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(54) **MONITORING CARDIAC BLOOD FLOW  
BALANCE RELATIONSHIP BETWEEN THE  
RIGHT AND LEFT HEART CHAMBERS AND  
CARDIAC REGULATION**

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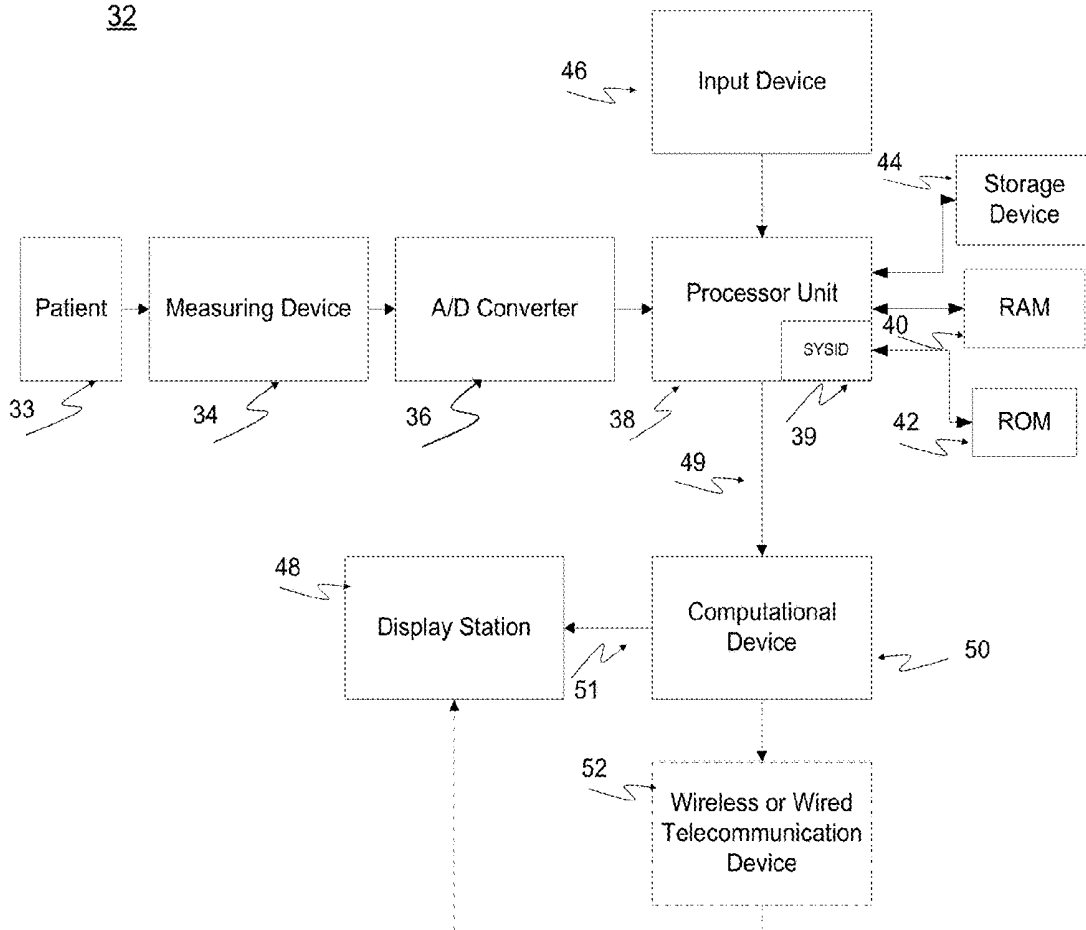
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(57) **ABSTRACT**

A system for measuring of cardiac blood flow balance parameter between the right chamber of the heart and the left chamber of the heart includes a sensor device for measuring one of blood pressure and blood flow rate and blood constituent concentration of a patient so as to generate an arterial pulse signal. A processing unit is responsive to the arterial pulse signal for generating a full arterial pulse signal, an arterio-venous pulse signal, and a balance parameter, including a balance ratio and a balance trend, and a pulse variability parameter. A computational device is responsive to the balance parameter for further generating a set of physiological parameters. A display station device is responsive to the set of physiological parameters from the computational device for displaying meaningful information.

32



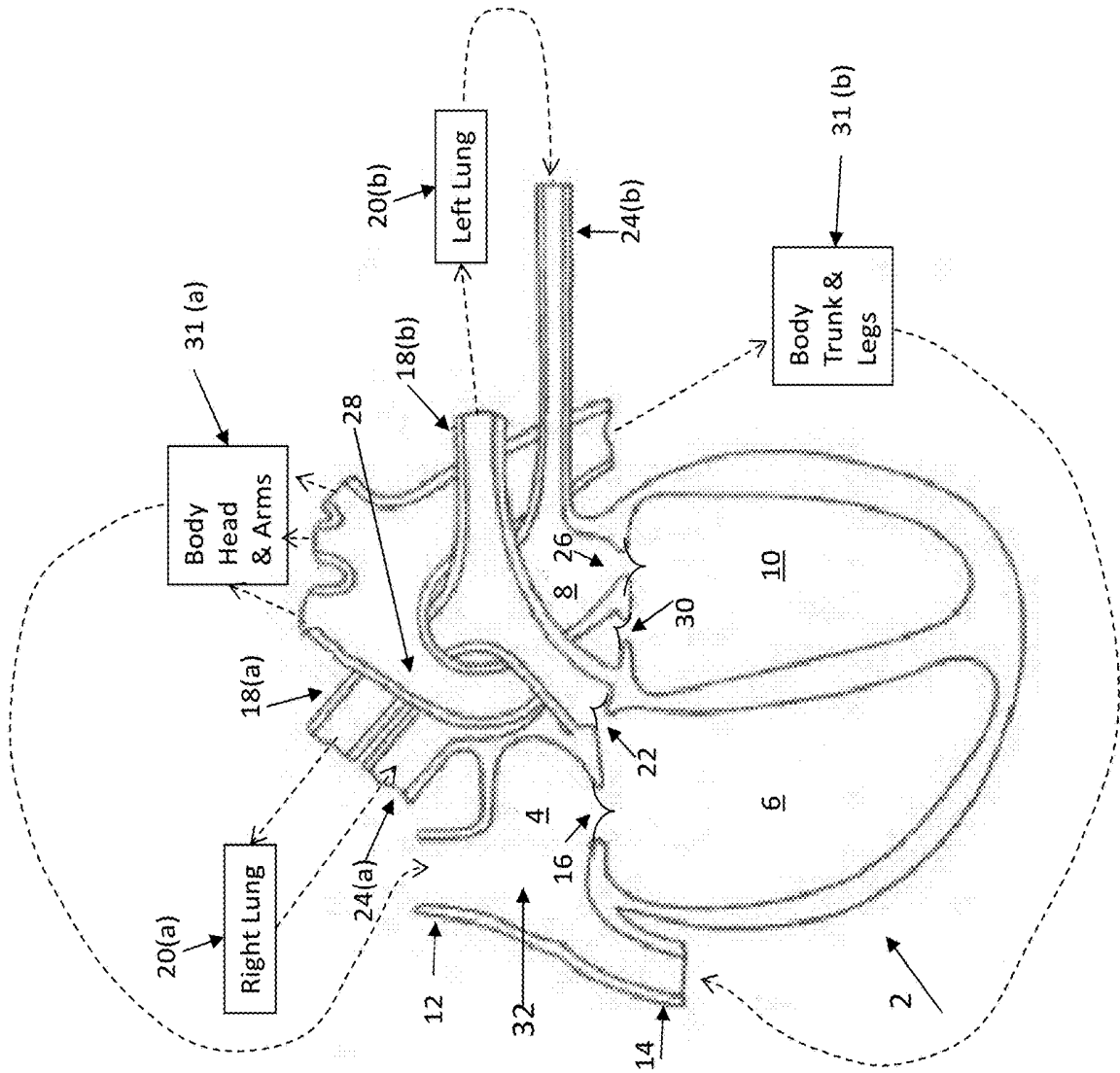


Fig. 1  
(Prior Art)

Fig. 2

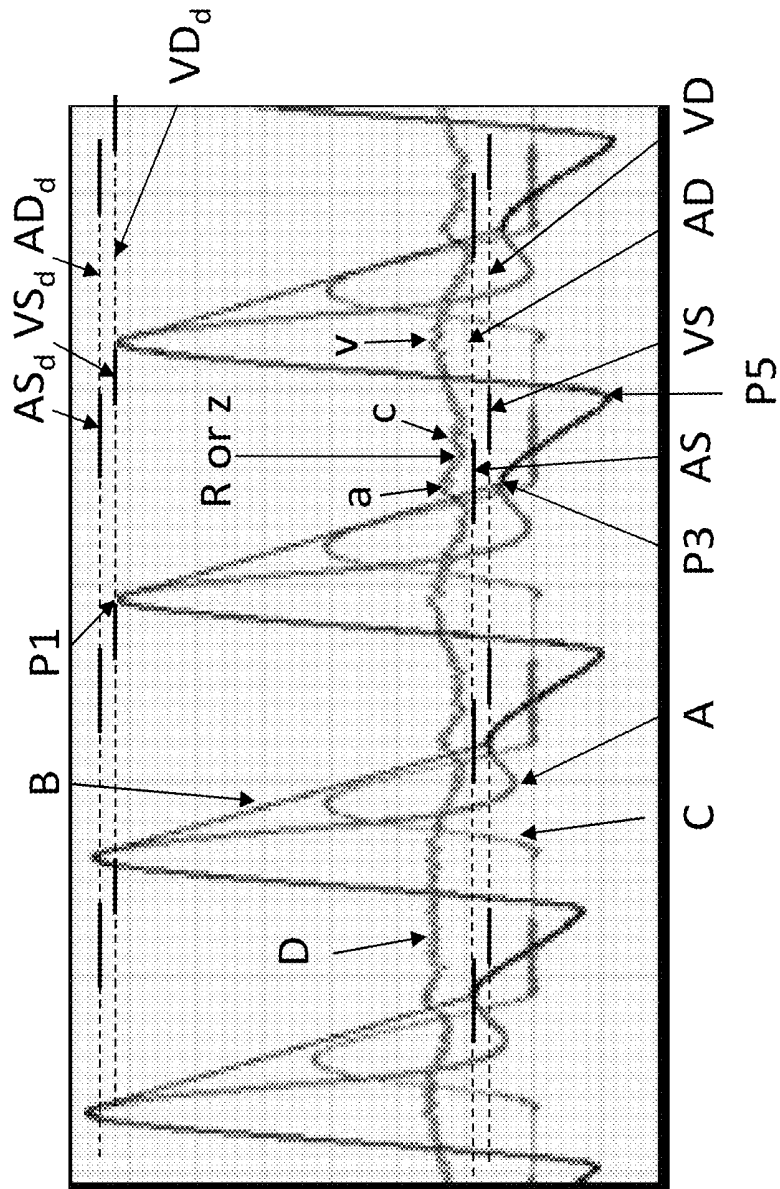


Fig. 3

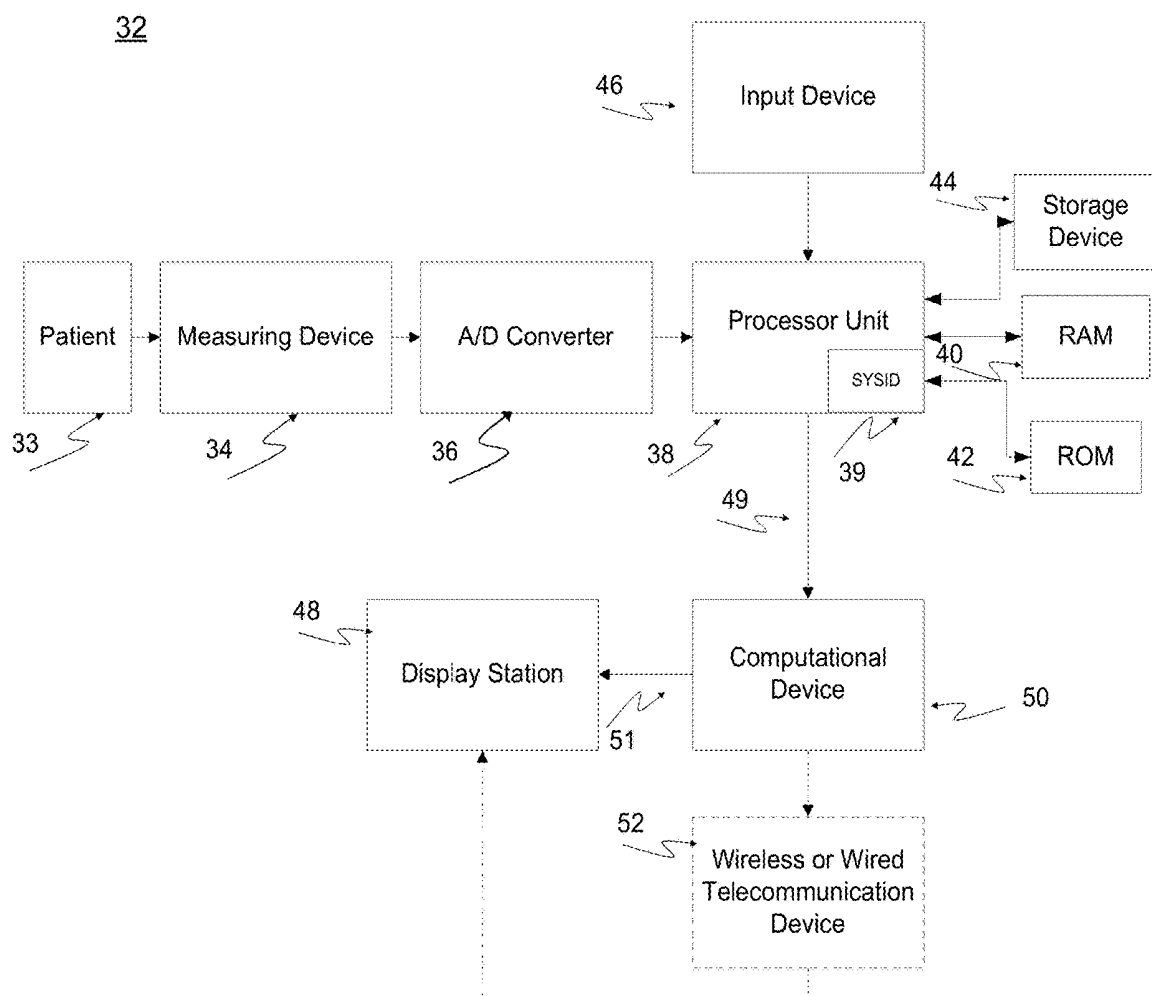
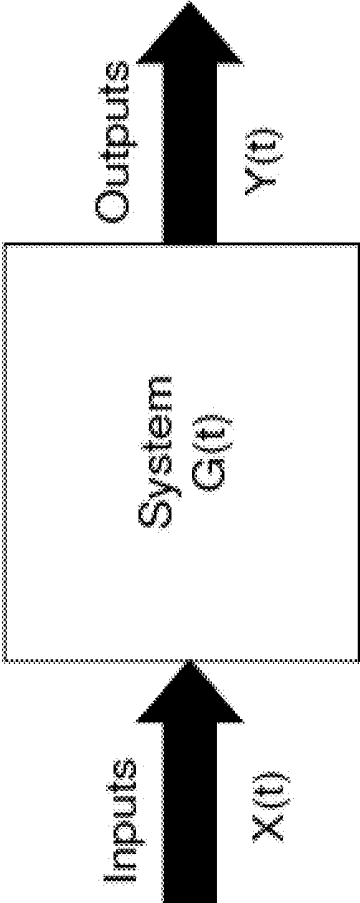


Fig 4.



$Y(t) = G(t) * X(t)$  Time Domain Convolution  
 $Y(f) = G(f) \cdot X(f)$  Frequency Domain Multiplication

Fig. 5

500

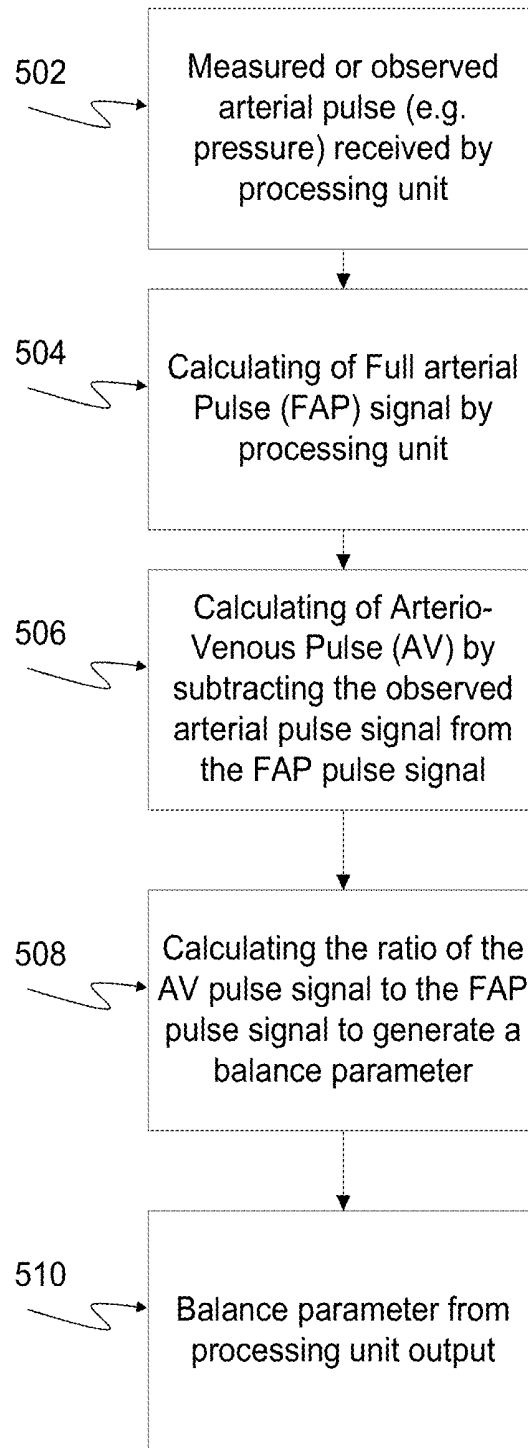


Fig 6

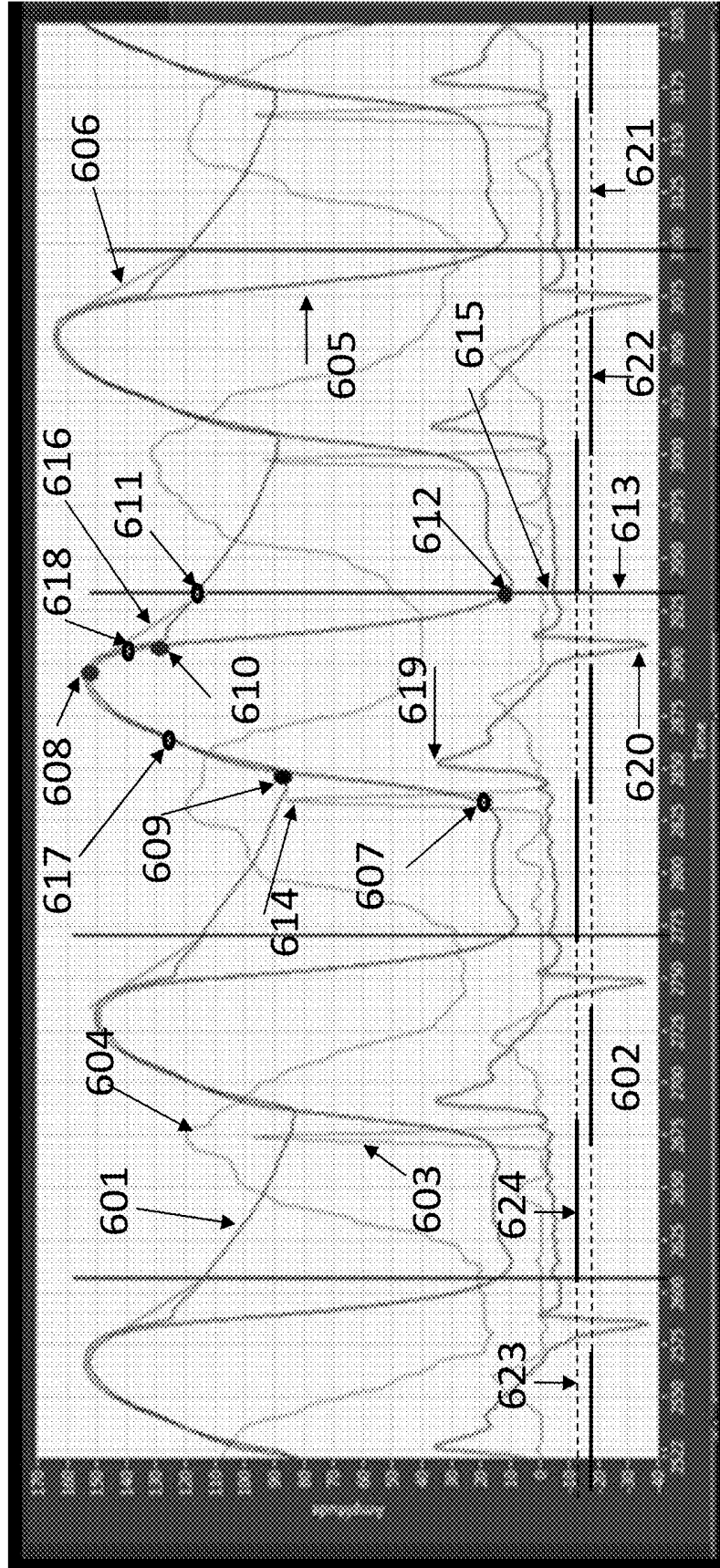


Fig 7

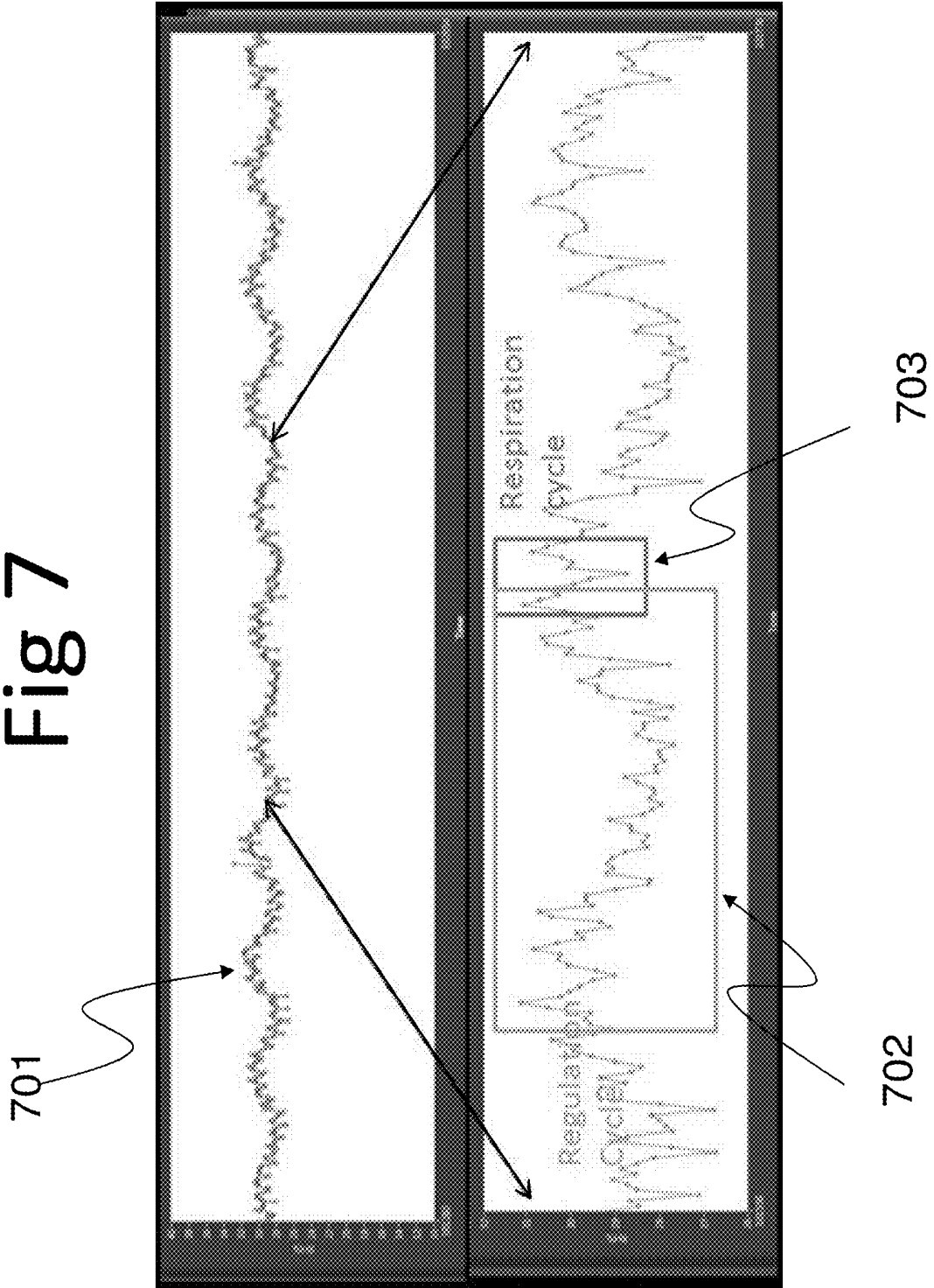


Fig 8(a)

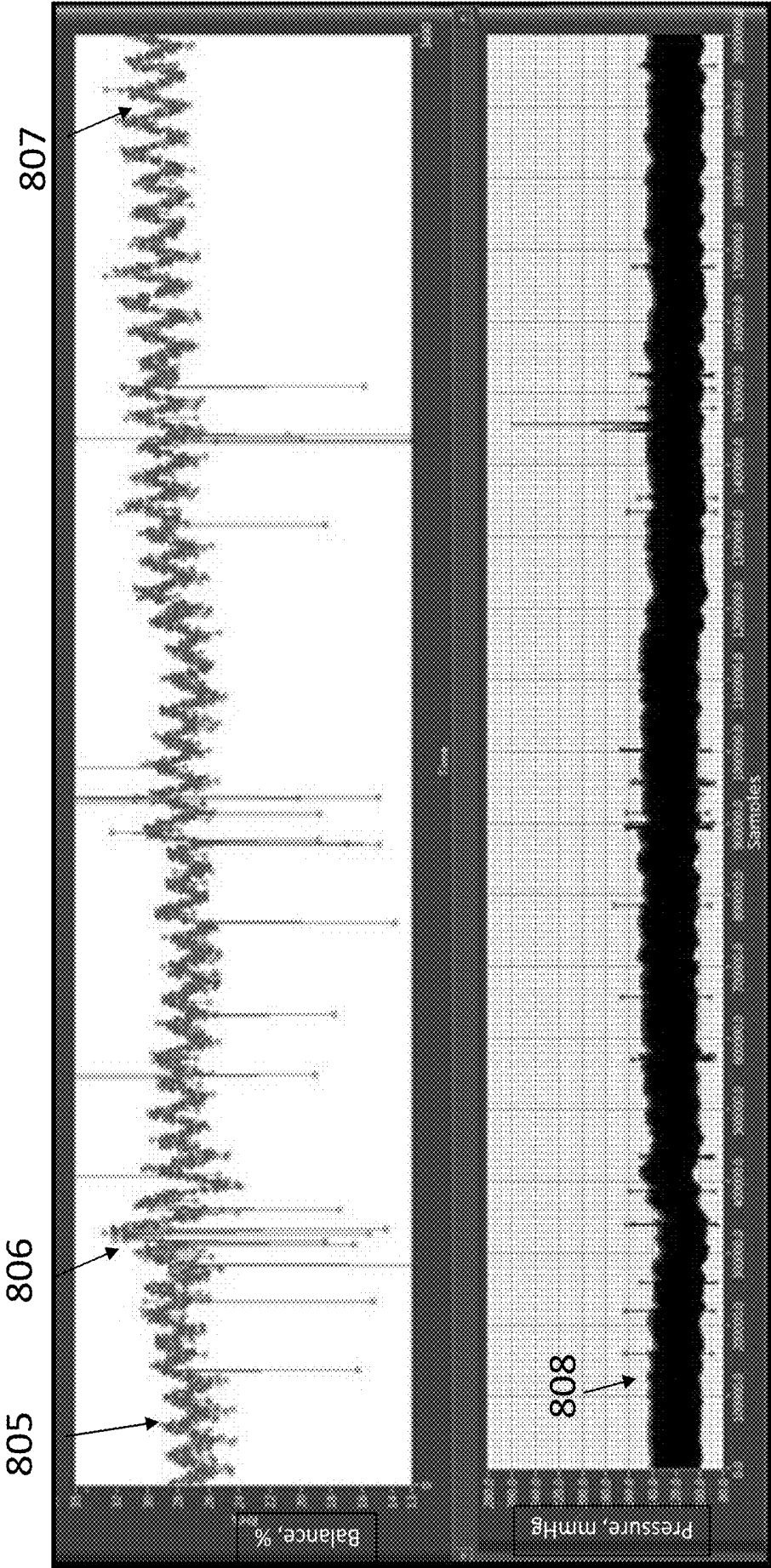
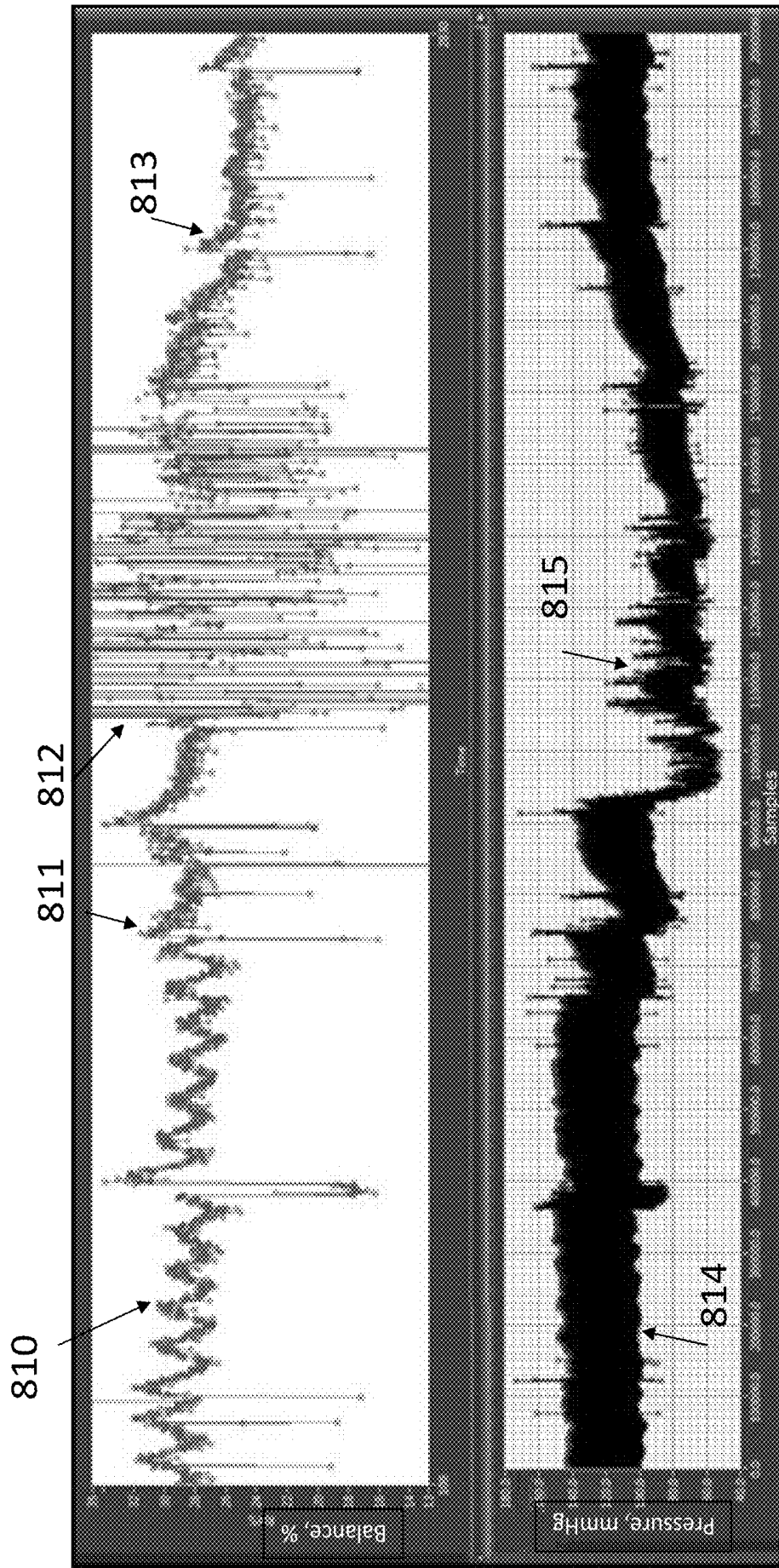


Fig 8(b)



900

Fig. 9

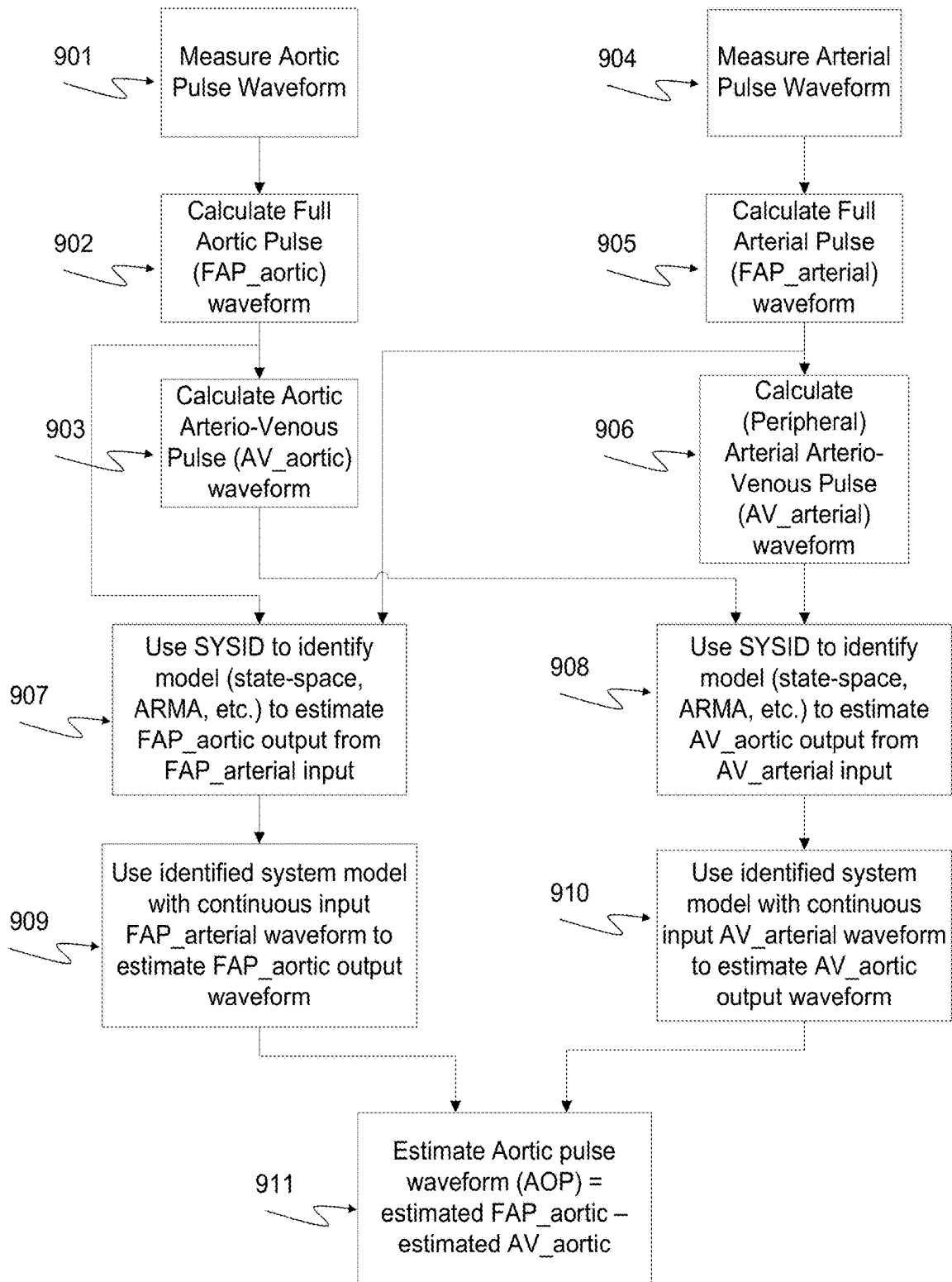
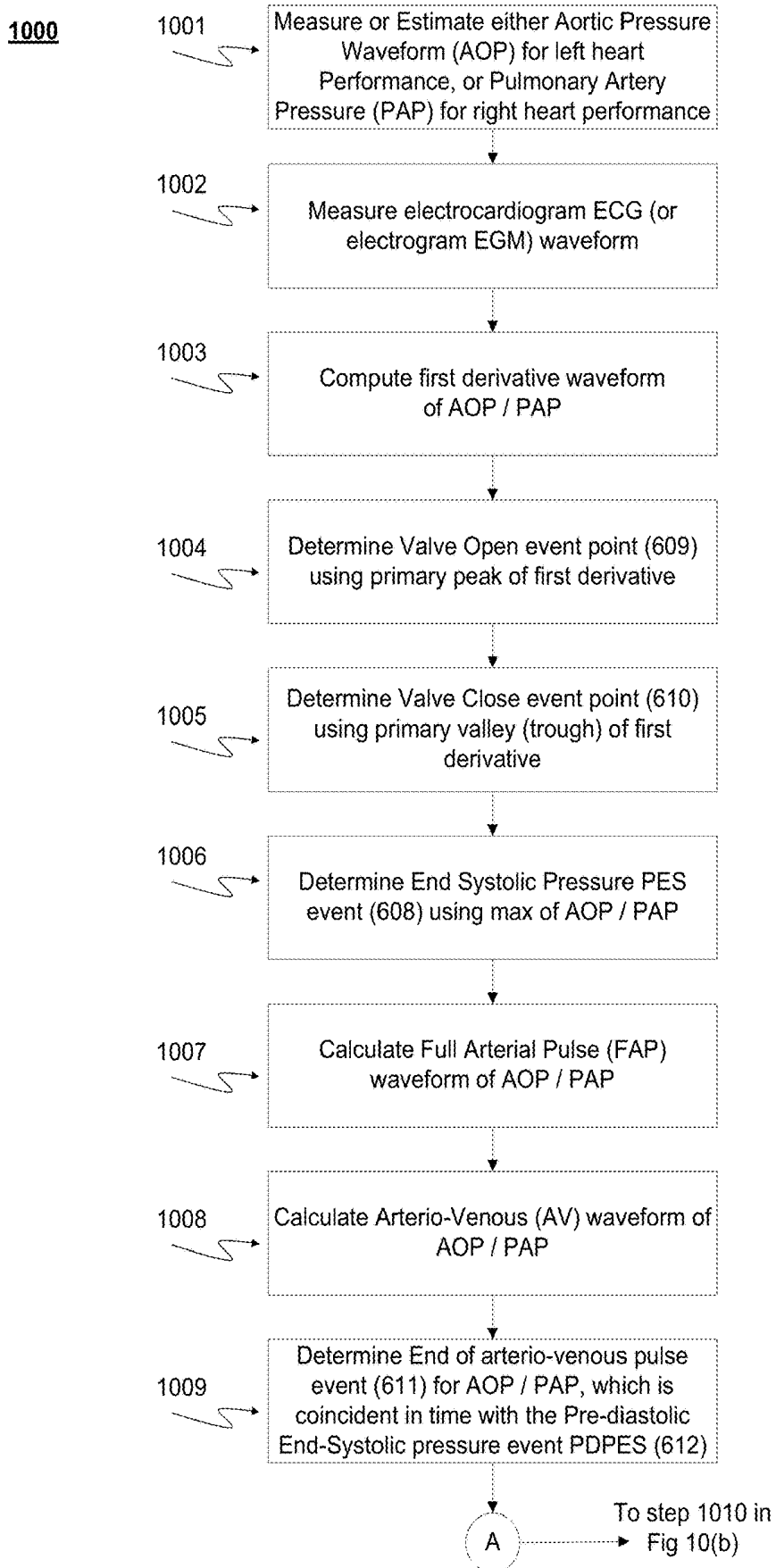
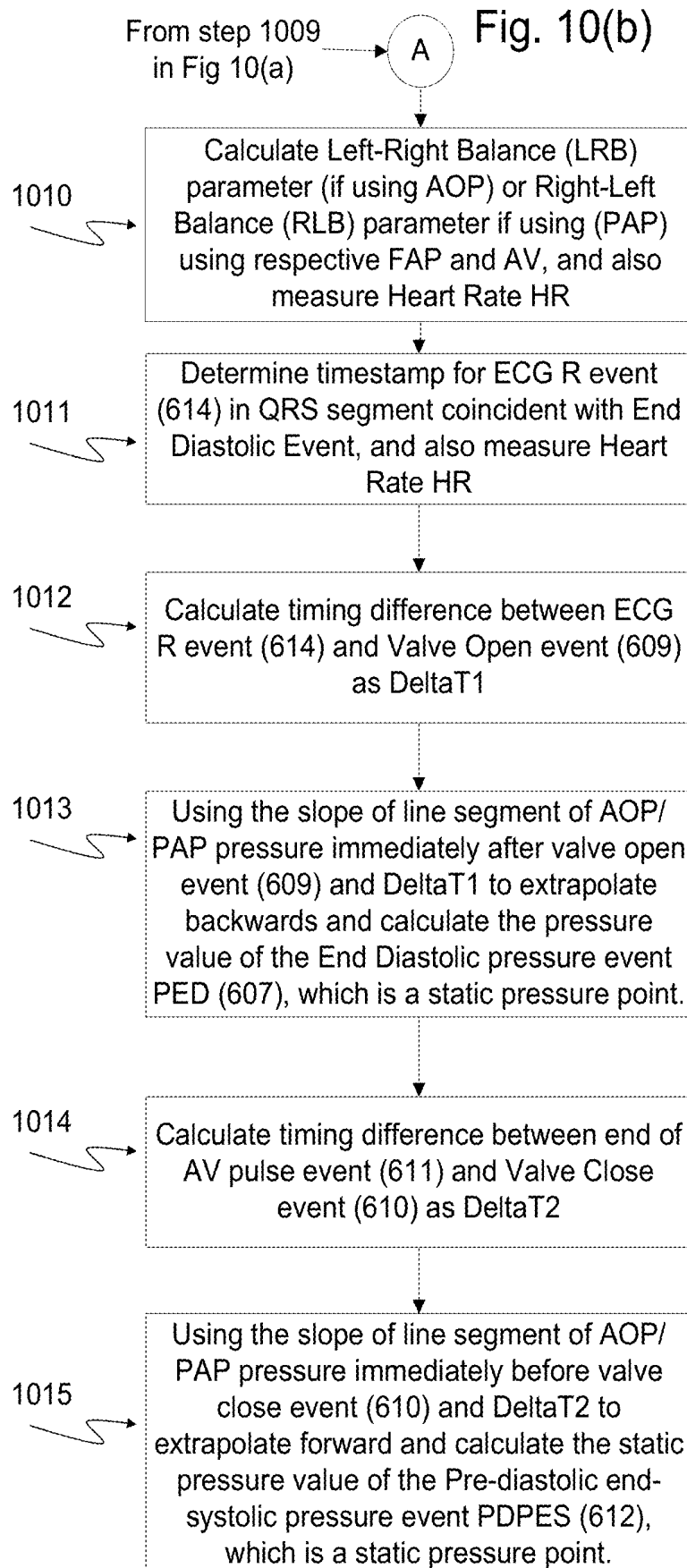


Fig. 10(a)





1100

Fig. 11

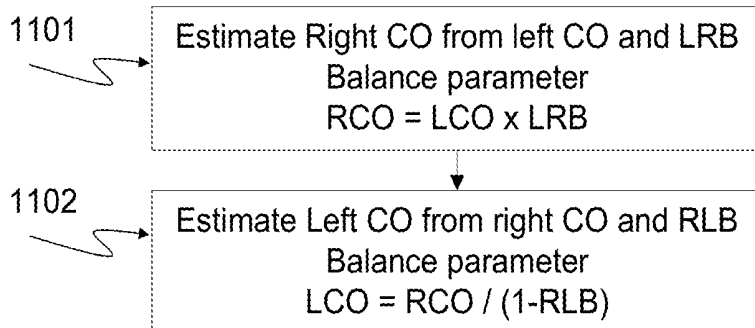


Fig. 12

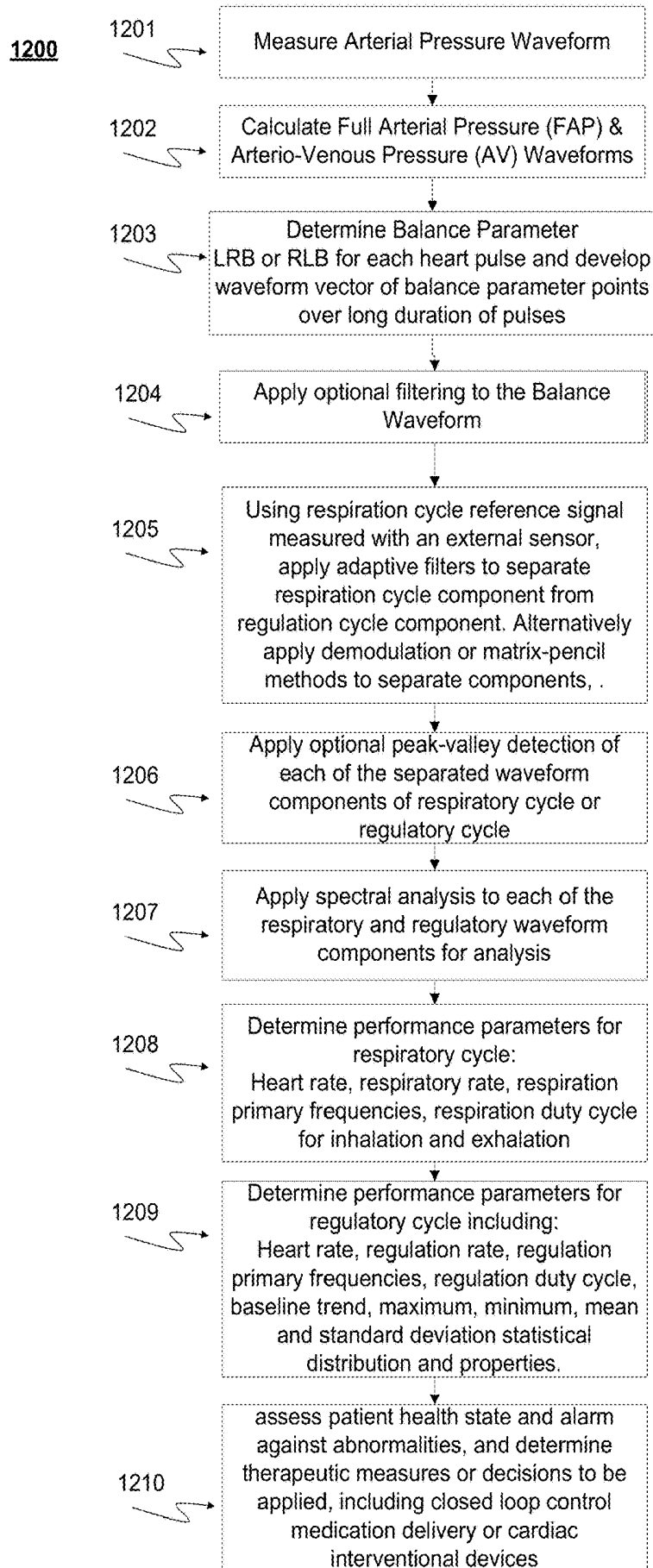


Figure 13

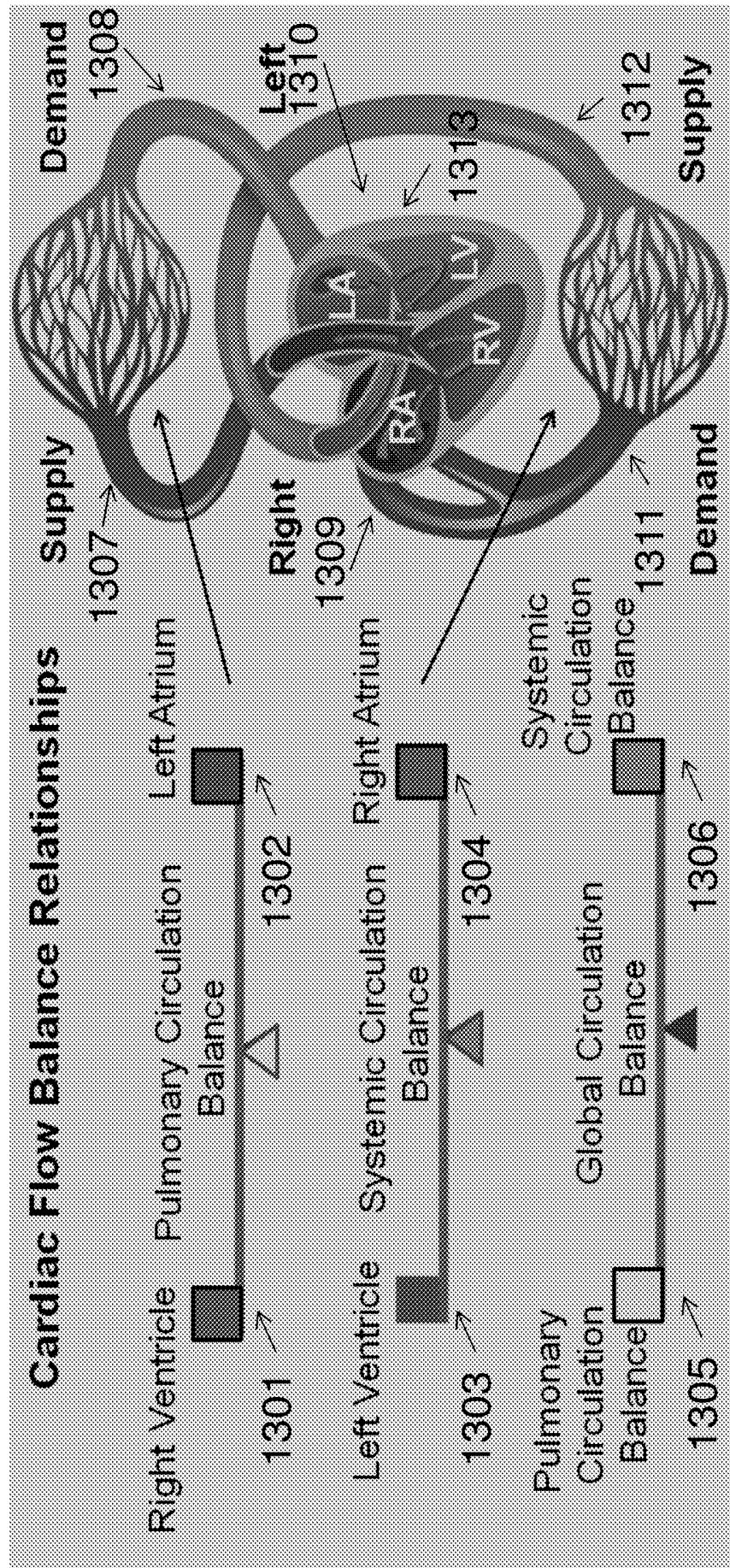


Figure 14

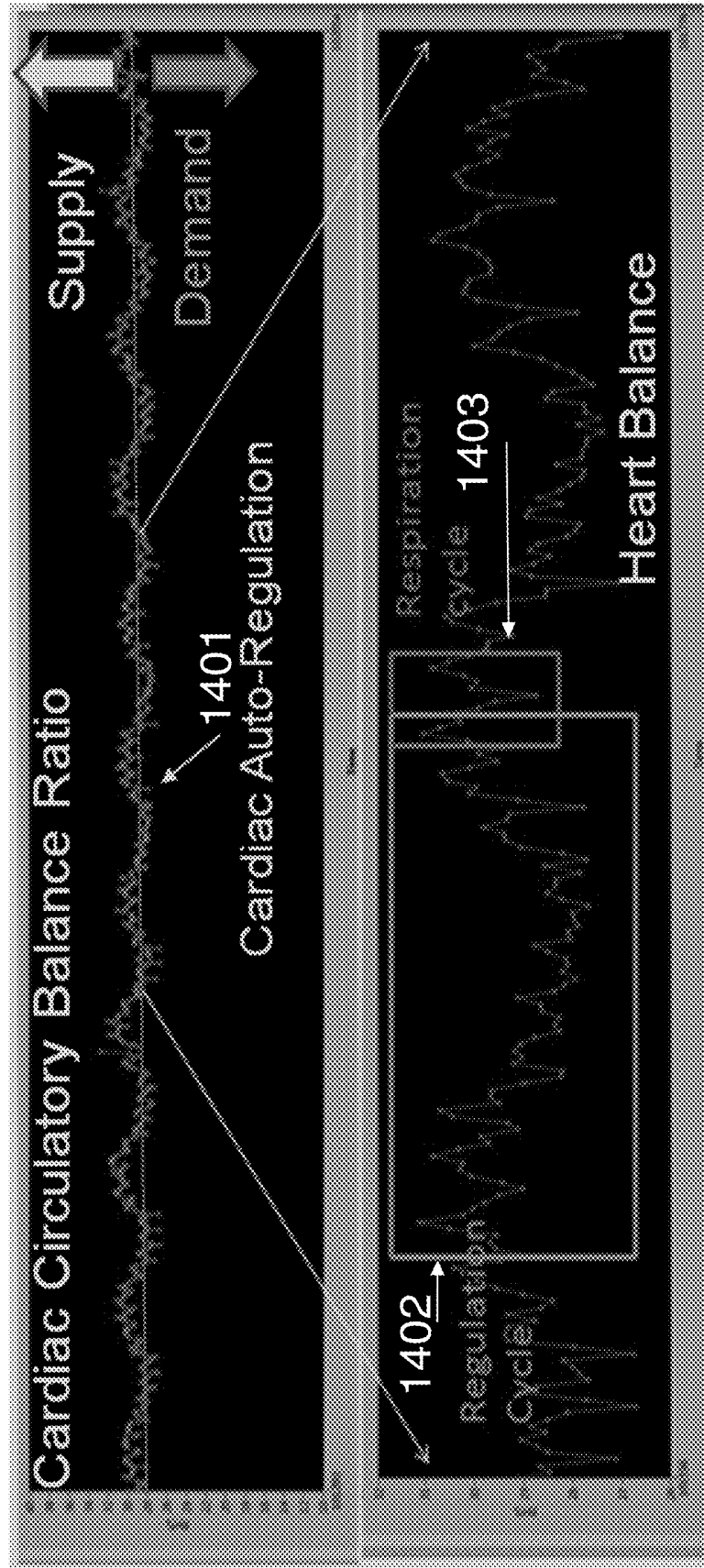


Figure 15

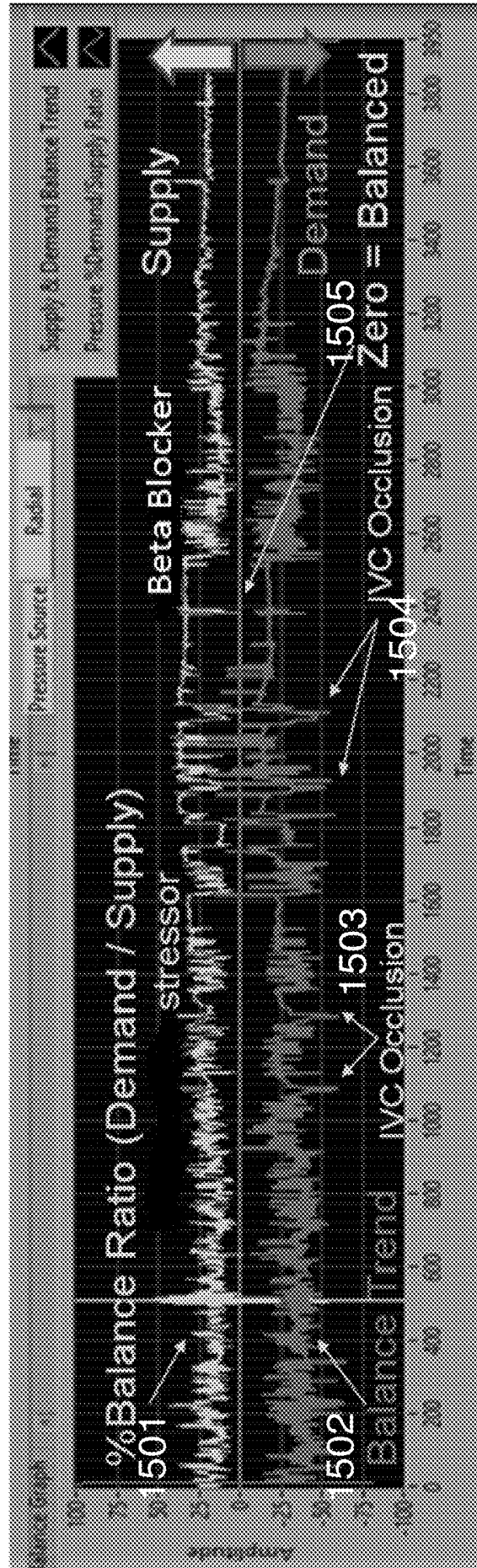
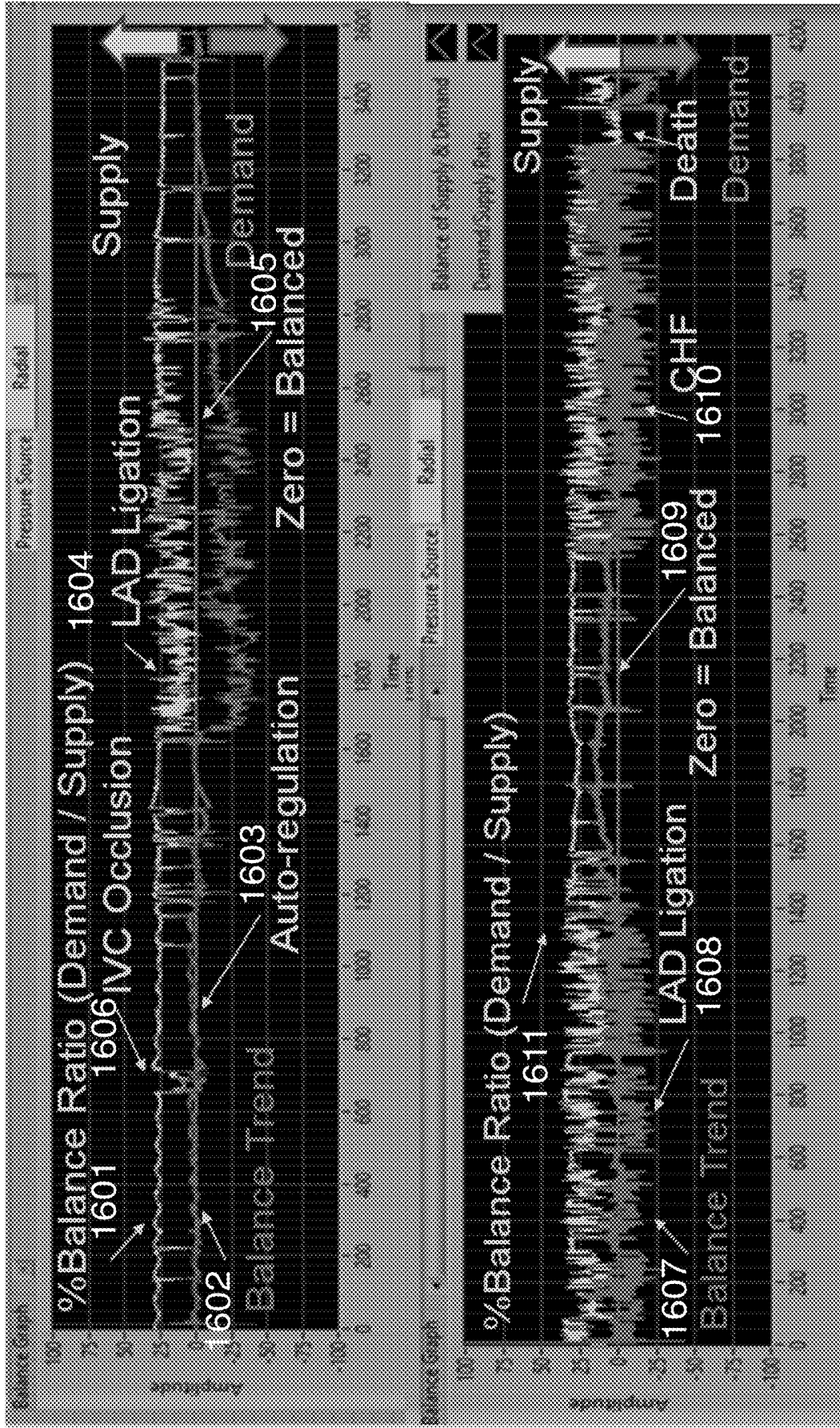
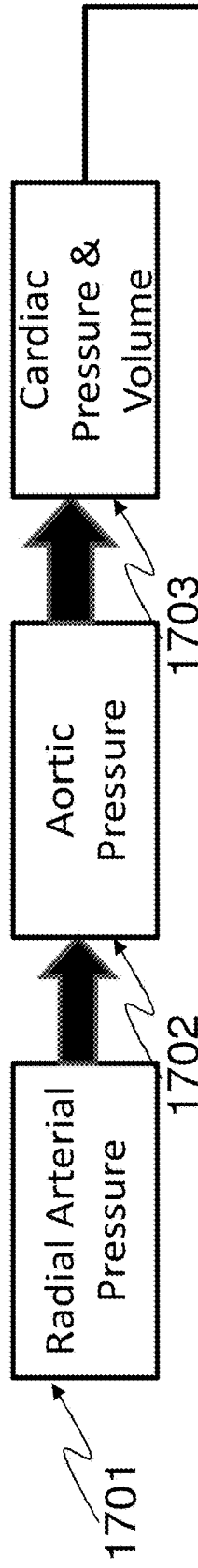


Figure 16



# Figure 17

## Step 1) Single-time invasive calibration with catheter



## Step 2) Non-invasive continuous hemodynamics monitoring

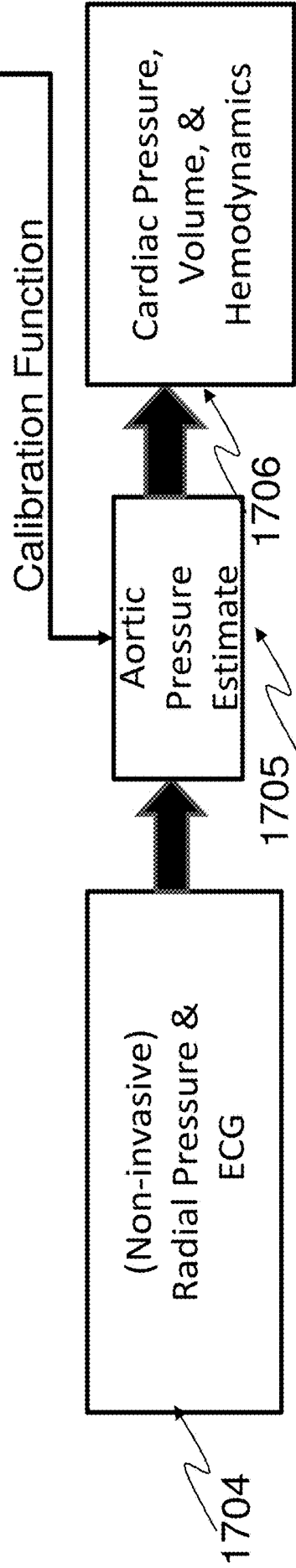


Figure 18

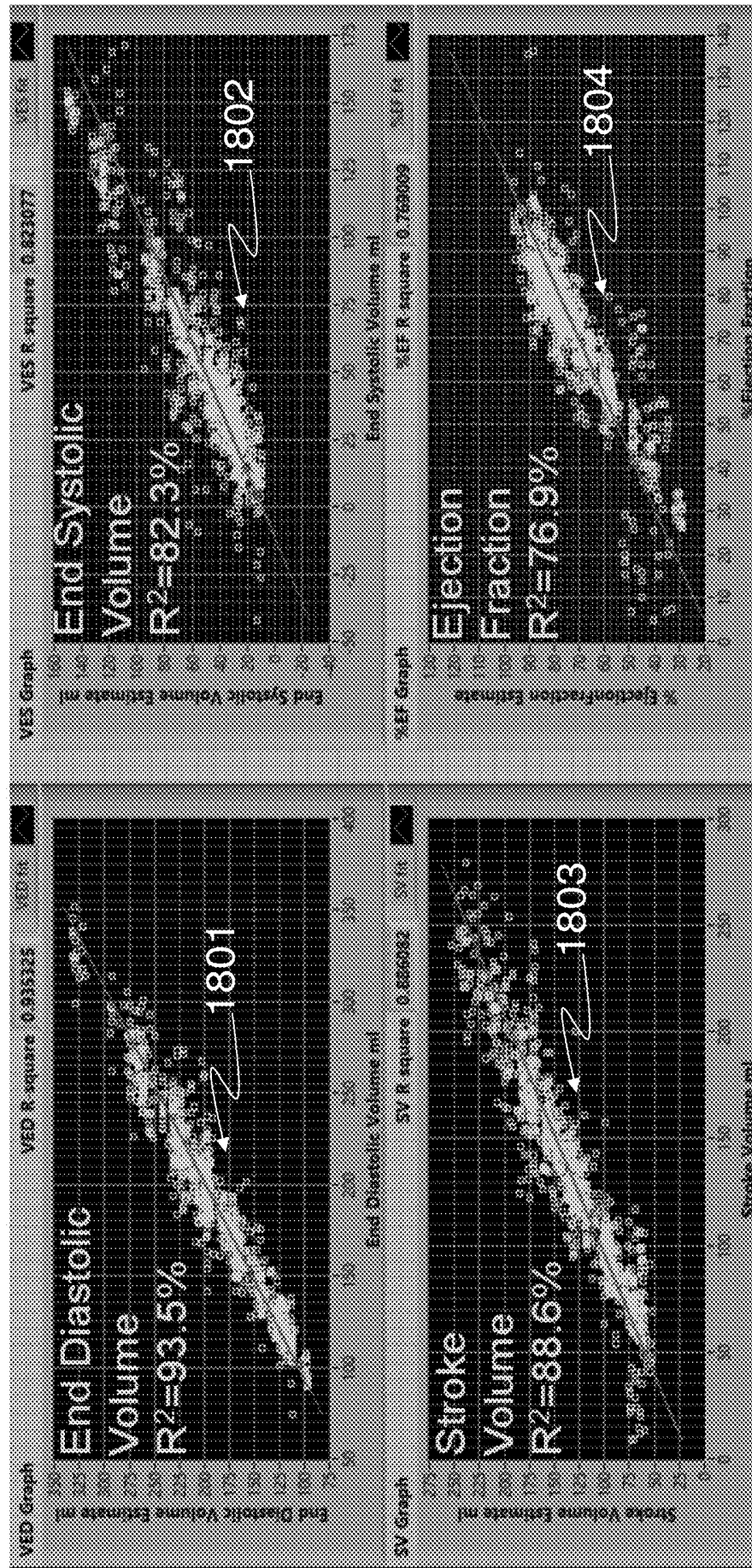


Figure 19

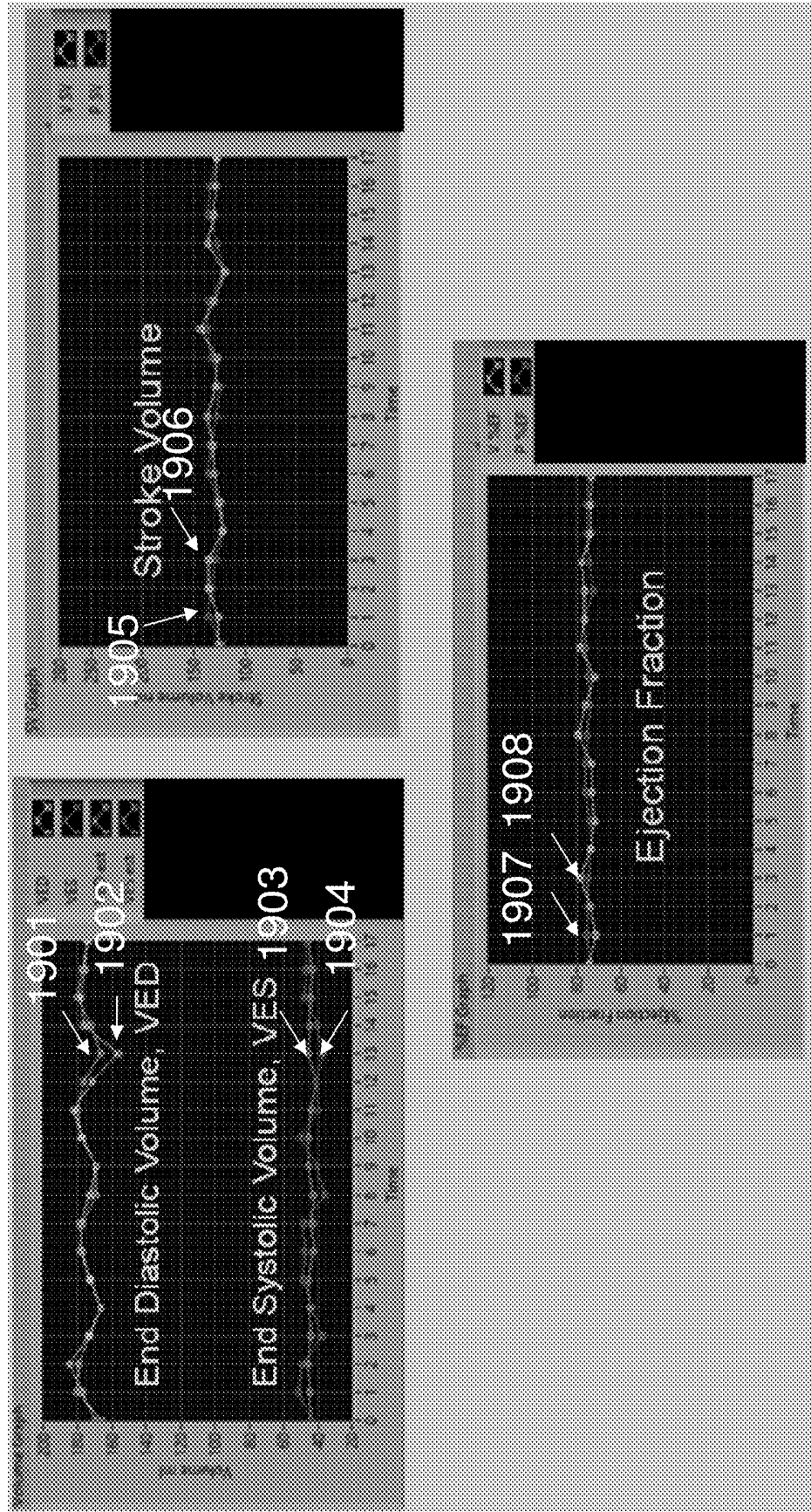
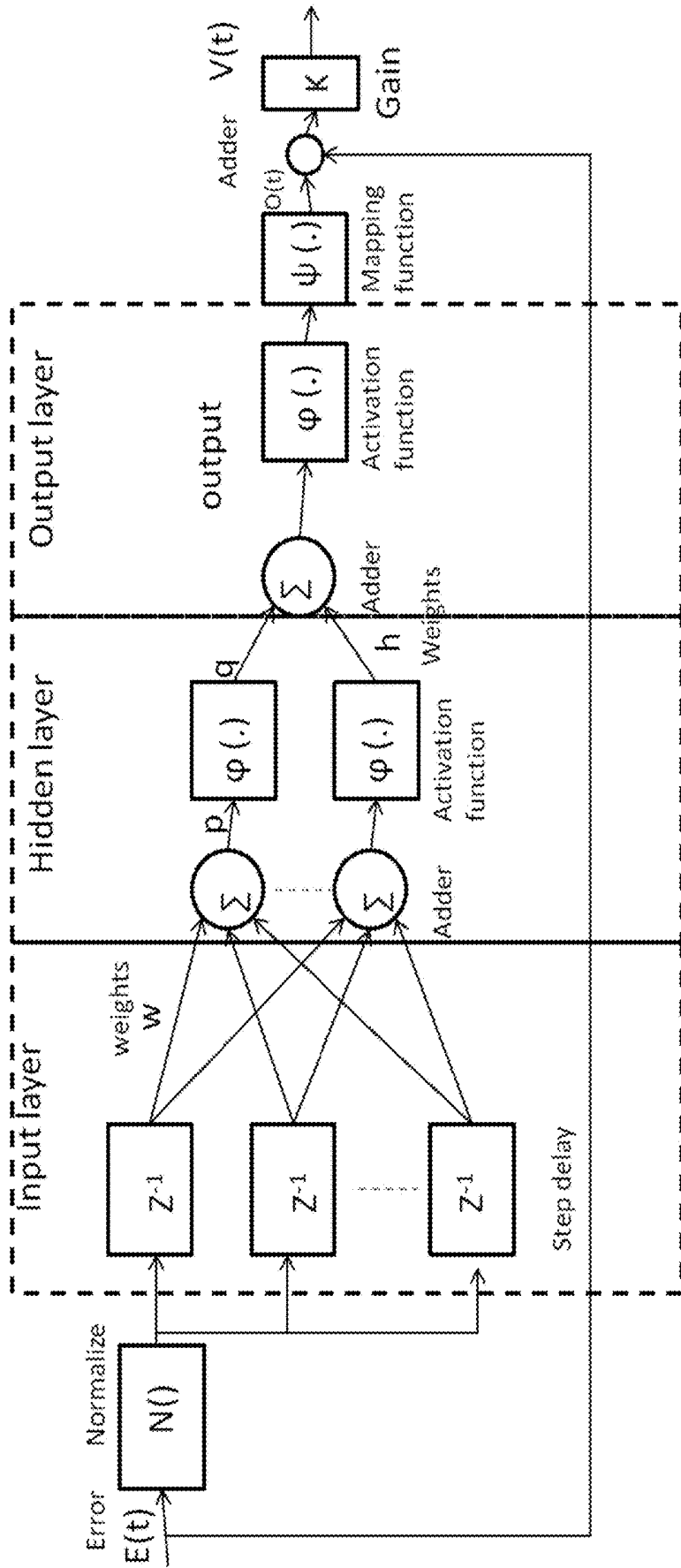


Figure 20. (Prior Art)

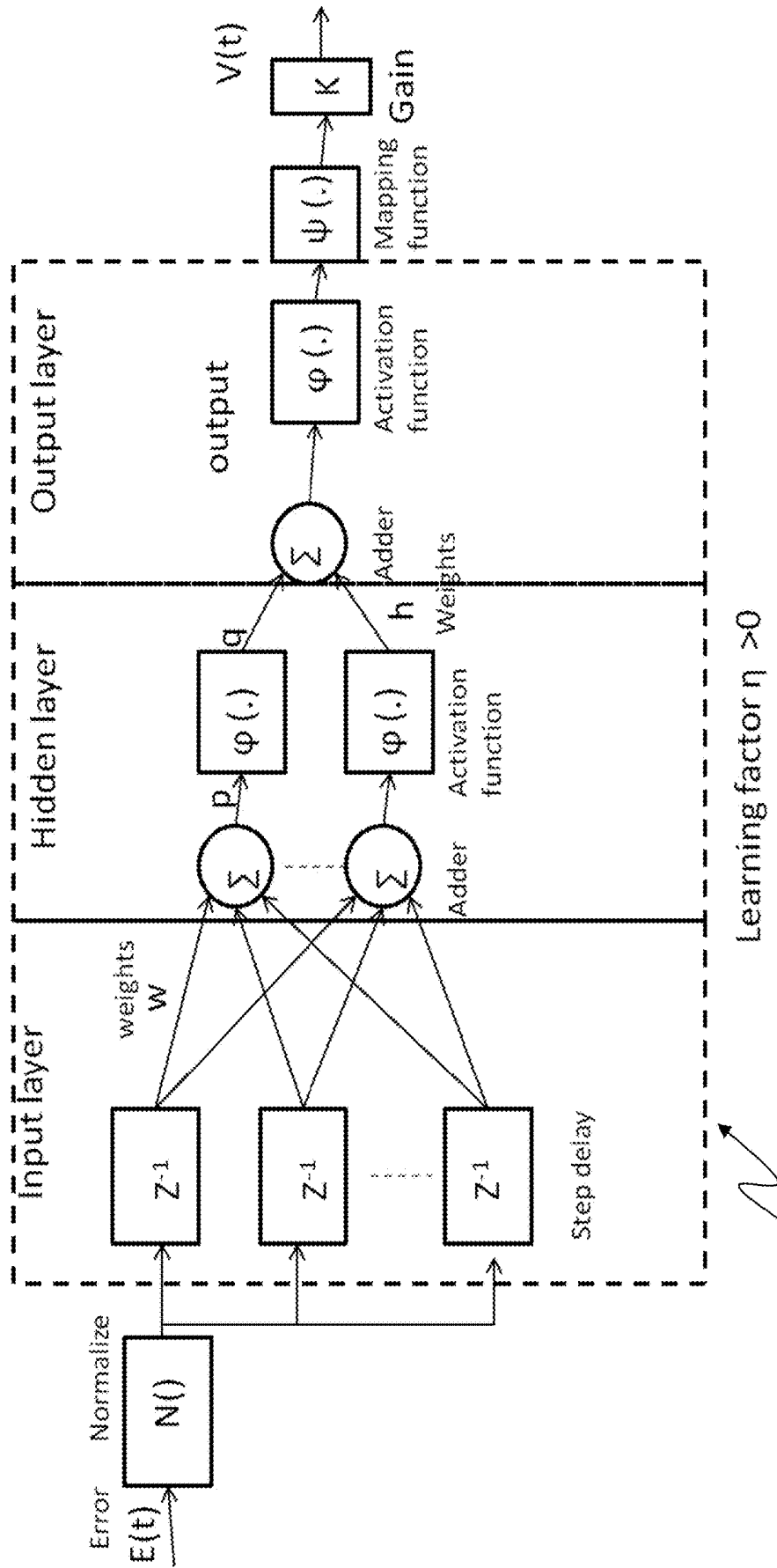


Learning factor  $\eta > 0$



2001

Figure 21



2201

Figure 22

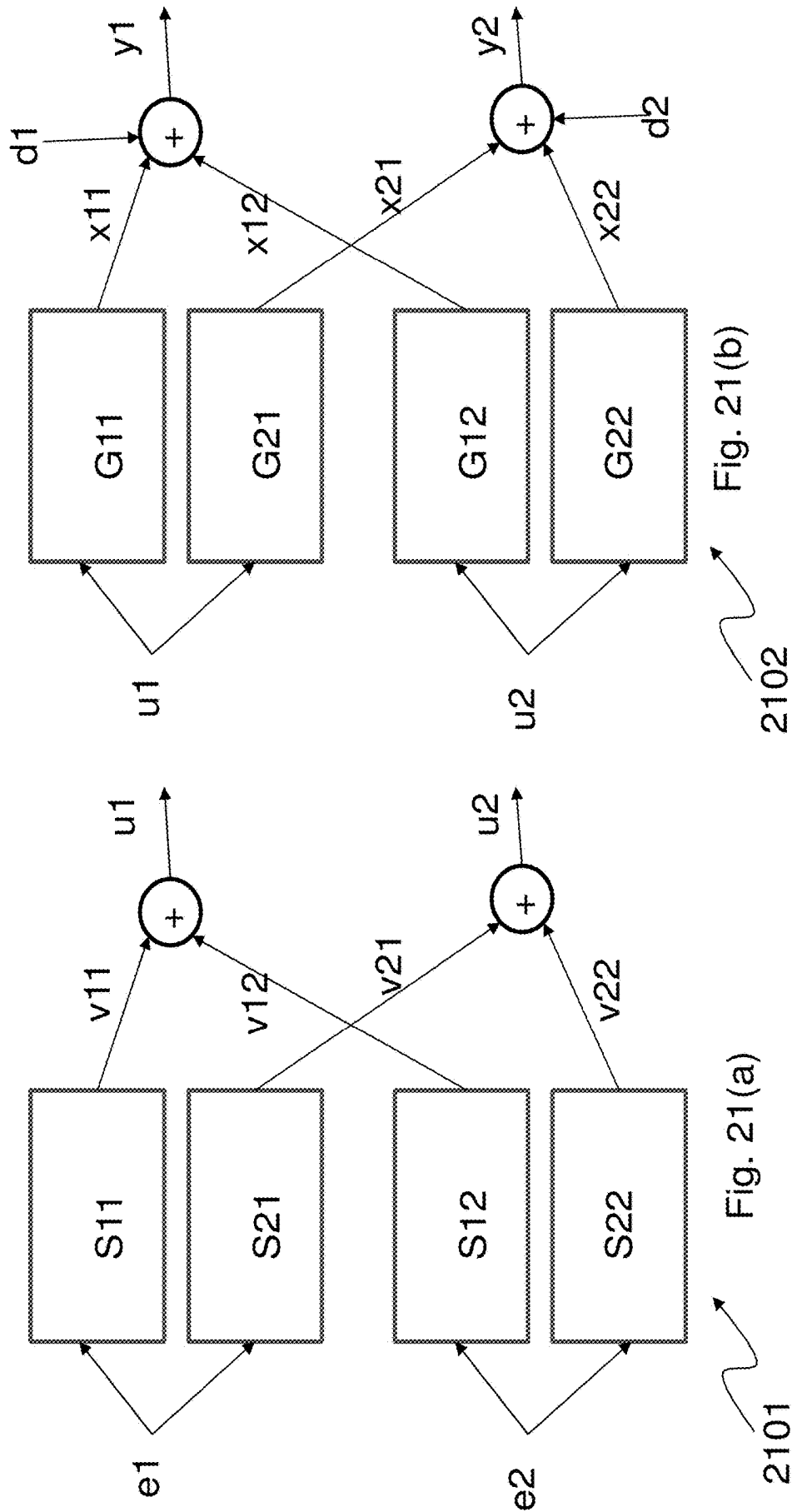
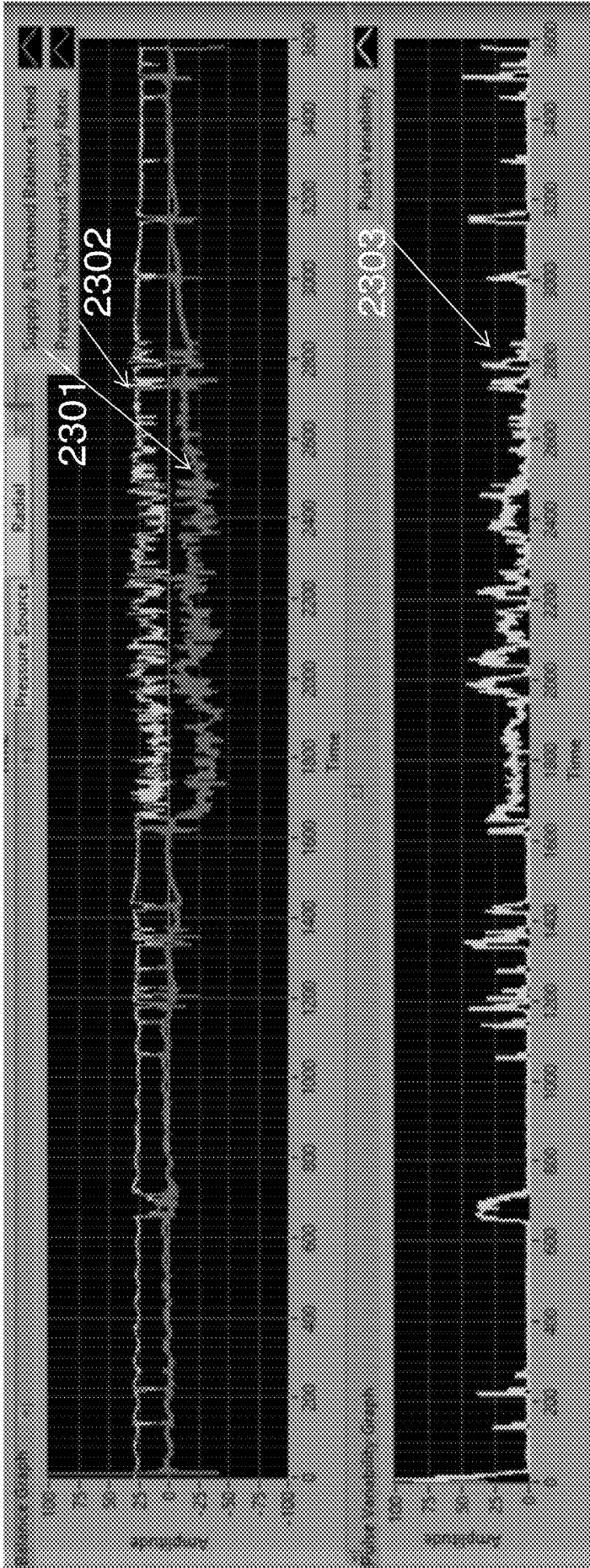


Figure 23



**MONITORING CARDIAC BLOOD FLOW  
BALANCE RELATIONSHIP BETWEEN THE  
RIGHT AND LEFT HEART CHAMBERS AND  
CARDIAC REGULATION**

CROSS-REFERENCE TO RELATED  
APPLICATIONS

[0001] This application is a non-provisional patent application which herein includes by reference and claims the benefits of provisional application 62/529,091 filed on Jul. 6, 2017, and provisional application Ser. No. 61/700,892 filed on Sep. 14, 2012, and non-provisional application US 2015/0018632 filed on Jul. 15, 2013, and non-provisional application US 2015/0025328 filed Jul. 19, 2013, and non-provisional application US 2015/0031969 filed Jul. 24, 2013.

BACKGROUND OF THE INVENTION

1. Technical Field

[0002] This invention relates generally to methods, devices, and systems used in the field of cardiovascular medicine for clinically assessing of a patient's health condition. More particularly, it relates to a new and improved system and method for monitoring cardiac blood flow balance between the right and left heart chambers based on arterial pulse waveforms representing blood pressure or flow or constituent concentration per cardiac cycle.

[0003] The description below is merely provided for general background information and for assisting in understanding the technical field to which the present invention is related. As is generally known in the art, the circulatory system in human beings is responsible for transporting oxygen and other nutrients to the cells of the human body. The circulatory system includes a heart, arteries, capillaries, and veins. In a healthy patient, the heart pumps blood with a certain pressure and volume so as to ensure that proper blood circulation is realized within the human body. Therefore, a discussion of the blood flow in the heart and congestive heart failure (CHF) will now be provided initially.

[0004] With attention directed to FIG. 1 of the drawings which is labeled "Prior Art", there is illustrated a diagram of the cardio-vascular system and its cycle. The heart 2 consists of four chambers; namely, the right atria 4, the right ventricle 6, the left atria 8 and the left ventricle 10. The right atria 4 receives carbon dioxide-filled blood which is returning from the body through the superior vena cava 12 and the inferior vena cava 14 where they join in the central venous. The right ventricle 6 receives the blood from the right atria 4 via a tricuspid valve 16 located between the two chambers. The right ventricle 6 pumps the blood via a pulmonary valve 22 into the pulmonary artery 18 which branches into the right pulmonary artery 18(a) for carrying the blood to the right lung 20(a), and the left pulmonary artery 18(b) for carrying the blood to the left lung 20(b). After receiving oxygen in the lungs 20(a) and 20(b), the oxygen-filled blood is returned to the left atria 8 of the heart 2 via the right pulmonary vein 24(a) and the left pulmonary vein 24(b). The blood is then passed into the left ventricle 10 via mitral valve 26. Next, the left ventricle 10 pumps the blood via the aortic valve 30 into the aorta 28 which branches into the ascending aorta for delivery throughout the upper body 31(a) heads and arms,

and the descending aorta 31(b) for delivery throughout the lower body including the trunk and legs via the network of arteries, capillaries, and finally returned to the heart 2 via the superior vena cava 12 and the inferior vena cava 14, which both merge into the center venous 32. Both the right atria 4 and the left atria 8 pump simultaneously, and both the right ventricle 6 and the left ventricle 10 pump simultaneously. In every pumping cycle, each chamber undergoes an expansion cycle called diastole followed by a contraction cycle called systole. On the ECG waveform, the Atrial diastolic phase extends between the S in the QRS complex until the start of the following P wave, while atrial systole is between the start of the P wave and extends through the following S in the QRS complex. Similarly, the Ventricular systolic phase is between the R in QRS-complex and the following end of the T-wave, while ventricular diastolic phase is between end of the T-wave and the following R in the QRS-complex. The atrial systolic phase occurs mostly during ventricular diastolic phase with a small fractional overlap to ventricular systole, while the atrial diastolic phase starts with the ventricular systolic phase but also extends and overlaps a substantial portion of the ventricular diastolic phase. Atrial diastolic phase has a longer duty cycle as compared to Atrial systolic phase.

[0005] The left ventricle (LV) and the right atrium (RA) are connected in a supply and demand relationship. Also the right ventricle (RV) and the left atrium (LA) are connected in a supply and demand relationship. A balance therefore must be achieved for each supply and demand relationship between the LV and RA (termed Left Right Balance (LRB)), and between the RV and LA (termed Right Left Balance (RLB)). Similarly a balance must be achieved across the two pairs of supply and demand relationships (LV, RA) and (RV, LA). Such a balance of the LRB and RLB balance parameters is termed the Systemic-Pulmonary circulation balance parameter (SPB) indicating a shift of the blood supply towards the systemic portion of the circulatory cycle or the pulmonary portion of the circulatory cycle. When the SPB is normal, this can be used to validate assumptions of the Fick equation that the left and right ventricular cardiac output on average are equivalent, as presumed in healthy individuals.

[0006] In healthy human beings, the right heart's cardiac output (RCO) blood flow is presumed equivalent to the left heart's cardiac output (LCO), on average, since the cycle must maintain steady blood flow through out the circulatory system. However, this is not true in patients with certain types of heart disease. For example, an infarction or arrhythmia on one side of the heart may lead to insufficiency in balance between the RCO and LCO. Thus, cardiac output (CO) is an important indicator not only for the diagnosis of the certain types of heart disease, but also for the continuous monitoring of a patient's health condition.

[0007] One basis for most common cardiac output-measurement systems is given by the well-known equation  $CO=HR \times SV$ , wherein CO is the cardiac output defined as blood flow in liters/min and is equal to the heart rate (HR) times the stroke volume (SV). The stroke volume is usually measured in liters, and the heart rate is usually measured in beats per minute. This equation explains that the amount of blood the heart pumps out over a unit of time (e.g., a minute) is simply equal to the amount it pumps out on every beat (stroke) times the number of beats per time unit.

[0008] If the right heart's cardiac output RCO (blood flow out of the right ventricle into the pulmonary artery) mostly

exceeds the left heart's cardiac output LCO (blood flow out of the left ventricle into the aorta) then blood accumulation in the pulmonary system (lungs) is evident, and which may lead to congestive heart failure (CHF). Similarly, if the left heart's cardiac output LCO mostly exceeds the right heart's cardiac output RCO then blood accumulation is evident in the body's vascular system and tissue fluids will accumulate leading to inflation in tissue or vascular diseases such as high blood pressure, and potentially may play a factor in the development of varicose veins. Therefore, a measure of the balance between the right heart's and left heart's cardiac outputs is necessary for monitoring the overall health of the cardiovascular and pulmonary systems.

**[0009]** In the inventor's view, the cardiac system is basically a control system with two primary objectives. The first primary objective is to maintain oxygen delivery to the tissue via adequate perfusion (blood flow and blood pressure), which is accomplished by varying the cardiac parameters such as heart rate, stroke volume, and contractility, and blood pressure. The heart's second primary objective is to maintain a target blood flow balance between the right and left heart chambers.

**[0010]** While the cardiac output balance between the right heart and the left heart chambers is an important clinical parameter for the purposes of diagnosing, treating, and monitoring cardiovascular disease, the conventional prior art techniques, such as thermodilution or dye dilution, for measuring this balance however requires intravascular or intramyocardial instrumentation which carries significant risks to the patient that includes increased perioperative morbidity and mortality, and increased long-term risks, such as stroke and pulmonary embolism. Additionally, intravascular instrumentation can only be performed by highly trained specialist which severely limits the availability of qualified physicians capable of implanting the device, thereby increasing the cost of the procedure.

**[0011]** Also, the estimation of the balance between right heart's and left heart's cardiac outputs by these conventional methods would necessitate measurement of both left and right cardiac outputs independently and then comparing (for example by subtraction or ratio or so any other means of comparison) both to obtain the measure of the balance. In addition, less invasive methods are also known in the prior art, but they have proved to be less accurate in their measurements. Further, the traditionally non-invasive methods are not able to distinguish between the left cardiac output and the right cardiac output.

## 2. Prior Art

**[0012]** In U.S. Pat. No. 5,211,177 to Chesney et al., there is disclosed a vascular impedance instrument which includes a transducer to obtain a digitized arterial blood pressure waveform. The digitized data is used to determine cardiac output and to subsequently obtain measurements of impedance parameters using the modified Windkessel model of the arterial system. The instrument is used as an aid in diagnosing, treating and monitoring patients with cardiovascular disease.

**[0013]** In U.S. Pat. No. 7,651,466 to Hatib, Feras et al., examines the pulse contour of arterial pressure pulses to map using an linear model (Windkessel method) into a flow pulse for purposes of estimating cardiac output. This does not take into account the relationships between the ventricular arterial supply and atrial venous demand on the shape of the

pressure waveform, but it analyzes the segment component of waveform prior to the onset of the interaction of the atrial venous demand on the ventricular arterial supply. Calibration of the model compliance parameters are required for estimating the dynamic relationship between blood pressure and flow rate.

**[0014]** In U.S. Pat. No. 8,282,564 to Parlikar, et. al., Similarly uses a lumped compliance model to model (Windkessel method) the relationship between arterial pressure and flow rate to estimate cardiac output, again the interaction effects between the ventricular arterial supply and atrial venous demand are not considered on the shape or morphology of the arterial pressure waveform.

**[0015]** In EU Patent No. EP1,848,330 B1 to Bennett, Tommy, et. al. Uses right ventricular pressure (RVP) pulse contour to estimate the stroke volume using a Windkessel pulse contour method. This method uses the full dynamic waveform information in the RVP to relate to the stroke volume information. It does not selectively choose specific landmarks on the pressure waveform to relate to volume information in a static method as in this invention.

**[0016]** In U.S. Pat. No. 5,368,040 to James Carney, describes analysis of ventricular pressure waveform and derives landmarks identified from first and second derivative inflection points or zero-crossing points or as related to features on an ECG waveform. It does not attempt to map these pressure values to volumetric information for purposes of estimating cardiac volumes or flow rates given pressure information.

**[0017]** The inventor of the instant invention, however, is not aware of system for measuring of a cardiac blood flow parameter between the left chamber of the heart and the right chamber of the heart that uses a sensor device to measure one of blood pressure and blood flow and blood constituent concentration of a patient so to generate an arterial pulse signal, a process unit for generating a full arterial pulse signal from the arterial pulse signal, the processor unit generating an arterio-venous pulse signal by subtracting the arterial pulse signal from the full arterial pulse signal, and the processor unit generating the balance parameter by calculating the ratio of the area under the curve of the arterio-venous pulse signal, measured by integration over the cardiac cycle, to the area under the curve of the full arterial pulse signal, measured by integration over the same cardiac cycle. The area under the curve can be calculated by the integration of the pulse waveform using any integration method, preferably trapezoidal approximation method. Whereas the DC offset can be either included in the integration, or normalized, or preferably removed by subtraction. In another embodiment the balance parameter is calculated using the ratio of the maximum peak value of the AV pulse to the maximum peak value of the FAP pulse over the same cardiac cycle. In another embodiment, the balance parameter is calculated using the sensitivity relationship between the area under the curve or the maximum peak values of the AV and FAP pulses, whereas sensitivity is preferably measured using the neural network sensitivity sensor (NNSS) or a partial or full derivative based method. The use of the word pulse throughout herein is used to more generally indicate the measured signal of type such as a blood pressure or blood flow rate or blood constituent concentration waveform.

**[0018]** In a preferred embodiment, a low pass filter, more preferably zero-phase or zero-delay low pass filter, can be

used to smooth out balance ratio parameter or balance sensitivity trend parameter prior to displaying it to the user, in order to provide a smooth update of true trend and avoid excessive noise spikes on per pulse basis.

**[0019]** The cardiac pulse variability can be measured using observed (measured) pulses, or AV pulses, or FAP pulses and indicated by measuring the sensitivity relationship or ratio relationship between the value for maximum peak and minimum valley for each of said pulses using its variable AC component. Whereas the sensitivity relationship is preferably evaluated using the Neural Network Sensitivity Sensor (NNSS) described herein, or using a partial derivative or full derivative based method. Said sensitivity or ratio measure of the pulse variability can be used with a low pass filter, preferably zero-phase or zero-delay low pass filter, to smooth out the indicated pulse variability measure before presentation or display. The pulse variability measure is indicative trend of cardiac output and blood flow and provides clinicians an indication of patient hydration status.

**[0020]** Therefore, it would be desirable to provide a system and method for measuring of a cardiac blood flow parameter between the left chamber of the heart and the right chamber of the heart useful in creating a clinical assessment of a patient's health condition. Further, it would be expedient that the system of the present invention be capable of performing as a clinically useful tool for the purpose of diagnosing, treating and monitoring the performance and function of a patient's liver, heart, lungs, brain, kidneys and other organs of the body. The present invention represents a significant improvement over the aforementioned prior art which is hereby incorporated by reference in their entirety.

#### BRIEF SUMMARY OF THE INVENTION

**[0021]** In broadest terms, a new and improved method is provided for the estimation of balance parameters between the right heart cardiac output and the left heart cardiac output based upon the analysis of the pulse wave of the arterial pulsation. The analysis includes the use of morphologic features of the Dicrotic Notch. The Dicrotic Notch is typically present throughout the Aortic arterial vasculature tree which is used to derive the Left-Right Balance (LRB) parameter. The Dicrotic Notch is also typically present throughout the Pulmonary Artery vasculature tree which is used to derive the Right-Left Balance (RLB) parameter.

**[0022]** The inventor has developed a new theory for explaining the morphology of the pulse wave of the arterial pulsation (arterial pulse pressure or flow waveforms) and specifically the occurrence of the Dicrotic Notch feature residing therein. The present invention is based upon the realization by the inventor that the morphology of an arterial pulse waveform lies in the circulatory nature of the blood in the vascular system. Most of the cardiac monitoring publications heretofore only analyzed either the left heart's performance or the right heart's performance without considering the interaction effects there between. In other words, the fact that the left and right heart cycles complement each other to complete the circulatory system was largely ignored.

**[0023]** The relationships between the right heart's cycle and the left heart's cycle cannot be ignored as there is a continuity of mass flow between them. Therefore blood flow, and subsequently characteristics of pressure behavior, on one side of the cycle must affect the other side of the cycle. The vascular pathways of arteries and veins are

interconnected by the capillary bed which forms the connectivity between the right and left heart chambers, and establishes the interdependency and continuity between them in terms of blood flow. While blood pressure between the right and left heart chambers is not directly interdependent, because of the capillary bed therebetween, the blood flow is directly interdependent as it has to be continuous between the left and the right heart chambers. This direct relationship in blood flow results, as will be explained later, in an indirect blood pressure interdependency consequently. Although blood pressure pulse is related to blood flow pulse by an arterial compliance factor, a balance parameter expressed as a ratio of two pressure components will normalize out the effects of the compliance and therefore the ratio will be equivalent, at least from a trend perspective, to a balance parameter that is derived from a blood flow pulse measurement. Similarly, the blood constituent concentration dynamic variation on a per cardiac cycle basis is directly related to blood flow as the concentration will change based on the speed of the mass flow per unit time by the sensor. Therefore balance parameter trends derived from blood constituent concentration will be consistent in behavior to balance parameter trends derived from blood flow.

**[0024]** Often the performance of the left side of the vascular system is analyzed separately from the right side, which yields weak conclusions that cannot be generalized. Traditional models that failed to take into account the simultaneous effects of the left and right heart cycles tend to bias the attribution of the dynamics it models to one side over the other, and therefore fail to represent the true dynamics fully. The venous side of the blood system is acted upon by the right atrium (RA) pull the blood up towards the heart. Therefore, the RA actually applies a pull force (demand) on the venous blood flow. On the left heart's side, the left ventricle (LV) actually applies a push force (supply) on the arterial blood flow to force the blood through the capillary system into the tissues.

**[0025]** These two events occur substantially in simultaneous timing (in normal subjects) so that the left side's (LV) push is aided by the right side's (RA) pull, in order to facilitate the movement of blood throughout the capillary system in the tissues. The LV and the RA are interconnected by the vascular tree (arteries, veins, and capillaries). While the blood pressure in the venous side is substantially directly independent of the blood pressure on the arterial side as they are connected via the capillaries, the blood flow between the arterial and venous side is substantially directly dependent, as conservation of mass laws and continuity of flow must apply. In other words, in healthy subjects, whatever blood mass flow the venous side demands the arterial side must supply and vice versa to complete the circulatory system. In subjects with circulatory system disorders, in addition to blood in the vascular system, some fluids are either stored or extracted from tissue fluids or the interstitial space. Blood pressure is an equilibrium point to the demand and supply of blood (or blood flow). It is therefore important to view pressure in the context of such supply and demand forces on blood flow. This invention exploits the relationships of blood supply and demand to each other, and uses them to derive information about the arterial supply of blood from venous pulse information and about venous demand of blood from arterial pulse information.

**[0026]** In view of the foregoing background, it is therefore an object of the present invention to provide a system and

method for measuring of cardiac blood flow balance parameter between the left chamber of the heart and the right chamber of the heart of improved design and performance. It is another object of the present invention to provide a system and method for clinically assessing of a patient's health condition based upon a balance parameter obtained by measured arterial pulse signal and assessing of the patient's health condition based upon the morphology of the balance signal. It is still another object of the present invention to provide a system and method for generating a full arterial pulse (FAP) signal, defined as the measured arterial pulse signal without the effect of atrial diastolic blood flow demand factor. The full arterial pulse (FAP) is defined by the measured arterial pulse signal without the effect of atrial diastolic blood flow demand, simulating an added component of arterio-venous (AV) pulse signal so as to result in a full arterial pulse signal (i.e. representing ventricular supply factor without atrial interaction effects). It is still yet another object of the present invention to provide a system and method for measuring of cardiac blood flow balance parameter which utilizes system identification techniques for modeling a transfer function relationship between a peripheral artery AV pulse or FAP signal and an aortic AV pulse or FAP signal. It is still yet another object of the present invention to provide a system and method for measuring of cardiac blood flow balance parameter which utilizes system identification techniques for modeling a transfer function relationship between a pulmonary artery peripheral branch AV pulse or FAP signal and the pulmonary artery AV pulse or FAP signal.

[0027] These and other objects, features and advantages of the invention are provided by a system for measuring of cardiac blood flow balance between the left chamber of the heart and the right chamber of the heart which includes a sensing device for measuring one of blood pressure and blood flow and blood constituent concentration of a patient so as to generate an arterial pulse signal. A system identification technique is responsive to the arterial pulse signal for generating a full arterial pulse signal.

[0028] Advantageously, a processor unit is provided and is configured to produce an arterio-venous pulse signal defined by subtracting the arterial pulse signal from the full arterial pulse signal. Further, the processor unit computes a balance parameter by calculating the ratio of the areas under the curve, by integration over the cardiac cycle, for the arterio-venous (AV) pulse signal to the full arterial pulse (FAP) signal. Whereas the DC offset can be either included, or normalized, or removed by subtraction. In a preferred embodiment, a simple linear transfer function is established between these two signals such as an auto-regressive moving average model is computed using system identification methods. In another preferred embodiment, a generalized state-space linear model of the balance parameter relationship between these two signals is computed using system identification methods.

[0029] In addition, a computational device generates a set of physiological parameters from the balance parameter. A display monitor receives the set of physiological parameters from the computational device for displaying meaningful information to indicate a clinical assessment of a patient's health condition.

[0030] These and other features and advantages of the disclosed system for measuring of cardiac blood flow balance parameter reside in the construction of parts and the

combination thereof, the mode of operation and use, as will become more apparent from the following description, reference being made to the accompanying drawings that form a part of this specification wherein like reference characters designate corresponding parts in the several views. The embodiments and features thereof are described and illustrated in conjunction with systems, tools and methods which are meant to exemplify and to illustrate, not being limiting in scope.

#### BRIEF DESCRIPTION OF THE DRAWINGS

[0031] FIG. 1 is a simplified diagram of a patient's cardiovascular system, useful in explaining the blood flow through the heart and labeled "Prior Art";

[0032] FIG. 2 depicts the various waveforms of a measured arterial pulse signal, a derived full arterial pulse signal, and a calculated arterio-venous pulse signal. It also depicts the simultaneously measured central venous pulse, and the systolic and diastolic phases of the atria and ventricles, as well as the delayed systolic and diastolic phases of the atria and ventricles as measured at the radial artery;

[0033] FIG. 3 is schematic block diagram of a system for producing a balance parameter and related information, constructed in accordance with the principles of the present invention;

[0034] FIG. 4 is a system transfer function which illustrates the time domain and frequency domain relationships between the input and output time series;

[0035] FIG. 5 is flow diagram of an exemplary embodiment of the software used by the microprocessor in FIG. 3 for generating the balance parameter and related information;

[0036] FIG. 6 are waveforms useful in explaining an embodiment of the invention on how cardiac volumetric and performance parameters can be determined using aortic pressure measurements allowing determination of the left ventricular pressure at certain key pressure points, which can be useful in mapping pressure information to volume information;

[0037] FIG. 7 is useful in explaining an embodiment of the invention on how to determine a cardiac auto-regulation rate and a respiratory rate determined from the cyclic variability of the balance parameter, computed for each cardiac cycle, and over a long term series of cardiac pulses;

[0038] FIGS. 8(a) depicts balance parameter waveforms useful in depicting the respiratory cycle and the normal cardiac auto-regulation function cycle, and FIG. 8(b) depicts disappearance of cardiac auto-regulation due to abnormal heart conditions, whereby its restoration is necessary for indicating restoration of normal heart condition;

[0039] FIG. 9 is a flow chart of an exemplary embodiment of the method for estimating aortic pulse waveform (AOP), defined by blood pressure, blood flow rate, or blood constituent concentration, based upon the estimated full aortic pulse and aortic arterio-venous pulse;

[0040] FIGS. 10(a) and 10(b) is a flow chart of an exemplary embodiment of the method of FIG. 6 used to determine ventricular pressure static points of end-diastolic pressure (PED), and pre-diastolic end-systolic pressure (PDPES) for use in computation of ventricular end-diastolic volume, end-systolic volume. It also describes the determination of PED and PDPES using aortic pressure alone without ventricular pressure information;

[0041] FIG. 11 is a flow chart of an exemplary embodiment of the method for estimating a second ventricular chamber's performance based upon information from a first ventricular chamber;

[0042] FIG. 12 is a flow chart of an exemplary embodiment of the method of FIG. 8 for using the balance parameter for measurement of the cardiac regulation cycle;

[0043] FIG. 13 depicts Cardiac Left/Right Blood Flow Circulation Demand & Supply Balance;

[0044] FIG. 14 depicts Cardiac Blood Flow Circulation Balance indicating Auto-Regulation of Supply and Demand;

[0045] FIG. 15 Balance parameter indicating vascular (IVC) occlusions, & medication effects on circulation;

[0046] FIG. 16 Balance parameter indicating LAD stenosis as an Early predictor of Congestive Heart Failure (CHF);

[0047] FIG. 17 depicts processing steps for a calibration process and an estimation process for estimating cardiac hemodynamics including cardiac left ventricular pressures and volumes noninvasively.

[0048] FIG. 18 and FIG. 19 depict a comparison between estimated and actual multiple cardiac hemodynamic parameters including end diastolic volume, end systolic volume, ejection fraction, and stroke volume. FIG. 18 presents the comparison on an x-y chart basis, and FIG. 19 depicts the comparison on a real-time waveform basis.

[0049] FIG. 20 Sensitivity sensing, S, between two input variables of a reference (target) variable R and a process variable P;

[0050] FIG. 21 A multi-input multi-output configuration for a sensitivity sensor, NNSS, between two input variables for MIMO structures;

[0051] FIG. 22 A multi-input multi-output configuration for a sensitivity relationship weighted final sensitivity;

[0052] FIG. 23 Pulse variability relationship to balance parameter;

#### DETAILED DESCRIPTION OF THE INVENTION

[0053] Before explaining the disclosed embodiments in detail, it is to be distinctly understood at the outset that the present invention shown in the drawings and predominantly described in detail in association with a system for measuring of cardiac blood flow balance parameter is not intended to serve as a limitation upon the scope or teachings thereof, but is to be considered merely for the purpose of convenience of illustration of one example of its application. In particular, the present invention may be applied to any other suitable physiological monitoring systems, such as oximetry systems, ECG systems, and any other suitable for monitoring of a patient's liver, heart, lungs, brain, kidneys and other organs of the body.

[0054] Historically, the dicrotic notch was thought to represent a reflection of a fast pressure pulse that is generated when the aortic valve opens above its cracking pressure, and blood pressure reflecting backward from the arterial side back toward the aorta. When this delayed and damped pressure component of the pulse wave overlaps with the original pressure wave component, then this would result in the combined components creating the elevation called the Dicrotic Notch. However, this was later largely dismissed and replaced by the valve recoil effect theory. The valve recoil effect theory hypothesized that the Dicrotic Notch was the result of the valve recoil after its passive closure due to pressure drop in the start of the ventricular diastole phase.

The valve recoil was believed to cause an elevation of the pressure post the initial decrease in pressure levels.

[0055] These theories however fail to explain why in some cases the Dicrotic Notch was completely missing in some patients, and in some other patients two dicrotic notches would appear. The fact that the Dicrotic Notch was not consistently seen in all subjects is another indication that this feature could not be entirely attributed to valve closure alone. The simultaneity of the valve closure event with the drop in pressure at the aorta and such event creating a change in the rate of reduction of blood pressure represented in the occurrence of the Dicrotic Notch has led to the false presumption that the Dicrotic Notch was formed due to that valve closure event alone. The inventor believes that the circulatory cycle must be analyzed completely to consider interdependencies between the arterial and venous side in terms of blood flow continuity and timing effects.

[0056] It has been observed by the inventor that the atrial diastolic blood flow demand on venous blood has an interaction effect on the ventricular systolic arterial supply of blood such that this interaction affects the blood flow and subsequently blood pressure. Such interaction and timing characteristics between the ventricular supply and atrial demand of blood is what primarily defines the dicrotic notch feature of the aortic and arterial pulse. When ventricles start systolic contraction to eject (supply) blood, the atria start diastolic expansion to dilate to fill with (demand) blood, however, the atrial dilation extends longer in duration to also overlap with ventricular diastolic expansion. During this phase and in the absence of further blood supply the level of steep decline of the pressure is due to atrial diastolic blood flow demand, which starts from demand zero level reaches a maximum demand peak (i.e. supply trough) and then reduces to demand zero level. When atrial diastolic blood flow demand ends, due to atrial systolic phase beginning, then the decay rate of the arterial side supply pressure returns to its natural decay rate due to ventricular supply pressure reduction alone at its natural decay rate. As the timing of this interaction and the level of supply and demand vary with patient's age or health state, there is sometimes observed the disappearance of the Dicrotic Notch or in other conditions the appearance of two Dicrotic Notches. This blood demand and supply interaction effect is realized across all the capillary interconnections between the venous vascular system and the arterial vascular system, and accumulatively the effect becomes stronger and more easily observed at larger arteries, especially at the aorta and pulmonary main arteries.

[0057] As was previously discussed, the pulse pressure at the aorta and other arterial peripheral branches exhibit "a forced" or "positively acted upon" loss of pressure that is subsequently followed by a rise in pressure, when this effect ends, resulting in the Dicrotic Notch. This is the primary effect creating the Dicrotic Notch. In other words, the two events (i.e. blood pressure drop followed by rise) create the feature of the Dicrotic Notch. The rise in the pressure post the valve's passive closure can not be fully attributed to the valve's recoil post the closure event alone. When the valve's recoil occurs, it represents a substantially smaller secondary pressure perturbation than is often observed in the aortic valve's pressure wave about the Dicrotic Notch defined by the primary effect.

[0058] The atrial diastolic blood flow demand (i.e. venous side "pull") on the blood flow overlaps simultaneously with

the ventricular systolic supply (i.e. arterial side push pressure) of the blood flow, during which the supply overwhelms the demand of blood and the pressure rises. The atrial diastolic blood flow demand then further overlaps with ventricular diastolic phase, which stops the blood supply due to heart valve closing effect, and allows the arterial pressure pulse to decay on its own natural decay rate. However, because of the continuous interaction with the atrial diastolic blood flow demand, which becomes the overwhelming dominant factor that reduces the blood pressure more rapidly at a forced decay rate that is steeper than the natural decay rate. This continues until the atrial systolic event start point, at which point the demand stops and the arterial decay rate returns to the natural decay rate. This interaction effects between demand and supply is substantially primarily responsible for the presence of the Dicrotic Notch. Such interaction profile is also varied or is affected by the blood viscosity or coagulation time, and can be used to estimate blood coagulation or coagulation variation. The venous side demand for blood flow overlapping with ventricular supply forces an increased blood flow through the capillaries greater than the rate it normally would occur at due to the arterial supply alone. It also lowers the realized pressure at the capillaries from the pressure levels that would be realized to accomplish the same flow rates due to arterial supply alone. This increased flow rate and at reduced pressure levels enhances the ability of blood cells to flow through the capillary bed more easily. In other words, the atrial diastolic blood flow demand assists the ventricular systolic blood flow supply in flowing the blood through the capillaries. The arterial pressure is affected by the presence or absence or interaction timing of the ventricular supply and atrial demand. The arterial supply pressure is lowered in the presence of the atrial (venous-side) demand pressure pulse. This venous side demand represented by the atrial diastolic blood flow demand pulse results in a reduction in the pressure on the arterial side at a faster rate (forced decay rate) than normal as compared to the case where the venous demand pulse was not present. Increased flow rate (due to venous side demand) also reduces the observed concentration at a fixed measurement point (per unit time) of blood constituents. Similarly, blood flow rate sensors also depict the dicrotic notch feature due to blood flow acceleration in the presence of atrial demand and the interaction of ventricular supply as discussed previously.

**[0059]** As defined herein, the term “arterio-venous pulse” or (AV pulse) refers to the portion of the arterial pulse (representing blood pressure or flow or constituent concentration) that depicts level reduction due to the atrial diastolic blood flow demand for blood (from the venous side), and the arterio-venous pulse is therefore measured on the arterial side and not the venous side. When the arterio-venous pulse demand or “pull” reaches its maximum, the arterial pulse supply or “push” reduces (dips) to a low level presenting the valley of the Dicrotic Notch, this forced decay rate of the arterial pulse pressure or flow is restored back to its natural (unforced) decay rate once the arterio-venous pulse is completed representing the end of venous demand and the remaining reduction rate of the arterial pulse pressure or flow occurs at its normal decay rate. The subsequent cessation or stopping of venous side demand for blood flow results in the arterial pressure increasing back to what it would have been normally at (unforced natural decay rate) due to reduced flow in the arterial side. The Dicrotic Notch

event itself is a representation of the arterio-venous pulse occurrence. The arterio-venous pulse demand “pull” accelerates the flow of blood by assisting the arterial pulse supply “push”.

**[0060]** Venous side pulsations simultaneously add a blood flow component (i.e. increase blood flow) due to its demand for blood, and force the blood to flow faster. The presence of the arterio-venous demand pulse reduces the overall pressure to flow the blood through the capillary channels from levels required to achieve the same flow rates in the absence of the arterio-venous demand pulse. This venous pulsation (blood flow pull or demand) is therefore responsible (primarily) for the creation of the arterial side Dicrotic Notch. Aortic valve recoil has a similar but smaller (secondary) effect on the aortic pressure and arterial pressure. Those two factors (venous demand, and aortic valve recoil) forming the dicrotic notch are discernible. The difference in the effects of the aortic valve recoil versus the venous side demand on the dicrotic notch can become more obvious or separable in the presence of cardiac abnormalities causing atrial filling timing to be not synchronized with ventricular ejection. In these cases, multiple notches can be observed along the dicrotic limb of the pressure wave.

**[0061]** Valve characteristics do not change significantly from heart beat cycle to cycle, but the venous side demand (due to atrial filling) is variable from heart beat cycle to cycle. Therefore, the variations in the arterio-venous pulse across heart cycle-to-cycle can be safely attributed to variations in venous side demand and arterial side supply. Because pressure measurements are used to quantify the arterio-venous pulse and pressure is measured on the arterial side, the area under the curve, by integration over the cardiac cycle, for the arterio-venous pulse is expected to be smaller than that for the arterial pulse itself as the capillary bed small channels exist between the arterial and the venous sides. On the other hand, if flow rate were measured on the arterial side, then the flow out of the arterial side must substantially equal the flow into the venous side, less a small component of fluid potentially transferred in/out of the tissue.

**[0062]** Therefore, an arterio-venous pulse measured from a blood flow waveform (or equivalent indirect methods such as blood constituent concentration variation related to flow) that is detected at the arterial side is expected to have an area under the curve that is substantially equivalent to the area under the curve for the venous flow waveform.

**[0063]** As a result of the foregoing discussion, it can be seen that the fundamental basis of the present invention is founded upon the development of new and novel theory for explaining the occurrence of the Dicrotic Notch. The present invention describes novel measures and physiologic parameters that are used as clinical indicators of cardiovascular and pulmonary performance and health status from the analysis of the pulse wave and its features, including the Dicrotic Notch. Furthermore, this invention describes methods for measurement of such novel parameters via a variety of sensory devices and algorithms or processing steps. In addition, this invention describes several utility applications for use of the derived novel indicator into clinical applications.

**[0064]** Referring now in detail to the various views of the drawings, there is illustrated in FIG. 2 a plurality of timing waveforms which are useful in explaining the present invention. The waveform A is the measured arterial pulse signal in a normal, healthy patient. The waveform B is the esti-

mated full arterial pulse (FAP) signal representing the arterial pulse in the hypothetical absence of venous (atrial) demand. The estimated full arterial pulse (FAP) signal is based on a linear or exponential line interpolation between the two points of primary peak point (x1, y1) and the secondary peak point (x2, y2) (of the arterial pulse signal A. The waveform C is the derived or calculated arterio-venous pulse (AV) signal obtained by subtracting the observed or measured arterial pulse signal A from the full arterial pulse signal B. The waveform D is the central venous pressure (CVP) waveform. A delay (D1) between features in the central venous waveform and the radial artery waveform will be present and substantially equivalent to the delay between the aortic waveform and the radial artery waveform (D2), because of equivalent vascular lengths. This delay can be measured between the start of ventricular systole and the start of the cycle of increasing arterial pressure in the radial artery. Ventricular systole begins substantially synchronously with the R event of the QRS complex of the electrocardiogram (ECG) or electrogram (EGM) waveform. This R event is also synchronous with the point z of the central venous waveform D. Other features of central venous waveform D are indicated as known in the art a-wave, c-wave, v-wave.

[0065] Indicated on the FIG. 2 is the period of the following phases: atrial diastole AD, atrial systole AS, ventricular diastole VD, and ventricular systole VS. It is important to note that the depicted indication of systolic and diastolic phases is from a volumetric perspective and not from a pressure perspective. The difference being the isovolemic periods. Also indicated is the delayed period of each of these phases when shifted in time by the previously indicated delay D1, and are indicated as follows: delayed atrial diastole ADd, delayed atrial systole ASd, delayed ventricular diastole VDD, and delayed ventricular systole VSd.

[0066] We notice that there is small overlap between (AS and VS) until ventricular valve open, however, there is a larger overlap between (AD and VS), (AS and VD), and (AD and VD). Similarly, there is small overlap between the delayed version of these phases (ASd and VSd) however, there is a larger overlap between (ADd and VSd), (ASd and VDD), and (ADd and VDD).

[0067] We note that in the interaction phase of (VSd and ADd) the ventricular supply overwhelms atrial demand and the pressure rises until the end of VSd, at which point the radial artery pressure reaches its maximum systolic value. Then the following phase is the interaction of VDD and ADd, however ventricular demand is blocked by the presence of valves, and the remaining effect on the radial pressure is the delayed atrial diastolic ADd demand. The pressure waveform depicts a natural slope of pressure decay once the pulse moves past the sensor, however, because of the contribution of the delayed atrial diastolic ADd demand, the radial pressure is forced to decay at a higher rate, until the next phase of interaction begins. Then the following phase is the interaction of VDD and ASd where atrial demand ends and the only remaining element is the natural decay rate of pressure, starting at the peak post the dicrotic notch. This natural rate of decay continues until the cycle restarts again with the interaction of the VSd and ADd. The period during the phase of interaction between VDD and ADd is the period of arterio-venous pulse waveform C, when atrial diastolic blood flow demand effect forces the slope of decay of the

radial pressure to be more negative than the natural rate of pulse decay during the ventricular diastolic phase as indicated during the interaction of the phases VDD and ASd.

[0068] Measurements obtained at the central venous or pulmonary vein are generally low pressure about 8-12 mmHg but are higher power and higher in signal to noise ratio than those obtained at smaller veins. The central venous pulse is also significantly modulated by the respiratory cycle.

[0069] While it is obvious to the reader that the calculation of the balance parameter can be performed either from the arterial or venous measurements, and yielding equivalent results of balance parameter detection and trending, the remaining discussion herein will focus on arterial side measurements because of their higher measurement power and resolution, and easier interpretation of the full arterial pulse and the arterio-venous pulse relative to equivalent information from the central venous pulse.

[0070] The Dicrotic Notch is present in the waveform A of the arterial pulse signal. The Dicrotic Notch is defined by a characteristic valley which resides between the peak (point 1) of the arterio-venous pulse (also the beginning of the diastolic phase of the ventricle), which is also followed by a secondary peak (point 3) which also marks the end of the arterio-venous pulse.

[0071] In the hypothetical absence of atrial (venous side) demand for blood flow and if the arterial blood flow was solely due to ventricular supply, the decrease from the peak (point P1) of the arterial pulse wave A (representing blood pressure pulse or blood flow pulse) is modeled and presumed to be substantially linear over time, for simplification, until it reaches the start of the settling phase. The settling phase of the pulse wave A is defined to start at the secondary peak (point P3) after the Dicrotic Notch valley, which also indicates the end of the arterio-venous pulse and the secondary valley (point P5) of the pulse wave which defines its end (end of the diastolic phase) and the beginning of a new pulse wave (start of the systolic phase).

[0072] The delays between the CVP waveform D and the arterio-venous waveform C observed at the radial artery are substantially equivalent to the delays between the aortic pressure and the radial pressure. It will also be noted of the symmetry in the shape of the AV waveform C obtained by subtracting the arterial pulse waveform A from the FAP waveform B.

[0073] In FIG. 3, there is shown a schematic block diagram of a system 32 for producing a balance parameter for use in one application, constructed in accordance with the principles of the present invention. The system 32 includes a sensing device or transducer 34, an A/D converter 36, a processor unit 38, a RAM 40, a ROM 42, and a storage device 44. The sensing device 34 is preferably a non-invasive arterial blood pressure or blood flow waveform measurement device operatively connected to a patient 33 for generating an arterial pulse signal. This arterial pulse signal is similar to waveform A in FIG. 2 and is fed as an input to the processor unit 38 via the A/D converter 36.

[0074] Further, the system 32 of the present invention includes a user input device 46 for inputting information to the processor unit, such as a keyboard or input terminal and a display station 48 for displaying useful information. The storage device can be any computer-readable media capable

of storing information that can be interpreted by the processor unit, such as a hard disc, floppy disc, or other digital storage apparatus.

**[0075]** The sampling rate of the A/D converter **36** is set to at least two times according to the Nyquist criteria, but is preferably set to five times or higher than the highest frequency component of interest in the pulse waveform of arterial blood pressure, or blood flow signal, or blood constituent concentration so as to satisfactorily capture the same. The A/D converter is used to digitize the analog arterial blood pressure or blood flow, or blood constituent concentration waveform signal generated by the sensing device. Alternatively, the processor unit could obtain the blood pressure or blood flow, or blood constituent concentration measurements in digital form directly from the sensing device so as to avoid the need of the A/D converter.

**[0076]** The sensing device **34** is preferably a non-invasive blood pressure, or blood flow, or blood constituent concentration waveform measurement device. A non-invasive blood pressure sensor such as a finger-cuff transducer unit using a counter pulsation technique in which the waveform is detected by balancing the air pressure in the finger cuff with the blood pressure in the patient's finger. A finger-cuff transducer unit of this type is commercially available from Ohmeda Monitoring Systems of Englewood, Colo. Also, tonometric blood pressure methods that press on the radial artery against the bone structure may be used to perform blood pressure measurements. A tonometric blood pressure of this type is the Vasotrac which is commercially available from Medwave. However, there exist many other prior art sensor types that can be used to measure either arterial and venous pulse pressure waveforms.

**[0077]** For blood flow measurements, this can be achieved by using non-invasive acoustic ultrasonic doppler frequency effect sensors, or using an electromagnetic wave (such as light or RF), or by directly using motion sensors, accelerometers, or using coriolis sensors, or by thermodilution methods, or dilution of concentration of a chemical such as lithium or dye or a radioactive material, or dilution of concentration of a blood constituent or component, or using differential pressure sensors measurements (for example Venturi flow sensing or orifice pinhole type sensors). Similarly, pulse flow can also be assessed using the dilution or concentration of a radiotracer chemical element such as those used in nuclear medicine, fluoroscopy, or in magnetic resonance functional imaging.

**[0078]** For blood constituent concentration measurements, this can be achieved by using a noninvasive optical plethysmography or using electrochemical sensors that have specificity and sensitivity to the target chemical analyte or blood constituent component, preferably with very fast response time.

**[0079]** For blood pressure measurements, this can be obtained using many sensor types including resistive or capacitive or inductive measurement elements, and any of the following broad categorizations of general types of sensing technologies including:

**[0080]** Invasive method include placement of a catheter to obtain pressure measurements consisting of:

**[0081]** Using implantable sensors that reside in a vascular branch or in a cardiac chamber or body organ.

**[0082]** Using catheter mounted pressure sensors, or via catheter fluidic channel pressure measurement over the

extended catheter using a pressure sensor that is either built into the catheter or external to it.

**[0083]** Minimally invasive methods include access to the artery or vein channel using a cannula and using a pressure sensor connected to the cannula channel to measure the vascular artery or vein pressure.

**[0084]** Non-invasive methods including:

**[0085]** Tonometric pressure pulse sensing methods

**[0086]** Optical plethysmography pulse sensing methods

**[0087]** RF pulse sensing methods

**[0088]** Acoustic pulse sensing methods such as those relating the Doppler frequency shift with the flow rate change.

**[0089]** Vibration pulse sensing methods such as placement of an accelerometer or force sensor in proximity to the vascular channel where pulse waveform or pulse velocity is intended to be measured.

**[0090]** Blood magneto-acoustic methods which use measurement of an electric current that is resultant from ion movement in a magnetic field.

**[0091]** Blood temperature dynamic variation can also be representative of the dynamics of the blood pulse when measured at a frequency high enough that is representative (at least twice as high based upon the Nyquist criteria, but preferably five times as high) of the dynamics of variations of the pulse. This is due to the thermal dilution gradient of the blood as it is accelerated past the sensor. This measurement can be also obtained more easily non-invasively, for example, using infrared sensors.

**[0092]** Blood constituents dynamic variations are also indicative of the pulsatile flow when measured at a frequency high enough that is representative (at least twice as high based upon the Nyquist criteria, but preferably five times as high) of the dynamics of variations of the pulse as represented in the arterial pulse.

**[0093]** The monitoring of the cardiac balance parameter in the present invention can be accomplished using pulsations measured from any artery, including, but not limited to, aortic, carotid, pulmonary, femoral, radial, renal, and hepatic arteries. Similarly, the monitoring of the cardiac balance parameter can be accomplished using pulsations measured from any vein including, but not limited to, central venous, superior vena cava, inferior vena cava, jugular, pulmonary, femoral, radial, renal, and hepatic veins. However, the signal power measured at a vein is much smaller than an artery. Nonetheless, the same primary components related to arterial supply and venous demand can be identified from venous measurements per cardiac cycle of any of blood pressure, blood flow, or blood constituent concentration and can be used for quantifying the balance parameter.

**[0094]** A balance parameter determined by measurements from the central venous, central venous branches, the aorta, or aortic arterial branch is called Left Right Balance (LRB), and a balance parameter determined from the pulmonary vein, pulmonary vein branches, pulmonary artery, or pulmonary arterial branches is called Right Left Balance (RLB).

**[0095]** In addition, the monitoring of the cardiac balance using a pulse waveform measured from any artery or vein can be accomplished with any of the aforementioned sensor types and sensory methods of measurement including invasive, minimally invasive, or noninvasive blood pulse monitoring, and including pulse waveforms that are indicative of

blood pressure, blood flow rate, or blood constituent or components dynamic variation over time.

**[0096]** Under the control of a software program stored in the ROM **42** or the storage device **44**, the processor unit **38** will derive the full arterial pulse (FAP) signal, as shown in the waveform B in FIG. 2, based upon the observed or measured arterial pulse signal A. Then, the processor unit **38** will calculate the arterio-venous (AV) pulse signal, as shown in the waveform C of FIG. 2, by subtracting the measured arterial pulse signal from the FAP signal. Finally, the processor unit will determine a balance parameter at its output **49** of the present invention by calculating the ratio of the area under the curve of the AV pulse signal, measured by integration over the cardiac cycle, to the FAP signal, measured by integration over the same cardiac cycle. Also the balance trend parameter is calculated using the relative sensitivity of the area under the curve of the AV pulse signal, measured by integration over the cardiac cycle, to the FAP signal, measured by integration over the same cardiac cycle. In another embodiment, the balance trend parameter is calculated using the relative sensitivity of the maximum peak value of the AV pulse to the maximum peak value of the FAP pulse. Whereas the balance trend is computed using a sensitivity model using the partial or full derivative method, or preferably using the neural network sensitivity sensor (NNSS) method as depicted in FIG. 20, relating two parameters, a reference (target) variable  $R(t)$  and a process variable  $P(t)$ , to each other with a sensitivity measure that describes the first differential or gradient between  $R(t)$  and  $P(t)$ . This balance ratio and balance trend parameters **49** are sent by the processor unit to a computational device **50** for generating a set of physiological parameters on its output **51** which are useful to indicate a clinical assessment of the patient's health condition.

**[0097]** In a preferred embodiment, a low pass filter, or more preferably, with zero-phase or zero-delay low pass filter, can be used to smooth out balance ratio parameter or balance sensitivity trend parameter prior to displaying it to the user, in order to provide a smooth update of true trend and avoid excessive noise spikes on per pulse basis.

**[0098]** The display station **48** receives the set of physiological parameters **51** from the computational device **50** for providing visual and audio monitoring of the patient's health condition and for alerting medical personnel of changes in the patient's health condition. As an option, a wired or wireless telecommunication system **52** shown in phantom may be coupled to receive the set of physiological parameters from the computational device **50** and for transmitting wirelessly the set of physiological parameters to the display station **48**, which can be located at a remote site away from the patient.

**[0099]** In the embodiment shown, the processor unit **38** is comprised of a general-purpose microprocessor which is adapted to execute software consisting of an operating system and one or more applications, as part of performing the functions described hereinbelow. Further, each of the various blocks illustrated in FIG. 3 is merely exemplary in nature and is intended to show one implementation of the system for providing the balance parameter using the arterial pulse signal and should not be construed as limiting. Indeed, the different embodiments may have widely varying software programs, data structures, applications, processes, and the like. Therefore, the various functions of each block may

in actual practice be combined, distributed, or otherwise arranged in any suitable fashion.

**[0100]** The processor unit **38** includes a System ID Processing Device **39** for establishing a transfer function relationship between FAP pulses of any artery in the left heart vasculature (as an input or inputs to the model) and the aortic artery main pulse which represents the primary source of such a pulse (as an output to the model). Similarly, the System ID Processing Device **39** can be used to establish a transfer function relationship between full arterial pulses (FAP) or arterio-venous (AV) pulses of any two arteries, defined in forward or backward input/output transfer function modeling steps. For example between the aorta and any of its arterial branches, or between the pulmonary artery and any of its arterial branches. Such transfer function relationships relate the pulses from a plurality of locations as input or output in order to use a more readily observable signal to estimate a more difficult to observe signal. For example, used in the right heart vasculature tree (as an input or inputs to the model) and the pulmonary artery main pulse as a primary source of such a pulse (as an output to the model). Preferably, linear state-space models can be used for establishing such transfer function relationship.

**[0101]** The aortic or pulmonary artery full arterial pulse (FAP) can be used as the source (input) producing the (peripheral) full arterial pulses (output) in the arterial branches, or the reverse (or inverse) model can be used. Similarly, the aortic or pulmonary artery arterio-venous pulse can be used as the source (input) producing the (peripheral) arterio-venous pulses (output) in the arterial branches, or the reverse (or inverse) model can be used. Whereas the estimated output arterio-venous pulse can be subtracted from the estimated output full arterial pulse to produce the estimated observed arterial pulse.

**[0102]** Similarly, the System ID Processing Device **39** can be used to establish a transfer function relationship between FAP pulses of any artery and the arterio-venous (AV) pulse from that same pulse, representing a model of the balance parameter. Preferably, a simple ratio of the AV pulse to the FAP pulse, i.e. a linear relationship of the first order can be determined. Also, a more generalized linear model of any order identified using system identification methods, preferably state-space model, can be used to describe the balance parameter as a system relationship. Such a generalized model will capture the relative amplitude ratio and timing delay (phase shift) information between the AV and FAP signals. Optionally, in a second step, given the balance parameter, or a model of it, and the input FAP, then the output AV pulse can be estimated.

**[0103]** Selection of either inputs or outputs allows for estimation or prediction of an unknown output, given an identified model parameters and measured inputs. A MISO (multiple inputs single output) type model mapping the relationship between a plurality of peripheral artery pulses (inputs, more easily measured) into the main primary source aortic or pulmonary artery pulse (output, more difficult to measure) for that vasculature is a preferred embodiment for enhancing accuracy and enabling easier prediction of the central pressure at the main vessel. For a single peripheral pulse, a SISO (single in single output) model is preferably used. Once a model is identified and the coefficients are defined, then the model can be used with the plurality of input peripheral pulses to estimate or predict the primary source arterial pulse.

**[0104]** Such stated pulse(s) can be produced by a measurement from different types of sensors including measurement of blood pressure, blood flow rate, blood temperature, or blood constituent concentration, such as hemoglobin, or glucose, or oxygen, or carbon dioxide content. The same principles of model identification can be applied using either the raw measured pulses or the estimated FAP pulses (full arterial pulse composed of the raw measured pulse plus the arterio-venous pulse) using any of these types of measurements.

**[0105]** Such models relating peripheral arterial (radial, femoral, renal, etc.) measurements and central arterial (aortic or pulmonary artery) measurements which can be identified using system identification methods can be used for estimation of such unknown blood constituent concentrations at a primary (central or main) blood vessel using only the measured peripheral blood constituent concentration. Such model based estimation methodology can lead to more accurate systemic estimation/measurement of such constituent concentration and therefore enable better medication delivery titration. Such medication delivery titration is desired for control or regulatory compensation of the desired constituent, especially with closed-loop feedback control systems, of which model free adaptive control or model predictive controllers (MPC) are two preferred embodiments. Similarly, a linear system identification, preferably state-space model, can be applied across peripheral Full Arterial Pulses (FAPs) such that an estimation of an output peripheral FAP from another input peripheral FAP can be obtained.

**[0106]** While there are available many tools and system identification strategies, the preferred embodiments of the present invention utilizes a direct system identification which uses state-space methods, and preferably in a multiple input single output (MISO) or single input single output (SISO) configuration, as more fully discussed below. The state-space MISO or SISO system identification allows implementation of system identification methods to the application of cardiovascular and pulmonary physiologic parameter estimation, and in the further improvement of application of the Balance parameter as input to such models to yield enhanced estimations to those parameters as model outputs. This invention applies the science of system identification to the realization of a feasible cardiovascular and pulmonary physiologic parameter estimation.

**[0107]** In FIG. 4, there is depicted a system transfer function  $G(t)$  which illustrates the relationships in the time and frequency domains between the inputs  $X(t)$  and outputs  $Y(t)$ . As can be seen, in the time domain the inputs  $X(t)$  are convoluted with the transfer function  $G(t)$  to produce the outputs  $Y(t)$ . On the other hand, in the frequency domain the inputs  $X(f)$  are simply multiplied in order to obtain the outputs  $Y(f)$ . Thus, the linear state-space model identification from frequency response function allows ease of adaptation of the system model using a change to its model order to optimize its approximation or estimation of the system outputs.

**[0108]** The state-space models (as described in reference 4, Linear Estimation, Kailath, Sayed, Hassibi. Prentice Hall, 2000, which is hereby incorporated by reference) are defined as

$$x(k+1)=Ax(k)+Bu(k)$$

$$y(k)=Cx(k)+Du(k)+n(k)$$

**[0109]** Where  $u(k)$ ,  $y(k)$ , and  $x(k)$  are time series of real numbers representing the input, output, and state variables, respectively, of the system, and  $n(k)$  is time series of real numbers representing the noise term which is assumed to be independent of the input sequence  $u(k)$ .  $A$ ,  $B$ ,  $C$ , and  $D$  indicate the coefficients vectors. The state variables represent the unforced dynamics of the system model.

**[0110]** By using the Fourier transform on both sides of both equations of the state-space model, we obtain the following, where  $\text{sup}$  indicates superscript, and  $\text{exp}$  indicates the exponent:

$$(\text{exp}.\text{sup}.jw)X(w)=AX(w)+BU(w)$$

$$Y(w)=CY(w)+DU(w)+N(w)$$

**[0111]** Where  $w$  is the frequency term,  $j$  indicates an imaginary number, and  $Y(w)$ ,  $U(w)$ ,  $X(w)$ ,  $N(w)$  are the frequency transformed output, input, noise, and state variables. The variables  $A$ ,  $B$ ,  $C$ ,  $D$  indicate the coefficients vectors. Where

$$G(\text{exp}.\text{sup}.jw)=G(z) \text{ at } z=\text{exp}.\text{sup}.jw=D+C(zI-A) \\ \text{sup}.-1)B \text{ at } z=\text{exp}.\text{sup}.jw \text{ and,}$$

$$Y(w)=G(\text{exp}.\text{sup}.jw)U(w)+N(w)$$

$Y(w)$  is the frequency response function (FRF) of the system.  $I$  represents the identity matrix and  $\text{sup}.-I$  represents a matrix inverse.

**[0112]** It will be noted that many other conventional System Identification methods exist and can provide substantially equivalent implementations to the state-space methods preferred in the present invention. These include (a) linear system identification (SYSID) methods, (b) nonlinear system identification methods, and (c) blind system identification methods. The background of each is discussed in details with algorithms describing prior art implementation details of such methods. The methods described apply for SISO (single input single output), MISO (multiple input single output), SIMO (single input (stimulus) multiple outputs (responses)), and MIMO (multiple inputs (stimuli) multiple outputs (responses)) configurations. This instant invention can equivalently, without loss of generality, use any of the other system identification methods mentioned below, and in any configuration of the inputs and outputs mentioned previously.

**[0113]** For example, as described in the references 1 through 6 listed in the Appendix, which are incorporated herein by reference, which describe linear systems estimation and system identification methods, the linear SYSID parametric and non-parametric methods include:

**[0114]** AR

**[0115]** ARX

**[0116]** ARMA

**[0117]** ARMAX

**[0118]** Generalized Linear

**[0119]** Output Error

**[0120]** Box-Jones

**[0121]** Continuous transfer function

**[0122]** Discrete transfer function

**[0123]** Impulse realization

**[0124]** User defined model

**[0125]** Principal components subspace identification

**[0126]** Discrete frequency transfer function from frequency response function

[0127] Continuous frequency transfer function from frequency response function

[0128] Maximum likelihood methods

[0129] The nonlinear SYSID methods include:

[0130] Neural Networks

[0131] Fuzzy Logic

[0132] Volterra Series

[0133] Weiner models (LMS, or recursive least square based)

[0134] Wavelets analysis

[0135] Nonlinear state-space models

[0136] The blind system identification methods include:

[0137] Laguerre model based

[0138] Deconvolution methods

[0139] The cardiovascular or pulmonary measured pulse waveform can refer to any of the following in continuous waveform or discrete values that are absolute or relative in value:

[0140] Blood pressure in any vascular location or cardiac chamber.

[0141] Blood flow rate in any vascular location.

[0142] Blood volume in any cardiac chamber.

[0143] Blood constituent concentration or chemical analyte concentration in any vascular location or cardiac chamber.

[0144] The system identification of the present invention can be either a SISO, if for example a single input channel was mapped to a single output channel, or, preferably, a MISO if the identified system was determined by mapping (or relating) multiple input channels (stimuli, measured cardiovascular or pulmonary parameters) into a single output channel (response, estimated cardiovascular or pulmonary parameter). Multiple inputs mapping provides greater informational content and therefore a better mapping accuracy to the output.

[0145] Alternatively, a MIMO configuration can be used where the input measured signals (measured cardiovascular or pulmonary parameters) are used to calculate multiple output responses (estimated cardiovascular or pulmonary parameters) in a single application of system identification methods rather than multiple applications of multiple sub-system identifications. Alternatively, a SIMO configuration can be applied where a single (measured cardiovascular or pulmonary parameter) is used as an input to determine system relationship with multiple outputs (estimated cardiovascular or pulmonary parameters).

[0146] FIG. 5 is flow diagram of an exemplary embodiment of the software used by the microprocessor in FIG. 3 for generating the balance parameter. The process 500 is performed by the processing unit 38. Initially, at step 502 the measured or observed arterial pulse waveform signal A of FIG. 2 is generated by the sensing device 34 and is received by the processing unit 38. At step 504, the processing unit calculates or derives the full arterial pulse (FAP) waveform signal B of FIG. 2 based upon the observed or measured arterial pulse waveform signal A. Next, in step 506 the processing unit calculates the arterial-venous (AV) pulse waveform signal C of FIG. 2 by subtracting the arterial pulse waveform signal from the FAP waveform signal. Finally, at step 510 the balance parameter 49 at the output of the processing unit is calculated by the ratio of the area under the curve of the AV pulse waveform signal, measured by integration over the cardiac cycle, to the FAP waveform signal, measured by integration over the same cardiac cycle,

in step 508. Alternatively, in a preferred embodiment, the ratio of the maximum peak value of the AV pulse waveform signal to the maximum peak value of the FAP pulse waveform signal over the same cardiac cycle is used. Whereas the DC offset can be either included in the integration, or normalized, or preferably removed by subtraction.

[0147] With attention directed to FIG. 6, there are shown a plurality of waveforms which are useful in explaining an embodiment of the present invention with respect to how certain key pressure points on the left ventricular pressure (LVP) or right ventricular pressure (RVP) signal can be determined based upon an aortic pressure or pulmonary artery measurement (or an estimate thereof). In particular, the key pressure points of interest include end-systolic pressure (PES), pre-diastole end-systolic pressure (PDPES), and end-diastolic pressure (PED). The timing timeline for atrial and ventricular systolic and diastolic phases are depicted to ease the interpretation of the full aortic pressure pulse, given the context of previous discussions in FIG. 2. It is important to note that the depicted indication of systolic and diastolic phases is from a volumetric perspective and not from a pressure perspective. The difference being the isovolemic periods. The atrial diastolic phase is 623. The atrial systolic phase is 624. The ventricular diastolic phase is 621. The ventricular systolic phase is 622. Adopting a pressure based convention for systolic and diastolic phase definition simply modifies the start and end of the defined phases.

[0148] PED substantially represents the pressure point on the ventricular pressure signal occurring pre iso-volemic contraction prior to the systolic contraction of the heart volume and after the diastolic expansion phase end. The end-systolic pressure (PES) and the PDPES pressure however are two separate pressure points on the ventricular pressure signal. The PDPES is indicative of the ventricular pressure value post the isovolemic relaxation phase and prior to diastolic expansion. The PED is indicative of the ventricular pressure value post diastolic expansion and pre-isovolemic contraction phase. While PED detection is generally easily detectable from the waveform of the ventricular pressure signal directly, the PDPES pressure point is not detectable directly therefrom since it is often affected by active forces of chamber muscle compression changing its relative timing.

[0149] The atrial diastolic blood flow demand creating filling of the atria and during the overlapping phase with ventricular isovolemic relaxation (indicated as part of ventricular volumetric diastole) is what creates the arterio-venous pulse, which is, as described herein, detectable at the aorta. Therefore, the timing signature marking the end of diastole for the atria must be detectable in the aortic pressure at the end of the arterio-venous pulse. This same event can be used to mark the end of the isovolemic relaxation event for the ventricle, after the systolic contraction phase end, at which point such pressure value in the ventricle (PDPES) represents substantially the post iso-volemic relaxation which occurs prior to diastolic expansion of the ventricular volume.

[0150] As can be seen in FIG. 6, the first waveform 605 is the measured ventricular pressure signal. The second waveform 601 is the measured aortic pressure (AOP) signal. The third waveform 603 is the measured electrocardiogram (ECG) signal, but can be equivalently replaced by an electrogram (EGM) signal. The fourth waveform 604 is a representation of the volume of blood in the ventricle.

Finally, the fifth waveform **602** is the first derivative of the aortic pressure (AOP) signal **601**.

[0151] The first key pressure point is PED which is designated by the dot **607** on the ventricular pressure signal **605**. The second key pressure point is PDPES which is designated by the dot **612** on the ventricular pressure signal **605**. The third key pressure point is the maximum ventricular pressure which is designated by the dot **608** on the ventricular pressure signal **605**. The fourth key pressure point is the point where ventricular pressure is equal to aortic pressure which is designated by the dot **609** located at a first intersection of the ventricular pressure signal **605** and the AOP signal **601**. The fifth key pressure point is the point where the ventricular pressure signal separates from and is no longer equal to the AOP signal which is designated by the dot **610** located at a second intersection of the ventricular pressure signal **605** and the AOP signal **601**.

[0152] Advantageously, the measurement of PDPES in this embodiment of the present invention is obtained from the measured waveform of the ventricular pressure signal **605** (or its estimate), and using event timing information derived from the waveform of the aortic pressure signal **601**. The derivative **602** of the aortic pressure signal depicts a small peak indicated by the dot **615**, at which point the ventricular chamber is at end systole, and the atrial chamber is at end diastole. At that event, the atrial diastolic blood flow demand (“pull effect”) of blood interaction with the ventricular blood flow supply (“push effect”) of blood reaches the phase end (end of arterio-venous pulse) where an aortic pressure first derivative **602** inflection point peak at that point indicates such event **615**.

[0153] Therefore, the PDPES (the dot **612**) from the ventricular pressure **605** can be measured (or estimated) at that event’s timing, which is also indicated with an inflection point **615** in the first derivative **602** of the aortic pressure **601**. It should be noted that the following observations can be made:

[0154] 1. PED can also be preferably detected at the ventricular pressure point when the ECG or equivalently EGM waveform triggers its R event in the QRS wave complex. PED alternatively can be measured from the ventricular pressure waveform directly using its 1st derivative valley.

[0155] 2. PES can be measured from the ventricular pressure waveform by finding the maximum value.

[0156] 3. The valve open event (the dot **609**) is detected preferably by the maximum peak **619** of the aortic pressure first derivative, or by the minimum valley of the aortic pressure, or by a peak of the first derivative of the ventricular pressure.

[0157] 4. The valve close event (the dot **610**) is detected preferably by the minimum valley **620** of the aortic pressure first derivative, or the valley of the aortic pressure at the dicrotic notch, or a valley of the first derivative of the ventricular pressure.

[0158] 5. The duration of the ventricular iso-volemic contraction event spans the duration of the end of P wave to R event in the QRS complex. Alternatively, preferably, it can be defined as spanning the QRS complex event duration.

[0159] 6. The duration of the ventricular iso-volemic relaxation event spans the duration of the electrical silence period of the heart.

[0160] The Balance parameter can be assessed from the aortic artery (or pulmonary artery) pulse signal using an estimation of the arterio-venous (AV) pulse signal, defined as the subtraction of the measured or observed arterial pulse from the full arterial pulse (FAP) signal at these two locations of the aorta or pulmonary artery. The aortic full arterial pulse (FAP\_aortic) (or equivalently the pulmonary artery full arterial pulse FAP\_pulmonary) is measured by extending a linear line **616** between the maximum pressure (peak-dot **608**) of the aortic pressure signal **601** until the end of the dicrotic notch, at which point it is advantageously substantially synchronous in timing with the PDPES (the dot **612**), which is detected from the aortic pressure signal (or its estimate) using an inflection peak (the dot **615**) of the first derivative of the aortic pressure signal as indicated in the waveform **602** of FIG. 6. An exponential line can also be used in place of the linear line **616** to model a non-constant rate of decrease of pressure as a linear line models a constant rate of decrease.

[0161] Still referring to FIG. 6, the small peak **615** of the first derivative (waveform **602**) of the aortic pressure signal **601** indicated at the vertical line **613**, corresponds with the pre-diastole end systolic event (PDPES) which is indicated on the ventricular pressure waveform **605** with the dot **612** and on the aortic pressure waveform **601** with the dot **611**. This PDPES event also corresponds with the ventricular volume being the smallest and is substantially coincident to the extended linear line **616** across the aortic pressure signal **601** used to define the FAP\_aortic pulse **606**. The vertical line **613** intersects the first derivative (waveform **602**) of aortic pressure signal **601** at an inflection point **615**.

[0162] Similarly, the end-diastolic event at which end-diastolic pressure (PED) is detected, defined by the ventricular pressure point post completion of the ventricular diastolic expansion, and prior to the onset of isovolemic contraction, is indicated as corresponding to the timing of the R-event **614** of the electrocardiogram (ECG) (or equivalently of an electrogram (EGM)) QRS-complex in the ECG (or equivalently EGM) waveform **603**. The PED value on the ventricular pressure waveform **605** is indicated by the dot **607**. The maximum ventricular pressure is also indicated on the waveform **605** by the dot **608**, which is equivalently the maximum aortic pressure on the waveform **601**. The dot **609** indicates the valve open pressure event, when both the ventricular pressure in the waveform **605** and the aortic pressure in the waveform **601** become equal. The dot **610** indicates the valve close pressure event, when the aortic pressure in the waveform **601** departs from the ventricular pressure in the waveform **605**.

[0163] With attention directed to FIGS. 10(a) and 10(b) of the drawings, there are shown, when joined together, a flow chart **1000** of an exemplary embodiment of the unique method of the present invention in FIG. 6 used to determine the PED and the PDPES values on the ventricular pressure signal for assessing the cardiac ventricular volumes and other cardiac performance parameters. Initially, in step **1001** a measurement (or an estimate thereof) of either the aortic pressure (AOP) waveform for the left heart performance or the pulmonary artery pressure (PAP) for the right heart is obtained, which is similar to the waveform **601** shown in FIG. 6. A measurement of the electrocardiogram (ECG) (or electrogram (EGM)) waveform is obtained in step **1002**, which is similar to the waveform **603** shown in FIG. 6. In step **1003**, the first derivative waveform of the waveform

obtained from step 1001 is computed, which is again similar to the waveform 602 depicted in FIG. 6. Next, in step 1004 the valve open event (the dot 609 in FIG. 6) is determined by using the maximum peak of the first derivative waveform obtained from the step 1003. Similarly, in step 1005 the valve close event (the dot 610 in FIG. 6) is determined by using the minimum valley of the first derivative waveform obtained from the step 1003.

[0164] Further, the end systolic pressure PES event is determined in step 1006 by using the maximum value of the waveform obtained from step 1001, which is similar to the dot 608 in FIG. 6. In step 1007, the full arterial pulse waveform of the aortic pressure AOP (FAP\_aortic) or the pulmonary artery pressure PAP (FAP\_pulmonary) from the step 1001 is calculated. In step 1008, the arterio-venous (AV) waveform of the aortic pressure AOP (AV\_aortic) or the pulmonary artery pressure PAP (AV\_pulmonary) from the step 1001 is calculated. In step 1009, the end (the dot 611) of the arterio-venous pulse event of the AV waveform in step 1008 is determined. This AV pulse event becomes zero for that heart cycle, which is substantially coincident in time with the pre-diastolic end-systolic pressure PDPES event, which is the dot 612 in FIG. 6, defined by the ventricular pressure point post completion of the isovolemic relaxation and prior to the onset of diastolic expansion.

[0165] Continuing in the process 1000, in step 1010 the Left-Right Balance (LRB) or Right-Left Balance (RLB) parameter is calculated dependent upon if AOP or PAP was used, respectively, in step 1001. This Balance parameter is based upon the respective FAP and AV generated using a transfer function, preferably using the ratio of the AV area under the curve, measured by integration over the cardiac cycle, to FAP area under the curve, measured by integration over the same cardiac cycle, or using the ratio of the maximum peak value of the AV pulse to the maximum peak value of the FAP pulse, or by some other relating transfer function, from the corresponding steps 1007 and 1008 as well as a measurement of the heart rate HR. Other relating transfer function includes the sensitivity relationship between AV and FAP pulses of a pulse waveform measure, preferably the area under the curve or maximum peak value, whereas sensitivity is preferably calculated using the NNSS method described herein or a partial derivative or full derivative based method. Whereas the DC offset can be either included, or normalized, or removed by subtraction. In step 1011, the timing of the R-wave event 614 in the QRS segment of the ECG (or EGM) waveform 603, which is substantially coincident with the end diastolic PED event (the dot 607), is determined as illustrated in FIG. 6. The heart rate is also measured in the step 1011. In step 1012, the time difference between the R-wave event 614 in the step 1011 and the valve open event 609 in step 1004 is calculated as a delta T1.

[0166] In step 1013, by using the slope of the line segment of the aortic pressure (AOP) (or pulmonary artery pressure (PAP)) pressure between the valve open event 609 and a short time after the valve opening (up to a period of a point approximately mid-way 617 of the maximum AOP (or PAP) pressure). The PED value of the ventricular pressure signal 605 can be extrapolated backwards from the AOP (or PAP) waveform signal 601 (or an estimate of it). The extrapolation period is defined by calculating the time between the ECG R event (614) and the valve open event (609) as delta T<sub>i</sub> of step 1012. The extrapolation slope is defined by measuring

the linear line fit across the AOP (or PAP) pressure points 609 and the selected point 617. This AOP (or PAP) line segment (x-values representing time, and y-values representing pressure values) is extrapolated backwards in time up to the x-axis point in time to estimate the PED event (the dot 607), as defined by the time occurrence of the R-event of the ECG (or EGM) waveform 603. The y-axis value based upon the result of this extrapolation is the PED pressure value at point 607, which is considered a first static pressure point. For a tiny fraction of a second during the heart cycle, at this first static pressure point, all positive and negative dynamic forces applied on the contained blood volume in the chamber are nulled, and the ventricle is in its largest volume. The first static pressure therefore represents the pressure only due to the volume of the blood in the ventricle at its largest volume.

[0167] In step 1014, the time difference between the end of the AV pulse event in the step 1009 and the valve close event in step 1005 is calculated as a delta T<sub>2</sub>. Next, in step 1015 by using the slope of the line segment of the AOP (or PAP) pressure between the valve close event 610 and a short time before the valve closing (up to a period of a point approximately mid-way 618 of the maximum AOP (or PAP) pressure). The PDPES value of the ventricular pressure can be extrapolated forward from the measured AOP (or PAP) pressure (or an its estimate of it). The extrapolation period is defined by calculating the time between the end of AV pulse event (611) and the valve close event (610) as delta T<sub>2</sub> of step 1014. The extrapolation slope is defined by measuring the linear line fit across the AOP (or PAP) pressure points 610 and the selected point 618. This AOP (or PAP) line segment (x-values representing time, and y-values representing pressure values) is extrapolated forward in time up to the x-axis point in time to estimate the PDPES event, as defined by the time occurrence of the small peak 615 in the first derivative (waveform 602) of the AOP (or PAP) pressure post the valve close event. The y-axis value based upon the result of this extrapolation is the PDPES pressure value at point 612, which is considered a second static pressure point. For a tiny fraction of a second during the heart cycle, at this second static pressure point, all positive and negative dynamic forces applied on the contained blood volume in the chamber are nulled, and the ventricle is in its smallest volume. The second static pressure therefore represents the pressure only due to the volume of the blood in the ventricle at its smallest volume.

[0168] In another aspect of the present invention, the inventor has realized that the variability of the Balance parameter (LRB or RLB) is a viable indicator of cardiac health state. The balance parameter is a very sensitive and specific measure for pulse variability, reflecting variation in supply and demand of blood, rather than the variation in the blood pressure or flow alone. It is also sensitive enough to detect respiration modulation effect of the cardiac cycle and advantageously allows the detection of respiration rate and parameters directly from the balance parameter variations.

[0169] With reference now to FIG. 7, there are shown a balance parameter 701 indication with a plurality of cardiac auto-regulation cycles modulating a plurality of respiratory cycles and cardiac cycles. Each point on the plot represents a single cardiac cycle. A single respiratory cycle portion of 701 is expanded and indicated in 703 boxed area. A single cardiac auto-regulation cycle of 701 is expanded and indicated in 702 boxed area. Thus, in this aspect of the present

invention, the respiration rate is detected from simple variation cycle of the balance parameter. Furthermore, the Balance parameter is sensitive enough to detect the cardiac auto-regulation cycle **702** which maintains the regulation of the balance between the left and right cardiac outputs. This longer period regulation cycle **702** was previously not measurable or realizable, but is now advantageously detected by the balance parameter of the instant invention. Characterization of auto-regulation function is therefore an important aspect of this invention. Detection of abnormalities, in the auto-regulation function and its cycle characteristics, alarming against auto-regulation abnormalities, and display and communication of auto-regulation information to users are all part of the instant invention.

[0170] With reference to FIG. **8(a)**, there is depicted the normal auto-regulation function to the cardiac cycle **805** as a vital indicator to the health state of the heart. Biased balance indication (i.e. too high, or too low) is an early indicator of heart disease. In FIG. **8(b)**, there is also depicted a perturbation at **806** causing the balance parameter to start an increasing trend **806** through **807**, which is not be easily detectable using blood pressure signal alone. As shown in FIG. **8(b)**, when the heart experiences abnormal ischemia **811** the auto-regulation periodic cycle pattern of the balance parameter disappears **813**, and when arrhythmia or fibrillation is experienced **812** a high variance of the balance parameter is indicated. Abnormal cardiac conditions such as arrhythmia or fibrillation are detected with the variance of the balance parameter above a threshold and can be indicated for an alarm. Restoration of auto-regulation periodic cycle is an essential indication of restoration of normal function to cardiac cycle. Absence of auto-regulation or other abnormal cardiac conditions are detected with the variance of the balance parameter below a threshold and can be indicated for an alarm. Abnormal cardiac condition detection and indication can be used for closed loop feedback in the delivery of therapeutic medications or to invoke therapy using a pacemaker, cardioverter, or a defibrillator.

[0171] With reference to FIG. **12**, there is illustrated a flow chart **1200** of an exemplary embodiment of the method of FIG. **7** and FIG. **8** for using the balance parameter for measuring the cardiac regulation cycle. By separating the respiration signal component **703** from the regulation signal component **702**, it is possible to obtain a better assessment of each of the individual components independently.

[0172] In the process **1200**, in the step **1201** an arterial pressure waveform (i.e., from an peripheral artery or pulmonary artery or aortic artery) is measured initially. In step **1202**, a full arterial pulse (FAP) and an arterio-venous (AV) pulse are calculated. In step **1203**, the balance parameter (LRB or RLB) is determined for each of heart pulse over a long duration of the heart pulses. In step **1204**, there is applied optionally a filtering to smooth out the balance parameter from step **1203**. In step **1205**, demodulation of the respiratory signal carrier is applied to obtain the regulatory signal carrier, or alternately demodulation of the regulatory signal carrier is applied to obtain the respiratory signal carrier.

[0173] However, this is difficult to achieve effectively since the respiration rate is variable. One preferred method is to use adaptive filters so as to filter out an undesired component (such as either respiration or regulation) in order to obtain the desired component (such as either regulation or respiration, respectively). Adaptive filters require a refer-

ence signal to indicate the component to be separated out. This reference signal is more easily provided as the respiratory signal which can be measured with a variety of methods, including motion sensitive accelerometers, or tension belt or modulation of the ECG waveforms, or nasal air flow sensor. Also, there are conventional tools known in the prior art, such as matrix-pencil methods for frequency estimation, can be used to separate very effectively the component frequencies representing the cardiac auto-regulation cycle from the respiration cycle.

[0174] In step **1206**, a peak-valley detection is optionally applied to the separated components from step **1205** to determine the end-of-cycle period, frequency, and duty cycle. Spectral frequency analysis (e.g. Fourier series) is performed in step **1207** to detect the primary regulation and respiration rate. In step **1208**, other respiratory parameters can also be derived such as the inhalation and exhalation duty cycle. The detected respiratory rate can be utilized to monitor and provide alert for respiratory disorders, including respiratory apnea in a highly sensitive, specific, and reliable method, that is not affected by additional noise sounds such as snoring or air flow measurement errors. This method can also be combined with anesthesia ventilators in order to determine patient restoration of respiratory function after respiratory depression with anesthesia or pain medications, to allow determination of optimal ventilator settings to wean patient of ventilator and to control the ventilator in closed loop to achieve patient weaning off ventilator. The respiration waveform can also be used to more generally assess patient respiratory function and health state. In step **1209**, by using the auto-regulation cycle rate, its period or frequency, or duty cycle, baseline trend, maximum, minimum, mean and standard deviation statistical distribution and properties, and heart rate can all be determined.

[0175] In step **1210**, the auto-regulation cycle and respiratory cycle characteristics and information can be used to assess patient health state and alarm against abnormalities, and determine therapeutic measures or decisions to be applied, including closed loop control of medication delivery or cardiac interventional devices. The balance parameter and auto-regulation performance parameters can be used to optimize the treatment protocol of renal dialysis such controlling the blood flow rate into the dialyzer to avoid abnormal cardiac balance and autoregulation performance.

[0176] The balance parameter and auto-regulation performance parameters can also be used in feedback closed loop control of dialysis devices to achieve optimal dialysis treatment control. The balance parameter and auto-regulation performance can also be used for detection and alarming of cardiac stress, patient pain, or surgical nociception, whereby marked trend variations in the balance levels and regulation cycle are associated with elevated pain levels or cardiac stress.

[0177] The balance parameter and auto-regulation performance parameters can be used to provide feedback control to ventricular assist devices and counter-pulsation devices to help optimize the blood flow for a more balanced auto-regulation of the heart cycle.

[0178] The balance parameter and auto-regulation performance parameters can be also used for guidance and optimization of lead placement locations of a pacemaker system to provide a more balanced auto-regulation performance of the heart cycle. The cardiac balance and autoregulation performance parameters can be used to evaluate blood flow

between the mother and the fetus and can be used to study the fetus cardiac regulation and balance as well as the mother's cardiac regulation and balance, and the interaction of both over the term of the pregnancy, to ensure that both mother and fetus are receiving optimal blood flow, and to ensure baby healthy rate of growth and nutrition with adequate blood flow. Abnormal fetal blood flow in relationship to mother's blood flow can be detected and alerted.

[0179] In FIG. 13, the balance parameter can be measured to indicate the balance between the two input parameters the supply 1307 to the pulmonary circulation of blood flow on the right side 1309 by the right ventricle (RV) 1301 and the demand 1308 for blood flow which is returned the left atrium (LA) 1302 on the left side 1310. The balance parameter can also be measured to indicate the balance between the two input parameters the supply 1312 to the systemic circulation of blood flow on the left side 1310 by the left ventricle (LV) (1303, 1313) and the demand 1311 for blood flow which is returned to the right atrium (RA) (1304, 1309). Furthermore, the balance between the two blood flow circulations, the pulmonary circulation 1305 and the systemic circulation 1306, can indicate the balance between the two input parameters said pulmonary circulation balance and said systemic circulation balance.

[0180] In FIG. 14, the balance parameter (balance ratio or balance trend) between the supply or demand for blood flow can be used to indicate the auto-regulation function of the heart 1401. The auto-regulation of the heart's supply and demand can be characterized by a regulation cycle 1402 with characteristic time period and amplitude and waveform morphology. The regulation cycle that is sinusoidal indicates healthy cardiac variability in shifting its bias of balance between the blood supply versus the blood demand sides of the balance parameter. The regulation cycle is also indicative of a characteristic respiration cycle 1403. The respiration cycle typically modulates the individual cardiac cycles, but the regulation cycle further regulates both the respiration cycle and the individual cardiac cycles. The cardiac auto-regulation cycle can therefore be used to measure global cardiac heart rate variability and blood flow variability. Also abnormal cardiac auto-regulation cycle can indicate abnormal auto-regulation or a disease state in either the supply side or the demand side of the cardiac cycle function, such as congestive heart failure, or regurgitating cardiac valve, or poor cardiac output, or constriction of a vein or artery, or cardiac muscle arrhythmia or fibrillation. Further testing can further narrow the source root cause of the cardiac imbalance or loss of auto-regulation sinusoidal pattern.

[0181] In FIG. 15, we depict two measures of the cardiac balance, the first is a balance ratio of the supply portion and demand portion of the blood flow pulses, and the second is balance trend of the supply portion and demand portion of the blood flow pulses. The balance ratio 1501 represents the ratio between the blood flow supply parameter represented by the area under the curve for the full arterial pulse and the demand parameter represented by the area under the curve for the arterio-venous pulse). This is an important parameter to view and normalize or track over time per patient, and to evaluate cardiac regulation characteristics relative to its historic performance for that patient. However, such a ratio may be different across patients due to different heart chamber size, and cardiac output and contractility performance variation, blood pressure and flow variation, and vascular system characteristics, and organ demand variation.

The cardiac balance trend 1502 provides a metric of the sensitivity (i.e. the partial derivative) between the blood flow supply parameter represented by the area under the curve for the full arterial pulse and the demand parameter represented by the area under the curve for the arterio-venous pulse). Both the balance ratio and the balance trend can be used to indicate circulatory occlusions such as inferior vena cava (IVC) (1503, 1504) or effects of blood pressure stressor medication or a beta blocker medication to reach is target or desired balance level. A typical desired balance level for the balance trend is the zero level 1505 indicating perfect balance between the supply and demand portions or parameters.

[0182] In FIG. 16, the balance ratio (1601, 1611) and balance trend (1602, 1607) are depicted to indicate detection of cardiac auto-regulation 1603, left anterior descending (LAD) occlusion (1604, 1608), inferior vena cava (IVC) occlusion 1606, and congestive heart failure 1610 clinical conditions. The zero-level line (1605, 1609) represents the ideal target for the balance trend parameter representing a perfect sensitivity or balance between the blood flow supply and demand without bias on either sides.

[0183] We measure the relative sensitivity between a reference (target) variable  $R(t)$  (as input  $u(t)$  or output  $y(t)$ ) and a process variable  $P(t)$  (as output  $y(t)$  or input  $u(t)$ ) using an accumulative sum of weighted or scaled terms for each of the time steps within a selected moving time window, each of the terms incorporates the partial derivative  $\partial y(n)/\partial u(n)$  as the gradient of  $y(t)$  with respect to  $u(t)$ , which represents the sensitivity of the output  $y(t)$  to variations of the input  $u(t)$ . Here, we define  $o(n)$  as the weighted sensitivity function of the process. Since the process is unknown, the sensitivity function is also unknown, but it is adaptively defined and resolved as the sensitivity function defines the relationship between the two inputs. An object of this invention is thus the use and application of this neural network sensitivity sensing (NNSS) function to resolve the sensitivity of clinical, physiologic, anatomic, metabolic, chemical, or physical variables to each other in clinical applications where it is often the case that the sensitivity between two variables is unknown, and this method allows us to resolve the sensitivity relationship between these two variables in a numerically stable way. This sensitivity function is based on the model free adaptive controller (MFA) that is defined in U.S. Pat. No. 6,055,524, which is hereby incorporated by reference. The '524 patent describes the use of a neural network that is capable of adapting to the sensitivity of the input functions specifically for use in control applications. In this invention, a similar neural network is defined but used solely for the purpose of open-loop sensitivity detection across multiple variables, but not for the purpose of closed-loop control as is claimed by the '524 patent and described in FIG. 20. The signals shown in FIG. 20 above are as follows:

[0184]  $r(t)$ -reference target signal

[0185]  $y(t)$ -Measured Variable or the Process Variable,  
 $y(t)=X(t)+d(t)$

[0186]  $X(t)$ -Process Output

[0187]  $u(t)$ -Controller Output

[0188]  $d(t)$ -Disturbance, the disturbance caused by noise or load changes that affect the sensitivity of the relationship to the reference  $r(t)$ .

[0189]  $e(t)$ -Error between the reference and measured Variable Since  $e(t)=r(t)-y(t)$

**[0190]** The objective of the neural network sensitivity sensor (NNSS) is to minimize the error between the reference and the measured variables. By reducing the error to zero, it is converging on the right sensitivity relationship between the two input variables. Linear and nonlinear relationships can be converging onto for detection of the sensitivity function. The gain of the process allows us to set the rate at which such sensitivity convergence takes place through the time updates of the convergence process.

**[0191]** The error reduction objective function can be set as

$$E(t)=(r(t)-y(t))/2$$

**[0192]** The activation functions are sigmoid defined herein as,

$$\text{Activation function } \phi(x)=1/(1+e^{-x})$$

**[0193]** Each output signal from the hidden layer is conveyed to the single neuron in the output layer via a path weighted by an individual weighting factor  $h_i$ , Where  $i=1, 2, \dots, N$ . These signals are summed in adder with  $h_0=1$ , the threshold signal for the output layer, and then filtered by activation function. The signal  $p(t)$  gets filtered by an activation function to produce  $q(t)$ .

**[0194]** The following function maps the range output of the neural network (0, 1) back to real space to produce the  $o(n)$  function Mapping function  $\psi(y)=\ln(y/(1-y))$

**[0195]** The following equations govern the update process for the neural network

$$p_j(n)=\sum w_{ij}(n)E_i(n)+1$$

$$q_j(n)=\phi(p_j(n))$$

$$o(n)=\psi(\phi(\sum E_j(n)q_j(n)+1))$$

$$o(n)=\sum h_j(n)q_j(n)+1$$

**[0196]** For single input single output (SISO) SSNN, the error is also feed forward to be added to  $o(t)$ , and multiplied by a Gain factor,  $K$ , as follows:  $v(t)=K(o(t)+e(t))$ .

**[0197]** For multiple inputs and multiple outputs (MIMO), or multiple inputs and single output (MISO) or single input and multiple outputs (SIMO) configurations, this feed forward operation occurs at the system level and not at the subsystem level as described below.

**[0198]** Where  $n$  denotes the  $n$ th iteration,  $o(t)$  is the continuous function of  $o(n)$ ,  $v(t)$  is the output of the neural network sensitivity sensor (NNSS),  $K>0$ , called sensitivity sensor (SS) gain, is a constant used to adjust the magnitude of the SS. This gain is used to adjust the NNSS solution convergence performance in a stable way. The real-time learning algorithm is developed to continuously update the values of the weighting factors of the NNSS as follows and as depicted in FIG. 21:

$$\Delta w_{ij}(n)=\eta K(\partial y(n)/\partial u(n))e(n)q_j(n)E_i(n)\sum h_k(n)$$

$$\Delta h_j(n)=\eta K(\partial y(n)/\partial u(n))e(n)q_j(n)$$

Where  $n>0$  is the learning rate or weight update rate.

**[0199]** Similarly the sensitivity due to MISO, MIMO, or SIMO configurations can be defined by same process between each of the input variables and each of the output variables, and the final sensitivity is a weighted sum of all the individual sensitivity relationships between each of the inputs and each of the outputs. The MIMO, MISO, or SIMO

sensitivity sensor is comprised of individually interconnected NNSS with combining of the outputs from different subsystems for each input.

**[0200]** In FIG. 22(a), where  $r$  is a reference which is compared with a difference operation to a measured process variable  $y$  to produce the residual error  $e$ . The error  $e1$  is the difference between a first process variable  $y1$  and its reference  $r1$ , and error  $e2$  is the difference between a second process variable  $y2$  and its reference  $r2$ . The error variables  $e1$  and  $e2$  are provided as inputs into sub-sensors  $S11$ ,  $S12$ ,  $S21$ ,  $S22$ , where  $v11$ ,  $v12$ ,  $v21$ , and  $v22$  are the sensors outputs, which get cross-combined in a sum Adder, optionally with a weighted sum, to produce the combined sensor outputs  $u1$  and  $u2$ .

**[0201]** In FIG. 22(b) depicts using two process inputs in a MIMO relationship with two process outputs, cross weighted adder is used. The input signals  $u1$ ,  $u2$  are passed as an input into a process MIMO process  $G$ , which is represented by four sub-processes  $G11$ ,  $G21$ ,  $G12$ ,  $G22$  that outputs  $x11$ ,  $x12$ ,  $x21$ ,  $x22$  and that combined, optionally via a weighed sum, with disturbance (noise)  $d1$ ,  $d2$  to produce the final MIMO NNSS output  $y1$  and  $y2$ , which is a measured variable.

**[0202]** In FIG. 23, depicts the relationship between blood flow supply and demand balance parameter trend **2301**, blood flow supply and demand balance parameter ratio **2302**, and the pulse variability **2303**. The balance ratio is a parameter that is typically greater than 0, but varies on its normal level from person to person. Variability in the balance ratio parameter can be used to indicate changes normal vs. abnormal for the same person. Population based normalization of the balance ratio can be done. The Balance trend parameter is typically centered around zero for a balanced blood flow supply and demand relationship. Variation from zero indicates bias either to the supply side ( $>0$ ) or the demand side ( $<0$ ). Population normalization in this case is not required to understand such bias relative to the zero line. The pulse variability indicate the variance of the heart rate within a moving time window across the heart rate time series, as known in the art. Waveform events indicated by Heart rate variability are also indicated in the balance parameter as shown graphically in **2303** and **2301** and **2302**. The pulse variability is also related to hydration status or cardiac output.

**[0203]** In a preferred embodiment, the cardiac balance is calculated using the Sensitivity between the Arterio-venous pulses signal, or their derived measures or components, and the full arterial pulses signal, or their derived measures or components. Said sensitivity is preferably calculated using the Neural Network Sensitivity Sensor (NNSS) method described herein or using a partial derivative or full derivative based method. Said derived measures or components include, but are not limited to, the maximum (peaks values) for each of the cardiac pulses signal, the integral (area under the curve) for the variable part for each of the cardiac pulses signal, the variance or variability for each of the pulses signal, and known in the art wavelet components of the said cardiac pulses.

**[0204]** The variance (or similarly standard deviation) of the Balance parameter has been advantageously found by the inventor to be a strong indicator of the health status of the subject's cardiac function. If the Balance parameter variance is too high as illustrated at **812** in FIG. 8(b), then it is an indicator of cardiac instability, potentially as an

indicator or measure of surgical nociception or pain, arrhythmia, ischemia or myocardial infarction, as some causes for high balance variance. These are typically associated with loss of the auto-regulation function indicated by the balance parameter at **811** and **813**. The absolute value of the balance parameter is used as an indicator of cardiac imbalance representing a biased performance toward the left or right heart blood pumping (cardiac output). Such an imbalance can lead to the accumulation of blood in parts of the circulatory system which can lead eventually to cardiac arrhythmia, or heart failure such as congestive heart failure.

**[0205]** If on the other hand the variance of the heart is too low from its normal auto-regulation cycle variance baseline, then this also is an indicator of hydration status or other cardiac problems including, for example hypovolemia, or low hydration or low cardiac output status, and is a strong predictor of heart failure. There exists a margin of normal variance of the Balance parameter about which the cardiac performance is automatically adaptively maintained by the heart during the auto-regulation cycle.

**[0206]** It is still another aspect of this invention to use both the Balance parameter as a mean target value set point and the Balance parameter variance (or standard deviation) margin as a second target set point in control of medication delivery that affect these cardiac performance parameters to the desired set target levels or margins. It is still yet another aspect of this invention is to use the measurement of Balance parameter variance at **812** as well as the absolute value of the Balance parameter at **807** for purposes of monitoring, predicting, and/or alerting against cardiac arrhythmias, fibrillation, ischemia, or heart failure such as congestive heart failure (CHF), or detection of onset of seizures and alarming against it.

**[0207]** In addition, valvular malfunction (regurgitation or insufficiency) will affect the appearance or timing of the Dicrotic Notch. It is thus envisioned that this invention can also use the RLB Balance and/or LRB Balance parameters to diagnose the presence or absence of as well as the degree or status of valvular regurgitation or insufficiency. Furthermore, another aspect of this invention is to use the balance parameter for indication of valve regurgitation or insufficiency.

**[0208]** Further, a cardiovascular control support system that provides pumping of a balloon in order to displace blood volume whereby the device is placed in a vascular artery or vein is used to achieve therapeutic or optimal Right-Left Balance levels. Another aspect of this invention is to use the Balance parameter as a process variable in a closed loop control system for controlling a counter pulsation (balloon) apparatus in order to achieve a desired set target value for cardiac parameters, including Balance parameter, blood pressure, and blood flow rate or cardiac output. A model predictive controller (MPC) or a model free adaptive controller can be preferably used for this purpose.

**[0209]** In yet another embodiment of the present invention, an estimated aortic pressure waveform can be derived by first estimating its components of FAP\_aortic and AV\_aortic and then subtracting the estimated AV\_aortic waveform from the estimated FAP\_aortic waveform. In FIG. 9, there is depicted a flow chart **900** of an exemplary technique for determining the estimated aortic pulse waveform, defined by one of aortic blood pressure, blood flow rate, and blood constituent concentration. Initially, at step **901** the aortic pulse is measured. Simultaneously, at step **904**

the (peripheral) arterial pulse (e.g., radial or femoral or carotid) is measured, defined by one of arterial blood pressure, blood flow rate, and blood constituent concentration. In step **902**, the aortic's full arterial pulse (FAP\_aortic) waveform is computed as defined herein. Similarly, the peripheral artery's full arterial pulse (FAP\_arterial) is computed in step **905**. Further, the aortic arterio-venous pulse (AV\_aortic) is computed in step **903**. The (peripheral) arterial arterio-venous pulse (AV\_arterial) is computed in step **906**.

**[0210]** In step **907**, the relationship or transfer function between the FAP\_arterial as input and the FAP\_aortic as output can be defined using many modeling and adaptive methods, including a system identification method, as mentioned herein, preferably using linear system identification producing a state-space linear model to define such transfer function. In one preferred embodiment, the mapping between FAP\_arterial and the FAP\_aortic, the transformation's transfer function (preferably state-space model's coefficients) can be identified easily. Such model creates a calibration transfer function when both FAP\_arterial and FAP\_aortic are available. In step **909**, such calibration transfer function can then be used with input FAP\_arterial to produce a more accurate estimate of the output FAP\_aortic. Similar or equivalent steps can be taken as above with FAP\_pulmonary derived from the pulmonary artery pulse replacing the FAP\_aortic, defined by one of pulmonary artery's blood pressure, blood flow rate, and blood constituent concentration.

**[0211]** Continuing in the process **900**, in the step **908** the (peripheral) arterial pulse (radial or femoral or carotid) arterio-venous pulse (AV\_arterial) as defined herein, is used to estimate the aortic's arterio-venous pulse (AV\_aortic) waveform as defined above. The relationship or transfer function between the AV\_arterial as input and the AV\_aortic as output can be defined using many modeling and adaptive methods, including a system identification method, as mentioned herein, preferably using linear system identification producing a state-space linear model to define such transfer function. In one preferred embodiment, the mapping between AV\_arterial and the AV\_aortic, the transformation's transfer function (preferably state-space model's coefficients) can be identified easily. Such model creates a calibration transfer function when both AV\_arterial and AV\_aortic are available. In step **910**, such calibration transfer function can then be used with input AV\_arterial to produce a more accurate estimate of the output AV\_aortic. Similar or equivalent steps can be taken as above with AV\_pulmonary derived from the pulmonary artery pulse replacing the AV\_aortic. Finally, in step **911** an estimate of the aortic pulse waveform can be derived by subtracting the estimated AV\_aortic waveform from the estimated FAP\_aortic waveform.

**[0212]** With reference to FIG. 11, there is shown a flow chart **1100** of an exemplary embodiment for estimating the other ventricular chamber's performance based upon information from the one ventricular chamber. The left heart's (VED-VES) is estimated, on average, using a Balance adjusted right cardiac output (RCO) divided by the heart rate (HR). If measured cardiac output is performed in the right heart, the volume difference (VED-VES)=RCO/HR. The parameter RCO can be adjusted using the Balance parameter value (Left-Right Balance (LRB) from any aortic branch

artery or Right-Left Balance (RLB) from any pulmonary branch artery) for the estimation of the left heart's cardiac output LCO.

[0213] In step 1102 of FIG. 11, the left ventricle's (LV) cardiac output is calculated as either  $LCO=(RCO/LRB)$  or  $LCO=RCO/(1-RLB)$ . Similarly, in step 1101 the right ventricle's (RV) cardiac output is calculated as either  $RCO=LCO*LRB$  or  $RCO=LCO*(1-RLB)$ . Both LRB and RLB are  $<1.0$  (i.e.  $<100\%$ ). The vascular vessel tree at the output of the left ventricle presents a narrowing diameter vessels which resulting in higher pressure requirement to pass fluid through it as opposed to the vascular tree at the input to the right atrium presenting and expanding diameter vessels which lowers the pressure measured at the right atrium, while maintaining average flow rate between the two chambers when the heart is in balance. The left cardiac pressure will have inefficiencies of transfer (e.g. backflow portion or tissue fluidic uptake). A higher RLB is associated with a lower LRB and is typically associated with left heart abnormal function. Similarly, a lower RLB is associated with a higher LRB and is typically associated with right heart abnormal function. A difference between the LCO and RCO is indicative of imbalance in RLB or LRB, which is indicative of cardiovascular disease state. In normal heart circulation, while the average left cardiac output out of the left ventricle is at higher pressure but it is generally at equivalent average flow rate as the right cardiac output from the right ventricle, which is at lower pressure.

[0214] Summarizing the above description of the present invention with respect to FIGS. 9 and 10, a plurality of peripherally measured arterial pulse in step 904 is used to derive a FAP\_arterial waveform in step 905, and a main artery (aortic or pulmonary artery) measured arterial pulse in step 901 is used to derive its FAP in step 902. Then, applying a calibration step, the plurality of input derived (peripheral)-arterial waveform(s) is/are mapped to output primary main artery FAP\_aortic (or FAP\_pulmonary) using a transfer function defined by a system identification method at respective steps 907 and 908. Preferably, the system identification method is a linear state-space model in a MISO (multi-input single-output) or SISO (single-input single-output) configuration.

[0215] Then, in the steps 909 and 910, there is provided continuously acquiring (peripheral) arterial pulse, deriving the FAP\_arterial and employing the previously identified transfer function to produce an estimate of the FAP\_aortic. In the step 911, there is provided using the continuously produced estimate of the FAP\_aortic to estimate the left ventricular (LV) pressure points of PED and PDPES obtained from the process 1000. Equivalently, there is provided using the continuously produced estimate of the FAP\_pulmonary to estimate the right ventricular (RV) pressure points of PED and PDPES.

[0216] In the process steps of FIG. 10(b), there is provided using the PED and PDPES estimated in conjunction with measured or estimated (adjusted) cardiac output and heart rate and Balance parameter to produce cardiac parameters including, but not limited to, absolute values of end-diastolic volume, end-systolic volume, and additional cardiac parameters known-in-the-art such as stroke volume, cardiac output, ejection fraction, systemic vascular resistance, contractility, and stroke work. Furthermore, there is produced an

estimate or measurement of the duration of the ventricular iso-volemic contraction period and the ventricular iso-volemic relaxation period.

[0217] In FIG. 17 and FIG. 18, depicts a step of single time calibration between an invasively measured cardiac blood pressure waveform, preferably aortic blood pressure 1702, and a more accessible peripherally measured blood pressure waveform, preferably radial arterial blood pressure 1701, and more preferably noninvasively measured blood pressure waveform from which we measure cardiac pressure and volumes. The details of which are described in patent application 2015/0025328 to Khair. A system identification method, preferably linear state space model is identified and used to map the peripherally measured blood pressure input waveform to the output invasively measured cardiac blood pressure waveform, preferably aortic blood pressure. After the calibration step, then we continuously measure the peripheral blood pressure waveform 1704 and use it as input to said identified model during the said calibration step to estimate the cardiac blood pressure waveform, preferably aortic 1705. From the estimated aortic blood pressure waveform, we can then continuously obtain estimated aortic blood pressure systolic and diastolic values and calculated cardiac chamber (left ventricle) systolic and diastolic pressures, and from which we can calculate diastolic volume 1801 and systolic volume 1802, and other cardiac hemodynamics including stroke volume 1803, cardiac output, ejection fraction 1804, and contractility.

[0218] FIG. 19, depicts a comparison between actual end diastolic volume 1901 and estimated end diastolic volume 1902, and actual end systolic volume 1903 and estimated end systolic volume 1904, and actual stroke volume 1905 and estimated stroke volume 1906, and actual ejection fraction 1907 and estimated ejection fraction 1908.

[0219] From the foregoing detailed description, it can thus be seen that the present invention provides a system for measuring of cardiac blood flow balance parameter between the right chamber of the heart and the left chamber of the heart which includes a sensor device for measuring one of blood pressure and blood flow rate and blood constituent concentration of a patient so as to generate an arterial pulse signal. A processing unit is responsive to the arterial pulse signal for generating a full arterial pulse (FAP) signal, an arterio-venous (AV) pulse signal, and a balance parameter. A computational device is responsive to the balance parameter for further generating a set of physiological parameters. A display station device is responsive to the set of physiological parameters from the computational device for displaying meaningful information.

[0220] In summary, we presented a system for measuring of cardiac blood flow balance parameter between the right chamber of the heart and the left chamber of the heart, comprising: sensing means for measuring one of blood pressure and blood flow rate and blood constituent concentration of a patient so as to generate a measured arterial pulse signal; processing means responsive to said arterial pulse signal for generating a full arterial pulse signal, defined as the measured arterial pulse signal substantially without the effect of atrial diastolic blood flow demand; said processing means for subtracting said arterial pulse signal from said full arterial pulse signal so as to generate an arterio-venous pulse signal; said processing means further calculating an area under the curve of said full arterial pulse (FAP) signal, measured by integration over the cardiac cycle, and an area

under the curve of said arterio-venous (AV) pulse signal, measured by integration over the same cardiac cycle, and a transfer function relationship using said arterio-venous pulse signal area under the curve and said full arterial pulse signal area under the curve to generate a balance parameter representation; computational means being responsive to said balance parameter representation for further generating a set of physiological parameters useful in clinical assessment of a patient's health condition. In an embodiment, said full arterial pulse signal is estimated from said measured arterial pulse signal using at least one of a linear line fit or exponential line fit including said measured arterial pulse signal peak and the peak following its dicrotic notch. In an embodiment, said transfer function is a partial derivative sensitivity relationship between said arterio-venous pulse signal and said full arterial pulse signal to generate a balance parameter representation. In an embodiment, said transfer function is a partial derivative sensitivity relationship calculated using a neural network between a plurality of the said arterio-venous pulse signal and a plurality of said full arterial pulse signal to generate a balance parameter representation. In an embodiment, said system having a display monitoring means responsive to said balance parameter and said set of physiological parameters from said processing means for displaying meaningful information. In an embodiment, said transfer function relationship is identified using a system identification method for generating said balance parameter. In an embodiment, said transfer function relationship is a linear state-space model for said balance parameter. In an embodiment, said set of physiological parameters are used to indicate a clinical assessment of a patient's health condition. In an embodiment, at least one of said balance parameter or said set of physiological parameters are used in at least one of the evaluation of a therapy process for the patient and in the controlling of said therapy process. In an embodiment, said set of physiological parameters are used to calculate oxygen consumption and oxygen delivery for the detection, diagnosis, prediction, or therapy of patients' disease state. In an embodiment, at least one of said balance parameter and said set of physiological parameters is used to control a medication delivery system for achieving target set points. In an embodiment, said set of physiological parameters are used to identify, using system identification models, transfer function relationship for blood arterial supply and demand for a plurality of organs in the body of the patient to assess each organ's health state, or to relatively compare their balance parameters. In an embodiment, said set of physiological parameters is used to provide monitoring of the patient's health condition, and alerting of abnormal events in patient's health condition. In an embodiment, further comprising wired or wireless telecommunication means for transmitting said balance parameter and said set of physiological parameters from said processing means to other processing, data storage, or display monitoring means. In an embodiment, said sensing means is attached to the patient using one of invasive, minimally invasive, and non-invasive techniques. In an embodiment, said blood flow measurement is obtained by one of acoustic sensors, optical sensors, motion sensors, differential pressure sensors, thermodilution or agent dilution sensors, and coriolis sensors. In an embodiment, said blood pressure measurement is obtained using one of invasive, minimally invasive, and non-invasive techniques. In an embodiment, said sensing means for measuring one of blood pressure and blood flow

rate and blood constituent concentration is obtained by measurements from an artery, including one of aortic, carotid, pulmonary, femoral, radial, renal and hepatic arteries. In an embodiment, said sensing means for measuring one of blood pressure and blood flow rate and blood constituent concentration is obtained by measurements from a vein, including one of central venous, superior vena cava, inferior vena cava, jugular, pulmonary, femoral, radial, renal, and hepatic veins generate a venous pulse signal.

**[0221]** In summary, we discussed a method for measuring of cardiac blood flow balance parameter between the right chamber of the heart and the left chamber of the heart, comprising the steps of: measuring with a sensor one of blood pressure and blood flow rate and blood constituent concentration of a patient so as to generate a measured arterial pulse signal; calculating a full arterial pulse signal from the measured arterial pulse signal, defined as the measured arterial pulse signal substantially without the effect of atrial diastolic blood flow demand; calculating an arterio-venous pulse signal by subtracting said arterial pulse signal from said full arterial pulse signal; generating a balance parameter by calculating a transfer function relationship using said full arterial pulse signal area under the curve, measured by integration over the cardiac cycle and said arterio-venous pulse signal area under the curve, measured by integration over the same cardiac cycle; and generating a set of physiological parameters in response to the balance parameter useful in clinical assessment of a patient's health condition. In an embodiment, wherein estimating said full arterial pulse signal from said measured arterial pulse signal using at least one of a linear line fit or exponential line fit including said measured arterial pulse signal's peak and the peak following its dicrotic notch. In an embodiment, wherein said transfer function is a partial derivative sensitivity relationship between said arterio-venous pulse signal and said full arterial pulse signal for generating a balance parameter representation. In an embodiment, said transfer function is a partial derivative sensitivity relationship calculated using a neural network between a plurality of the said arterio-venous pulse signal and a plurality of said full arterial pulse signal for generating a balance parameter representation. In an embodiment, displaying said balance parameter and set of physiological parameters from said processing means for displaying meaningful information. In an embodiment, identifying said transfer function relationship using a system identification method for generating said balance parameter. In an embodiment, wherein said transfer function relationship is a linear state-space model for said balance parameter. In an embodiment, wherein the step of measuring includes the step of attaching the sensor to the patient using one of invasive, minimally invasive, and non-invasive techniques. In an embodiment, wherein the step of measuring the blood flow is obtained by one of acoustic sensors, motion sensors, differential pressure sensors, thermodilution or agent dilution sensors, and coriolis sensors. In an embodiment, wherein the step of measuring the blood pressure is obtained using one of invasive, minimally invasive, and non-invasive techniques. In an embodiment, wherein the step of measuring one of blood pressure, blood flow rate, and blood constituent concentration is obtained by measurements from an artery, including one of aortic, carotid, pulmonary, femoral, radial, renal, and hepatic arteries. In an embodiment, wherein the step of measuring one of blood pressure, and blood flow rate, and blood

constituent concentration is obtained by measurements from a vein, including one of central venous, superior vena cava, inferior vena cava, jugular, pulmonary, femoral, radial, renal, and hepatic veins generate a venous pulse signal. In an embodiment, wherein the derived cardiac balance parameter is used to optimize or control a cardiac pacemaker in one of location of leads placement, pacing power level, pacing pulse shape, pacing period, or pacing duty cycle. In an embodiment, wherein the derived cardiac balance parameter is used to detect and optionally alert against cardiac fibrillation or arrhythmia condition. In an embodiment, further comprising the steps of providing a processor unit and configuring the processor unit to produce a transfer function which computes one of the estimated full arterial pulse signal or the arterio-venous pulse signal based upon the measured arterial pulse signal. In an embodiment, wherein said processor unit employs a system identification technique to define said transfer function. In an embodiment, wherein said system identification technique uses a linear state-space model identification. In an embodiment, wherein said balance signal provides physiological information of the heart. In an embodiment, wherein said balance signal provides physiological information of the brain. In an embodiment, wherein said balance signal provides physiological information of the lungs. In an embodiment, wherein said balance signal provides physiological information of the kidney. In an embodiment, wherein said balance signal provides physiological information of the patient's body organ. In an embodiment, at least two said balance parameters are relatively compared to provides physiological information of the patient.

**[0222]** In summary, we presented a method for generating a sensitivity parameter between a first input parameter and a second input parameter, comprising the steps of: Obtaining said first input parameter and said second input parameter representing one of a process variable and a process variable target or reference; Comparing said first input parameter and said second input parameter to each other to obtain a difference error between them; Presenting said error into the input taps of an input layer of a neural network; Multiplying said input taps values by weights representing a partial derivative between said first input parameter and said second input parameter to produce weighted input values; Summing the weighted input value of said input layer and presenting summed weighted input value into the taps of the hidden layer of said neural network; Mapping said summed weighted input values using an activation function for presenting said hidden layer; Multiplying said hidden taps values by weights representing a partial derivative between said first input parameter and said second input parameter to produce weighted hidden values; Summing said weighted hidden values and of said hidden layer and presenting summed weighted hidden values into the taps of the output layer of said neural network; Mapping said summed weighted hidden values using an output layer activation function for presenting said output layer;

**[0223]** Presenting said output layer activation function output to a mapping function for desired range of neural network output to generate a sensitivity measurement between said two input parameters; and Optionally multiplying said sensitivity measurement with a gain value for sensitivity scaling. In an embodiment, wherein the sensitivity parameter represents a blood flow balance parameter, and

one of the first input parameter and the second input parameter represent one of a blood flow supply and a blood flow demand.

**[0224]** While there has been illustrated and described what is at present considered to be a preferred embodiment of the present invention, it will be understood by those skilled in the art that various changes and modifications may be made, and equivalents may be substituted for elements thereof without departing from the true scope of the invention. In addition, many modifications may be made to adapt a particular situation or material to the teachings of the invention without departing from the central scope thereof. Therefore, it is intended that this invention not be limited to the particular embodiment disclosed as the best mode contemplated for carrying out the invention, but that the invention will include all embodiments falling within the scope of the appended claims.

## APPENDIX

### References

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- [0229]** 5. Multivariable System Identification For Process Control, Y. Zhu, Elsevier Science; 1 edition (October, 2001)
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What is claimed is:

1. A system for measuring of cardiac blood flow balance parameter between the right chamber of the heart and the left chamber of the heart, comprising:

sensing means for measuring one of blood pressure and blood flow rate and blood constituent concentration of a patient so as to generate a measured arterial pulse signal;

processing means responsive to said arterial pulse signal for generating a full arterial pulse signal, defined as the measured arterial pulse signal substantially without the effect of atrial diastolic blood flow demand;

said processing means for subtracting said arterial pulse signal from said full arterial pulse signal so as to generate an arterio-venous pulse signal;

said processing means further calculating an area under the curve of said full arterial pulse (FAP) signal, measured by integration over the cardiac cycle, and an area under the curve of said arterio-venous (AV) pulse signal, measured by integration over the same cardiac cycle, and a transfer function relationship using said arterio-venous pulse signal area under the curve and said full arterial pulse signal area under the curve to generate a balance parameter representation;

computational means being responsive to said balance parameter representation for further generating a set of physiological parameters useful in clinical assessment of a patient's health condition.

2. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said full arterial pulse signal is estimated from said measured arterial pulse signal using at least one of a linear line fit or exponential line fit including said measured arterial pulse signal peak and the peak following its dicrotic notch.

3. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said transfer function is a partial derivative sensitivity relationship between said arterio-venous pulse signal and said full arterial pulse signal to generate a balance parameter representation.

4. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said transfer function is a partial derivative sensitivity relationship calculated using a neural network between a plurality of the said arterio-venous pulse signal and a plurality of said full arterial pulse signal to generate a balance parameter representation.

5. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, having a display monitoring means responsive to said balance parameter and said set of physiological parameters from said processing means for displaying meaningful information.

6. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said transfer function relationship is identified using a system identification method for generating said balance parameter.

7. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said transfer function relationship is a linear state-space model for said balance parameter.

8. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said set of physiological parameters are used to indicate a clinical assessment of a patient's health condition.

9. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein at least one of said balance parameter or said set of physiological parameters are used in at least one of the evaluation of a therapy process for the patient and in the controlling of said therapy process.

10. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said set of physiological parameters are used to calculate oxygen consumption and oxygen delivery for the detection, diagnosis, prediction, or therapy of patients' disease state.

11. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein at least one of said balance parameter and said set of physiological parameters is used to control a medication delivery system for achieving target set points.

12. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said set of physiological parameters are used to identify, using system identification models, transfer function relationship for blood arterial supply and demand for a plurality of organs in the body of the patient to assess each organ's health state, or to relatively compare their balance parameters.

13. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said set of physiological parameters is used to provide monitoring of the patient's health condition, and alerting of abnormal events in patient's health condition.

14. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein further comprising wired or wireless telecommunication means for transmitting said balance parameter and said set of physiological parameters from said processing means to other processing, data storage, or display monitoring means.

15. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said sensing means is attached to the patient using one of invasive, minimally invasive, and non-invasive techniques.

16. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said blood flow measurement is obtained by one of acoustic sensors, optical sensors, motion sensors, differential pressure sensors, thermodilution or agent dilution sensors, and coriolis sensors.

17. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said blood pressure measurement is obtained using one of invasive, minimally invasive, and non-invasive techniques.

18. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said sensing means for measuring one of blood pressure and blood flow rate and blood constituent concentration is obtained by measurements from an artery, including one of aortic, carotid, pulmonary, femoral, radial, renal and hepatic arteries.

19. A system for measuring of cardiac blood flow balance parameter as claimed in claim 1, wherein said sensing means for measuring one of blood pressure and blood flow rate and blood constituent concentration is obtained by measurements from a vein, including one of central venous, superior vena cava, inferior vena cava, jugular, pulmonary, femoral, radial, renal, and hepatic veins generate a venous pulse signal.

20. A method for measuring of cardiac blood flow balance parameter between the right chamber of the heart and the left chamber of the heart, comprising the steps of:

measuring with a sensor one of blood pressure and blood flow rate and blood constituent concentration of a patient so as to generate a measured arterial pulse signal;

calculating a full arterial pulse signal from the measured arterial pulse signal, defined as the measured arterial pulse signal substantially without the effect of atrial diastolic blood flow demand;

calculating an arterio-venous pulse signal by subtracting said arterial pulse signal from said full arterial pulse signal;

generating a balance parameter by calculating a transfer function relationship using said full arterial pulse signal area under the curve, measured by integration over the cardiac cycle and said arterio-venous pulse signal area under the curve, measured by integration over the same cardiac cycle; and

generating a set of physiological parameters in response to the balance parameter useful in clinical assessment of a patient's health condition.

21. A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein estimating said full arterial pulse signal from said measured arterial pulse signal using at least one of a linear line fit or exponential line fit including said measured arterial pulse signal's peak and the peak following its dicrotic notch.

22. A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein said transfer

function is a partial derivative sensitivity relationship between said arterio-venous pulse signal and said full arterial pulse signal for generating a balance parameter representation.

**23.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein said transfer function is a partial derivative sensitivity relationship calculated using a neural network between a plurality of the said arterio-venous pulse signal and a plurality of said full arterial pulse signal for generating a balance parameter representation.

**24.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, displaying said balance parameter and set of physiological parameters from said processing means for displaying meaningful information.

**25.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, identifying said transfer function relationship using a system identification method for generating said balance parameter.

**26.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein said transfer function relationship is a linear state-space model for said balance parameter.

**27.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein the step of measuring includes the step of attaching the sensor to the patient using one of invasive, minimally invasive, and non-invasive techniques.

**28.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein the step of measuring the blood flow is obtained by one of acoustic sensors, motion sensors, differential pressure sensors, thermolulution or agent dilution sensors, and coriolis sensors.

**29.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein the step of measuring the blood pressure is obtained using one of invasive, minimally invasive, and non-invasive techniques.

**30.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein the step of measuring one of blood pressure, blood flow rate, and blood constituent concentration is obtained by measurements from an artery, including one of aortic, carotid, pulmonary, femoral, radial, renal, and hepatic arteries.

**31.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein the step of measuring one of blood pressure, and blood flow rate, and blood constituent concentration is obtained by measurements from a vein, including one of central venous, superior vena cava, inferior vena cava, jugular, pulmonary, femoral, radial, renal, and hepatic veins generate a venous pulse signal.

**32.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein the derived cardiac balance parameter is used to optimize or control a cardiac pacemaker in one of location of leads placement, pacing power level, pacing pulse shape, pacing period, or pacing duty cycle.

**33.** A method for measuring of cardiac blood flow balance parameter as claimed in claim 20, wherein the derived cardiac balance parameter is used to detect and optionally alert against cardiac fibrillation or arrhythmia condition.

**34.** A method for clinically assessing a patient's health condition as claimed in claim 20, further comprising the steps of providing a processor unit and configuring the

processor unit to produce a transfer function which computes one of the estimated full arterial pulse signal or the arterio-venous pulse signal based upon the measured arterial pulse signal.

**35.** A method for clinically assessing of a patient's health condition as claimed in claim 20, wherein said processor unit employs a system identification technique to define said transfer function.

**36.** A method for clinically assessing of a patient's health condition as claimed in claim 20, wherein said system identification technique uses a linear state-space model identification.

**37.** A method for clinically assessing of a patient's health condition as claimed in claim 20, wherein said balance signal provides physiological information of the heart.

**38.** A method for clinically assessing of a patient's health condition as claimed in claim 20, wherein said balance signal provides physiological information of the brain.

**39.** A method for clinically assessing of a patient's health condition as claimed in claim 20, wherein said balance signal provides physiological information of the lungs.

**40.** A method for clinically assessing of a patient's health condition as claimed in claim 20, wherein said balance signal provides physiological information of the kidney.

**41.** A method for clinically assessing a patient's health condition as claimed in claim 20, wherein said balance signal provides physiological information of the patient's body organ.

**42.** A method for clinically assessing a patient's health condition as claimed in claim 20, at least two said balance parameters are relatively compared to provides physiological information of the patient.

**43.** A method for generating a sensitivity parameter between a first input parameter and a second input parameter, comprising the steps of:

Obtaining said first input parameter and said second input parameter representing one of a process variable and a process variable target or reference;

Comparing said first input parameter and said second input parameter to each other to obtain a difference error between them;

Presenting said error into the input taps of an input layer of a neural network;

Multiplying said input taps values by weights representing a partial derivative between said first input parameter and said second input parameter to produce weighted input values;

Summing the weighted input value of said input layer and presenting summed weighted input value into the taps of the hidden layer of said neural network;

Mapping said summed weighted input values using an activation function for presenting said hidden layer;

Multiplying said hidden taps values by weights representing a partial derivative between said first input parameter and said second input parameter to produce weighted hidden values;

Summing said weighted hidden values and of said hidden layer and presenting summed weighted hidden values into the taps of the output layer of said neural network;

Mapping said summed weighted hidden values using an output layer activation function for presenting said output layer;

Presenting said output layer activation function output to a mapping function for desired range of neural network

output to generate a sensitivity measurement between said two input parameters; and

Optionally multiplying said sensitivity measurement with a gain value for sensitivity scaling.

**44.** A method for generating a sensitivity parameter as claimed in claim **43**, wherein the sensitivity parameter represents a blood flow balance parameter, and one of the first input parameter and the second input parameter represent one of a blood flow supply and a blood flow demand.

\* \* \* \* \*

专利名称(译)	监测左右心腔之间的心脏血流平衡关系和心脏调节		
公开(公告)号	<a href="#">US20200008686A1</a>	公开(公告)日	2020-01-09
申请号	US16/028403	申请日	2018-07-05
发明人	KHAIR, MOHAMMAD MOHAMMAD		
IPC分类号	A61B5/02 A61B5/026 A61B5/028 A61B8/06 A61B5/0275 A61B5/0215 A61B5/022 A61B5/00 A61B5/08 A61B5/20		
CPC分类号	A61B5/02241 A61B5/201 A61B5/0261 A61B5/7278 A61B5/028 A61B8/06 A61B5/02755 A61B5/6876 A61B5/0215 A61B2560/0475 A61B5/0002 A61B5/742 A61B5/746 A61B5/7264 A61B5/4839 A61B5/08 A61N1/36571 A61B5/4866 A61B5/02028 A61B5/4064 A61B5/7203 A61B5/029 A61B5/7242 A61N1/36564 A61N1/36585		
外部链接	<a href="#">Espacenet</a> <a href="#">USPTO</a>		

摘要(译)

一种用于测量心脏的右室和心脏的左室之间的心脏血液流动平衡参数的系统，包括用于测量患者的血压和血流速度以及血液成分浓度之一以产生血流的传感器装置。动脉脉搏信号。处理单元响应于动脉脉冲信号以生成完整的动脉脉冲信号，动静脉脉冲信号以及包括平衡比和平衡趋势的平衡参数以及脉动可变性参数。计算设备响应于平衡参数以进一步生成一组生理参数。显示站设备响应于来自计算设备的一组生理参数以显示有意义的信息。

