



US006884221B2

(12) **United States Patent**
Narimatsu et al.

(10) **Patent No.:** **US 6,884,221 B2**
(45) **Date of Patent:** **Apr. 26, 2005**

(54) **CIRCULATORY-ORGAN EVALUATING APPARATUS**

(75) Inventors: **Kiyoyuki Narimatsu**, Komaki (JP);
Toshihiko Ogura, Komaki (JP); **Akira Tampo**, Komaki (JP)

(73) Assignee: **Colin Medical Technology Co Corporation**, Komaki (JP)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 123 days.

(21) Appl. No.: **10/370,617**

(22) Filed: **Feb. 24, 2003**

(65) **Prior Publication Data**

US 2003/0236464 A1 Dec. 25, 2003

(30) **Foreign Application Priority Data**

Jun. 20, 2002 (JP) 2002-180370

(51) Int. Cl.⁷ **A61B 5/02**; A61B 5/00

(52) U.S. Cl. **600/485**; 600/483; 600/481;
600/300

(58) Field of Search 600/481, 483,
600/485, 486, 490, 492-496, 500-503,
508-526

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,265,011 A 11/1993 O'Rourke
5,743,856 A 4/1998 Oka et al.
6,196,974 B1 3/2001 Miwa

6,322,516 B1 11/2001 Masuda et al.
6,440,079 B1 * 8/2002 Ogura et al. 600/492
6,612,993 B1 * 9/2003 Narimatsu 600/500
6,659,958 B1 * 12/2003 Narimatsu et al. 600/485
6,666,827 B1 * 12/2003 Narimatsu 600/490
6,702,754 B1 * 3/2004 Ogura et al. 600/500
6,712,768 B1 * 3/2004 Ogura et al. 600/494
6,719,704 B1 * 4/2004 Narimatsu et al. 600/500
6,746,405 B1 * 6/2004 Narimatsu 600/494

FOREIGN PATENT DOCUMENTS

EP 1 050 267 A1 11/2000
JP A 9-140679 6/1997
JP 11-045286 2/1999
JP 2001-069659 3/2001

OTHER PUBLICATIONS

Masuda, Y. "Basis of Pulse Wave, Waveform of Pressure Wave (Pulse Wave)," *Pulse Wave Velocity*, May 1, 2002, pp. 18-25, in Japanese and English translation pp 1-7.

* cited by examiner

Primary Examiner—Robert L. Nasser

Assistant Examiner—Navin Natnithithadha

(74) Attorney, Agent, or Firm—Olliff & Berridge

(57) **ABSTRACT**

A circulatory-organ evaluating apparatus for evaluating a circulatory organ of a living subject, including a display device, and a control device which controls the display device to simultaneously and graphically display a blood pressure, an augmentation index, and pulse-wave-velocity-related information of the subject.

6 Claims, 10 Drawing Sheets

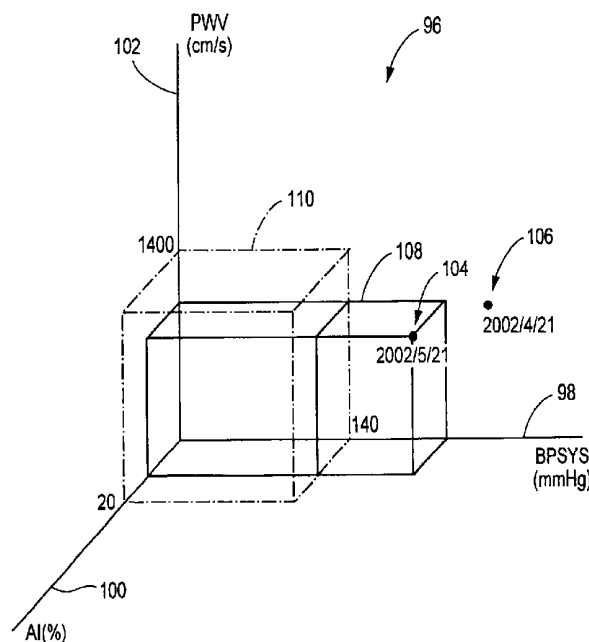
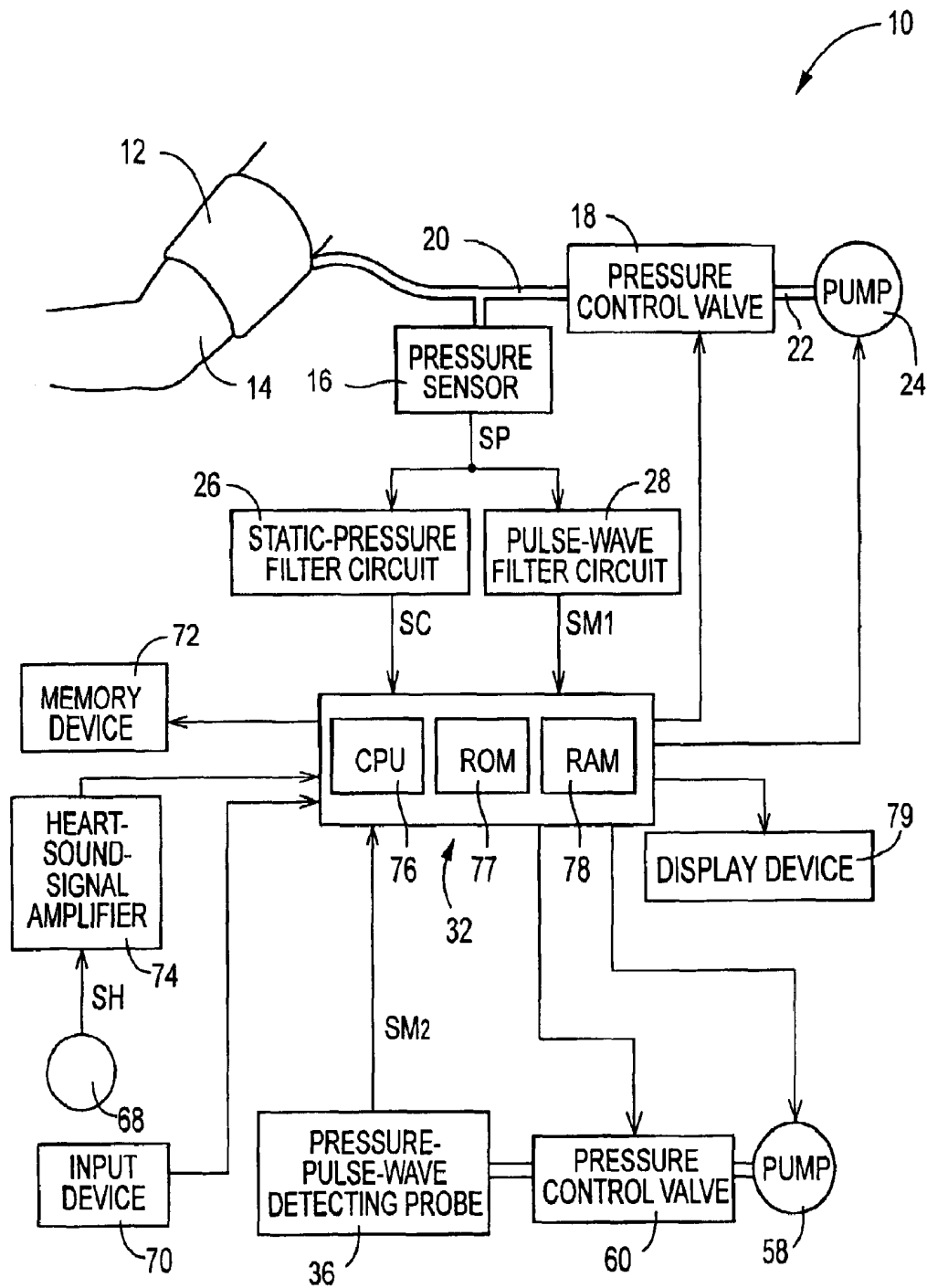


FIG. 1



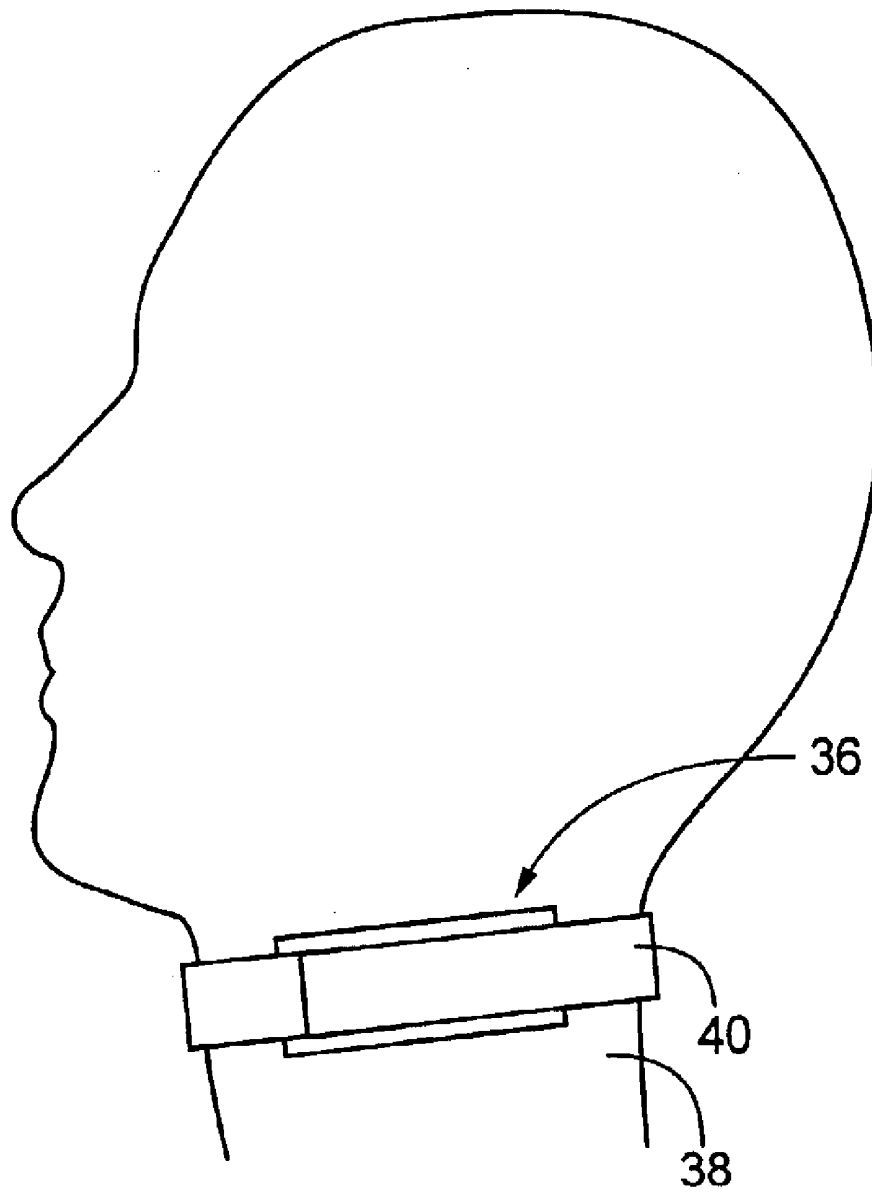
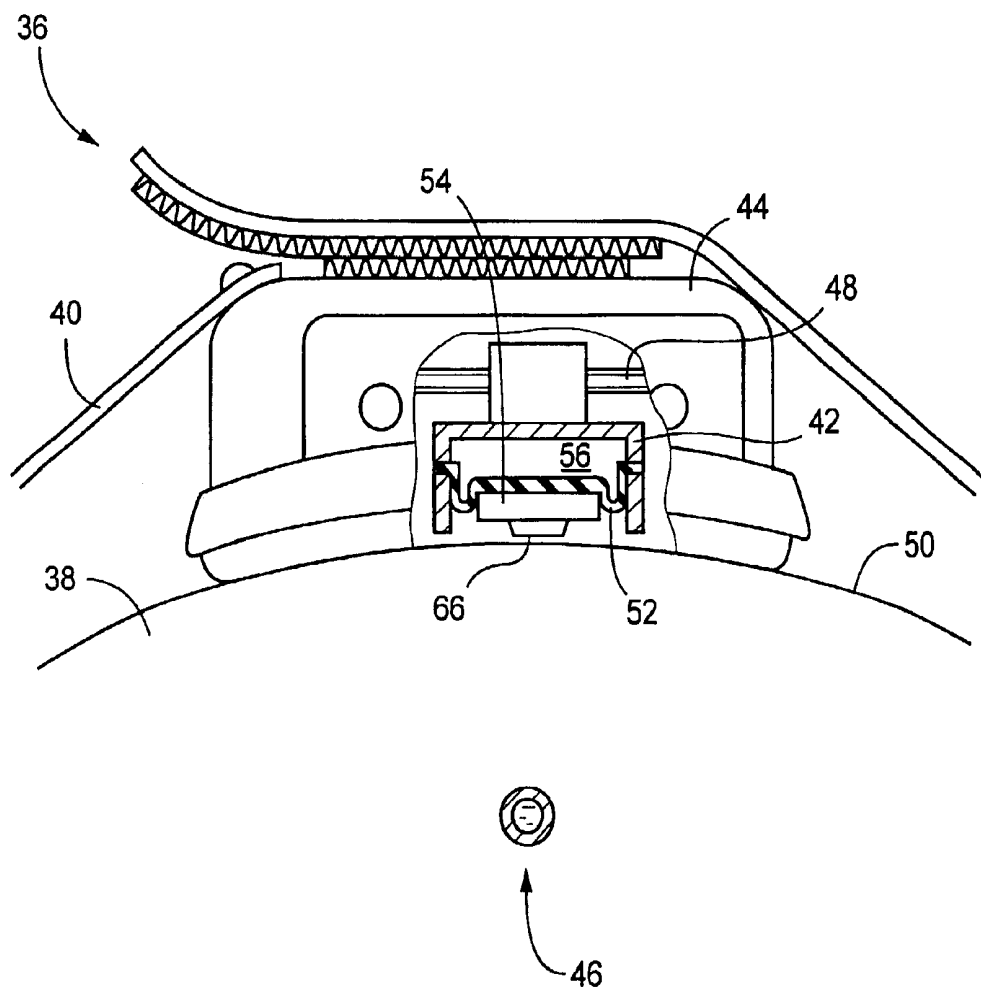


FIG. 2

FIG. 3



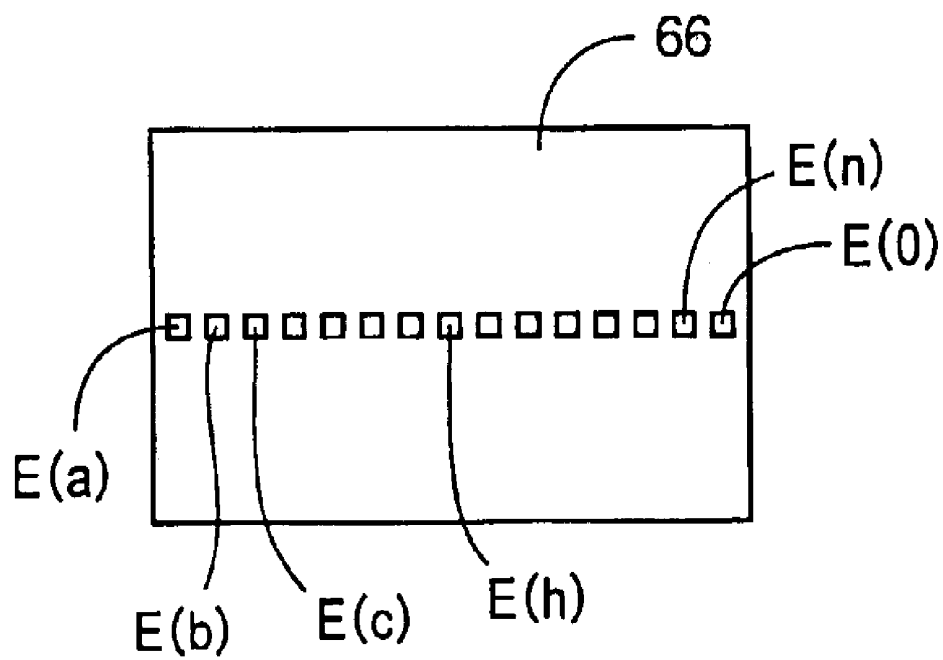


FIG. 4

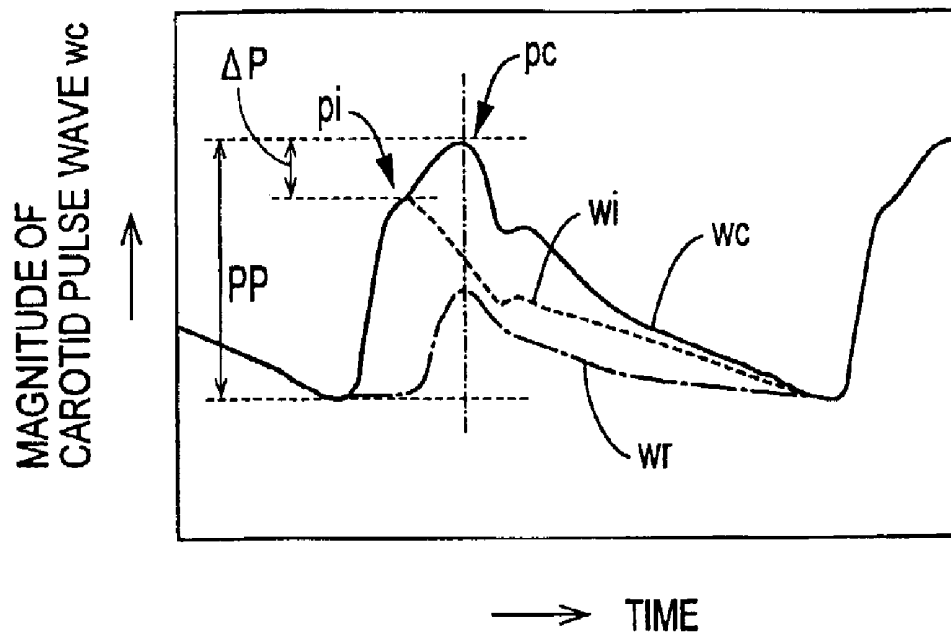
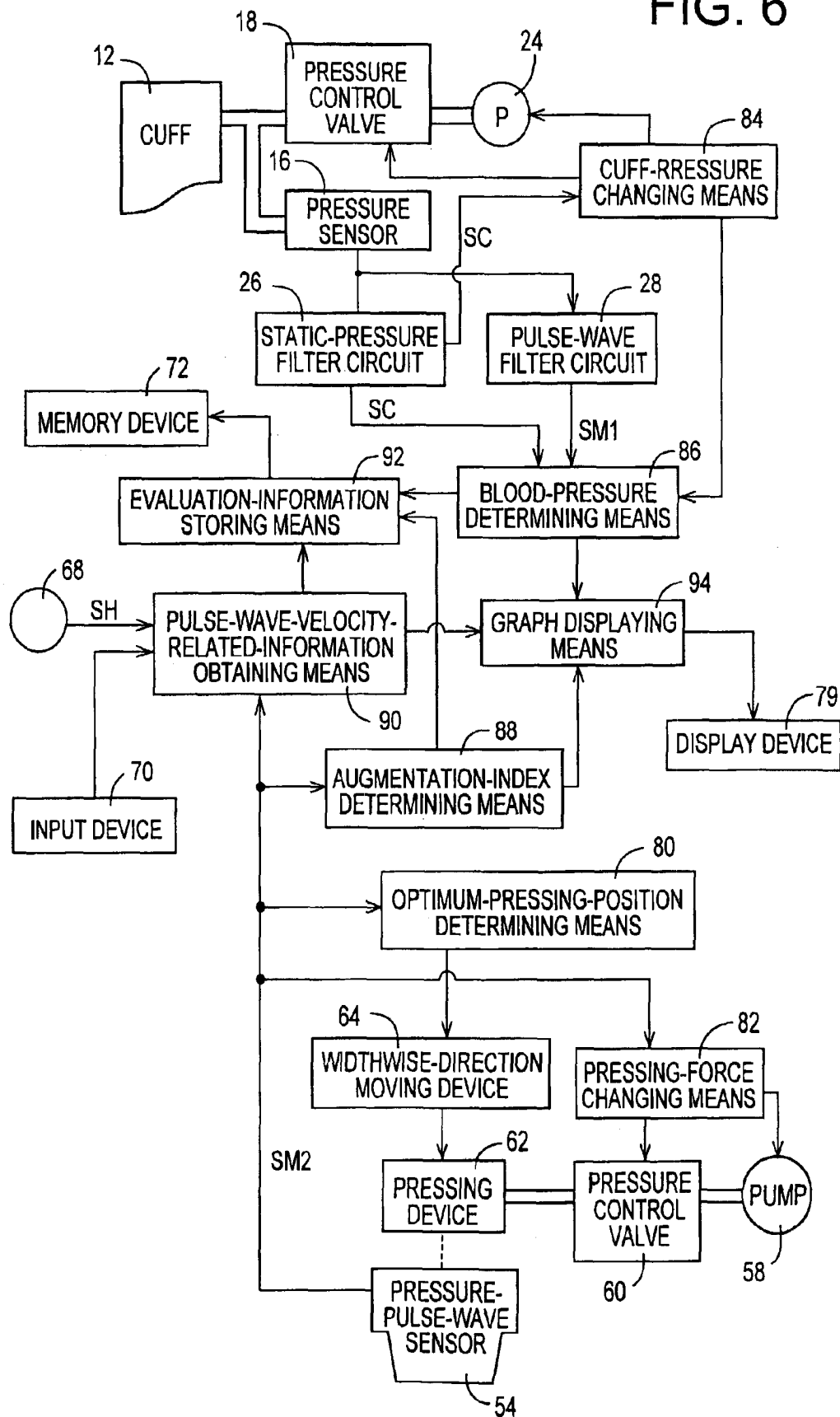


FIG. 5

FIG. 6



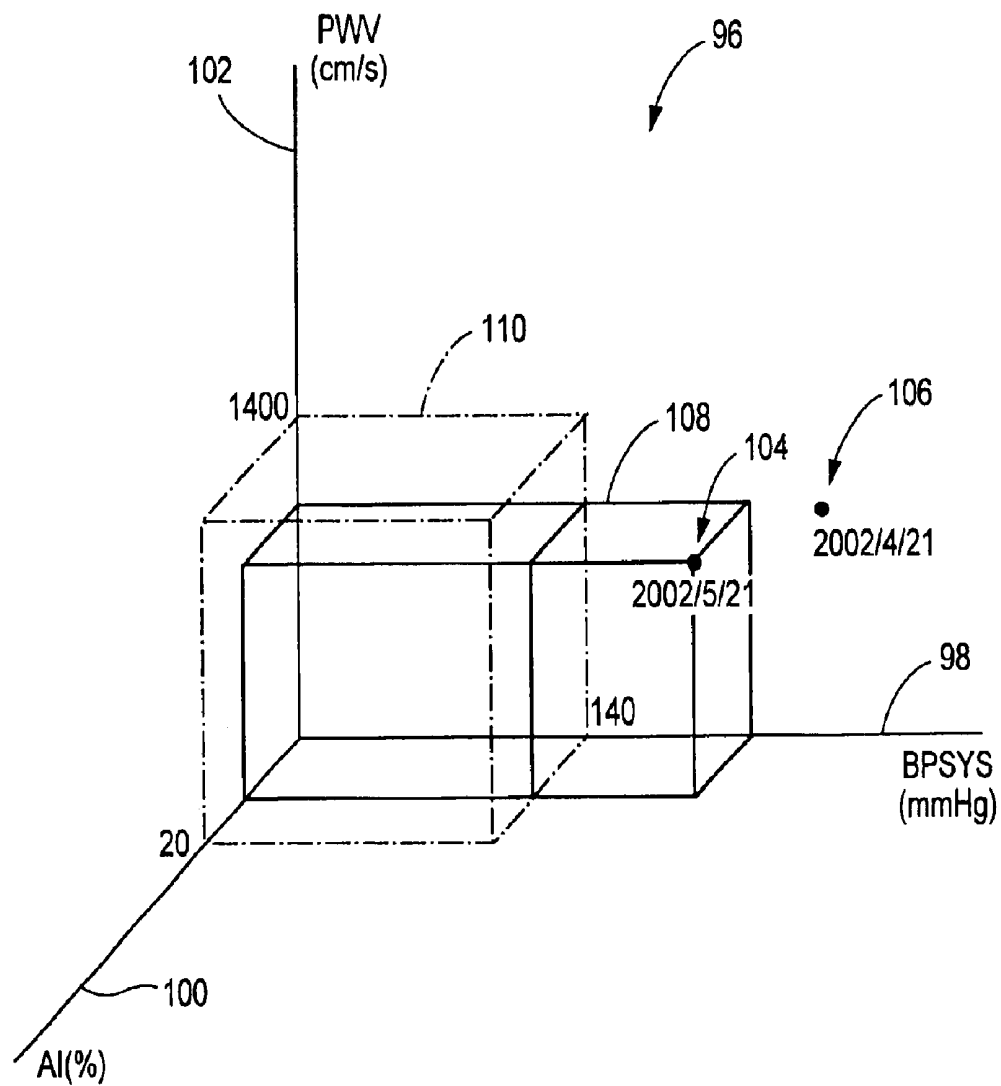


FIG. 7

FIG. 8

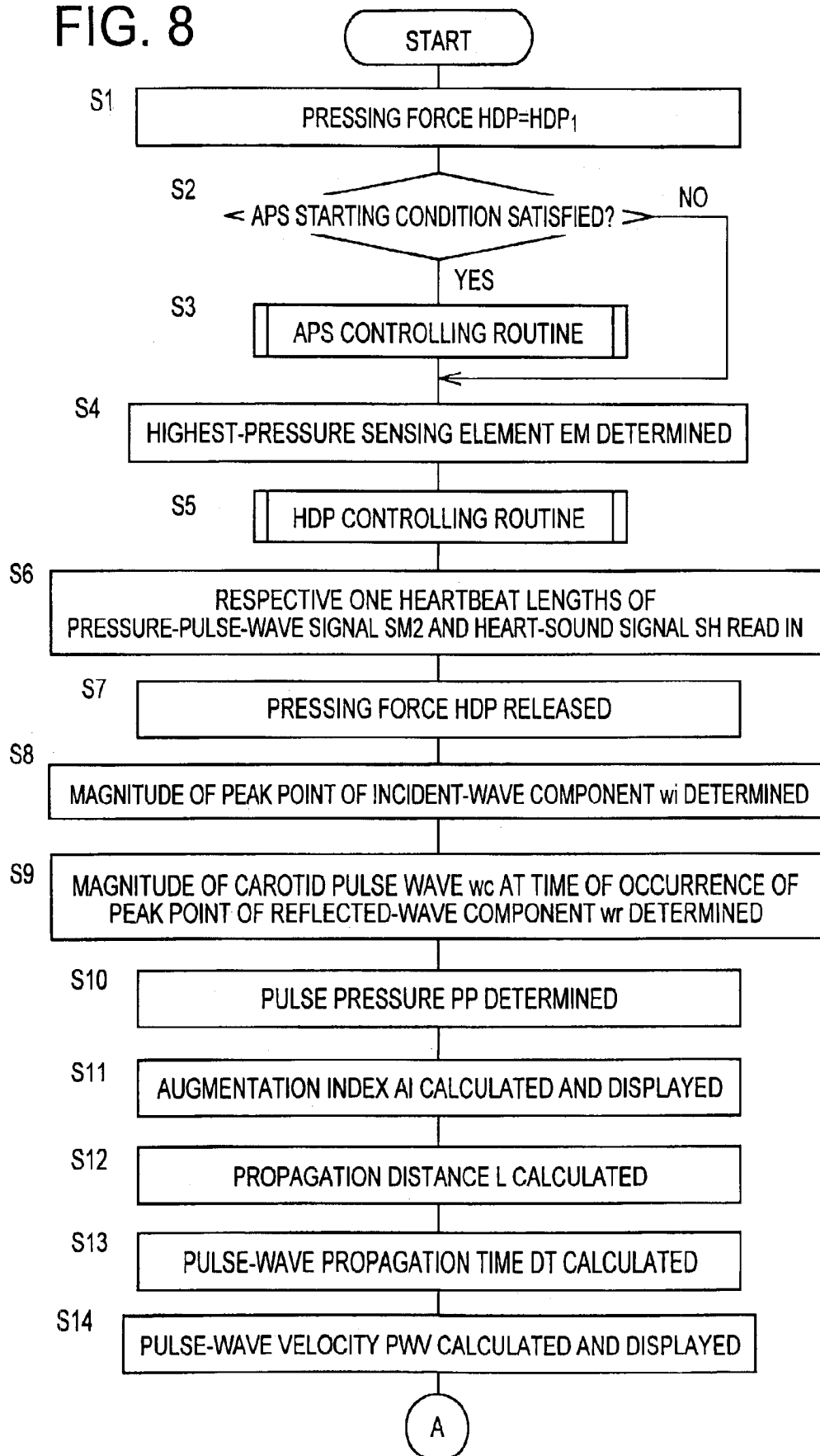


FIG. 9

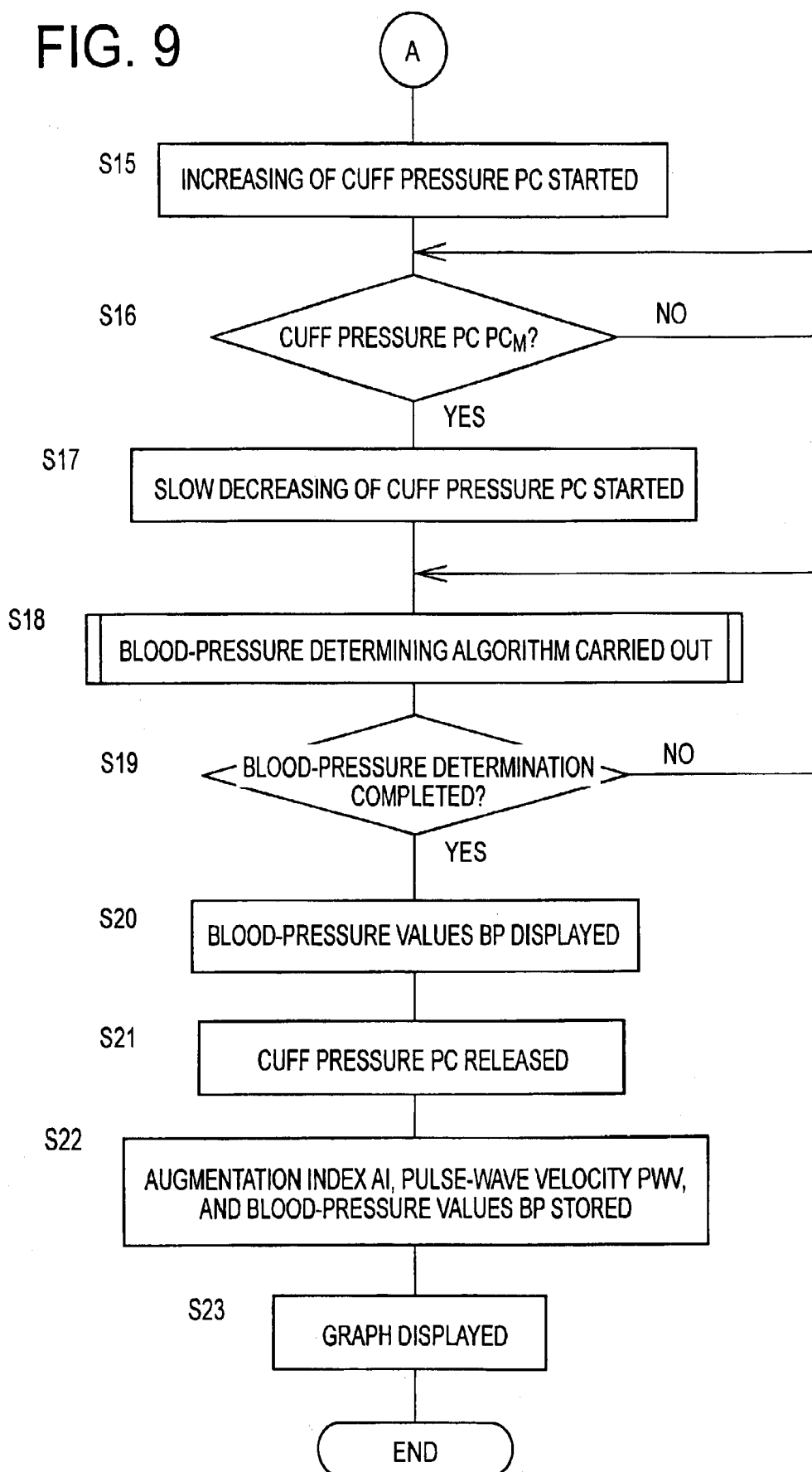
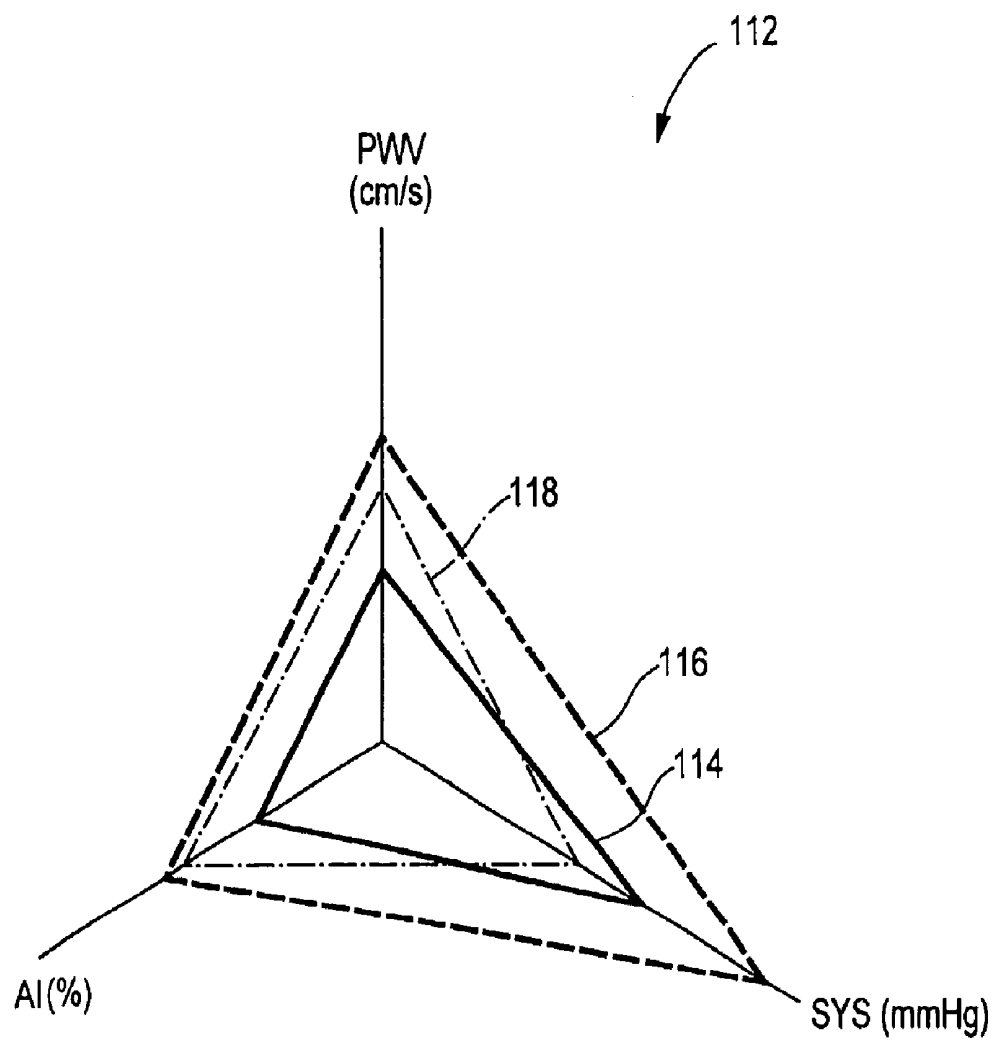


FIG. 10



CIRCULATORY-ORGAN EVALUATING APPARATUS

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to a circulatory-organ evaluating apparatus which evaluates a circulatory organ of a living subject.

2. Related Art Statement

The dynamic function of circulatory organ of a living person is primarily defined by the pumping function of the person's heart, and also is influenced by the degree of contraction or dilation of walls of blood vessels (e.g., the degree of hardness of arteries) or the diameter of blood vessels.

Therefore, the dynamic function of circulatory organ is evaluated by measuring, e.g., the person's blood pressure. If the person is diagnosed with hypertension, the person will undergo a medical treatment such as administration of antihypertensive drug. In addition, since the blood pressure is influenced by the degree of hardness of arteries, there will be some cases where pulse-wave-velocity-related information that is related to a velocity at which a pulse wave propagates between the person's two portions is obtained so as to evaluate the degree of hardness of arteries as well. Those evaluations may be made by, e.g., an apparatus disclosed by Japanese Patent Document No. 9-140679 or its corresponding U.S. Pat. No. 5,743,856.

Meanwhile, there are some indexes used for evaluating the circulatory organ of living person. Those indexes include, in addition to blood pressure and pulse-wave-velocity-related information, augmentation index, AI. The augmentation index indicates a proportion of a reflected-wave component of a pulse wave to an incident-wave component of the same. In order to accurately evaluate the circulatory organ, it is preferred to simultaneously measure augmentation index in addition to blood pressure and pulse-wave-velocity-related information, because the increased number of physical parameters obtained from the person can be used to evaluate the circulatory organ.

However, in the case where the circulatory organ is evaluated by measuring the three indexes, i.e., blood pressure, pulse-wave-velocity-related information, and augmentation index, if respective measured values of the three indexes are just displayed, it is difficult to speculate how pulse-wave-velocity-related information or augmentation index influences blood pressure, or conversely if blood pressure influences pulse-wave-velocity-related information or augmentation index. Thus, there are some cases where it is difficult to evaluate the circulatory organ if the three measured values are just displayed. In particular, in the case where a patient is informed of a therapy to be taken or of therapeutic effects, it is difficult for the patient to evaluate the circulatory organ based on only the measured values.

SUMMARY OF THE INVENTION

It is therefore an object of the present invention to provide a circulatory-organ evaluating apparatus which permits easy evaluation of the circulatory organ of a living subject.

The above object has been achieved by the present invention. According to the present invention, there is provided a circulatory-organ evaluating apparatus for evaluating a circulatory organ of a living subject, comprising a display device; and a control device which controls the

display device to simultaneously and graphically display a blood pressure, an augmentation index, and pulse-wave-velocity-related information of the subject.

According to this invention, a person can graphically recognize, on the display device, the blood pressure, augmentation index, and pulse-wave-velocity-related information of the subject. Thus, the person is permitted to easily evaluate the circulatory organ of the subject.

According to a preferred feature of the present invention, the display device displays a multi-dimensional graph representing at least the blood pressure, the augmentation index, and the pulse-wave-velocity-related information of the subject. According to this feature, since the person can graphically recognize, from the single multi-dimensional graph, the three parameters, i.e., the blood pressure, the augmentation index, and the pulse-wave-velocity-related information, the person can easily evaluate the circulatory organ based on those increased number of parameters. However, the display device may display respective graphs of the blood pressure, the augmentation index, and the pulse-wave-velocity-related information, independent of each other.

According to another feature of the present invention, the multi-dimensional graph includes a standard figure representing a standard blood pressure, a standard augmentation index, and standard pulse-wave-velocity-related information. According to this feature, the person can more easily evaluate the circulatory organ by graphically comparing the measured values of the blood pressure, the augmentation index, and the pulse-wave-velocity-related information, with the standard figure.

According to another feature of the present invention, the multi-dimensional graph represents respective time-wise changes of the blood pressure, the augmentation index, and the pulse-wave-velocity-related information of the subject. According to this feature, since the person can graphically recognize the respective time-wise changes of the blood pressure, the augmentation index, and the pulse-wave-velocity-related information, the person can easily evaluate a time-wise change of the circulatory organ. Therefore, the person can easily evaluate the effect of medication or the improvement of subject's habits.

BRIEF DESCRIPTION OF THE DRAWINGS

The above and optional objects, features, and advantages of the present invention will be better understood by reading the following detailed description of the preferred embodiments of the invention when considered in conjunction with the accompanying drawings, in which:

FIG. 1 is a diagrammatic view showing a circuitry of a circulatory-organ evaluating apparatus to which the present invention is applied;

FIG. 2 is a view showing a state in which a pressure-pulse-wave detecting probe of the apparatus of FIG. 1 is worn on a cervical portion of a living subject;

FIG. 3 is an enlarged view of the pressure-pulse-wave detecting probe of FIG. 2, a portion of the probe being cut away;

FIG. 4 is a view for explaining a state in which an array of pressure sensing elements is provided in a press surface of a pressure-pulse-wave sensor shown in FIG. 3;

FIG. 5 is a view showing an example of a carotid pulse wave, w_c , represented by a pressure-pulse-wave signal, $SM2$, supplied from one of the pressure sensing elements of the pressure-pulse-wave sensor of FIG. 3;

FIG. 6 is a diagrammatic view for explaining essential control functions of an electronic control device of the apparatus of FIG. 1;

FIG. 7 is a view showing a three-dimensional scatter diagram as an example of a three-dimensional graph which is displayed on a display device by a graph displaying means shown in FIG. 6;

FIG. 8 is a flow chart for explaining a portion of the control functions of a CPU (central processing unit) of the control device, shown in the diagrammatic view of FIG. 6;

FIG. 9 is a flow chart for explaining another portion of the control functions of the CPU, shown in the diagrammatic view of FIG. 6; and

FIG. 10 is a view showing a radar chart as another example of the three-dimensional graph which is displayed on the display device by the graph displaying means shown in FIG. 6.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

Hereinafter, there will be described an embodiment of the present invention in detail by reference to the drawings. FIG. 1 is a diagrammatic view showing a circuitry of a circulatory-organ evaluating apparatus 10 to which the present invention is applied.

In FIG. 1, reference numeral 12 designates an inflatable cuff which includes a belt-like cloth bag and a rubber bag accommodated in the cloth bag and which is adapted to be wound around a brachial portion 14 of a patient as a living subject. The cuff 12 is connected via a piping 20 to a pressure sensor 16 and a pressure control valve 18. The pressure control valve 18 is connected via a piping 22 to an air pump 24. The pressure control valve 18 adjusts a pressure of a pressurized air supplied from the air pump 24, and supplies the pressure-adjusted air to the cuff 12, or discharges the pressurized air from the cuff 12, so as to control an air pressure in the cuff 12.

The pressure sensor 16 detects the air pressure in the cuff 12, and supplies a pressure signal, SP, representing the detected air pressure, to a static-pressure filter circuit 26 and a pulse-wave filter circuit 28. The static-pressure filter circuit 26 includes a low-pass filter which extracts, from the pressure signal SP, a cuff-pressure signal, SC, representing a static component of the detected air pressure, i.e., a pressing pressure of the cuff 12 (hereinafter, referred to as the cuff pressure, PC). The filter circuit 26 supplies the cuff-pressure signal SC to an electronic control device 32 via an A/D (analog-to-digital) converter, not shown. The pulse-wave filter circuit 28 includes a band-pass filter which extracts, from the pressure signal SP, a cuff-pulse-wave signal, SM1, representing a cuff pulse wave as an oscillatory component of the detected air pressure. The filter circuit 28 supplies the cuff-pulse-wave signal SM1 to the control device 32 via an A/D converter, not shown. The cuff pulse wave represented by the cuff-pulse-wave signal SM1 is a brachial pulse wave, wb, produced from a brachial artery, not shown, of the patient that is pressed by the cuff 12.

The present circulatory-organ evaluating apparatus 10 includes a pressure-pulse-wave detecting probe 36, shown in FIG. 2, that functions as a carotid-pulse-wave detecting device. The pressure-pulse-wave detecting probe 36 is worn on a cervical portion 38 of the patient, as shown in FIG. 2, with the help of a band 40. As shown in detail in FIG. 3, the pressure-pulse-wave detecting probe 36 includes a container-like sensor housing 42; a case 44 which accommodates the sensor housing 42; and a feed screw 48 which

is threadedly engaged with the sensor housing 42 and is rotated by an electric motor, not shown, provided in the case 44 so as to move the sensor housing 42 in a widthwise direction of a carotid artery 46. With the help of the band, the pressure-pulse-wave detecting probe 36 is detachably attached to the cervical portion 38, such that an open end of the sensor housing 42 is opposed to a body surface 50 of the cervical portion 38.

In addition, the pressure-pulse-wave detecting probe 36 includes a pressure-pulse-wave sensor 54 which is secured via a diaphragm 52 to an inner wall of the sensor housing 42, such that the sensor 54 is movable relative to the housing 42 and is advanceable out of the open end of the same 42. The sensor housing 42, the diaphragm 52, etc. cooperate with one another to define a pressure chamber 56, which is supplied with a pressurized air from an air pump 58 via a pressure-control valve 60, as shown in FIG. 1, so that the pressure-pulse-wave sensor 54 is pressed against the body surface 50 with a pressing force corresponding to the air pressure in the pressure chamber 56.

The sensor housing 42 and the diaphragm 52 cooperate with each other to provide a pressing device 62 which presses the pressure-pulse-wave sensor 54 against the carotid artery 46, and the feed screw 48 and the not-shown motor cooperate with each other to provide a widthwise-direction moving device 64 which moves the pressure-pulse-wave sensor 54 in the widthwise direction of the carotid artery 46 and thereby changes a pressing position where the sensor 54 is pressed on the body surface 50.

The pressure-pulse-wave sensor 54 has a pressing surface 66, and a number of semiconductor pressure sensing elements (hereinafter, referred to as the "pressure sensing elements") E which are arranged in the pressing surface 66 at a regular interval in the widthwise direction of the carotid artery 46, i.e., in the direction of movement of the sensor 54 parallel to the feed screw 48, over a length greater than the diameter of the carotid artery 46. For example, as shown in FIG. 4, fifteen pressure sensing elements E(a), E(b), . . . , E(o) are arranged at a regular interval of, e.g., 0.6 mm.

The pressure-pulse-wave detecting probe 36, constructed as described above, is pressed against the body surface 50 of the cervical portion 38 right above the carotid artery 46, so that the pressure-pulse-wave sensor 54 detects a pressure pulse wave (i.e., a carotid pulse wave, wc) which is produced from the carotid artery 46 and is transmitted to the body surface 50, and supplies a pressure-pulse-wave signal, SM2, representing the detected carotid pulse wave wc, to the control device 32 via an A/D converter, not shown. An example of the carotid pulse wave wc represented by the pressure-pulse-wave signal SM2 continuously supplied from the pressure-pulse-wave sensor 54 is indicated at solid line in FIG. 5.

Back to FIG. 1, the circulatory-organ evaluating apparatus 10 further includes a heart-sound microphone 68, an input device 70, and a memory device 72. The heart-sound microphone 68 is attached, with an adhesive tape, not shown, or the like, to a chest, not shown, of the patient. The microphone 68 incorporates a piezoelectric element, not shown, which converts heart sounds produced from the patient's heart, into an electric signal, i.e., a heart-sound signal, SH. A heart-sound-signal amplifier 74 includes four sorts of filters, not shown, which cooperate with one another to attenuate a low-pitch component having a great energy and thereby amplifies and filters a high-pitch component of the heart-sound signal SH supplied from the microphone 68. The heart-sound signal SH amplified and filtered by the

amplifier 74 is supplied to the control device 32 via an A/D converter, not shown.

The input device 70 includes a keyboard, not shown, which is operated by an operator such as a doctor to input an identification code and a stature, T, of the patient. The input device 70 supplies respective signals representing the input-
5 subject's identification code and stature T, to the control device 32.

The memory device 72 is provided by a well-known memory such as a magnetic disk, a magnetic tape, or a non-volatile semiconductor memory, and stores, in a memory area allotted for each of a plurality of patients, a blood pressure, BP, an augmentation index, AI, and a pulse-wave velocity, PWV, of the each patient that are determined by the electronic control device 32.

The electronic control device 32 is provided by a so-called microcomputer including a CPU 76, a ROM (read only memory) 77, a RAM (random access memory) 78, and an I/O (input-and-output) port, not shown. The CPU 76 processes signals according to the control programs pre-stored in the ROM 77 by utilizing the temporary-storage function of the RAM 78, and supplies drive signals via the I/O port to the air pumps 24, 58 and the pressure control valves 18, 60 so as to control the cuff pressure PC and the pressure in the pressure chamber 56. Moreover, the CPU 76 determines, based on the cuff-pulse-wave signal SM1, the pressure-pulse-wave signal SM2, the cuff-pressure signal SC, and the heart-sound signal SH, each supplied to the control device 32, a blood pressure BP, an augmentation index AI, and a pulse-wave velocity PWV of each patient, and controls a display device 79 and the memory device 72.

FIG. 6 is a diagrammatic view for explaining essential control functions of the electronic control device 32 of the circulatory-organ evaluating apparatus.

An optimum-pressing-position determining device or means 80 judges whether a prescribed pressing-position changing condition is satisfied, i.e., whether one (hereinafter, referred to as the "highest-pressure sensing element, EM") of the pressure sensing elements E of the pressure-pulse-wave sensor 54 that detects the highest pressure of the respective pressures detected by all the elements E is positioned in one of prescribed opposite end portions of the array of pressure sensing elements E. Each of the prescribed opposite end portions of the array of elements E may be a range having a prescribed length including a corresponding one of the opposite ends of the array of elements E, or a range accommodating a prescribed number of elements E including a corresponding one of the respective elements E located at the opposite ends of the array. When this pressing-position changing condition is satisfied, the determining means 80 carries out the following pressing-position changing operation: After the pressing device 62 once moves the pressure-pulse-wave sensor 54 away from the body surface 50, the widthwise-direction moving device 64 moves the pressing device 62 and the sensor 54 over a prescribed distance, and then the pressing device 62 presses the sensor 54 with a prescribed, considerably small, first pressing force HDP1 that would be smaller than an optimum pressing force HDPO, described later. In this state, the determining means 80 judges again whether the prescribed pressing-position changing condition is satisfied. The determining means 80 repeats carrying out the above-described operation and judgment till the pressing-position changing condition is not satisfied any longer, preferably till the highest-pressure sensing element EM is positioned in a prescribed middle portion of the array of elements E. The

length, or element number, employed for defining each of the opposite end portions of the array of elements E is prescribed based on the diameter of the artery (i.e., the carotid artery 46) to be pressed by the pressure-pulse-wave sensor 54, and may be one fourth of the diameter.

A pressing-force changing device or means 82 changes, after the optimum-pressing-position determining means 80 positions the pressure-pulse-wave sensor 54 at the optimum pressing position, a pressing force HDP (i.e., a hold-down pressure) applied by the pressing device 62 to the sensor 54, within a prescribed pressing-force range, either stepwise in response to each heartbeat of the patient or continuously at a prescribed, considerably low rate. Based on the carotid pulse wave wc obtained during the changing of the pressing force HDP, the changing means 82 determines an optimum pressing force HDPO and maintains the pressing force applied by the pressing device 62 to the sensor 54, at the thus determined optimum pressing force HDPO. Here, the optimum pressing force HDPO is so determined that a pulse pressure PP of the carotid pulse wave wc detected by the highest-pressure sensing element EM pressed with the pressing force HDP (i.e., a pressure difference obtained by subtracting a pressure corresponding to a minimum magnitude, from a pressure corresponding to a maximum magnitude, of one heartbeat-synchronous pulse of the carotid pulse wave wc) may not be smaller than a predetermined lower-limit pulse pressure PPL. The lower-limit pulse pressure PPL is experimentally determined, in advance, as a value which assures that a clear carotid pulse wave wc can be detected. If a pulse pressure PP is too small, a clear carotid pulse wave wc cannot be obtained.

A cuff-pressure changing device or means 84 operates, based on the cuff-pressure signal SC supplied from the static-pressure filter circuit 26, the pressure control valve 18 and the air pump 24 so as to quickly increase the cuff pressure PC to a prescribed target pressure PC_M (e.g., 180 mmHg) that would be higher than a systolic blood pressure BP_{SYS} of the patient and, subsequently, slowly decrease the cuff pressure at a rate of, e.g., 2 or 3 mmHg/sec. After a blood-pressure determining device or means 86, described below, determines blood-pressure values BP of the patient, the changing means 84 releases the cuff pressure to an atmospheric pressure.

The blood-pressure determining means 86 determines, based on the cuff-pressure signal SC continuously supplied from the static-pressure filter circuit 26, and the cuff-pulse-wave signal SM1 continuously supplied from the pulse-wave filter circuit 28, each during the slow decreasing of the cuff pressure PC under the control of the cuff-pressure changing means 84, a systolic blood pressure BP_{SYS} , a mean blood pressure BP_{MEAN} , and a diastolic blood pressure BP_{DIA} of the patient, according to well-known oscillometric method.

An augmentation-index determining device or means 88 first determines respective times of occurrence of respective peak points, pi and pr , of an incident-wave component, wi , and a reflected-wave component, wr , which are contained in a heartbeat-synchronous pulse of the carotid pulse wave wc continuously detected by the highest-pressure sensing element EM of the pressure-pulse-wave sensor 54 in the state in which the pressing force HDP applied to the sensor 54 is maintained at the optimum pressing force HDPO. Then, the determining means 88 determines a pressure difference, ΔP , by subtracting a pressure corresponding to a magnitude of the carotid pulse wave wc at the time of occurrence of the peak point pi of the incident-wave component from a pressure corresponding to a magnitude of the carotid pulse

wave *wc* at the time of occurrence of the peak point *pr* of the reflected-wave component *wr*. In addition, the determining means **88** determines a pulse pressure *PP* of the heartbeat-synchronous pulse of the carotid pulse wave *wc*. And, the determining means **88** calculates an augmentation index *AI* by substituting the following Expression 1 with the thus determined pressure difference ΔP and pulse pressure *PP*:

$$AI = (\Delta P / PP) \times 100(\%) \quad (\text{Expression 1})$$

Here, the manner in which the time of occurrence of peak point *pi* of incident-wave component *wi* of the carotid pulse wave *wc* is explained. The carotid pulse wave *wc* contains the incident-wave component *wi*, indicated at broken line in FIG. 5, and the peak point *pi* of the incident-wave component *wi* corresponds to an inflection point or a local-maximum point of the composite carotid pulse wave *wc* (i.e., observed wave) that occurs between a rising point (i.e., a minimum point) and a peak point, *pc*, of the composite wave *wc*. In the example shown in FIG. 5, the peak point *pi* of the incident-wave component *wi* corresponds to an inflection point of the observed wave *wc*. To this end, the determining means **88** subjects the continuously obtained pressure-pulse-wave signal *SM2* to a mathematical treatment commonly used to identify an inflection or local-maximum point; such as a differentiation treatment or a filter treatment.

The time of occurrence of peak point *pr* of the reflected-wave component *wr* is generally defined as a time of occurrence of the first local-maximum point following the peak point *pi* of the incident-wave component *wi*. Therefore, in the case, shown in FIG. 5, where a peak point *pi* of an incident-wave component *wi* does not coincide with a peak point *pc* of a carotid pulse wave *wc*, the time of occurrence of peak point *pc* of the carotid pulse wave *wc* is determined as the time of occurrence of peak point *pr* of reflected-wave component *wr*. On the other hand, in the case where a peak point *pi* of an incident-wave component *wi* is so large that the peak point *pi* of incident-wave component *wi* also defines a peak point *pc* of a carotid pulse wave *wc*, the time of occurrence of the first local-maximum point following the peak point *pi* of incident-wave component *wi* is determined as the time of occurrence of peak point *pr* of reflected-wave component *wr*.

A pulse-wave-velocity-related-information obtaining device or means **90** first determines a prescribed point (e.g., a starting point of second heart sound, *II*) of a heart-sound waveform represented by the heart-sound signal *SH* continuously supplied from the heart-sound microphone **68**, and a prescribed point (e.g., a dicrotic notch) of the carotid pulse wave *wc*, represented by the pressure-pulse-wave signal *SM2*, that corresponds to the prescribed point of the heart-sound waveform, and then determines a time difference between a time of detection of the prescribed point of the heart-sound waveform and a time of detection of the prescribed point of the carotid pulse wave *wc*. This time difference is a pulse-wave propagation time, *DT*, that is needed for the carotid pulse wave *wc* to propagate from the patient's aortic valve to the cervical portion **38** where the pressure-pulse-wave detecting probe **36** is worn. In addition, the obtaining means **90** determines, according to a pre-determined relationship between stature *T* and propagation distance *L*, represented by the following Expression 2, and based on the patient's stature *T* supplied from the input device **70**, a propagation distance *L* between the patient's aortic valve and the cervical portion **38** where the probe **36** is worn, and finally determines a pulse-wave velocity *PWV* (cm/sec) according to the following Expression 3 and based

on the thus obtained propagation distance *L* and pulse-wave propagation time *DT*:

$$L = aT + b \quad (\text{Expression 2})$$

(*a* and *b* are experimentally determined constants)

$$PWV = L / DT \quad (\text{Expression 3})$$

An evaluation-information storing device or means **92** stores, in a certain memory area of the memory device **72**, a set of circulatory-organ evaluation information including the blood-pressure values *BP* determined by the blood-pressure determining means **86**, the augmentation index *AI* determined by the augmentation-index determining means **88**, and the pulse-wave velocity *PWV* obtained by the pulse-wave-velocity-related-information obtaining means **90**, such that the set of evaluation information is associated with the identification code identifying each patient and a date (or a date and a time) when the set of evaluation information is obtained.

A graph displaying device or means **94** operates the display device **79** to display a three-dimensional graph having a blood-pressure axis, an augmentation-index axis, and a pulse-wave-velocity-related-information axis. In addition, the graph displaying means **96** operates the display device **79** to display, in the three-dimensional graph, a symbol representing the blood-pressure values *BP* determined by the blood-pressure determining means **86**, the augmentation index *AI* determined by the augmentation-index determining means **88**, and the pulse-wave velocity *PWV* obtained by the pulse-wave-velocity-related-information obtaining means **90**. If the memory device **72** stores one or more past sets of evaluation information that had been obtained from the patient, the graph displaying means **94** operates the display device **79** to additionally display, in the three-dimensional graph, one or more symbols representing the one or more past sets of evaluation information.

FIG. 7 shows an example of the three-dimensional graph displayed by the display device **79** under control of the graph displaying means **94**, that is, a three-dimensional scatter diagram **96** as a sort of three-dimensional graph. The three-dimensional scatter diagram **96** has a systolic-blood-pressure axis **98**, an augmentation-index axis **100**, and a pulse-wave-velocity axis **102**, and shows a circle and a day **104** representing a systolic blood pressure value *BP_{SYS}*, an augmentation index *AI*, and a pulse-wave velocity *PWV* that are measured in the current measuring operation carried out on the day, and a circle and a day **106** representing a systolic blood pressure value *BP_{SYS}*, an augmentation index *AI*, and a pulse-wave velocity *PWV* that were measured in the past measuring operation carried out on the day and are stored in the memory device **72**. Thus, a person such as a doctor can recognize respective time-wise changes of the systolic blood pressure value *BP_{SYS}*, augmentation index *AI*, and pulse-wave velocity *PWV* of the patient. In addition, the diagram **96** shows a measurement figure **108** representing the three values measured in the current measurement, and having a rectangular parallelepiped shape.

The three-dimensional scatter diagram **96** additionally shows a standard figure **110** representing respective upper-limit values of respective normal ranges (i.e., respective standard values) of the systolic blood pressure value *BP_{SYS}*, augmentation index *AI*, and pulse-wave velocity *PWV*, and having a cubic shape. In the example shown in FIG. 7, the respective upper-limit values of respective normal ranges of the systolic blood pressure value *BP_{SYS}*, augmentation index

AI, and pulse-wave velocity PWV are 140 mmHg, 20%, and 1,400 cm/s, respectively.

By comparing the measurement figure 108, or the position of the circle 104, with the standard figure 110 in the three-dimensional scatter graph 96 shown in FIG. 7, a person can easily recognize that the patient's augmentation index AI and pulse-wave velocity PWV are normal but the patient's systolic blood pressure BP_{sys} is too high. In addition, by comparing the circle 104 with the circle 106, the person can recognize that the patient's systolic blood pressure BP_{sys} is being improved.

FIGS. 8 and 9 are flow charts for explaining the control functions of the CPU 76, shown in the diagrammatic view of FIG. 6.

In FIG. 8, first, the CPU carries out Steps S1 through S3 (hereinafter, the term "Step(s)" is omitted) corresponding to the optimum-pressing-position determining means 80. First, at S1, the CPU operates the pressing device 62 to change the pressure in the pressure chamber 56, thereby changing the pressing force HDP applied to the pressure-pulse-wave sensor 54, to the prescribed first pressing force HDPI. The first pressing force HDPI is experimentally determined, in advance, such that an S/N ratio of the carotid pulse wave wc detected by each of the pressure sensing elements E is so great as to be able to determine, with sufficiently high accuracy, a magnitude of a peak point pc of a heartbeat-synchronous pulse of the carotid pulse wave wc.

Subsequently, at S2, the CPU judges whether the pressing-position changing condition (i.e., the APS starting condition) is satisfied, i.e., whether a current highest-pressure sensing element EM as one of the pressure sensing elements E of the pressure-pulse-wave sensor 54 is positioned in one of the prescribed opposite end portions of the array of elements E. If a negative judgment is made at S2, the control jumps to S4, described later.

On the other hand, if a positive judgment is made at S2, that is, if the current position of the pressure-pulse-wave sensor 54 relative to the carotid artery 46 is not appropriate, the control goes to S3, i.e., an APS controlling routine in which the pressure-pulse-wave sensor 54 is positioned at an optimum pressing position where the current highest-pressure sensing element EM is located at substantially the middle of the array of elements E. More specifically described, the CPU carries out the following pressing-position changing operation: After the pressing device 62 once moves the pressure-pulse-wave sensor 54 away from the body surface 50, the widthwise-direction moving device 64 moves the pressing device 62 and the sensor 54 over a prescribed distance, and then the pressing device 62 presses again the sensor 54 with the first pressing force HDPI. In this state, the CPU judges again whether the prescribed pressing-position changing condition is satisfied. The CPU repeats those actions and judgment till the pressing-position changing condition is not satisfied any longer, more specifically described, till the current highest-pressure sensing element EM is positioned in the prescribed middle portion of the array of elements E.

After the pressure-pulse-wave sensor 54 is positioned at the optimum pressing position at S3, or if a negative judgment is made at S2, the control goes to S4 to determine the current highest-pressure sensing element EM as a final highest-pressure sensing element EM of the sensor 54. S4 is followed by S5 corresponding to the pressing-force determining means 82, i.e., an HDP controlling routine. More specifically described, the CPU operates the pressing device 62 so that the pressing force HDP applied to the pressure-pulse-wave sensor 54 is continuously increased from the

first pressing force HDPI. During this increasing of the pressing force HDP, the CPU determines an optimum pressing force HDPO at which a pulse pressure PP of a heartbeat-synchronous pulse of the carotid pulse wave wc detected by the highest-pressure sensing element EM determined at S4 exceeds the prescribed lower-limit pulse pressure PPL, and maintains the pressing force HDP applied to the sensor 54, at the thus determined optimum pressing force HDPO.

Then, the control goes to S6 where the CPU reads in respective one-heartbeat lengths of the pressure-pulse-wave signal SM2 supplied from the highest-pressure sensing element EM determined at S4 and the heart-sound signal SH supplied from the heart-sound microphone 68. Then, the control goes to S7 to stop the air pump 58 and operate the pressure control valve 60 so that the pressing force HDP applied to the sensor 54 is decreased down to an atmospheric pressure.

Subsequently, the control proceeds with S8 through S11 corresponding to the augmentation-index calculating means 88. First, at S8, the CPU subjects, to a fourth-order differentiation, a heartbeat-synchronous pulse of the carotid pulse wave wc that is represented by the one-heartbeat length of the pressure-pulse-wave signal SM2 read in at S6, more specifically described, a portion of the carotid pulse wave wc that is located between a rising point and a peak point pc of the pulse, so as to identify an inflection point or a local-maximum point located between the rising point and the peak point pc. The CPU determines a magnitude of the thus identified inflection point or local-maximum point as a magnitude of the peak point pi of the incident-wave component wi.

Subsequently, at S9, the CPU determines a magnitude of the carotid pulse wave wc at the time of occurrence of the peak point of reflected-wave component wr. More specifically described, in the case where the peak point pi of incident-wave component wi, determined at S8, does not coincide with a maximum point of the heartbeat-synchronous pulse of the carotid pulse wave wc, a magnitude of the maximum point of the heartbeat-synchronous pulse of the carotid pulse wave wc is determined as the magnitude of the carotid pulse wave wc at the time of occurrence of the peak point of the reflected-wave component wr. On the other hand, in the case where the peak point pi of the incident-wave component wi, determined at S8, coincides with the maximum point of the heartbeat-synchronous pulse of the carotid pulse wave wc, a magnitude of the first local-maximum point following the peak point pi of the incident-wave component wi is determined as the magnitude of the carotid pulse wave wc at the time of occurrence of the peak point of reflected-wave component wr.

Then, at S10, the CPU determines a pulse pressure PP of the heartbeat-synchronous pulse of the carotid pulse wave wc, read in at S6. S10 is followed by S11 to calculate a pressure difference ΔP by subtracting the magnitude of the carotid pulse wave wc at the time of occurrence of the peak point of the incident-wave component wi, determined at S8, from the magnitude of the carotid pulse wave wc at the time of occurrence of the peak point of the reflected-wave component wr, determined at S9. Additionally, the CPU calculates an augmentation index AI by substituting the augmentation-index calculating expression shown in the form of Expression 1, with the thus calculated pressure difference ΔP and the pulse pressure PP determined at S10, and operates the display device 79 to display the thus calculated augmentation index AI in digits.

Then, the control goes to S12 through S14 corresponding to the pulse-wave-velocity-related-information obtaining

11

means 90. At S12, the CPU calculates a propagation distance L by substituting Expression 2 with the stature T that had been inputted through the input device 72 before the current operation of the CPU is started. Subsequently, at S13, the CPU identifies a starting point of a second heart sound II based on the one-heartbeat length of the heart-sound signal SH, read in at S6, and a dirotic-notch point of a heartbeat-synchronous pulse of the carotid pulse wave wc based on the one-heartbeat length of the pressure-pulse-wave signal SM2, read in at S6, and calculates, as a pulse-wave propagation time DT, a time difference between a time of detection of the starting point of the second heart sound II and a time of detection of the dirotic-notch point of the carotid pulse wave wc. Then, at S14, the CPU calculates a pulse-wave velocity PWV by substituting Expression 3 with the propagation distance L and the propagation time DT respectively calculated at S12 and S13, and operates the display device 79 to display the thus calculated pulse-wave velocity PWV in digits.

Next, S15 and the following steps shown in FIG. 9 will be described. At S15, the CPU starts the air pump 24 and operate the pressure control valve 18 so as to start quickly increasing the cuff pressure PC. Subsequently, at S16, the CPU judges whether the cuff pressure PC has exceeded a target pressure PC_M pre-set at 180 mmHg. S16 is repeated until a positive judgment is made, while the cuff pressure PC is quickly increased. Meanwhile, if a positive judgment is made at S16, the control goes to S17 to stop the air pump 24 and operate the pressure control valve 18 so as to start slowly decreasing the cuff pressure PC at a rate of about 3 mmHg/sec.

Next, the control goes to S18 to S20 corresponding to the blood-pressure determining means 86. At S18, the CPU determines, based on change of respective amplitudes of successive heartbeat-synchronous pulses of the brachial pulse wave wb represented by the cuff-pulse-wave signal SM1 continuously obtained during the slow decreasing of the cuff pressure PC, a systolic blood pressure BP_{SYS} , a mean blood pressure BP_{MEAN} , and a diastolic blood pressure BP_{DIA} of the patient, according to well-known oscillometric blood-pressure determining algorithm. Then, at S19, the CPU judges whether the determination of the blood-pressure values BP has completed at S18.

S18 and S19 are repeated till a positive judgment is made at S19. Thus, the blood-pressure determining algorithm is continued. Meanwhile, if a positive judgment is made at S19, the control goes to S20 to operate the display device 79 to display the thus determined systolic, mean, and diastolic blood-pressure values BP_{SYS} , BP_{MEAN} , BP_{DIA} in digits. Then, the control goes to S21 to operate the pressure control valve 18 to decrease the cuff pressure PC to an atmospheric pressure. In the present flow chart, S15 through S17 and S21 correspond to the cuff-pressure changing means 84.

Then, the control goes to S22 corresponding to the evaluation-information storing means 92. At S22, the CPU stores, in a certain memory area of the memory device 72, the augmentation index AI calculated at S11, the pulse-wave velocity PWV calculated at S14, and the systolic blood pressure BP_{SYS} calculated at S18, as a set of evaluation information obtained, from the patient identified by the identification code inputted before the current operation of the CPU is started, together with data indicative of a date when the set of evaluation information is obtained from the patient.

Then, the control goes to S23 corresponding to the graph displaying means 94. At S23, the CPU operates the display device 79 to display the three-dimensional scatter diagram

12

96, as shown in FIG. 7, and additionally display, in the graph 96, the circle 104 and the measurement figure 108 each of which represents the augmentation index AI calculated at S11, the pulse-wave velocity PWV calculated at S14, and the systolic blood pressure BP_{SYS} calculated at S18; the standard figure 110; and the circle 106 representing the past set of evaluation information that has been stored, in the memory device 72, for the patient identified by the identification code inputted before the current operation of the CPU is started.

In the above-described embodiment, a person such as a doctor can graphically recognize, from the three-dimensional scatter diagram 96 displayed on the display device 79, the respective measured values of the systolic blood pressure BP_{SYS} , the augmentation index AI, and the pulse-wave velocity PWV of the patient. Thus, the person can easily make a total evaluation of the circulatory organ of the patient based on the three parameters.

Moreover, in the above-described embodiment, the three-dimensional scatter diagram 96 shows the standard figure 110 representing the respective upper limits of the respective normal ranges of systolic blood pressure BP_{SYS} , augmentation index AI, and pulse-wave velocity PWV. Therefore, the person can more easily evaluate the circulatory organ by comparing the measurement figure 108 or the circle 104 representing the respective measured values of the systolic blood pressure BP_{SYS} , the augmentation index AI, and the pulse-wave velocity PWV of the patient, with the standard figure 110.

Moreover, in the above-described embodiment, the three-dimensional scatter diagram 96 shows the circles 104, 106 that cooperate with each other to represent the respective timewise changes of the systolic blood pressure BP_{SYS} , the augmentation index AI, and the pulse-wave velocity PWV of the patient. Thus, the person can graphically recognize the respective timewise changes of the systolic blood pressure BP_{SYS} , the augmentation index AI, and the pulse-wave velocity PWV of the patient, and accordingly can easily recognize the timewise change of dynamic function of the circulatory organ of the patient. Therefore, the person can easily evaluate the effect of medication or the improvement of patient's habits.

While the present invention has been described in its preferred embodiment by reference to the drawings, it is to be understood that the invention may otherwise be embodied.

For example, in the illustrated embodiment, the three-dimensional scatter diagram 96 is used as a sort of three-dimensional graph. However, other sorts of three-dimensional graphs may be used; such as a radar chart 112 as shown in FIG. 10. In FIG. 10, a solid-line triangle 114 represents respective values of systolic blood pressure BP_{SYS} , augmentation index AI, and pulse-wave velocity PWV of a patient, measured in a current measuring operation; a broken-line triangle 116 represents respective values of systolic blood pressure BP_{SYS} , augmentation index AI, and pulse-wave velocity PWV of the same patient, measured in the preceding or last measuring operation; and a one-dot-chain-line triangle 118 represents respective upper-limit values (i.e., respective standard values) of systolic blood pressure BP_{SYS} , augmentation index AI, and pulse-wave velocity PWV.

Also, in the illustrated embodiment, the three-dimensional scatter diagram 96 represents the systolic blood pressure BP_{SYS} , the augmentation index AI, and the pulse-wave velocity PWV of the subject. However, the systolic blood pressure BP_{SYS} may be replaced with the mean or diastolic blood pressure BP_{MEAN} , BP_{DIA} , and the pulse-wave

13

velocity PWV may be replaced with the pulse-wave propagation time DT.

Also, in the illustrated embodiment, the display device **79** displays the three-dimensional scatter diagram **96** as the three-dimensional graph. However, it is possible to obtain 5 another or other sorts of circulatory-organ evaluation information (e.g., heart rate, HR) and display a four- or higher-dimensional graph additionally representing the other sorts of evaluation information.

Also, the illustrated circulatory-organ evaluating apparatus **10** has the functions of measuring the blood pressure BP, the augmentation index AI, and the pulse-wave velocity PWV of living subject. However, the illustrated apparatus **10** may be so modified as not to have those functions. In the 10 latter case, other devices may measure those parameters, and supply the respective measured values of the parameters to the thus modified apparatus **10**.

Also, the illustrated apparatus **10** substantially simultaneously measure the blood pressure BP, the augmentation index AI, and the pulse-wave velocity PWV of living subject. However, it is not essentially required to substantially simultaneously measure those parameters, so long as the respective measured values of those parameters are 20 simultaneously displayed.

Also, in the illustrated embodiment, the blood pressure BP is measured at the brachial portion **14**, the augmentation index AI is measured at the cervical portion **38**, and the pulse-wave velocity PWV is measured at the portion between the subject's heart (i.e., aortic valve) and the cervical portion **38**. However, those parameters may be 25 measured at other portions of the subject. For example, blood pressure BP may be measured at a femoral portion or an ankle; augmentation index AI may be measured at a brachial portion **14**; and pulse-wave velocity PWV may be measured at a portion between a brachium and an ankle. 35

Also, the generally-used augmentation-index calculating expression (i.e., Expression 1) employs the pulse pressure PP as its denominator. However, though the denominator may be replaced with the magnitude of the carotid pulse wave wc at the time of occurrence of peak point of the incident-wave component wi, a value calculated according to the thus modified expression adequately reflects a degree of arteriosclerosis of the subject. Therefore, the pulse pressure PP in Expression 1 may be replaced with the magnitude of the carotid pulse wave wc at the time of occurrence of 40 peak point of the incident-wave component wi. 45

The present invention may be embodied with other various changes without departing from the spirit of the invention.

14

What is claimed is:

1. A circulatory-organ evaluating apparatus for evaluating a circulatory organ of a living subject, comprising:

a display device which displays a three-dimensional graph representing a blood pressure, an augmentation index, and a pulse-wave-velocity-related information of the subject; and

a control device which controls the display device to simultaneously and graphically display the blood pressure, the augmentation index, and the pulse-wave-velocity-related information of the subject.

2. A circulatory-organ evaluating apparatus recited in claim 1, wherein the three-dimensional graph includes a standard figure representing a standard blood pressure, a standard augmentation index, and standard pulse-wave-velocity-related information.

3. A circulatory-organ evaluating apparatus recited in claim 1, wherein the three-dimensional graph represents respective time-wise changes of the blood pressure, the augmentation index, and the pulse-wave-velocity-related information of the subject.

4. A circulatory-organ evaluating apparatus according to claim 3, further comprising a memory device which stores at least one past set of evaluation information including at least a blood pressure, an augmentation index, and pulse-wave-velocity-related information that were measured from the subject in at least one past measuring operation, wherein the control device controls, based on the set of evaluation information stored in the memory device, the display device to display the three-dimensional graph representing the respective time-wise changes of the blood pressure, the augmentation index, and the pulse-wave-velocity-related-information information of the subject.

5. A circulatory-organ evaluating apparatus according to claim 1, further comprising an input device which is operable by an operator to input identification information identifying the subject.

6. A circulatory-organ evaluating apparatus according to claim 1, further comprising:

a blood-pressure measuring device which measures the blood pressure of the subject;

an augmentation-index determining device which determines the augmentation index of the subject; and a pulse-wave-velocity-related-information obtaining device which obtains the pulse-wave-velocity-related information of the subject.

* * * * *

专利名称(译)	循环器官评估装置		
公开(公告)号	US6884221	公开(公告)日	2005-04-26
申请号	US10/370617	申请日	2003-02-24
[标]申请(专利权)人(译)	日本柯林股份有限公司		
申请(专利权)人(译)	COLIN CORPORATION		
当前申请(专利权)人(译)	欧姆龙保健CO., LTD.		
[标]发明人	NARIMATSU KIIYOYUKI OGURA TOSHIHIKO TAMPO AKIRA		
发明人	NARIMATSU, KIIYOYUKI OGURA, TOSHIHIKO TAMPO, AKIRA		
IPC分类号	A61B5/021 A61B5/026 A61B5/0285 A61B5/0205 A61B5/022 A61B5/0245 A61B5/025 A61B10/00 A61B5/02 A61B5/00		
CPC分类号	A61B5/021 A61B5/0285 A61B5/6822 A61B5/743 A61B5/7445 A61B2562/0247 A61B2562/043		
代理机构(译)	OLIFF & 贝里奇		
优先权	2002180370 2002-06-20 JP		
其他公开文献	US20030236464A1		
外部链接	Espacenet USPTO		

摘要(译)

一种用于评估生物体的循环器官的循环器官评估装置，包括显示装置，以及控制该显示装置同时且图形地显示血压，增强指数和脉搏波速度的控制装置。该主题的相关信息。

