the pulse period change of the patient calculated before depressurization of the cuff,

an upper end of the threshold value range comprises a product of: the pulse rate or the pulse period change

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calculated before depressurization of the cuff; and
 the discretionary constant plus one, and
 a lower end of the threshold value range comprises a
 product of: the pulse rate or the pulse period change
 calculated before depressurization of the cuff; and
 one minus the discretionary constant;
 a rest/unrest condition determination section that deter-
 mines whether the patient was in a rest condition or an
 unrest condition during blood pressure measurement by
 comparing at least one of the pulse rate of the patient
 calculated during and before depressurization of the cuff
 and the pulse period change of the patient calculated
 during and before depressurization of the cuff with the
 threshold value based on data of the pulse period mea-
 sured in the past measurement of the patient; and
 a display unit that displays the measured blood pressure
 and whether the blood pressure was measured during the
 rest condition or the unrest condition of the patient.
 9. The blood pressure measurement device according to
 claim 8, wherein the threshold calculation section adjusts the
 threshold value depending on a measurement condition of the
 patient.

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10. The blood pressure measurement device according to
 claim 8,
 wherein the rest/unrest condition determination section
 determines whether the patient was in the rest condition
 or the unrest condition during blood pressure measure-
 ment by examining both the pulse rate of the patient
 calculated during and before depressurization of the cuff
 and the pulse period change of the patient calculated
 during and before depressurization of the cuff, and if at
 least one of the pulse rate of the patient calculated during
 and before depressurization of the cuff and the pulse
 period change of the patient calculated during and
 before depressurization of the cuff is found to be differ-
 ent from a pulse rate of past measurement of the patient
 and a pulse period change of past measurement of the by
 more than a corresponding threshold value, the patient is
 determined to have been in the unrest condition during
 blood pressure measurement, and
 wherein the pulse rate of past measurement of the patient
 and the pulse period change of past measurement of the
 patient are based on the pulse period measured in the
 past measurement of the patient.

* * * * *

专利名称(译)	具有确定患者休息状态的功能的血压测量装置		
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摘要(译)

一种方法包括对血压测量装置的袖带加压，所述血压测量装置缠绕在患者的测量部位周围，在袖带减压期间和之前测量患者的血压和脉冲周期，基于计算脉冲周期的阈值范围。如果测量的脉冲周期被确定为在过去测量的脉冲周期的阈值范围内，则确定患者的过去测量的脉冲周期的数据，确定测量的脉冲周期是否在过去测量的脉冲周期的阈值范围内。在减压期间和之前，显示患者处于休息状态，并且如果在减压期间或之前确定测量的脉冲周期不在过去测量的脉冲周期的阈值范围内，则显示患者在不安的情况。

