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(54) **PHYSIOLOGICAL PARAMETER TRACKING SYSTEM**

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USPC 600/310, 322, 323, 326, 328, 330, 331, 600/334, 335, 336, 340, 344, 473, 476; 356/41

See application file for complete search history.

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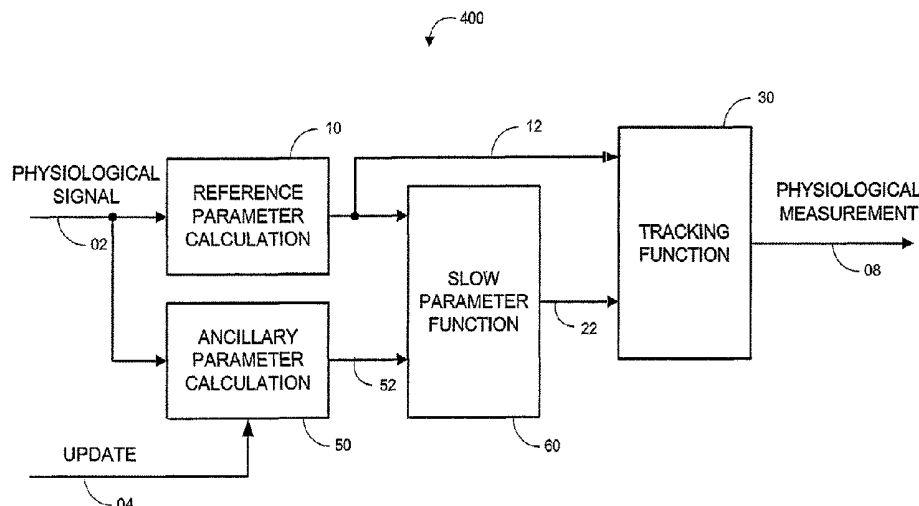
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(57) **ABSTRACT**

A physiological parameter tracking system has a reference parameter calculator configured to provide a reference parameter responsive to a physiological signal input. A physiological measurement output is a physiological parameter derived from the physiological signal input during a favorable condition and an estimate of the physiological parameter according to the reference parameter during an unfavorable condition.

19 Claims, 5 Drawing Sheets



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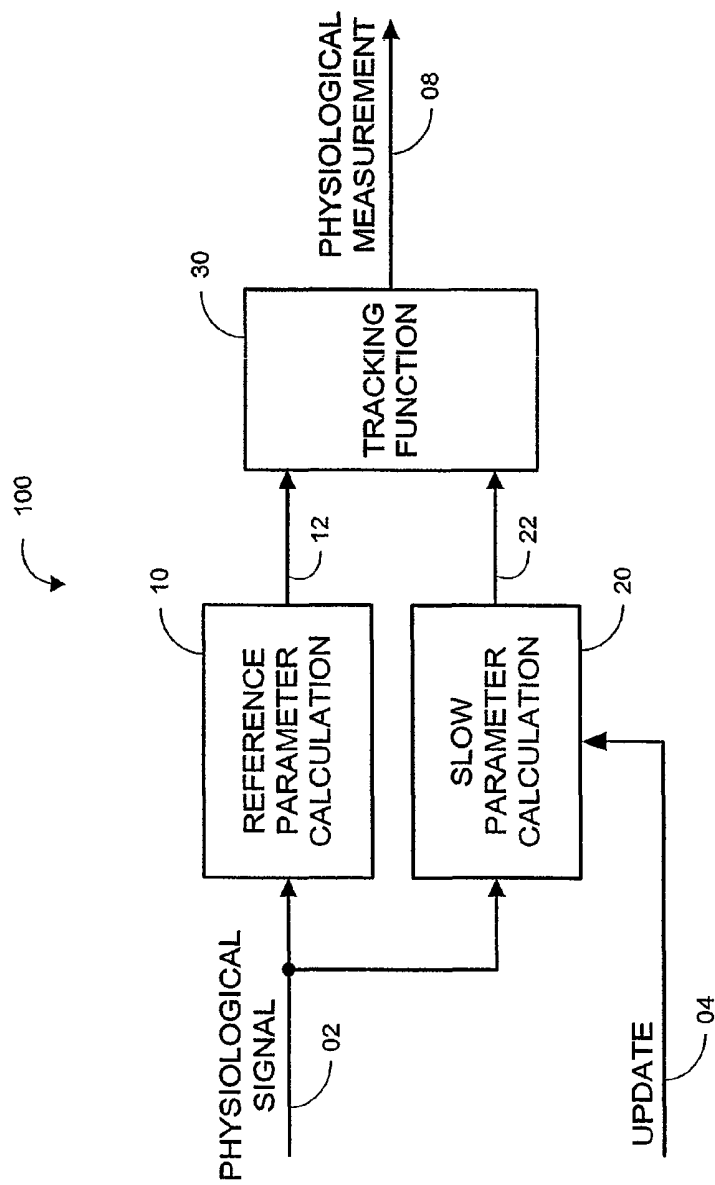


FIG. 1

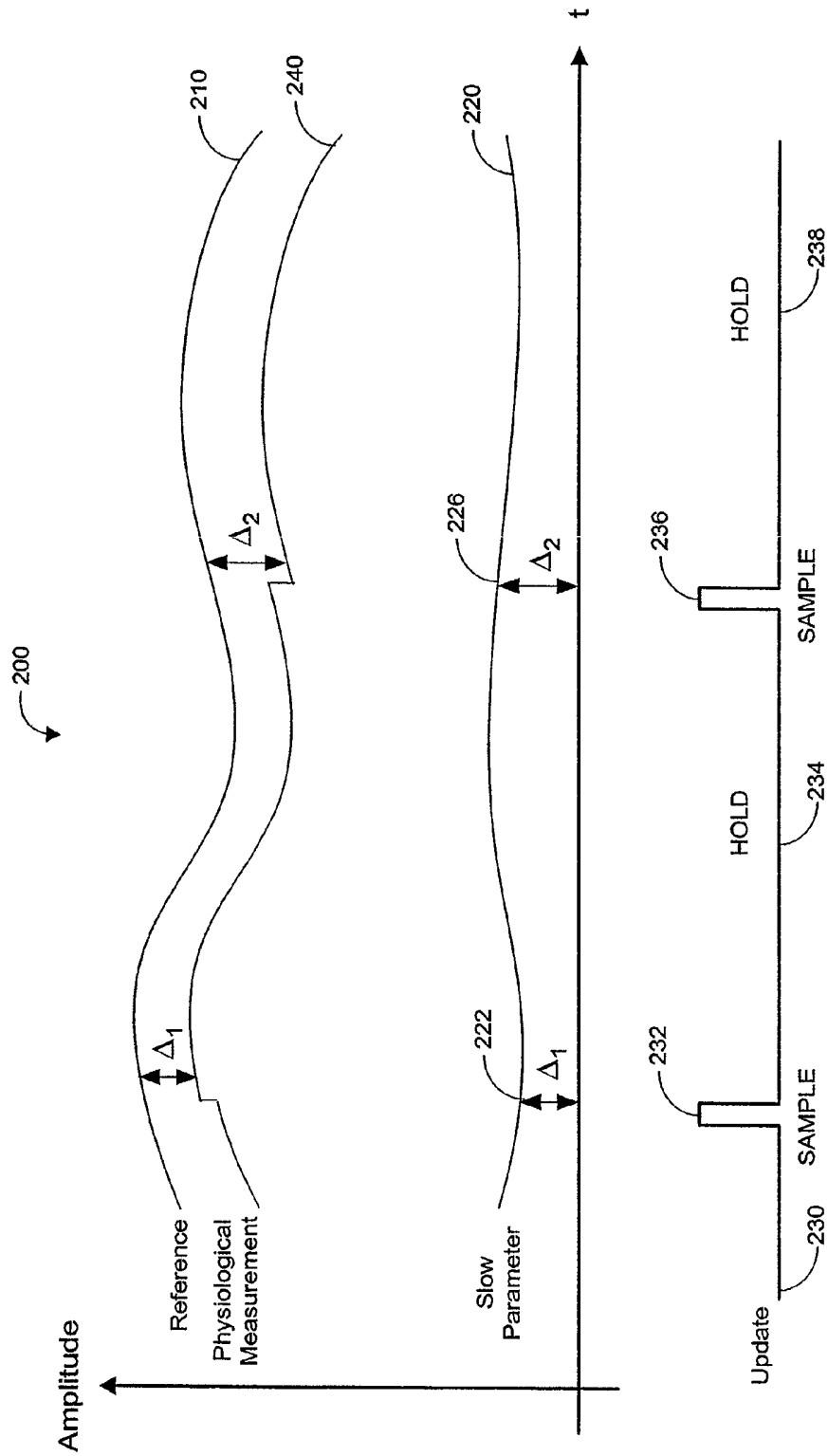


FIG. 2

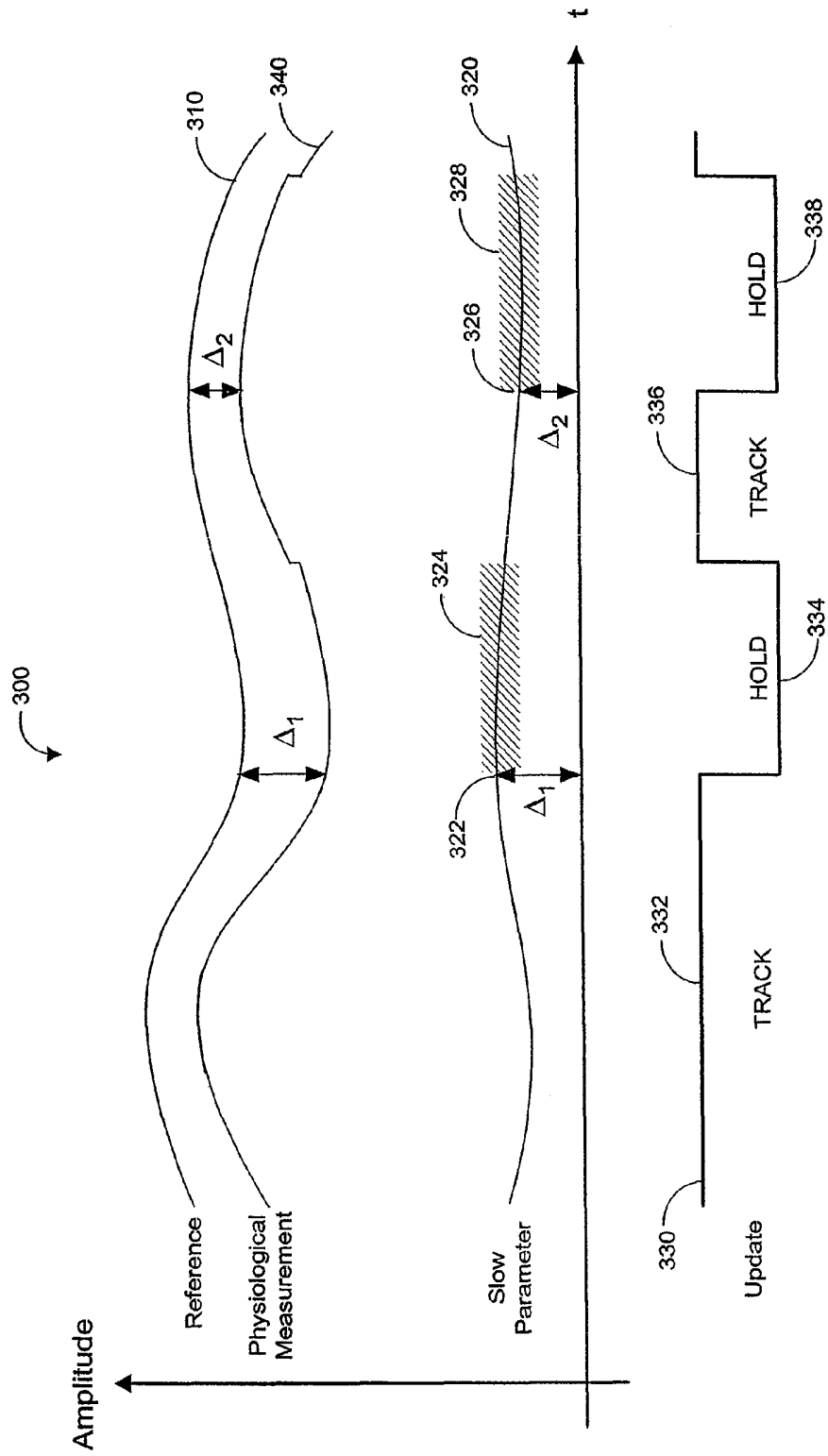


FIG. 3

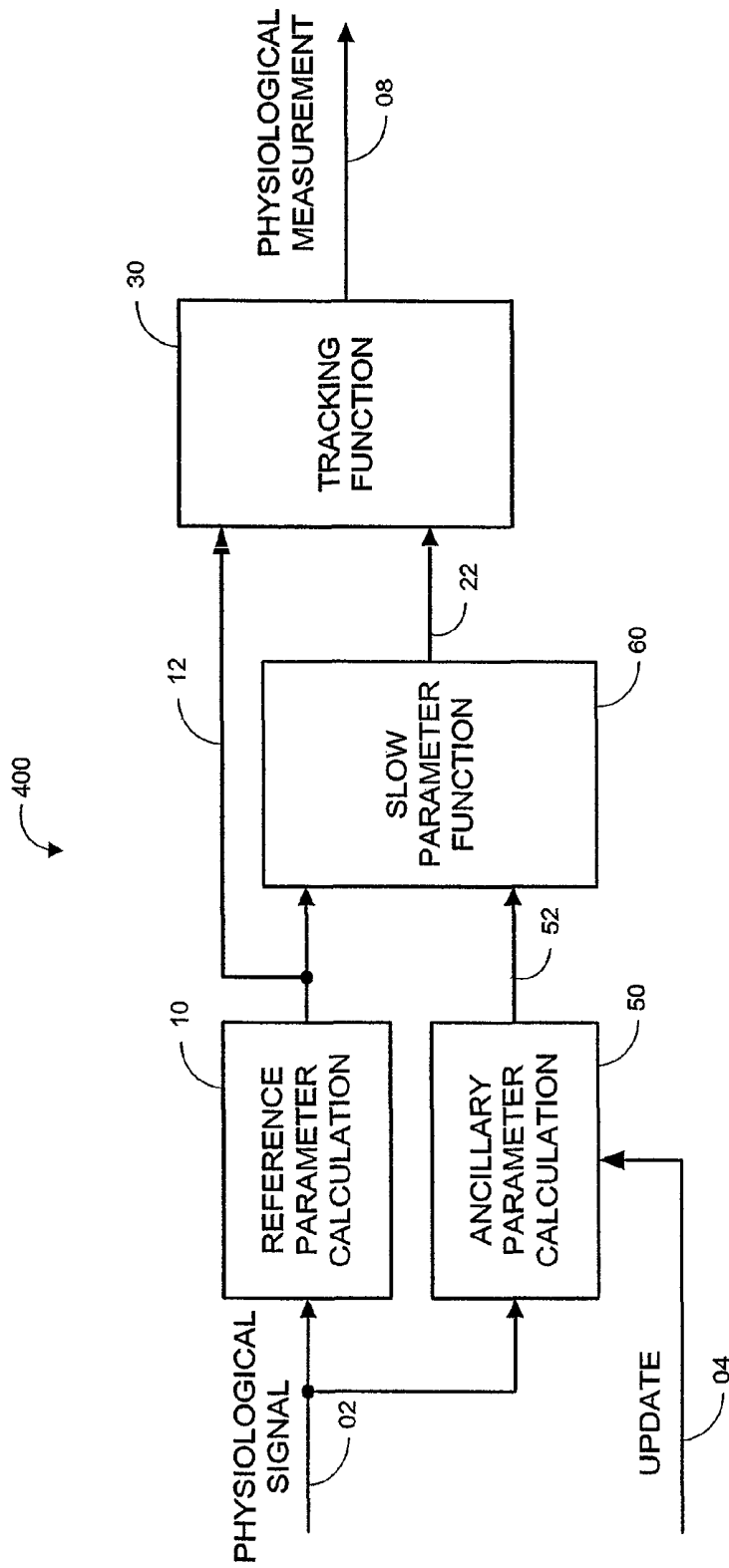


FIG. 4

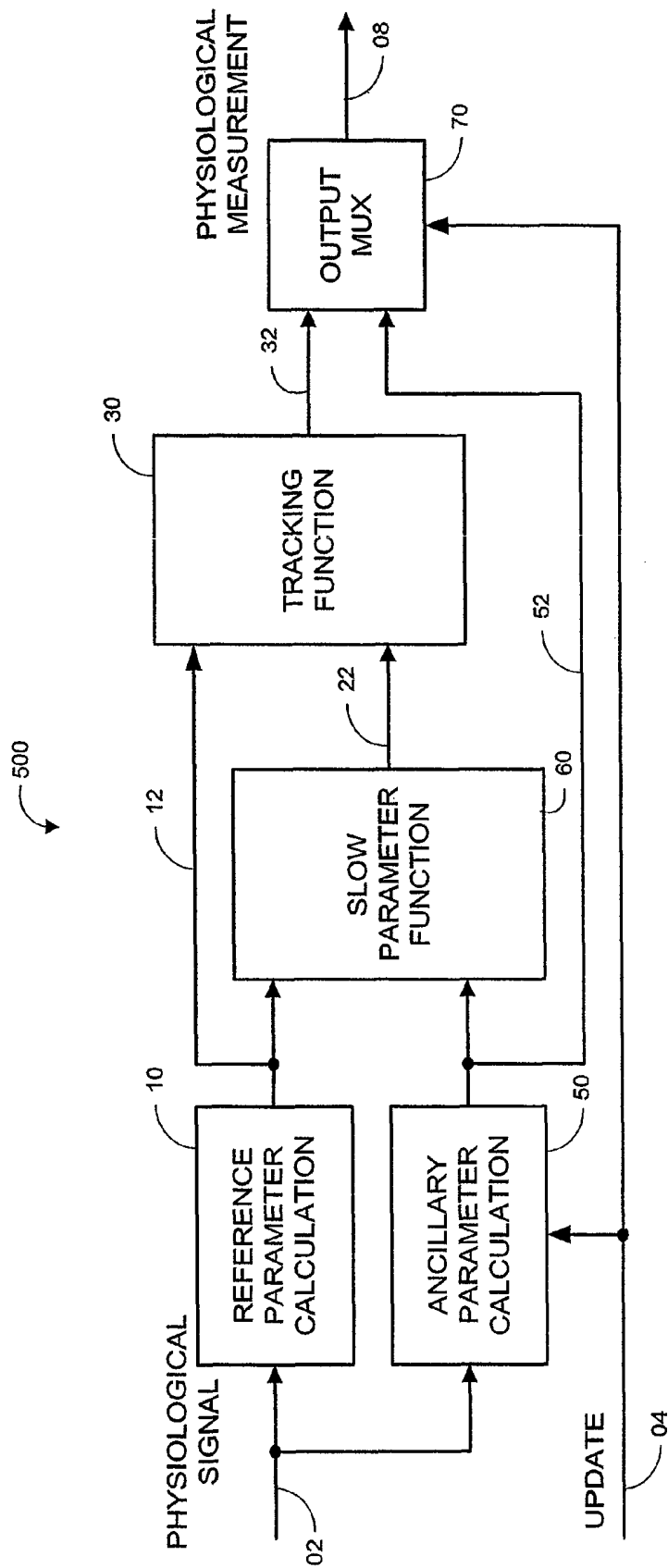


FIG. 5

PHYSIOLOGICAL PARAMETER TRACKING SYSTEM

CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims priority benefit under 35 U.S.C. §120 from, and is a continuation of U.S. patent application Ser. No. 11/834,602, filed Aug. 6, 2007, entitled "Physiological Parameter Tracking System", which is a continuation of U.S. patent application Ser. No. 10/930,048, filed Aug. 30, 2004, entitled "Physiological Parameter Tracking System", which claims priority benefit under 35 U.S.C. §119(e) from U.S. Provisional Patent Application No. 60/498,749, filed Aug. 28, 2003, entitled "Physiological Parameter Tracking System". The present application incorporates the foregoing disclosures herein by reference.

BACKGROUND OF THE INVENTION

Oxygen transport from the lungs to body tissue can be monitored by measuring various physiological parameters. For example, oxygen saturation of arterial blood (S_aO_2) is a measure of the ratio of oxyhemoglobin (HbO_2) concentration to the sum of HbO_2 and deoxyhemoglobin (Hb) concentrations in the arterial blood. Because HbO_2 is the major oxygen carrying component of blood, S_aO_2 is indicative of oxygen delivery to body tissues. As another example, oxygen saturation of venous blood (S_vO_2) is a similar measure of HbO_2 and Hb concentrations in venous blood and is indicative of oxygen consumption by body tissues. Measurements of the concentrations of carboxyhemoglobin ($HbCO$) and methemoglobin ($MetHb$) are indicative of abnormal hemoglobin constituents that interfere with oxygen transport.

Pulse oximetry is a noninvasive, easy to use, inexpensive procedure for measuring the oxygen saturation level of arterial blood. Pulse oximeters perform a spectral analysis of the pulsatile component of arterial blood in order to determine oxygen saturation ($S_{pa}O_2$), which is an estimate of S_aO_2 . A pulse oximetry system has a sensor and a monitor. The sensor has emitters that typically consist of a red light emitting diode (LED) and an infrared LED that project light through blood vessels and capillaries underneath a tissue site, such as a fingernail bed. A sensor also has a detector that typically is a photodiode positioned opposite the LEDs so as to detect the emitted light as it emerges from the tissue site. A pulse oximetry sensor is described in U.S. Pat. No. 6,088,607 entitled "Low Noise Optical Probe," which is assigned to Masimo Corporation, Irvine, Calif. and incorporated by reference herein.

SUMMARY OF THE INVENTION

One aspect of a physiological parameter tracking system comprises a physiological signal and first, second, third and fourth calculators. The physiological signal has at least first and second intensity signal components received from a light-sensitive detector that detects light of at least first and second wavelengths transmitted through body tissue carrying pulsing blood. The first calculator is configured to output a reference parameter responsive to the physiological signal. The second calculator is configured to output an ancillary parameter responsive to the physiological signal. The third calculator is configured to output a slow parameter that is a function of the reference parameter and the ancillary parameter. The slow parameter is a function of time that is slowly

varying relative to the reference parameter and the ancillary parameter. A fourth calculator is configured to output a physiological measurement responsive to the reference parameter and the slow parameter. In an embodiment, the fourth calculator provides a physiological measurement that is at least in part a function of the reference parameter and the slow parameter. In an embodiment, the physiological measurement is a function of the reference parameter and the slow parameter during a first time interval and is the ancillary parameter during a second time interval. In an embodiment, the first time interval includes a period when calculations of the ancillary parameter are unfavorable. In an embodiment, the second time interval includes a period when calculations of the ancillary parameter are favorable.

Another aspect of a physiological parameter tracking system comprises inputting a physiological signal, deriving a physiological measurement from the physiological signal during a favorable condition, estimating the physiological measurement during an unfavorable condition and outputting a combination of the derived physiological measurement and the estimated physiological measurement. In an embodiment, estimating comprises calculating a slow parameter that is physiologically related to the reference parameter and the physiological measurement and tracking the reference parameter with the slow parameter. In an embodiment, outputting comprises selecting between estimated physiological measurement and derived measurement according to the favorable condition and the unfavorable condition. In an embodiment, the favorable condition and the unfavorable conditions relate to power consumption goals. In an embodiment, the favorable condition and the unfavorable conditions relate to the quality of the physiological signal.

A further aspect of a physiological parameter tracking system comprises a physiological signal input, a reference parameter calculator and a physiological measurement means for outputting and estimating. The physiological signal input has at least first and second intensity signal components received from a light-sensitive detector that detects light of at least first and second wavelengths transmitted through body tissue carrying pulsing blood. The reference parameter calculator is configured to output a reference parameter responsive to the physiological signal. The physiological measurement means outputs a physiological parameter derived from the physiological signal input during a favorable condition and estimates the physiological parameter according to the reference parameter during an unfavorable condition. In an embodiment, a slow parameter means relates the reference parameter to the physiological parameter during the unfavorable condition. In an embodiment, an update means selects a first time period for outputting the derived physiological parameter and a second time period for outputting the estimated physiological parameter.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a block diagram of a slow parameter calculation embodiment of a physiological parameter tracking system;

FIG. 2 is a graph illustrating operation of a physiological parameter tracking system in a sample and hold (S/H) mode;

FIG. 3 is a graph illustrating operation of a physiological parameter tracking system in a track and hold (T/H) mode;

FIG. 4 is a block diagram of an ancillary calculation embodiment of a physiological parameter tracking system for operation in a S/H mode; and

FIG. 5 is a block diagram of an ancillary calculation embodiment of a physiological parameter tracking system for operation in a T/H mode.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Overview

FIGS. 1, 4 and 5 illustrate embodiments of a physiological parameter tracking system that advantageously provide a clinically accurate physiological measurement by tracking a reference parameter based upon a slowly varying ("slow") parameter. As such, it is not necessary to continuously or frequently perform complex calculations to derive the physiological measurement. That is, the physiological measurement is a relatively simple function of the reference parameter and the slow parameter. Slow parameter calculations are performed only when conditions are favorable, or alternatively, suspended when conditions are not favorable, as indicated by an update command. The update command may be responsive to conditions such as power consumption goals or the quality of a physiological signal input to name a few.

In one embodiment, the slow parameter is HbCO or MetHb, and the reference parameter is $S_{pa}O_2$. Accordingly, the physiological measurement is $S_{pa}O_2$ corrected for the presence of one or both of these abnormal hemoglobin constituents. In another embodiment, the slow parameter is $\Delta_{av}=S_{pa}O_2-S_vO_2$, a measure of oxygen consumption at a tissue site, and the reference parameter is $S_{pa}O_2$. Accordingly, the physiological measurement is an estimate of S_vO_2 .

Slow Parameter Calculation

FIG. 1 illustrates a slow parameter calculation embodiment of a physiological parameter tracking system 100 in which the slow parameter 22 is derived from and responsive to a physiological signal 02. The physiological parameter tracking system 100 has a physiological signal 02 input, a reference parameter calculation 10, a slow parameter calculation 20 and a tracking function 30 and generates a physiological measurement 08 output. The reference parameter calculation 10 generates a reference parameter 12 from the physiological signal 02. The slow parameter calculation 20 generates the slow parameter 22 from the physiological signal 02 input. The tracking function 30 generates the physiological measurement 08 from the reference parameter 12 and the slow parameter 22.

As shown in FIG. 1, the physiological signal 02 is responsive to a physiological condition. In one embodiment, the physiological signal 02 originates from an optical sensor (not shown) attached to a tissue site. The sensor transmits multiple wavelengths of optical energy $\lambda_1, \lambda_2, \dots, \lambda_n$ into the tissue site and detects corresponding optical energy emerging from the tissue site. The reference parameter calculation 10 may include pulse oximetry algorithms that operate on the physiological signal 02 to generate arterial oxygen saturation, $S_{pa}O_2$, as the reference parameter 12. A pulse oximetry signal processor and algorithms are described in U.S. Pat. No. 5,632,272 entitled Signal Processing Apparatus which is assigned to Masimo Corporation, Irvine, Calif. and incorporated by reference herein.

Also shown in FIG. 1, the slow parameter calculation 20 generates a slow parameter 22 from the physiological signal input 02 according to an update command 04. As an example, the slow parameter calculation 20 may include algorithms that operate on the physiological signal 02 to generate a measure of the concentration of abnormal hemoglobin, such as HbCO or MetHb. Multiple wavelength

signal processing for measuring abnormal hemoglobin constituents, for example, is described in U.S. Provisional Patent App. No. 60/426,638 entitled "Parameter Compensated Physiological Monitor," U.S. Provisional Patent App. No. 60/428,419 entitled "Blood Parameter Measurement System," and U.S. Pat. No. 6,229,856 entitled "Method and Apparatus for Demodulating Signals in a Pulse Oximetry System, which is assigned to Masimo Corporation, Irvine, Calif., all incorporated by reference herein.

Further shown in FIG. 1, the update command 04 may operate in a sample and hold (S/H) mode. That is, when the update command 04 is asserted, the slow parameter calculation 20 is triggered and the resulting slow parameter 22 value is held until a subsequent calculation. Operation of a physiological parameter tracking system having a S/H update is described with respect to FIG. 2, below. Alternatively, the update command 04 may operate in a track and hold (T/H) mode. That is, while the update command 04 is asserted, the slow parameter calculation 20 continues to generate values for the slow parameter 22. When the update command 04 is not asserted, the last generated value of the slow parameter 22 is held until the update command 04 is once more asserted. Operation of a physiological parameter tracking system having a T/H update is described with respect to FIG. 3, below.

Tracking Examples

FIG. 2 is an amplitude versus time graph 200 illustrating operation of a physiological parameter tracking system utilizing a S/H update. The graph 200 illustrates a reference curve 210 corresponding to a reference parameter 12 (FIG. 1) and a slow parameter curve 220 corresponding to a slow parameter 22 (FIG. 1). Below the graph 200 is a timing diagram 230 corresponding to the update command 04 (FIG. 1). A physiological measurement curve 240 corresponds to the physiological measurement 08 (FIG. 1).

As shown in FIG. 2, the physiological measurement curve 240 tracks the reference curve 210 according to a tracking function 30 (FIG. 1), which in this illustration is the difference between the reference parameter 12 (FIG. 1) and the slow parameter 22 (FIG. 1). A slow parameter 220 value is calculated at sample times 232, 236 and maintained throughout hold periods 234, 238. In particular, during a first sample time 232, a slow parameter value 222 of Δ_1 is calculated, and during a second sample time 236, a slow parameter value 226 of Δ_2 is calculated. As a result, during a first hold period 234, the physiological measurement curve 240 tracks the reference curve 210 by a difference of Δ_1 . Likewise, during a second hold period 238, the physiological measurement curve 240 tracks the reference curve 210 by a difference of Δ_2 . In this manner, the physiological measurement 240 is advantageously displayed with clinical accuracy utilizing only occasional computational resources and reducing power consumption accordingly.

FIG. 3 is an amplitude versus time graph 300 illustrating operation of a physiological parameter tracking system utilizing a T/H update. The graph 300 illustrates a reference curve 310 corresponding to a reference parameter 12 (FIG. 1) and a slow parameter curve 320 corresponding to a slow parameter 22 (FIG. 1). Below the graph 300 is a timing diagram 330 corresponding to the update command 04 (FIG. 1). A physiological measurement curve 340 corresponds to the physiological measurement 08 (FIG. 1).

As shown in FIG. 3, the physiological measurement curve 340 tracks the reference curve 310 according to a tracking function 30 (FIG. 1), which, again, is the difference between the reference parameter 12 (FIG. 1) and the slow parameter 22 (FIG. 1). Slow parameter 320 values are calculated

throughout track periods 332, 336, and the last computed values are maintained throughout the corresponding hold periods 334, 338. In particular, during a first track period 332, the physiological measurement curve 340 is the reference curve 310 minus the slow parameter curve 320. At the end of the first track period 332, a slow parameter value 332 of Δ_1 is maintained throughout the first hold period 334. As a result, during the first hold period 334, the physiological measurement curve 340 is the reference curve 310 minus Δ_1 and does not depend on the slow parameter curve 320. That is, during the first hold period 332, the physiological measurement curve 340 tracks the reference curve 310 by a difference of Δ_1 .

The "track" periods 332, 336 are so named because the slow parameter calculation 20 (FIG. 1) in response to the update timing 330 operates in a manner roughly analogous to a conventional track/hold amplifier when its output tracks the input. These are not to be confused with the periods when the physiological measurement curve 340 is "tracking" the reference parameter curve 310, which actually is during the hold periods 334, 338, when the slow parameter 22 (FIG. 1) output is held constant.

Also shown in FIG. 3, during a second track period 336, the physiological measurement curve 340 is again the reference curve 310 minus the slow parameter curve 320. At the end of the second track period 336, a slow parameter value 326 of Δ_2 is maintained throughout the second hold period 338. As a result, during the second hold period 338, the physiological measurement curve 340 is the reference curve 310 minus Δ_2 and does not depend on the slow parameter curve 320. That is, during the second hold period 338, the physiological measurement curve 340 tracks the reference curve 310 at a difference of Δ_2 .

Further shown in FIG. 3, the hold periods 334, 338 may correspond to slow parameter drop-out periods 324, 328, i.e. periods when the slow parameter cannot be accurately calculated. In this manner, the physiological measurement 340 is advantageously displayed with clinical accuracy even when noise or other signal corruption prevents measurement of the slow parameter 320.

Ancillary Parameter Calculation

FIG. 4 illustrates an ancillary parameter calculation embodiment of a physiological parameter tracking system 400 in which the slow parameter 22 is derived from an ancillary parameter 52 in S/H mode. The ancillary parameter 52, in turn, is derived from a physiological signal 02. That is, unlike the slow parameter calculation embodiment 100 (FIG. 1), the slow parameter 22 is only indirectly derived from and responsive to the physiological signal 02. The physiological parameter tracking system 400 has a physiological signal 02 input, a reference parameter calculation 10 and a tracking function 30, and, accordingly, generates a physiological measurement 08, similarly as described with respect to FIG. 1, above. However, in the ancillary calculation embodiment 400, the slow parameter 22 is a function 60 of the reference parameter 12 and/or an ancillary parameter 52. An ancillary parameter calculation 50 generates the ancillary parameter 52 from the physiological signal input 02 according to a S/H update command 04 input, such as described with respect to FIG. 2, above.

As an example, the ancillary parameter calculation 50 may include algorithms that operate on the physiological signal 02 to intermittently calculate venous oxygen saturation, $S_{pv}O_2$, as determined by a S/H update command 04. A corresponding slow parameter function 60 is the difference between an $S_{pa}O_2$ reference parameter 12 and the $S_{pv}O_2$ ancillary parameter 52 to yield a Δ_{av} slow parameter 22.

Then, the tracking function 30 is a difference between the $S_{pa}O_2$ reference parameter 12 and the sampled Δ_{av} slow parameter 22 to generate a $S_{pv}O_2$ physiological measurement 08. That is, the physiological measurement 08 in this example advantageously provides a continuous measurement of venous saturation $S_{pv}O_2$ utilizing intermittent calculations of $S_{pv}O_2$. Apparatus and methods for determining $S_{pv}O_2$ from mechanical or ventilator induced perturbation of the venous blood volume are described in U.S. Pat. No. 5,638,816 entitled "Active Pulse Blood Constituent Monitoring" and U.S. Pat. No. 6,334,065 entitled "Stereo Pulse Oximeter," which are assigned to Masimo Corporation, Irvine, Calif. and are incorporated by reference herein.

FIG. 5 illustrates an ancillary parameter calculation embodiment of a physiological parameter tracking system 500 in which the slow parameter 22 is derived from an ancillary parameter 52 in T/H mode. The ancillary parameter 52, in turn, is derived from a physiological signal 02. The physiological parameter tracking system 500 has a physiological signal 02 input, a reference parameter calculation 10, an ancillary parameter calculation 50, a slow parameter function 60 and a tracking function 30, and, accordingly, generates a physiological measurement 08, similarly as described with respect to FIG. 4, above. However, in this ancillary calculation embodiment 500, the update command 04 operates in a track and hold mode, as described with respect to FIG. 3, above. Accordingly, the ancillary calculation embodiment 500 also has an output multiplexer 70 having the tracking function output 32 and the ancillary parameter 52 as inputs and the physiological measurement 08 as an output, as controlled by the update command 04 input. As such, the physiological measurement 08 is the ancillary parameter 52 during a track period 332, 336 (FIG. 3) of the update command 04 and is a function of the ancillary parameter 52 and the reference parameter 10 during a hold period 334, 338 (FIG. 3) of the update command 04. That is, the physiological measurement 08 is advantageously the ancillary parameter 52 except during a hold period, when the physiological measurement 08 tracks the reference parameter 12 according to the maintained value of the slow parameter 22.

As an example, the ancillary parameter calculation 50 may continuously calculate venous oxygen saturation, $S_{pv}O_2$, as determined by the update command 04 during track periods, and this calculation is provided as the physiological measurement 08. However, during hold periods of the update command 04, the physiological measurement 08 becomes $S_{pv}O_2$, i.e. the $S_{pa}O_2$ reference parameter 12 minus a maintained value of the Δ_{av} slow parameter 22. The physiological measurement 08 in this example advantageously provides a measurement of venous saturation that is continuous through drop-out periods.

A physiological parameter tracking system has been disclosed in detail in connection with various embodiments. These embodiments are disclosed by way of examples only and are not to limit the scope of the claims that follow. One of ordinary skill in the art will appreciate many variations and modifications.

What is claimed is:

1. A method of monitoring a patient's condition by electronically tracking one or more physiological parameters of the patient through processing signals received from a noninvasive sensor, said electronically tracking including transforming said signals into measurements of said one or more physiological parameters, the method comprising: receiving a physiological signal from a noninvasive physiological sensor that detects light of at least first

and second wavelengths transmitted through body tissue carrying pulsing blood;
electronically processing said signal in a pulse oximetry signal processor, including:

calculating a first blood parameter responsive to the physiological signal, the first blood parameter having a first rate of amplitude change over time;
responsive to an update command:

in a sample period of the update command calculating a sampled value of a second blood parameter responsive to the physiological signal, the second blood parameter having a second rate of amplitude change over time, wherein the first blood parameter and the second blood parameter are different blood parameters, and

in a subsequent hold period of the update command following the sample period, maintaining the sampled value of the second blood parameter calculated during the sample period,

calculating a third blood parameter based at least in part on a difference between a current value of the first blood parameter and the sampled value of the second blood parameter; and

determining a physiological measurement of oxygen saturation in the body tissue at least partly by tracking the first blood parameter by the third blood parameter such that, in the sample period, the physiological parameter represents the sampled value of the second blood parameter and, in the hold period, the physiological parameter represents an estimate of the second blood parameter based on the current value of the first blood parameter and the sampled value of the second blood parameter, wherein determining the physiological measurement using the sample period and the hold period utilizes less computational resources and consumes less power than determining the physiological measurement without the sample period and the hold period; and

displaying the physiological measurement.

2. The method of claim 1, wherein the calculating said second blood parameter comprises suspending calculations during a first time interval comprising the hold period and performing calculations during a second time interval comprising the sample period as indicated by the update command.

3. The method of claim 2, wherein the determining said physiological measurement further comprises calculating the physiological measurement, during the first time interval, as a function of the first blood parameter and a difference corresponding to the measured value of the third blood parameter.

4. The method of claim 2, wherein the determining said physiological measurement further comprises outputting the second blood parameter during the second time interval.

5. The method of claim 1, further comprising asserting the update command wherein, upon assertion, a value of the third blood parameter is repeatedly generated.

6. The method of claim 1, further comprising asserting the update command wherein, upon assertion, a value of the third blood parameter is held for a predetermined duration.

7. The method of claim 1, wherein the calculating said second blood parameter further comprises repeatedly calculating venous oxygen saturation.

8. The method of claim 1, comprising providing said noninvasive physiological sensor.

9. The method of claim 1, comprising providing a patient monitor including an input to receive said signal and a signal processor to execute said processing.

10. The method of claim 1, further comprising asserting the update command responsive to power consumption goals or quality of a physiological signal input.

11. A patient monitor configured to receive a signal from a non-invasive optical sensor, the monitor comprising:

an input configured to receive a physiological signal from a noninvasive physiological sensor that detects light of at least first and second wavelengths transmitted through body tissue carrying pulsing blood, said physiological signal responsive to said detection of said light; and

a pulse oximetry signal processor configured to receive data responsive to said physiological signal and configured to:

calculate a first blood parameter responsive to the physiological signal, the first blood parameter having a first rate of amplitude change over time,
responsive to an update command:

in a sample period of the update command calculate a sampled value of a second blood parameter responsive to the physiological signal, the second blood parameter having a second rate of amplitude change over time, wherein the first blood parameter and the second blood parameter are different blood parameters, and

in a subsequent hold period of the update command following the sample period, maintain the sampled value of the second blood parameter calculated during the sample period,

calculate a third blood parameter based at least in part on a difference between a current value of the first blood parameter and the sampled value of the second blood parameter, and

determine a physiological measurement of oxygen saturation in the body tissue at least partly as a difference between the first blood parameter and the third blood parameter such that, in the sample period, the physiological parameter represents the sampled value of the second blood parameter and, in the hold period, the physiological parameter represents an estimate of the second blood parameter based on the current value of the first blood parameter and the sampled value of the second blood parameter; and
a display configured to display the physiological measurement.

12. The patient monitor of claim 11, wherein the signal processor is also configured to suspend calculations during a first time interval comprising the hold period and to perform calculations during a second time interval comprising the sample period as indicated by the update command.

13. The patient monitor of claim 12, wherein the signal processor is also configured to calculate the physiological measurement as a function of the first blood parameter and the third blood parameter during the first time interval.

14. The patient monitor of claim 12, wherein the signal processor is also configured to output the second blood parameter during the second time interval.

15. The patient monitor of claim 11, wherein the signal processor is also configured to generate a measure of the concentration of abnormal hemoglobin.

16. The patient monitor of claim 11, wherein the signal processor is also configured to assert the update command wherein, upon assertion, a value of the third blood parameter is repeatedly generated.

17. The patient monitor of claim 11, wherein the signal processor is also configured to assert the update command wherein, upon assertion, a value of the third blood parameter is held for a predetermined duration.

18. The patient monitor of claim 11, wherein the signal processor is also configured to repeatedly calculate venous oxygen saturation.

19. The patient monitor of claim 11, wherein the signal processor is configured to assert the update command responsive to power consumption goals or quality of a physiological signal input.

* * * * *

专利名称(译)	生理参数跟踪系统		
公开(公告)号	US9788768	公开(公告)日	2017-10-17
申请号	US13/777936	申请日	2013-02-26
[标]申请(专利权)人(译)	梅西莫股份有限公司		
申请(专利权)人(译)	Masimo公司		
当前申请(专利权)人(译)	Masimo公司		
[标]发明人	AL ALI AMMAR DIAB MOHAMED K WEBER WALTER M		
发明人	AL-ALI, AMMAR DIAB, MOHAMED K. WEBER, WALTER M.		
IPC分类号	A61B5/1455 A61B5/00		
CPC分类号	A61B5/1455 A61B5/14551 A61B5/14552 A61B5/72 A61B2560/0204		
代理机构(译)	KNOBBE , MARTENS , 奥尔森 & BEAR LLP		
优先权	11/834602 2013-02-26 US 10/930048 2007-08-07 US 60/498749 2003-08-28 US		
其他公开文献	US20130274572A1		
外部链接	Espacenet USPTO		

摘要(译)

生理参数跟踪系统具有参考参数计算器，其被配置为响应于生理信号输入提供参考参数。生理测量输出是从在有利条件期间输入的生理信号导出的生理参数和在不利条件期间根据参考参数估计生理参数。

