



(19) **United States**

(12) **Patent Application Publication** (10) **Pub. No.: US 2004/0044276 A1**

Arnold

(43) **Pub. Date:**

Mar. 4, 2004

(54) **METHOD AND APPRATUS FOR MEASURING PULSUS PARADOXUS**

(57)

ABSTRACT

(76) Inventor: **Donald Arnold**, Nashville, TN (US)

Correspondence Address:
KNOBBE MARTENS OLSON & BEAR LLP
2040 MAIN STREET
FOURTEENTH FLOOR
IRVINE, CA 92614 (US)

(21) Appl. No.: **10/228,641**

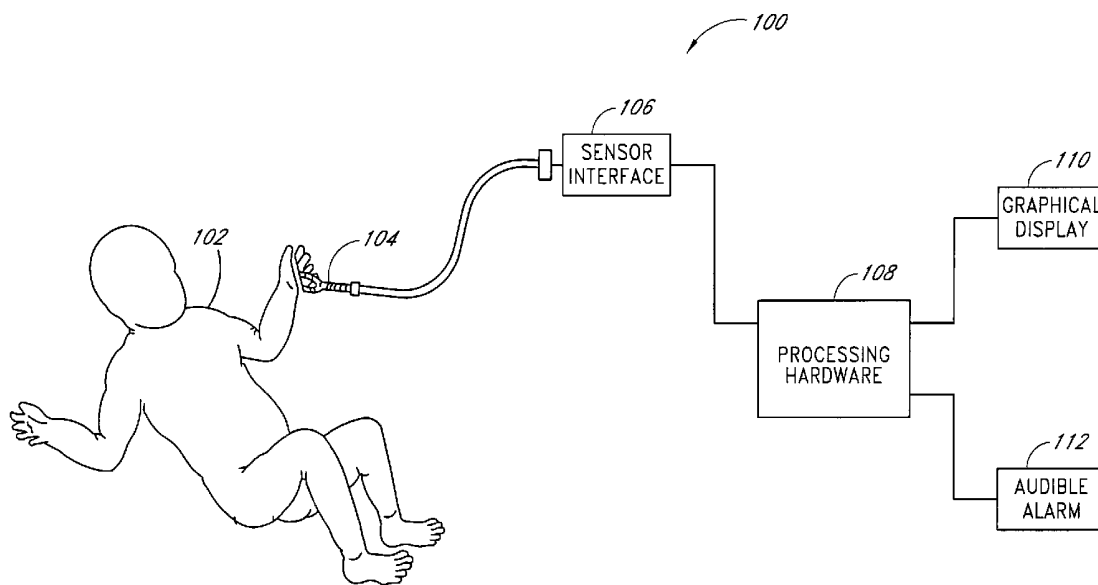
(22) Filed: **Aug. 27, 2002**

Publication Classification

(51) **Int. Cl.⁷** **A61B 5/00**

(52) **U.S. Cl.** **600/323**

A method and apparatus are disclosed of utilizing a source of arterial and/or arteriolar pulse waveform data from a patient for the purpose of measuring pulsus paradoxus. The arterial pulse waveform data source described is a pulse oximeter plethysmograph but can be any similar waveform data source, including intra-arterial transducer, blood pressure transducer, or plethysmograph. Through incorporation of the measurements of values, such as the area under the pulse waveform curve, that are time-domain functions of a change in height of the pulse waveform over at least a partial duration of the waveform, embodiments of the present invention represent a significant improvement upon previously described methods of measuring pulsus paradoxus and, further, produce improved accuracy in measurement of the multiple contributing variables generating pulsus paradoxus, in particular the contribution of diastolic events, to the extent that the physical signs so measured can be regarded as "Revised Pulsus Paradoxus."



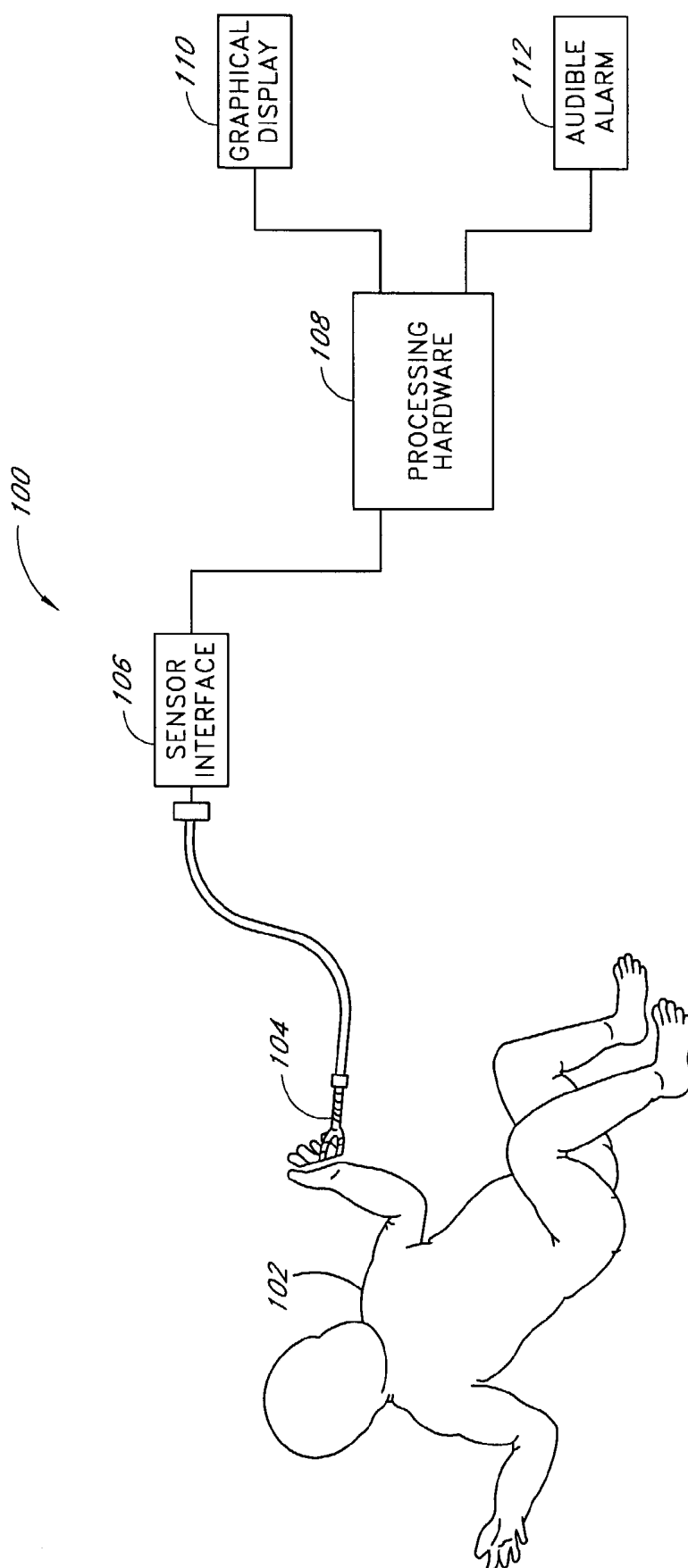


FIG. 1

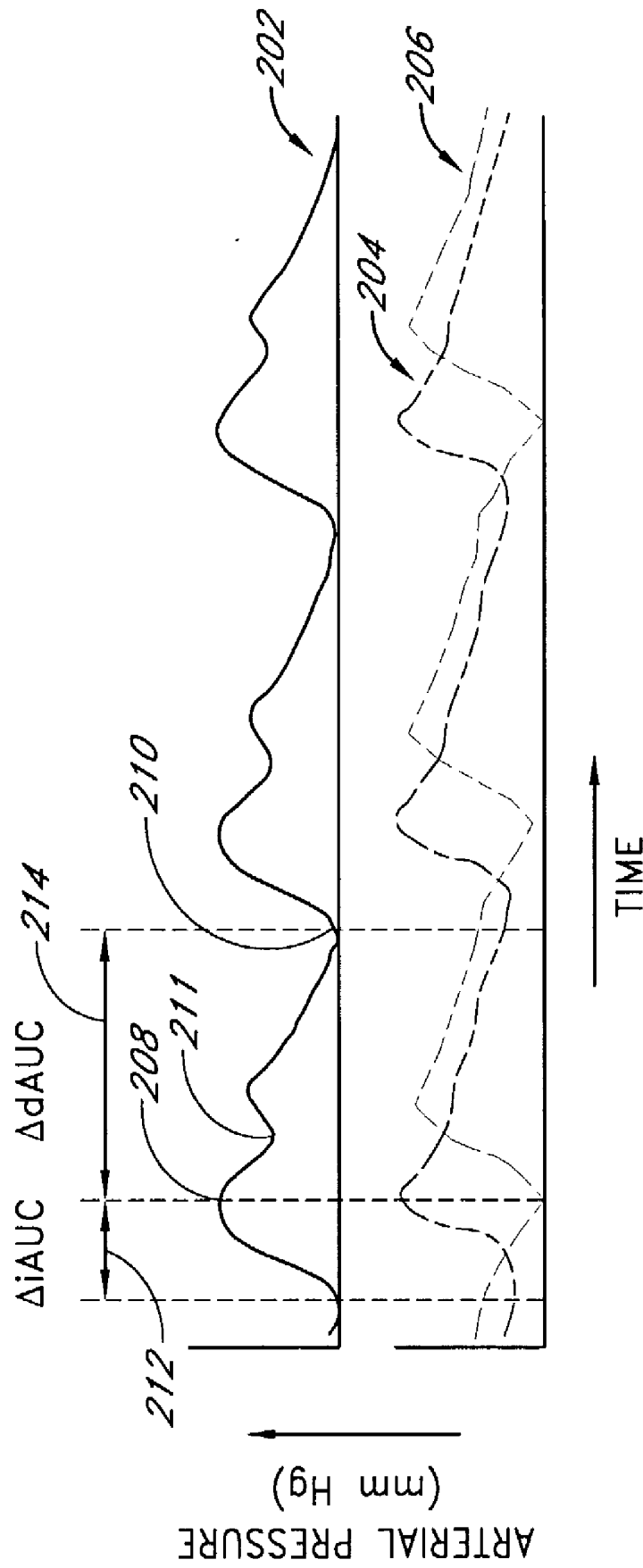


FIG. 2

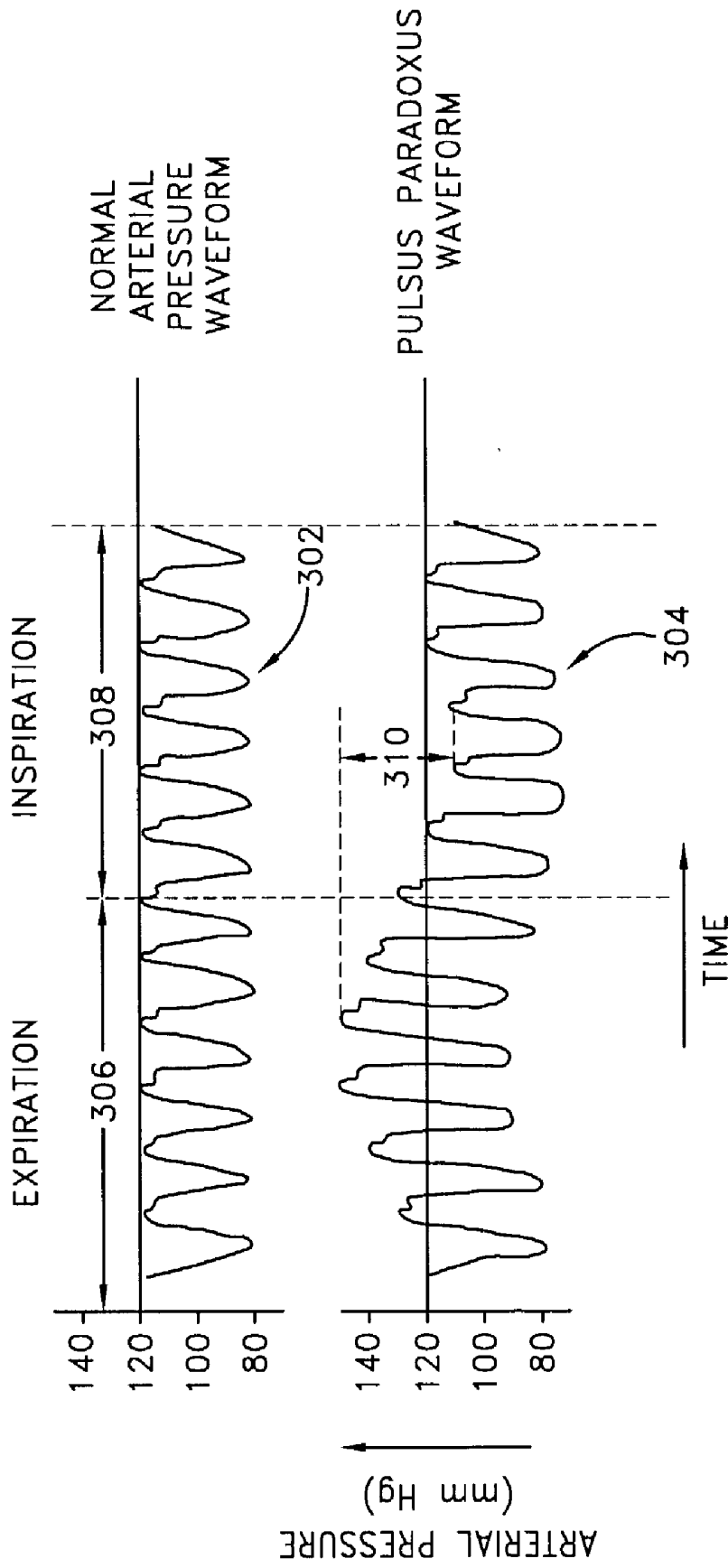


FIG. 3

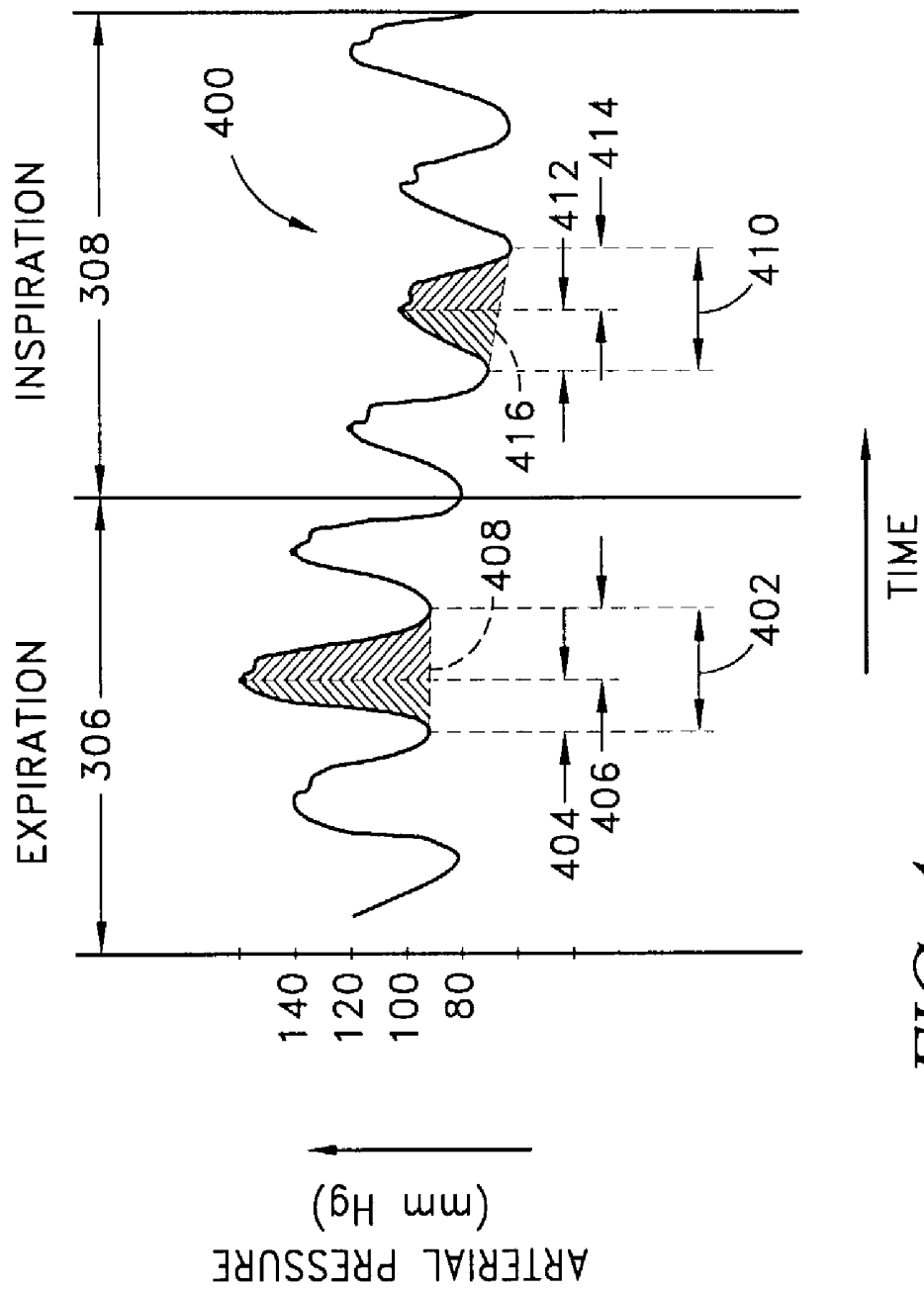


FIG. 4

METHOD AND APPRATUS FOR MEASURING PULSUS PARADOXUS

FIELD OF THE INVENTION

[0001] Embodiments of the invention arise in the field of medical diagnosis, prognosis, and monitoring of patients afflicted with a variety of diseases and clinical syndromes such as asthma, croup, tension pneumothorax, pericardial tamponade, pulmonary embolus, hypovolemic shock, and sleep apnea. In particular, the invention relates to an improved non-invasive system and method for monitoring the presence and degree of pulsus paradoxus within a patient. The invention is particularly useful for rapidly assessing the status of a patient in acute respiratory distress in order to more accurately and objectively determine the severity of the patient's condition.

BACKGROUND OF THE INVENTION

[0002] Pulsus paradoxus is a medical term referring to a quantifiable, exaggerated decrease in arterial blood pressure during inspiration. In normal patients, this decrease is in the range of about 2-5 mm Hg; whereas, in a patient suffering from certain medical conditions, pulsus paradoxus can reach 10 mm Hg or higher. Pulsus paradoxus has been noted in a variety of medical conditions including, but not limited to, upper airway obstruction, bronchial asthma, tricuspid atresia, mitral stenosis, conditions of decreased left ventricular compliance, croup, tension pneumothorax, pericardial tamponade, pulmonary embolism, hypovolemic shock, and sleep apnea.

[0003] Asthma and Related Diseases

[0004] Asthma accounts for almost two million Emergency Department admissions annually in the U.S., and it is estimated that 29 million people will be diagnosed with asthma in the next two decades. Hospitalizations for childhood asthma have increased 3% to 5% annually, and mortality from asthma has increased 10% annually since 1977. Other forms of lung disease, including but not limited to chronic obstructive pulmonary disease ("COPD") and emphysema, place a heavy burden on patients and on the system of medical care. Early recognition and accurate assessment of the severity of airway obstruction and of the response to therapy are fundamental to the improvement of health for persons with these disorders.

[0005] Common measures used currently to assess the severity of asthma are clinical assessment, arterial blood gas analysis, spirometry, and pulse oximetry; however, all are subject to certain shortcomings. Clinical assessment scores, for example, exhibit marked interobserver variability and have been incompletely validated. Arterial blood gas analysis is an invasive and painful technique and is often complicated by therapeutic administration of O₂ and β -adrenergic drugs and is therefore unreliable as an indicator of asthma severity. Tests of forced expiratory flow, as in spirometry, are effort dependent, typically cannot be used with children, and can actually exacerbate the underlying disease process. In part because physicians' ability to accurately assess pulmonary signs can be unreliable, numerous clinical scoring systems and management guidelines have been established for diseases such as asthma.

[0006] Despite the publication of the National Heart Lung and Blood Institute ("NHLBI") Guidelines for Emergency

Department Asthma Management in 1991, the asthma mortality rate in children ages 5-17 nearly tripled between 1980 and 1996. Many experts are at a loss to explain the rising mortality of asthmatic patients in view of the improving quality of acute pharmacological management of asthma and the enhanced sophistication of emergency physicians and pre-hospital care systems. One explanation lies in the observation that there has been little change in how the acute asthmatic patient is evaluated. A recent development in assessing acute asthma has been the use of pulse oximetry (SPO₂) which measures the degree of oxygen saturation of hemoglobin non-invasively and empirically. Despite the ubiquitous availability of pulse oximetry, (-adrenergic drugs, used widely, can result in ventilation-perfusion mismatch, leading to a fall in SPO₂ even though the patient is improving. Finally, changes in SPO₂ reflect atelectasis and intrapulmonary shunting and do not directly provide information regarding airflow obstruction and ventilation. An easily recognized, sensitive and objective parameter, by which practitioners of all levels of expertise could quickly recognize an asthma exacerbation, would help in more accurate diagnoses.

[0007] Easily measured, objective and accurate indices of severity for acute exacerbation of bronchiolitis, croup, emphysema and COPD are also not available. Additionally, medical emergencies such as cardiac tamponade, hypovolemia, and pulmonary embolism are difficult to diagnose and/or to quantify in severity. Finally, the response of these disorders to treatment is likewise difficult to objectively measure.

[0008] Traditional Methods of Pulsus Paradoxus Measurement

[0009] Although measurement of pulsus paradoxus is recommended by numerous authoritative medical practice guidelines (for example, the previously cited NHLBI Guidelines for Emergency Department Asthma Management), pulsus paradoxus is rarely recorded in clinical practice. Resistance by physicians to the application of pulsus paradoxus for the objective assessment of disease severity, and asthma in particular, is largely due to the difficulty in measuring pulsus paradoxus in a rapidly breathing patient by currently employed methods.

[0010] One conventional method of measuring pulsus paradoxus is through the use of intra-arterial catheters. Although intra-arterial catheters can often give reliable measurements of pulsus paradoxus, placement of these catheters is painful and associated with significant risk. Furthermore, placement of intra-arterial catheters should only be done by highly trained medical personnel using sophisticated monitoring equipment, preferably in a hospital setting. Consequently, this method is not favored for general use.

[0011] Another method for measuring pulsus paradoxus is with the use of a sphygmomanometer, commonly referred to as a blood pressure cuff. This technique involves inflating the sphygmomanometer to above systolic pressure and slowly deflating it. As the elevated systolic pressure occurring during expiration is approached, heart sounds will be heard intermittently (during expiration only), and as the lower systolic pressure occurring during inspiration is approached, heart sounds will be heard continuously. The difference between these points at which heart sounds are

heard, first intermittently and then continuously, is a patient's pulsus paradoxus. However, this traditional method of measurement is difficult in the clinical setting in which noise, rapid respiratory rates and patient movement are the norm, and provides measurement of pulsus paradoxus at only a single point in time. Moreover, this process is ergonomically very difficult to perform and multiple operator efforts are typically required. As a result, the method is often inaccurate and unreliable. Manually derived pulsus paradoxus also correlates poorly with pulsus paradoxus calculated from intra-arterial pressure and, despite the NHLBI's recommendations, actual measurement of pulsus paradoxus is rare.

[0012] A third method of measuring pulsus paradoxus has been developed through photoplethysmographic techniques. In the field of photoplethysmography, pulses of light having different wavelengths are transmitted through or reflected by a patient's tissue to non-invasively determine various blood analyte values. More particularly, a photoplethysmographic device known as a pulse oximeter is employed to determine pulse rates and blood oxygen levels. Pulse oximeters typically include a probe that is attached to a patient's appendage (e.g., finger, ear lobe or nasal septum). The probe directs light signal pulses generated by a plurality of emitters through the appendage, wherein portions of the light signals are absorbed by the tissue. The intensity of light transmitted by the tissue is monitored by one or more detectors which output signals indicative of the light absorbency characteristics of the tissue. Because the blood analytes of interest each differentially absorb more light at one wavelength than at other wavelengths, the ratio of detector output signals can be used to compute the blood analyte concentrations.

[0013] By way of primary example, it is known that oxyhemoglobin (O₂Hb) absorbs light more readily in the infrared region than in the red region, whereas reduced hemoglobin (RHb), or deoxyhemoglobin, more readily absorbs light in the red region than in the infrared region. As such, oxygenated blood with a high concentration of oxyhemoglobin and a low concentration of reduced hemoglobin will tend to have a high ratio of optical transmissivity in the red region to optical transmissivity in the infrared region. The relative transmissivity of blood at red and infrared center wavelengths can be employed as a measure of blood oxygen saturation (SpO₂).

[0014] It is also recognized that concentrations of other related blood constituents (e.g., carboxyhemoglobin (COHb) and methemoglobin (MetHb)) can be measured with a similar approach since such analytes also have unique light absorbency characteristics at different corresponding wavelengths. The determination of such additional constituents can serve to enhance the measurement of blood oxygen saturation.

[0015] As will be appreciated by one skilled in the art, the detector output signal in pulse oximeters contains non-pulsatile and pulsatile components. The non-pulsatile component is influenced by the absorbency of tissue, venous blood, capillary blood, non-pulsatile arterial blood, the intensity of the light signals, ambient environmental light, and the sensitivity of the detector. The pulsatile component reflects the expansion of the arteriolar bed with arterial blood, and the varying amplitude of this pulsatile component depends upon the blood volume change per pulse as a result

of arteriolar inflow. As such, the pulsatile component isolates the optical absorption attributable to the arterial blood component of the vascular bed and provides a basis for monitoring changes in the concentration of the noted blood analytes, oxyhemoglobin and deoxyhemoglobin. This feature of all plethysmographic oximeters, isolation of the pulsatile component of the arteriolar vascular bed and the waveform signal so generated, can be used to determine pulsus paradoxus for a particular patient.

[0016] Despite the problems inherent in these methods of detecting pulsus paradoxus, the advantages of measuring and monitoring pulsus paradoxus are significant. These measurements provide valuable insight into how troubled the act of breathing is for a given patient, and can help physicians detect, assess and treat numerous respiratory ailments. The NHLBI has recognized the advantages of measuring pulsus paradoxus and has recommended that pulsus paradoxus be measured on all asthmatic patients, despite inherent inaccuracies of the sphygmomanometric technique. Moreover, the NHLBI has advised that any patient with a pulsus paradoxus of 12 mm Hg or greater be hospitalized.

SUMMARY OF THE INVENTION

[0017] What is needed are new methods and systems for reliably measuring pulsus paradoxus and aiding in the examination, diagnosis and treatment of patients suffering from pulsus paradoxus-associated diseases. In particular, there is a need for an accurate, non-effort dependent, real-time, non-invasive apparatus and method for accurately measuring pulsus paradoxus. Embodiments of the present invention comprise a system and method to provide a sensitive, objective, accurate, real-time, painless and non-invasive parameter by which to assess these disorders.

[0018] In accordance with one aspect of the present invention, a method for measuring pulsus paradoxus in a mammal is provided. The method comprises obtaining data indicative of pulsatile cardiovascular behavior from the mammal, determining area under a curve for each of two or more measured pulse waveforms, and comparing the calculated areas to determine the presence or absence and magnitude of pulsus paradoxus.

[0019] In accordance with another aspect of the present invention, a system for measuring pulsus paradoxus in a mammal is provided. The system provided comprises a means for obtaining pulse waveform data indicative of pulsatile cardiovascular behavior from the mammal, a means for determining area under a curve from at least two pulse waveforms, and a means for comparing the calculated areas to determine the presence or absence and magnitude of pulsus paradoxus.

[0020] In accordance with another aspect of the invention, a system for measuring pulsus paradoxus in a mammal is provided. This system comprises a pulse oximeter that can obtain pulse waveform data indicative of pulsatile cardiovascular behavior from the mammal, a measurer that determines area under the curve from at least two pulse waveforms, and an analyzer that compares the calculated areas to determine the presence, absence and magnitude of pulsus paradoxus.

[0021] In accordance with another aspect of the invention, a method for measuring and/or detecting pulsus paradoxus

in a mammal is provided. The method comprises obtaining data indicative of pulsatile cardiovascular behavior from the mammal, the data comprising at least a first and a second pulse waveform; determining a first value that is a time-domain function of a change in height of the first pulse waveform over at least a partial duration of the first pulse waveform; determining a second value that is a time-domain function of a change in height of the second pulse waveform over at least a partial duration of the second pulse waveform; comparing the first value and the second value to measure pulsus paradoxus.

[0022] Time-domain functions of a change in height of a pulse waveform over at least a partial duration of the pulse waveform include area under the curve (AUC), slope, average slope, maximal slope, and other functions. In some embodiments, there is generally no need to obtain frequency-domain information, such as that provided by Fourier transform or power spectral analysis, in order to make a determination of the presence and/or magnitude of pulsus paradoxus.

[0023] In accordance with other aspects of the invention, the method comprises obtaining data indicative of pulsatile cardiovascular behavior from a mammal, the data comprising a first plethysmographic or pressure amplitude obtained at a first time interval after an onset of a first pulse, and a second plethysmographic or pressure amplitude obtained at a second time interval after the onset of first pulse; determining a first value that is a time-domain function of a difference between the first and second plethysmographic or pressure amplitudes; obtaining a third plethysmographic or pressure amplitude obtained at the first time interval after an onset of a second pulse, and a fourth plethysmographic or pressure amplitude obtained at the second time interval after the onset of the second pulse; determining a second value that is a time-domain function of a difference between the third and fourth plethysmographic or pressure amplitudes; and comparing the first value and the second value to measure pulsus paradoxus.

[0024] As is well known in the field of plethysmography, a "plethysmographic amplitude" represents the magnitude of some measure of plethysmography, such as light transmission or absorption, and can reflect a magnitude of pressure in or volume of one or more vessels or chambers. Other factors, such as oxyhemoglobin (O₂Hb), reduced hemoglobin (RHb), or deoxyhemoglobin, may also influence plethysmographic amplitude, as discussed previously. As used herein, "pressure amplitude" means blood pressure or a magnitude of absolute or relative blood pressure based on a system that uses a proxy for blood pressure, such as plethysmography.

[0025] Note that in obtaining a plethysmographic or pressure amplitude (or magnitude) at a time interval after an onset of a first pulse, this time interval may be of a fixed or predetermined duration, such as 0.2 sec or 0.3 msec, or it may be a relative time interval, such as the first half of systole or the 25-75% interval of the cardiac cycle or pulse duration. Also, this time interval can be set to zero (0), such that the plethysmographic or pressure amplitude at a time interval of zero after an onset of a pulse may itself equal zero, or any other baseline value of plethysmographic or baseline pressure amplitude that exists at the onset of the pulse.

BRIEF DESCRIPTION OF THE DRAWINGS

[0026] These and other aspects of this invention will be readily apparent from the detailed description below and the appended drawings, which are meant to illustrate and not to limit the invention, and in which:

[0027] **FIG. 1** depicts a pulse oximeter system connected to a mammalian patient.

[0028] **FIG. 2** depicts an arterial pressure waveform, in accordance with one aspect of the present invention.

[0029] **FIG. 3** depicts both a normal arterial pressure waveform and a pulsus paradoxus waveform, in accordance with another aspect of the present invention.

[0030] **FIG. 4** illustrates measurements made upon a waveform through a preferred embodiment of the present invention.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

[0031] Embodiments of the present invention will be more completely understood through the following detailed description, which should be read in conjunction with the attached drawings. In this description, like numbers refer to similar elements within various embodiments of the present invention.

[0032] Pulsus paradoxus represents a quantifiable, exaggerated decrease in arterial blood pressure during inspiration. Although frequently described in the medical literature as merely a decrease in systolic blood pressure, this physiologic phenomenon actually represents a decrease in left ventricular stroke volume ("LVS_V") during the inspiratory phase of the respiratory cycle and/or an augmentation of LVS_V during the expiratory phase of the respiratory cycle. This subtle distinction is important in development of the most accurate means by which to measure pulsus paradoxus.

[0033] The mechanisms of pulsus paradoxus are incompletely understood and can differ among various diseases. In severe acute asthma, for example, airway obstruction results from bronchospasm, mucous membrane edema and mucus plugging. Overcoming this obstruction to maintain airflow and ventilation necessitates the generation of increased negative intrapleural pressure on inspiration and increased positive intrapleural pressure on expiration. These exaggerated swings of intrapleural pressure result in increased left ventricular afterload during inspiration and decreased left ventricular afterload during expiration and concomitant decreased LVS_V and increased LVS_V during inspiration and expiration, respectively. Air-trapping and hyperinflation result in alveolar distension, pulmonary arteriolar vasoconstriction, and increased impedance to right ventricular ejection during inspiration. This results in decreased left ventricular filling and further impaired left ventricular stroke output. Diminished left ventricular filling also results from shift of the interventricular septum leftward as a result of augmented filling of the right ventricle. Conversely, these events are influenced inversely during expiration and result in increased LVS_V during expiration. Each of these alterations contribute to paradoxical pulse in asthma and other respiratory and cardiovascular disease states.

[0034] More generally, pulsus paradoxus is caused by variably altering intrapleural pressure, left ventricular com-

pliance, pulmonary hyperinflation, or systemic or pulmonary venous return. Any of these events, individually or collectively, lead to an accentuation of the decrease in LVSV during the inspiratory phase respiratory cycle and/or an augmentation of LVSV during the expiratory phase of the respiratory cycle, and thus the phenomenon of pulsus paradoxus. These changes in LVSV and resulting pulsus paradoxus have been demonstrated to result from the following physiologic events:

- [0035] 1. Increasingly negative intrapleural pressure during inspiration resulting in increased transmural (absolute intracardiac minus intrapleural) pressure.
- [0036] 2. Impairment to left ventricular ejection during inspiration resulting from increased afterload (as the left ventricular must create greater contractile force to overcome the increasingly negative intrapleural pressure).
- [0037] 3. Mechanical impairment to left ventricular filling (decreased compliance) due to shift of the interventricular septum leftward (resulting from increased venous return to the right heart as a result of increased negative intrapleural pressure during inspiration), resulting in decreased left ventricular stroke volume during inspiration.
- [0038] 4. Increased impedance to right ventricular ejection during inspiration due to increased alveolar volume and increased resistance to flow through alveolar capillaries, resulting in decreased pulmonary venous return to the left atrium and ventricle and decreased LVSV.

[0039] Typical methods of detecting pulsus paradoxus have numerous, significant disadvantages. For example, using intra-arterial catheters to detect pulsus paradoxus is painful, associated with significant risk and should only be done by highly trained medical personnel using sophisticated monitoring equipment, preferably in a hospital setting. Determining pulsus paradoxus through the use of a sphygmomanometer is often difficult and frequently results in unreliable readings. Through photoplethysmography, pulsus paradoxus can be determined in non-invasive fashion, but results can not be reliable and provide limited information to a physician or caregiver.

[0040] Previous attempts to quantify pulsus paradoxus from oximeter plethysmographic and other waveforms representing pulsatile arterial flow have done so by measuring changes in waveform height, a one-dimensional parameter, similar to the traditional methods described above, or by measuring variation of the plethysmograph baseline. Because pulsus paradoxus represents changes in LVSV and flow, these methods lack the sensitivity and accuracy of a method based upon two or three-dimensional parameters. Moreover, the existing techniques only measure changes in the height of the systolic waveform and do not adequately measure diastolic changes occurring during the cardiac cycle that contribute to pulsus paradoxus. The multiple physiologic events contributing to pulsus paradoxus and the corollary, that the diastolic and systolic contributions to pulsus paradoxus vary depending on the disease process involved, make evident the need to consider these diastolic contributions when measuring pulsus paradoxus. In the preferred embodiment, a method and system are provided

which utilizes changes in area under a measured waveform curve ("AUC"), a two-dimensional parameter, to determine pulsus paradoxus. Preferably, the provided method and system also accurately incorporate diastolic changes in order to make a more sensitive and accurate assessment of the physiologic changes contributing to pulsus paradoxus.

[0041] FIG. 1 depicts a pulse oximeter system, generally designated as 100, connected to a mammalian patient 102. Preferably, pulse oximeter system 100 includes an optical transducer or probe 104, a sensor interface 106, processing hardware 108, a graphical display 110 and an audible alarm 112. In the preferred embodiment, the probe 104 is attached to the patient 102 on a digit, earlobe or other site overlying an arteriolar vascular bed. As illustrated in FIG. 1, the probe 104 can be attached to the fingers of the patient 102, but other locations are also possible. The probe 104 typically comprises an exterior housing that applies active elements of the probe 104 to tissue of the patient 102 containing an arterial or arteriolar vascular bed that is to be monitored. In the preferred embodiment, the probe 104 also contains one or more light-emitting diodes ("LEDs") and one or more light detectors to monitor the level of light transmitted through or reflected from the patient's 102 vascular tissue. The light detectors typically measure the changing optical absorption of light transmitted from LEDs at approximately 660 nm and 940 nm which results from volume expansion of the arteriolar and capillary beds during pulsatile cardiac blood flow. The probe transmits signals describing the detected light to at least one sensor interface 106. In addition, although a single probe 104 and a single sensor interface 106 are illustrated, one skilled in the art will recognize that alternative embodiments employing multiple probes 104 and/or multiple sensor interfaces 106 are also possible. One exemplary design of a typical pulse oximeter system is disclosed in U.S. Pat. No. 6,385,471 by Mortz, the entire disclosure of which is hereby incorporated by reference.

[0042] The signal received by the sensor interface 106 from the probe 104 is traditionally an analog signal and is preferably processed by additional analog circuitry and converted by an analog-to-digital converter circuit into a set of digital measurements before being transmitted to the processing hardware 108.

[0043] The processing hardware 108 can comprise a specific device, such as a digital signal processor specifically programmed or hardwired to implement the features of embodiments of the present invention. The processing hardware 108 can also comprise a general purpose computer, incorporating elements such as an Intel Pentium™ processor, physical memory, input and output devices, etc., programmed with software implementing features embodied by the present invention. The primary function of the processing hardware 108 is to measure the maximal change in area under a continuous optical plethysmographic waveform curve corresponding to arterial pulsation, to display data related to this measurement on the graphic display 110, and to trigger an audible alarm 112 to alert physicians or other caregivers to the presence and severity of pulsus paradoxus. The operation of the processing hardware 108 is described in additional detail below and, for the purpose of this disclosure, it is assumed that the other elements disclosed in FIG. 1 are the conventional components found in existing pulse oximeter systems.

false positive reports (e.g., those due to arrhythmia) and increasing the system's overall accuracy.

[0051] Advantageously, embodiments of the system and methods provided herein reliably reflect corresponding increases or decreases in airway obstruction and pulsus paradoxus more accurately and less invasively than previous methods. Whether by pulse oximeter plethysmography, finger plethysmograph or other waveform, the preferred embodiment involves placing a non-invasive and painless transducer on a finger, ear lobe or other body area. The necessary calculations are conducted in real time, allowing for nearly instantaneous measurement of pulsus paradoxus, rapid notification for medical personnel, and quicker diagnosis and treatment. Furthermore, states of low perfusion, such as hypovolemia, will have little or no influence on the accuracy of pulsus paradoxus determined by embodiments of the present invention because the measurement of pulsus paradoxus is based upon relative changes Δ AUC during the respiratory cycle and not upon absolute values of AUC or other waveform indices.

[0052] In addition, the two dimensional parameters measured (Δ AUC, Δ iAUC and Δ dAUC) are far more accurate for determining the presence and/or magnitude of pulsus paradoxus than manual measurement or methods utilizing only change in waveform height. The decrease in LVSV that results in pulsus paradoxus is a change in volume, a three-dimensional entity. The two-dimensional mathematical waveform indices, including changes in area under the waveform curve which we utilize to measure pulsus paradoxus, accurately represent the true changes occurring in LVSV and, thus, pulsus paradoxus. Previous developers have not recognized the importance of utilizing this two-dimensional parameter, area under the curve, to measure pulsus paradoxus.

[0053] Furthermore, numerous advantages not realized under previous techniques are also achieved. For example:

[0054] 1. The preferred embodiment does not require simultaneous determination of the phases of the respiratory cycle.

[0055] 2. The preferred embodiment is based upon the percent change in AUC and, thus requires less intensive use of processing hardware and/or software, allowing less sophisticated and/or less expensive components to be used.

[0056] 3. The preferred embodiment is not significantly influenced by changes in waveform baseline occurring as a result of movement of the patient 102.

[0057] 4. Although the timing of events causing pulsus paradoxus during the respiratory cycle is altered during positive pressure ventilation, the preferred embodiment nonetheless provides accurate measurement of pulsus paradoxus during positive pressure ventilation.

[0058] In addition to the diagnosis of pulsus paradoxus, embodiments of the present invention allow for accurate and timely recognition of airway obstruction (e.g., from mucus plugs), mechanical failure (equipment failure), loss of the airway (tracheostomy tube displacement) or other conditions associated with pulsus paradoxus such as cardiac tamponade. Consequently, the methods and systems pro-

vided herein will aid in the rapid diagnosis and treatment of patients currently or potentially suffering from Sudden Infant Death Syndrome or obstructive sleep apnea. Technology dependent patients dependent on home assisted ventilation and artificial airways (tracheostomy) will also benefit from this monitoring.

[0059] Given the embodiments of the invention described herein, it should be apparent to those skilled in the art that the foregoing is merely illustrative and not limiting, having been presented by way of example only. Numerous modifications and other embodiments are within the scope of one of ordinary skill in the art and are contemplated as falling within the scope of the invention as defined by the appended claims and equivalents thereof.

I claim:

1. A method for measuring pulsus paradoxus in a mammal, the method comprising:

obtaining data indicative of pulsatile cardiovascular behavior from the mammal, the data comprising at least a first and a second pulse waveform;

determining a first AUC from at least one component of the first pulse waveform;

determining a second AUC from at least one component of the second pulse waveform;

comparing the first AUC and the second AUC to measure pulsus paradoxus.

2. The method of claim 1, further comprising determining the presence or absence of pulsus paradoxus.

3. The method of claim 2, further comprising correlating a magnitude of the difference between the first AUC and second AUC with a magnitude of pulsus paradoxus.

4. The method of claim 1, wherein the comparing step is performed by a computer.

5. The method of claim 1, wherein the data indicative of pulsatile cardiovascular behavior is obtained from plethysmography.

6. The method of claim 5, wherein the data indicative of pulsatile cardiovascular behavior is obtained from pulse oximetry.

7. The method of claim 6, wherein the data indicative of pulsatile cardiovascular behavior is obtained from reflective pulse oximetry.

8. The method of claim 6, wherein the data indicative of pulsatile cardiovascular behavior is obtained from transmissive pulse oximetry.

9. The method of claim 1, wherein the at least one component of the first pulse waveform comprises a systolic portion of the first pulse waveform, and the at least one component of the second pulse waveform comprises a systolic portion of the second pulse waveform.

10. The method of claim 9, wherein the at least one component of the first pulse waveform further comprises a diastolic portion of the first pulse waveform, and the at least one component of the second pulse waveform further comprises a diastolic portion of the second pulse waveform.

11. The method of claim 1, wherein the comparing step comprises determining a difference between the first AUC and second AUC.

12. The method of claim 11, wherein an absolute value of the difference between the first AUC and second AUC is

maximal for any two of a set of pulse waveforms obtained over a duration of at least one respiratory cycle.

13. The method of claim 11, further comprising providing a determination of the presence or absence of pulsus paradoxus in the mammal, the determination based upon whether a magnitude of the difference between the first AUC and second AUC is greater than a threshold value.

14. The method of claim 13, wherein pulsus paradoxus is determined to exist when the magnitude of the difference between the first AUC and second AUC is greater than 10 percent.

15. The method of claim 14, wherein pulsus paradoxus is determined to exist when the magnitude of the difference between the first AUC and second AUC is greater than 20 percent.

16. The method of claim 15, wherein pulsus paradoxus is determined to exist when the magnitude of the difference between the first AUC and second AUC is greater than 50 percent.

17. The method of claim 1, further comprising identifying and suppressing artifacts present within the data indicative of pulsatile cardiovascular behavior.

18. The method of claim 12, further comprising calculating an average for the difference in AUC.

19. The method of claim 18, further comprising calculating a standard deviation from the average difference in AUC.

20. The method of claim 19, further comprising calculating a variance from the standard deviation of the difference in AUC.

21. The method of claim 12, further comprising calculating a weighted signal average for the difference in AUC.

22. A system for measuring pulsus paradoxus in a mammal, the system comprising:

means for obtaining data indicative of pulsatile cardiovascular behavior from the mammal, the data comprising at least a first and a second pulse waveform;

means for determining a first AUC from at least one component of the first pulse waveform, and for determining a second AUC from at least one component of the second pulse waveform;

means for comparing the first AUC and the second AUC to measure pulsus paradoxus.

23. The system of claim 22, wherein the means for obtaining data indicative of pulsatile cardiovascular behavior comprises a plethysmograph.

24. The system of claim 23, wherein the plethysmograph comprises a reflective plethysmograph.

25. The system of claim 23, wherein the plethysmograph comprises a transmissive plethysmograph.

26. The system of claim 23, wherein the means for obtaining data indicative of pulsatile cardiovascular behavior comprises a pulse oximeter.

27. A system for measuring pulsus paradoxus in a mammal, the system comprising:

a pulse oximeter that obtains data indicative of pulsatile cardiovascular behavior from the mammal, the data comprising at least a first and a second pulse waveform;

a measurer that measures a first AUC from at least one component of the first pulse waveform, and measures a second AUC from at least one component of the second pulse waveform;

an analyzer that compares the first AUC and the second AUC to measure pulsus paradoxus.

28. The system of claim 27, wherein a computer comprises the analyzer.

29. A method for measuring pulsus paradoxus in a mammal, the method comprising:

obtaining data indicative of pulsatile cardiovascular behavior from the mammal, the data comprising at least a first and a second pulse waveform;

determining a first value that is a time-domain function of a change in height of the first pulse waveform over at least a partial duration of the first pulse waveform;

determining a second value that is a time-domain function of a change in height of the second pulse waveform over at least a partial duration of the second pulse waveform;

comparing the first value and the second value to measure pulsus paradoxus.

30. The method of claim 29, further comprising determining the presence or absence of pulsus paradoxus.

31. The method of claim 29, further comprising correlating a magnitude of the difference between the first value and second value with a magnitude of pulsus paradoxus.

32. The method of claim 29, wherein the first and second values are slopes.

33. The method of claim 32, wherein the first and second values are average slopes.

34. The method of claim 32, wherein the first and second values are maximal slopes.

35. The method of claim 29, wherein the first value is a first AUC and the second value is a second AUC.

36. The method of claim 29, wherein the partial duration of the first pulse waveform is systole.

37. A method for measuring pulsus paradoxus in a mammal, the method comprising:

obtaining data indicative of pulsatile cardiovascular behavior from the mammal, the data comprising a first plethysmographic or pressure amplitude obtained at a first time interval after an onset of a first pulse, and a second plethysmographic or pressure amplitude obtained at a second time interval after the onset of the first pulse;

determining a first value that is a time-domain function of a difference between the first and second plethysmographic or pressure amplitudes;

obtaining a third plethysmographic or pressure amplitude obtained at the first time interval after an onset of a second pulse, and a fourth plethysmographic or pressure amplitude obtained at the second time interval after the onset of the second pulse;

determining a second value that is a time-domain function of a difference between the third and fourth plethysmographic or pressure amplitudes; and

comparing the first value and the second value to measure pulsus paradoxus.

38. The method of claim 37, further comprising determining the presence or absence of pulsus paradoxus.

39. The method of claim 37, further comprising correlating a magnitude of the difference between the first value and second value with a magnitude of pulsus paradoxus.

40. The method of claim 37, wherein the first and second values are slopes.

41. The method of claim 40, wherein the first and second values are average slopes.

42. The method of claim 40, wherein the first and second values are maximal slopes.

43. The method of claim 40, wherein the first value is a first AUC and the second value is a second AUC.

44. The method of claim 37, wherein the partial duration of the first pulse waveform is systole.

45. A method for measuring pulsus paradoxus in a mammal, the method comprising:

obtaining data indicative of pulsatile cardiovascular behavior from the mammal, the data comprising a first plethysmographic or pressure amplitude obtained at a time interval after an onset of a first pulse, and a second

plethysmographic or pressure amplitude obtained at said time interval after an onset of a second pulse;

comparing the first plethysmographic or pressure amplitude and the second plethysmographic or pressure amplitude to measure pulsus paradoxus.

46. The method of claim 45, wherein the first plethysmographic or pressure amplitude is a minimum for at least two pulses in a respiratory cycle and the second plethysmographic or pressure amplitude is a maximum for the at least two pulses in the respiratory cycle.

47. A method for measuring pulsus paradoxus, comprising:

comparing plethysmographic amplitude as a function of time during inspiration with plethysmographic amplitude as a function of time during expiration.

* * * * *

专利名称(译)	测量脉冲矛盾的方法和装置		
公开(公告)号	US20040044276A1	公开(公告)日	2004-03-04
申请号	US10/228641	申请日	2002-08-27
[标]申请(专利权)人(译)	阿诺德DONALD		
申请(专利权)人(译)	阿诺德DONALD		
当前申请(专利权)人(译)	PRECISION奇脉, INC.		
[标]发明人	ARNOLD DONALD		
发明人	ARNOLD, DONALD		
IPC分类号	A61B5/00		
CPC分类号	A61B5/14551		
其他公开文献	US6869402		
外部链接	Espacenet USPTO		

摘要(译)

公开了一种利用来自患者的动脉和/或小动脉脉搏波形数据源以测量脉搏矛盾的方法和装置。所描述的动脉脉搏波形数据源是脉搏血氧计体积描记器，但可以是任何类似的波形数据源，包括动脉内换能器，血压换能器或体积描记器。通过结合诸如脉冲波形曲线下面积的值的测量值，其是在波形的至少部分持续时间内脉冲波形的高度变化的时域函数，本发明的实施例表示在先前描述的测量脉搏矛盾的方法方面的显著改进，并且进一步提高了产生脉搏悖论的多个贡献变量的测量精度，特别是舒张事件的贡献，达到如此测量的体征可被视为“修改了Pulsus Paradoxus。”

