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(54) **PERFUSION INDEX SMOOTHING**

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(73) Proprietor: **Masimo Corporation
Irvine, CA 92618 (US)**

(72) Inventor: **AL-ALI, Ammar
Tustin, 92782 (US)**

(74) Representative: **Vossius & Partner
Patentanwälte Rechtsanwälte mbB
Siebertstrasse 3
81675 München (DE)**

(56) References cited:
**EP-A- 0 870 466 WO-A-2004/034898
WO-A-2006/097866 US-A1- 2003 163 032
US-B1- 6 334 065**

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Description

BACKGROUND OF THE DISCLOSURE

Field of the Disclosure

[0001] The present disclosure relates in general to patient monitoring and in particular to oximeter patient monitors capable of determining perfusion index measurements.

Description of the Related Art

[0002] Oximeter systems providing measurements of a monitored patient have become the standard of care in many patient care settings, including surgical, post surgical, neonatal, general ward, home care, physical training, and the like. In general, oximeter systems accept one or more noninvasive signals from an optical sensor or probe capable of emitting multiple wavelengths of light into a tissue site and capable of detecting light after attenuated by the tissue site. The optical sensor generally outputs intensity signal data. Fig. 1 illustrates a photoplethysmograph intensity signal 100 output by an oximeter sensor. An oximeter does not directly detect absorption, and hence does not directly measure a standard plethysmograph waveform. However, the standard plethysmograph can be derived by observing that the detected intensity signal 100 is merely an out of phase version of an absorption profile known to one of skill in the art. That is, the peak detected intensity 102, generally corresponds to a minimum absorption, and minimum detected intensity 104, generally corresponds to a maximum absorption. Further, a rapid rise in absorption during an inflow phase of the plethysmograph is reflected in a rapid decline 106 in intensity. Likewise, a gradual decline in absorption during the outflow phase of the plethysmograph is reflected in a gradual increase 108 in detected intensity.

[0003] Fig. 2 illustrates a flow calculator 200 which receives a processed signal 202 responsive to at least one of the intensity signals output from the sensor. In an embodiment, the flow calculator outputs an indication of blood flow, such as, for example, a perfusion index (PI) 204. In an embodiment, the PI 204 comprises a relative indication of pulse strength at a monitoring site. For example, the PI 204 may be defined as the ratio of the wavelength's (A) AC signal to the DC signal, or the percentage of pulsatile signal to non-pulsatile signal, according to the following:

$$PI = (\lambda_{\max} - \lambda_{\min})/\lambda_{DC}$$

where λ_{\max} is the maximum value, λ_{\min} is the minimum value, and λ_{DC} is the average value of the signal 202.

[0004] Once calculated, the PI 204 may advantageously

be displayed in a wide number of ways, including rising LEDs or other display elements, text, graphics, or other visual elements including color, flashing, and the like, trended data, trace data, or the like. Fig. 3A illustrates a display output for an oximeter patient monitor 302 including a textual PI display 304 (shown as "3.25 PI") ranging from 1.0 to 20. Fig. 3B illustrates a display output for a handheld oximeter patient monitor 322 including a PI bar 324 ranging from, for example, "<.1%" to ">5%" with steps of "<.1%", ".25%", ".5%", "1%", "1.25%", "1.5%", "1.75%", "2%", "3%", and ">5%." An artisan will recognize from the disclosure herein that various steps and a wide variety of scalars or other mathematical mapping can be used to make PI numbers more readily understandable to a caregiver. However, using the foregoing scale, the PI bar 324 can be used as a diagnostic tool during low perfusion for the accurate prediction of illness severity, especially in neonates. Moreover, the rate of change in the PI bar 324 can be indicative of blood loss, sleep arousal, severe hypertension, pain management, the presence or absence of drugs, or the like. According to one embodiment, a measurement below about "1.25%" may indicate medical situations in need of caregiver attention, specifically in monitored neonates. Because of the relevance of about "1.25%", the PI bar 324 may advantageously include level indicia that switch sides of the PI bar 324, thus highlighting any readings below about that threshold. Moreover, behavior of the PI bar 324, as discussed below, may advantageously draw attention to monitored values below such a threshold.

[0005] The PI bar 324 may advantageously activate LEDs from a bottom toward a top such that the bar "fills" to a level proportional to the measured value. In one embodiment, the PI bar 324 shows a static value of perfusion for a given time period, such as, for example, one or more pulses. In another embodiment, or functional setting, the PI bar 324 may advantageously pulse with a pulse rate, may hold the last reading and optionally fade until the next reading, may indicate historical readings through colors or fades, traces or the like. Additionally, the PI bar 324 may advantageously change colors, flash, increasingly flash, or the like to indicate worsening measured values of perfusion.

[0006] As discussed above, the monitors 302, 322 may include output functionality that outputs, for example, trend perfusion data, such that a caregiver can monitor measured values of perfusion over time. Alternatively or additionally, the monitors 302, 322 may display historical trace data on an appropriate display indicating the measured values of perfusion over time. In an embodiment, the trend data is uploaded to an external computing device through, for example, a multipurpose sensor connector or other input output systems such as USB, serial or parallel ports, computing networks, or the like. Additional information regarding the calculation and use of PI is disclosed in the '065 patent, assigned to Masimo Corporation ("Masimo") of Irvine, California.

[0007] WO 2004/034898 A2 discloses a method for

the presentation of information concerning variations of the arterial filling with blood (perfusion) of organs of living beings on the user surface of a display screen. In said method the data required for the presentation (perfusion index) is derived, using an algorithm, from measuring values of a non-invasive photometric measuring process for determining the arterial oxygen saturation of the blood. A first perfusion index is defined as a reference value and the subsequent perfusion indices are determined as relative deviations with respect to the reference value, said relative deviations being presented in the form of analog, graphic elements on the user surface as information concerning the variations of the perfusion.

SUMMARY OF THE DISCLOSURE

[0008] Aspects of the present disclosure include a system which provides a more accurate indication of PI. In one embodiment, the wavelength used to determine PI is in the infrared (IR) spectrum. IR wavelengths are more independent of saturation than are wavelengths in the Red (R) spectrum. As a result, when the saturation changes, the PI calculated from a R wavelength tends to vary greater than the PI calculated from a IR wavelength. Although the IR intensity signal data is less responsive to saturation than various other intensity channels, the IR intensity signal remains somewhat susceptible to motion-induced noise, distortion and the like. In general, PI measurements during lower signal quality or distortion generally trends the PI too high. As understood by an artisan from the disclosure herein, higher PI measurements should correlate to better perfusion through the tissue site, and lower PI measurements should correlate to lower perfusion through the tissue site. Thus, noise may cause a patient monitor to give a caregiver the indication that a patient has better perfusion through a measurement site than is actually the case.

[0009] Based at least thereon, embodiments of the present disclosure seek to reduce an error between a calculated measurement of PI and actual conditions. In an embodiment, a baseline perfusion index ("baseline PI") is determined and used to improve measurements during motion conditions. For example, a smoother may advantageously determine a baseline PI during strong signal quality, high signal confidence conditions, and use that baseline PI or a statistical combination of baseline PIs instead of or in addition to the current PI that is being influenced by noise. In an embodiment, the smoother selects the lower PI value as the output PI value. This is because motion noise tends to increase PI, thus the lower PI value tends to be the more accurate value.

[0010] In an embodiment, PI is calculated using at least two different calculation techniques. The two different calculations are analyzed and a PI indication is determined. In an embodiment, the lower of the two PI values is chosen. In an embodiment, the PI values are averaged or statistically analyzed to determine a more accurate value for PI.

[0011] The present invention relates to a method of smoothing a perfusion index measurement according to claim 1 and to an oximeter according to claim 3. The present disclosure further discloses the following methods and oximeter which do not fall within the scope of the claims.

[0012] A method of determining an indication of perfusion index in a patient monitor is disclosed. The method includes the steps of receiving plethysmographic data, calculating two or more indications of perfusion index using at least two different calculation techniques, and determining a final indication of perfusion index based on the indications of perfusion index. In an example, the final indication of perfusion index is determined by selecting the lowest indication of perfusion index. In an example, the final indication of perfusion index is determined by statistical analysis. In an example, the final indication of perfusion index is determined by averaging the indications of perfusion index. perfusion index is disclosed. The method includes the steps of receiving plethysmographic data and determining an indication of perfusion index by utilizing at least one calculation techniques to determine a resulting indication of perfusion index. In an example, determining the indication of perfusion index includes utilizing the calculation technique that will result in the lowest perfusion index value.

[0013] A disclosed oximeter includes an input capable of receiving intensity signal data responsive to intensity signals acquired from a detector capable of detecting light attenuated by body tissue, a first calculator configured to calculate a first indication of perfusion index using a first calculation technique, a second calculator configured to calculate a second indication of perfusion index using a second calculation technique and a processor configured to utilize at least one of the first and second calculators to determine a resulting indication of perfusion index. In an example, the processor is configured to select the calculator which calculates the lowest indication of perfusion index. In an example, the processor utilizes both calculation techniques. In an example, the processor is further configured to select the lowest indication of perfusion index as the resulting indication of perfusion index.

[0014] A method of determining an indication of a physiological condition using data responsive to intensity signals acquired from a detector capable of detecting light attenuated by body tissue is disclosed, The method includes the steps of determining one or more indications of pulse information from data responsive to intensity signals acquired from a detector capable of detecting light attenuated by body tissue, determining a first indication of amplitude data for a single pulse based on the one or more indications of pulse information, determining a second indication of amplitude data for a plurality of pulses based on the one or more indications of pulse information, determining a final indication of amplitude data based on the first and second indications and outputting the final indication. In an example, the amplitude data is

used to determine an indication of perfusion. In an embodiment, determining a final indication includes selecting the lowest indication of amplitude from the first and second indications of amplitude. In an example, determining a final indication includes averaging the first and second indications of amplitude. In an embodiment, determining a final indication is based on a statistical analysis of the first and second indications of amplitude. In an example, the combination includes an averaged combination of pulses. In an example, the combination includes a maximum and minimum amplitude of a plurality of pulses. In an example, the combination includes a statistical combination of pulses.

[0015] For purposes of summarizing the disclosure, certain aspects, advantages and novel features of the disclosure have been described herein. Of course, it is to be understood that not necessarily all such aspects, advantages or features will be embodied in any particular embodiment of the disclosure.

BRIEF DESCRIPTION OF THE DRAWINGS

[0016] A general architecture that implements the various features of the disclosure will now be described with reference to the drawings. The drawings and the associated descriptions are provided to illustrate embodiments of the disclosure and not to limit its scope.

Fig. 1 illustrates a graph showing an intensity "plethysmograph" oximetry waveform.

Fig. 2 illustrates a flow calculator capable of determining measurements for a perfusion index.

Figs. 3A - 3B illustrate patient monitors capable of calculating and displaying the measurements of **Fig. 2**.

Fig. 4 illustrates a simplified block diagram of a smoothed perfusion index generator, according to an embodiment of the disclosure.

Fig. 5 illustrates a flow chart of a smoothing process, according to an embodiment of the disclosure.

Fig. 6 illustrates another embodiment of a simplified block diagram of a smoothed perfusion index generator.

Fig. 7 illustrates a simplified block diagram of an embodiment of a PI determination system using multiple PI calculation techniques.

Fig. 8 illustrates a flow chart of a PI decision process, according to another embodiment of the disclosure.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

[0017] **Fig. 4** illustrates a simplified block diagram of a smoothed perfusion index generator 400, according to an embodiment of the disclosure. As shown, the generator 400 includes a flow calculator 402, a smoother 404 and optional indications of noise, including a measure of distortion 406, signal quality 408, and waveform quality

410. In an embodiment, a data signal 420 responsive to an intensity signal is input into the flow calculator 402, and a current value 422 of PI is calculated. A skilled artisan will recognize a number of calculations that can determine values of perfusion, including, for example, the foregoing ratio of the AC signal to the DC signal. The current value 422 of PI, which in an embodiment is subject to noise and distortion in the data signal 420, is input into the smoother 404, which reduces an error between the current value 422 of PI and actual perfusion conditions. For example, the smoother 404 determines a baseline PI, and depending upon an indication of some or all of an amount of distortion, noise, signal quality, and/or waveform quality in the data signal 422, substitute or combine the baseline PI for or with the current value 422 to generate an output PI 430.

[0018] In an embodiment, the wavelength used to determine PI is in the infrared (IR) spectrum. IR wavelengths are more independent of saturation than are wavelengths in, for example, the Red (R) spectrum. As a result, when the saturation changes, the PI calculated from a R wavelength tends to vary greater than the PI calculated from a IR wavelength.

[0019] In an embodiment, the distortion signal 406 may comprise a Boolean value indicating whether the data signal 422 includes, for example, motion-induced noise. Although an artisan will recognize from the disclosure herein a number of methodologies for deriving the distortion signal 406, derivation of a Boolean distortion signal is disclosed in U.S. Pat. No. 6,606,511. Alternatively, or in addition to, the signal quality signal 408 may comprise a Boolean value indicating whether the data signal 422 meets various waveform criteria. Although an artisan will recognize from the disclosure herein a number of methodologies for deriving the signal quality signal 408, derivation of a Boolean distortion signal is disclosed in the '511 patent. Alternatively, or in addition to, a feature extractor 440 may advantageously produce waveform quality outputs 410, indicative of waveform quality or waveform shape. Although an artisan will recognize from the disclosure herein a number of methodologies for deriving the waveform quality signal 410, derivation thereof is disclosed in the '065 patent.

[0020] Thus, the smoother 404 accepts one or more different indicators of the quality of the data signal 420, and determines how to smooth the PI output to reduce errors between data trends and actual perfusion conditions. In an embodiment, the smoothing may advantageously comprise statistical weighting, other statistical combinations, or simply passing the PI 422 through to the output, depending upon one or more of the quality signals 406, 408, 410, or logical combinations thereof. In an embodiment, the smoother 404 selects the lowest PI value to pass to the output PI 422. This is because motion noise tends to increase PI, thus the lower PI value tends to be the more accurate value.

[0021] Upon the output of a smoothed PI measurement, a monitor may advantageously audibly and/or vis-

ually presents the measurement to a caregiver, and when the measurement meets certain defined thresholds or behaviors, the monitor may advantageously audibly and/or visually alert the caregiver. In other embodiments, the monitor may communicate with other computing devices to alert the caregiver, may compare longer term trend data before alarming, or the like.

[0022] Fig. 5 illustrates a flow chart of a smoothing process 500, according to an embodiment of the disclosure. In block 502, data is acquired responsive to one or more of multiple channel intensity signals. In block 504, a determination is made whether distortion is present, whether signal quality is good, whether there is motion-induced-noise or the like. When acceptable noise levels exist, in block 506, the baseline PI becomes the current PI and in block 508, the baseline PI is output. However, when unacceptable signal quality or the like is detected, the current PI is compared to the baseline PI. When the current PI is smaller than the baseline PI, in block 506, the baseline PI becomes the current PI. However, when the current PI is greater than or equal to the baseline PI, the baseline PI is output in block 510. Thus, the smoothing process 500 determines when a baseline PI can be acquired in quality monitoring conditions, and then uses that baseline PI to avoid drift or other errors that may deteriorate the accuracy of the PI output. An artisan will recognize from the disclosure herein that the baseline PI could be reset periodically, some or all iterations, when other occurrences or measurements suggest a reinitialization, when various trends occur, or the like.

[0023] Fig. 6 illustrates another embodiment of a simplified block diagram of a smoothed perfusion index generator 600. As shown, the generator 600 includes a smoother 604 and optional indications of noise, including a measure of distortion 606, signal quality 608, waveform quality 610, and Pleth Feature Extractor similar to those described with respect to Fig. 4. In an embodiment, a data signal 620 responsive to an intensity signal is input into the PI Analyzer 602, and a PI value 422 is outputted for use by the smoother 604. In an embodiment, the PI Analyzer employs multiple different calculation techniques to determine a more accurate indication of PI. In an embodiment, based on the indications of distortion signal quality, waveform quality and/or other indications of signal quality, a specific calculation technique is used to determine a PI output value 622.

[0024] Fig. 7 illustrates a simplified block diagram of an embodiment of a PI determination system 700 using multiple PI calculation techniques. As shown, pleth data 720 is input into the system. The pleth data 720 is then routed to at least two different PI calculators 702, 704. In an embodiment, more than two different types of calculation techniques can be used. The at least two PI calculators 702, 704 output PI indications for input into the PI selector 706. The PI selector 706 determines a PI value to output. In an embodiment, the PI selector chooses the lowest PI value. In general, the lower PI value should be the more accurate value, particularly in the presence

of motion induced noise. This is because motion induced noise tends to raise PI values. In an embodiment, the PI selector 706 averages or otherwise analyzes the PI values using statistical modeling in order to determine a more accurate value of PI.

[0025] In an embodiment, one of the PI calculators 702, 704 determines a PI value based on pulse by pulse determination of PI, such as, for example, using the following PI formula as described above:

$$PI = (\lambda_{\max} - \lambda_{\min})/\lambda_{DC}$$

[0026] In an embodiment, one of the PI calculators 702, 704 determines a PI value based on a fixed or variable interval of pleth data including more than one pulse, in effect calculating a bulk PI.

[0027] Fig. 8 illustrates a flow chart of a PI decision process 800, according to another embodiment of the disclosure. At block 802, the process 800 receives pleth data 802. The process then moves to block 804 where the process 800 calculates at least two different PI indications, such as, for example, as described with respect to Fig. 7. The process 800 then moves to block 806 where an indication of PI is determined based on the different calculations, again, such as, for example, as described with respect to Fig. 7. The process 800 then repeats itself for each PI value determination.

[0028] While the above perfusion index determination system has been described in certain embodiments herein, other embodiments of the present disclosure will be known to those of skill in the art from the descriptions herein. Moreover, the described embodiments have been presented by way of example only, and are not intended to limit the scope of the disclosure. Moreover, those of skill in the art understand that information and signals can be represented using a variety of different technologies and techniques. For example, data, instructions, commands, information, signals, bits, symbols, and chips that can be referenced throughout the above description may be represented by voltages, currents, electromagnetic waves, magnetic fields or particles, optical fields or particles, or any combination thereof.

[0029] Those of skill in the art further appreciate that the various illustrative logical blocks, modules, circuits, and algorithm steps described in connection with the embodiments disclosed herein can be implemented as electronic hardware, computer software, or combinations of both. To illustrate this interchangeability of hardware and software, various illustrative components, blocks, modules, circuits, and steps have been described above generally in terms of their functionality. Whether such functionality is implemented as hardware or software depends upon the particular application and design constraints imposed on the overall system. Skilled artisans can implement the described functionality in varying ways for each particular application, but such implemen-

tation decisions should not be interpreted as causing a departure from the scope of the present disclosure.

[0030] The various illustrative logical blocks, modules, and circuits described in connection with the embodiments disclosed herein can be implemented or performed with a general purpose processor, a digital signal processor (DSP), an application specific integrated circuit (ASIC), a field programmable gate array (FPGA) or other programmable logic device, discrete gate or transistor logic, discrete hardware components, or any combination thereof designed to perform the functions described herein. A general purpose processor can be a microprocessor, but in the alternative, the processor can be any conventional processor, controller, microcontroller, or state machine. A processor can also be implemented as a combination of computing devices, e.g., a combination of a DSP and a microprocessor, a plurality of microprocessors, one or more microprocessors in conjunction with a DSP core, or any other such configuration.

[0031] The steps of a method or algorithm described in connection with the embodiments disclosed herein can be embodied directly in hardware, in a software module executed by a processor, or in a combination of the two. A software module can reside in RAM memory, flash memory, ROM memory, EPROM memory, EEPROM memory, registers, hard disk, a removable disk, a CD-ROM, or other form of storage medium known in the art. A storage medium is coupled to the processor, such that the processor can read information from, and write information to, the storage medium. In the alternative, the storage medium can be integral to the processor. The processor and the storage medium can reside in an ASIC. The ASIC can reside in a user terminal, physiological monitor and/or sensor. The processor and the storage medium can reside as discrete components in a user terminal, physiological monitor and/or sensor.

[0032] Although the foregoing disclosure has been described in terms of certain preferred embodiments, other embodiments will be apparent to those of ordinary skill in the art from the disclosure herein. Additionally, other combinations, omissions, substitutions and modifications will be apparent to the skilled artisan in view of the disclosure herein. Moreover, it is contemplated that various aspects and features of the disclosure described can be practiced separately, combined together, or substituted for one another, and that a variety of combination and subcombinations of the features and aspects can be made and still fall within the scope of the disclosure. Furthermore, the systems described above need not include all of the modules and functions described in the preferred embodiments.

Claims

1. A method of smoothing a perfusion index measurement, said method being performed by an oximeter system and comprising the following steps:

receiving intensity signal data responsive to intensity signals acquired from a detector capable of detecting light attenuated by body tissue; calculating a current value of perfusion index and inputting said current value into a smoother module, said smoother module further accepting one or more indicators of the quality of the intensity signal data and performing the following steps:

when said one or more indicators indicate a good quality of the received intensity signal data, a baseline perfusion index becomes the current value of perfusion index and the baseline perfusion index is output, and

when said one or more indicators indicate a poor quality of the received intensity signal data, the current value of the perfusion index is compared to the baseline perfusion index, and

when said current value of perfusion index is smaller than the baseline perfusion index, the baseline perfusion index becomes the current value of perfusion index and the baseline perfusion index is output, and

when said current value of perfusion index is greater than or equal to the baseline perfusion index, the baseline perfusion index is output.

2. The method of Claim 1, wherein the signal is responsive to an IR intensity signal.
3. An oximeter comprising:

a flow calculator (402) configured to receive intensity signal data responsive to intensity signals acquired from a detector capable of detecting light attenuated by body tissue, and configured to calculate a current perfusion index value; a memory storing a baseline perfusion index; a smoother module (404) including a processor configured to accept one or more indicators of the quality of the intensity signal data and to perform the following steps:

when said one or more indicators indicate a good quality of the received intensity signal data, the baseline perfusion index becomes the current perfusion index value and the baseline perfusion index is output, and

when said one or more indicators indicate a poor quality of the received intensity signal data, the current perfusion index value is

compared to the baseline perfusion index, and

when said current perfusion index value is smaller than the baseline perfusion index, the baseline perfusion index becomes the current perfusion index value and the baseline perfusion index is output,
when said current perfusion index value is greater than or equal to the baseline perfusion index, the baseline perfusion index is output.

4. The oximeter of Claim 3, further comprising a monitor configured to alert a caregiver when said output meets defined thresholds or behaviors.

Patentansprüche

1. Verfahren zum Glätten einer Perfusionsindexmessung, wobei das Verfahren durch ein Oximetersystem ausgeführt wird und folgende Schritte aufweist:

Empfangen von Intensitätssignaldaten ansprechend auf von einem Detektor, der in der Lage ist, von Körpergewebe abgeschwächtes Licht zu detektieren, erfasste Intensitätssignale, Berechnen eines aktuellen Werts des Perfusionsindex und Eingeben des aktuellen Werts in ein Glättungsmodul, wobei das Glättungsmodul ferner einen oder mehrere Indikatoren zur Qualität der Intensitätssignaldaten akzeptiert und folgende Schritte ausführt:

wenn der eine oder die mehreren Indikatoren eine gute Qualität der empfangenen Intensitätssignaldaten angeben, wird ein Grundlinien-Perfusionsindex zum aktuellen Wert des Perfusionsindex und wird der Grundlinien-Perfusionsindex ausgegeben, und
wenn der eine oder die mehreren Indikatoren eine schlechte Qualität der empfangenen Intensitätssignaldaten angeben, wird der aktuelle Wert des Perfusionsindex mit dem Grundlinien-Perfusionsindex verglichen, und

wenn der aktuelle Wert des Perfusionsindex kleiner als der Grundlinien-Perfusionsindex ist, wird der Grundlinien-Perfusionsindex zum aktuellen Wert des Perfusionsindex und wird der Grundlinien-Perfusionsindex ausgegeben, und
wenn der aktuelle Wert des Perfusionsindex größer oder gleich dem Grundlinien-Perfusionsindex ist, wird der Grundlinien-Perfusionsindex ausgegeben.

2. Verfahren nach Anspruch 1, wobei das Signal auf ein IR-Intensitätssignal anspricht.

3. Oximeter, welches Folgendes aufweist:

eine Flussberechnungseinrichtung (402), die dafür ausgelegt ist, Intensitätssignaldaten ansprechend auf von einem Detektor, der in der Lage ist, von Körpergewebe abgeschwächtes Licht zu detektieren, erfasste Intensitätssignale zu empfangen, und dafür ausgelegt ist, einen aktuellen Perfusionsindexwert zu berechnen, einen Speicher, der einen Grundlinien-Perfusionsindex speichert,
ein Glättungsmodul (404), das einen Prozessor aufweist, der dafür ausgelegt ist, einen oder mehrere Indikatoren zur Qualität der Intensitätssignaldaten zu akzeptieren und die folgenden Schritte auszuführen:

wenn der eine oder die mehreren Indikatoren eine gute Qualität der empfangenen Intensitätssignaldaten angeben, wird der Grundlinien-Perfusionsindex zum aktuellen Perfusionsindexwert und wird der Grundlinien-Perfusionsindex ausgegeben, und
wenn der eine oder die mehreren Indikatoren eine schlechte Qualität der empfangenen Intensitätssignaldaten angeben, wird der aktuelle Perfusionsindexwert mit dem Grundlinien-Perfusionsindex verglichen, und
wenn der aktuelle Wert des Perfusionsindex kleiner als der Grundlinien-Perfusionsindex ist, wird der Grundlinien-Perfusionsindex zum aktuellen Wert des Perfusionsindex und wird der Grundlinien-Perfusionsindex ausgegeben,
wenn der aktuelle Wert des Perfusionsindex größer oder gleich dem Grundlinien-Perfusionsindex ist, wird der Grundlinien-Perfusionsindex ausgegeben.

4. Oximeter nach Anspruch 3, welches ferner eine Überwachungseinrichtung aufweist, die dafür ausgelegt ist, einen Pflegedienstleister zu warnen, wenn die Ausgabe definierte Schwellenwerte oder Verhaltensweisen erfüllt.

Revendications

1. Procédé de lissage d'une mesure d'indice de perfusion, ledit procédé étant effectué par un système d'oxymètre et comprenant les étapes suivantes :

réception de données de signaux d'intensité en réponse à des signaux d'intensité acquis à partir

d'un détecteur capable de détecter une lumière atténuée par un tissu corporel ;
calcul d'une valeur actuelle d'un indice de perfusion et entrée de ladite valeur actuelle dans un module de lissage, ledit module de lissage acceptant en outre un ou plusieurs indicateurs de la qualité des données de signaux d'intensité et effectuant les étapes suivantes :

lorsque ledit un ou plusieurs indicateurs indiquent une bonne qualité des données de signaux d'intensité reçus, un indice de perfusion de base devient la valeur actuelle de l'indice de perfusion et l'indice de perfusion de base est émis, et

lorsque ledit un ou plusieurs indicateurs indiquent une qualité médiocre des données de signaux d'intensité reçus, la valeur actuelle de l'indice de perfusion est comparée à l'indice de perfusion de base, et

lorsque ladite valeur actuelle de l'indice de perfusion est inférieure à l'indice de perfusion de base, l'indice de perfusion de base devient la valeur actuelle de l'indice de perfusion et l'indice de perfusion de base est émis, et

lorsque ladite valeur actuelle de l'indice de perfusion est supérieure ou égale à l'indice de perfusion de base, l'indice de perfusion de base est émis.

2. Procédé selon la revendication 1, dans lequel le signal est envoyé en réponse à un signal d'intensité IR.

3. Oxymètre comprenant :

un calculateur de débit (402) conçu pour recevoir des données de signaux d'intensité en réponse à des signaux d'intensité acquis à partir d'un détecteur capable de détecter une lumière atténuée par un tissu corporel, et conçu pour calculer une valeur actuelle de l'indice de perfusion ;

une mémoire stockant un indice de perfusion de base ;

un module de lissage (404) comprenant un processeur conçu pour accepter ou un plusieurs indicateurs de la qualité des données de signaux d'intensité et pour effectuer les étapes suivantes :

lorsque ledit un ou plusieurs indicateurs indiquent une bonne qualité des données de signaux d'intensité reçus, l'indice de perfusion de base devient la valeur actuelle de l'indice de perfusion et l'indice de perfusion de base est émis, et

lorsque ledit un ou plusieurs indicateurs in-

diquent une médiocre qualité des données de signaux d'intensité reçus, la valeur actuelle de l'indice de perfusion est comparée à l'indice de perfusion de base, et

lorsque ladite valeur actuelle de l'indice de perfusion est inférieure à l'indice de perfusion de base, l'indice de perfusion de base devient la valeur actuelle de l'indice de perfusion et l'indice de perfusion de base est émis, et

lorsque ladite valeur actuelle de l'indice de perfusion est supérieure ou égale à l'indice de perfusion de base, l'indice de perfusion de base est émis.

4. Oxymètre selon la revendication 3, comprenant en outre un moniteur conçu pour alerter un soignant lorsque ladite sortie atteint des seuils ou des comportements définis.

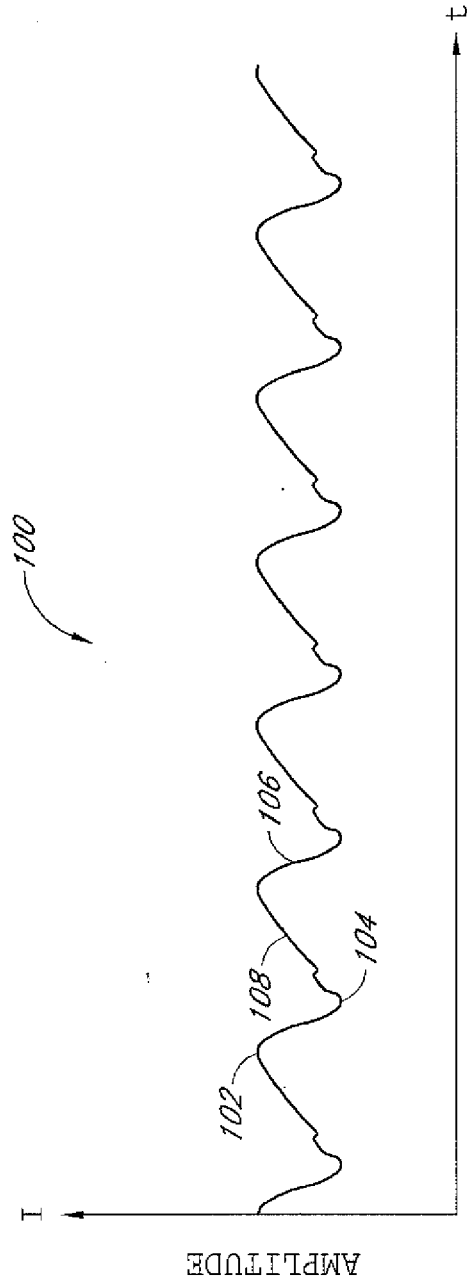


FIG. 1

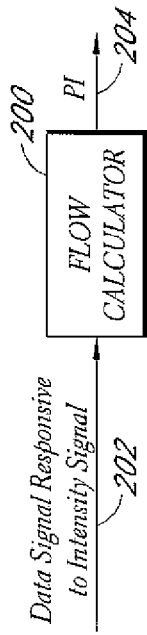


FIG. 2

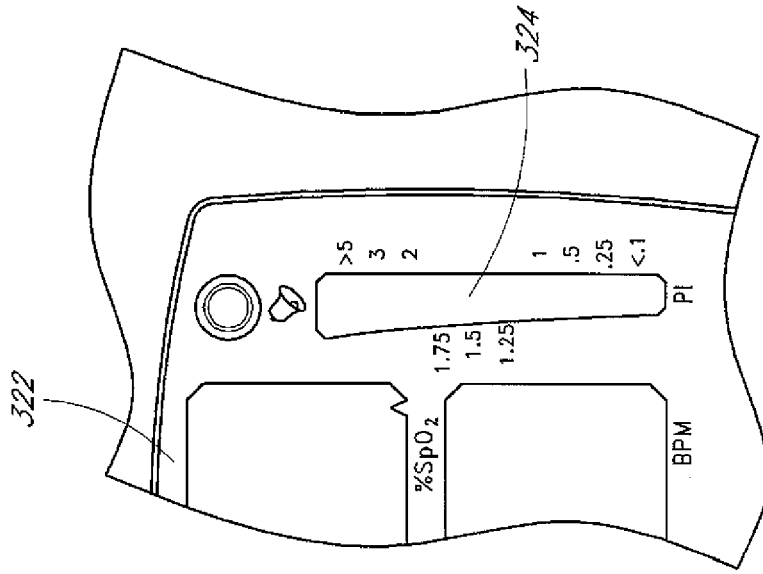


FIG. 3B

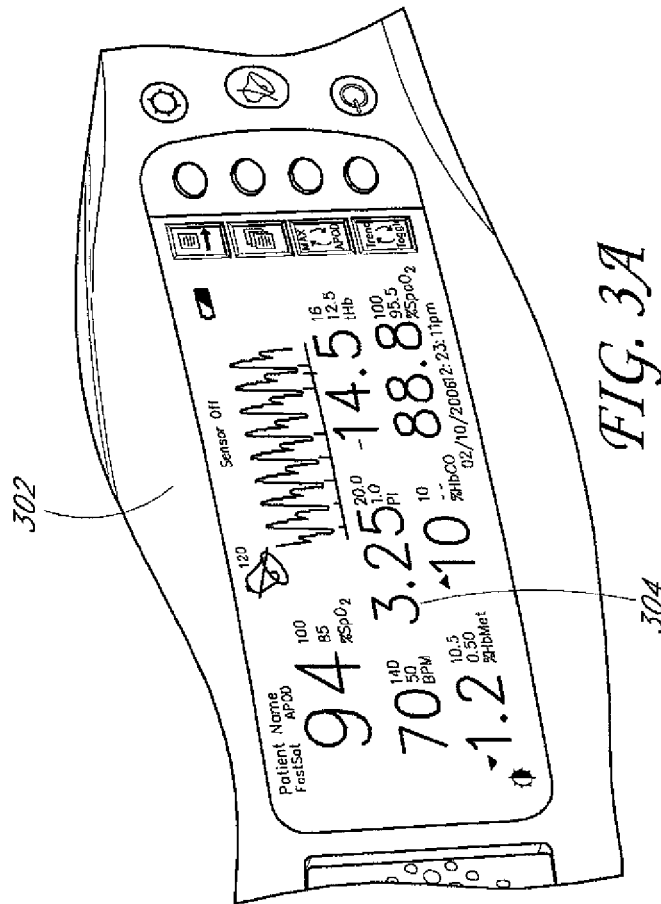


FIG. 3A

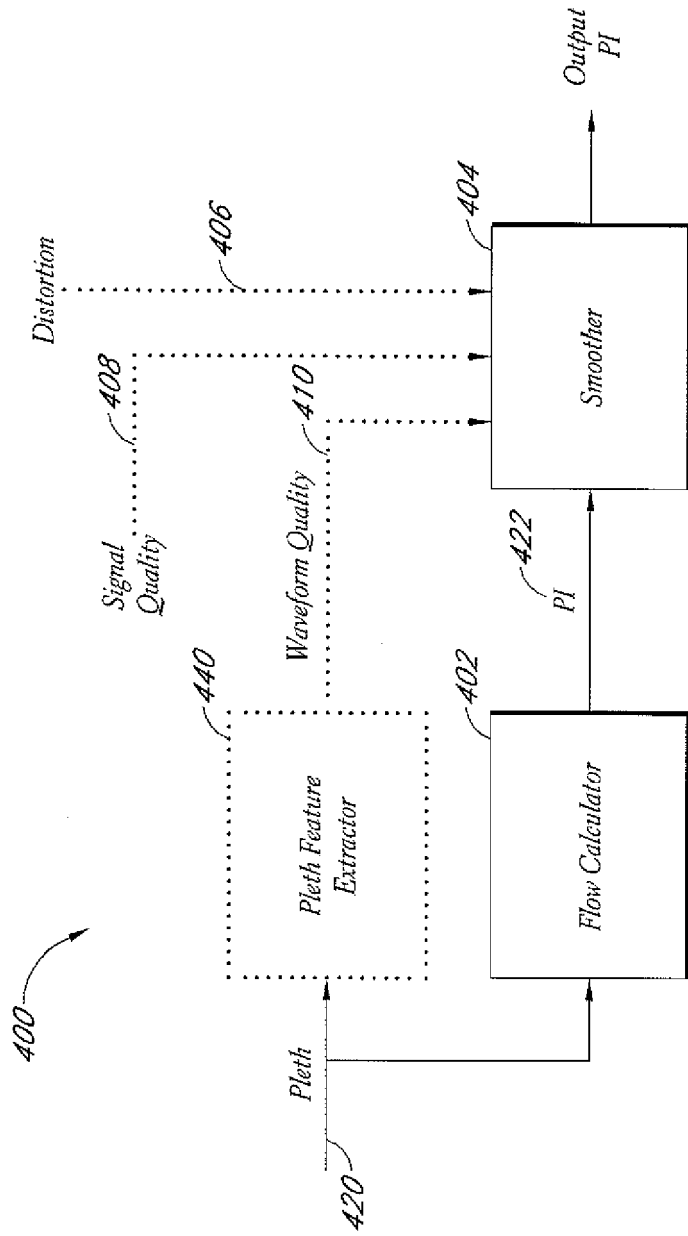


FIG. 4

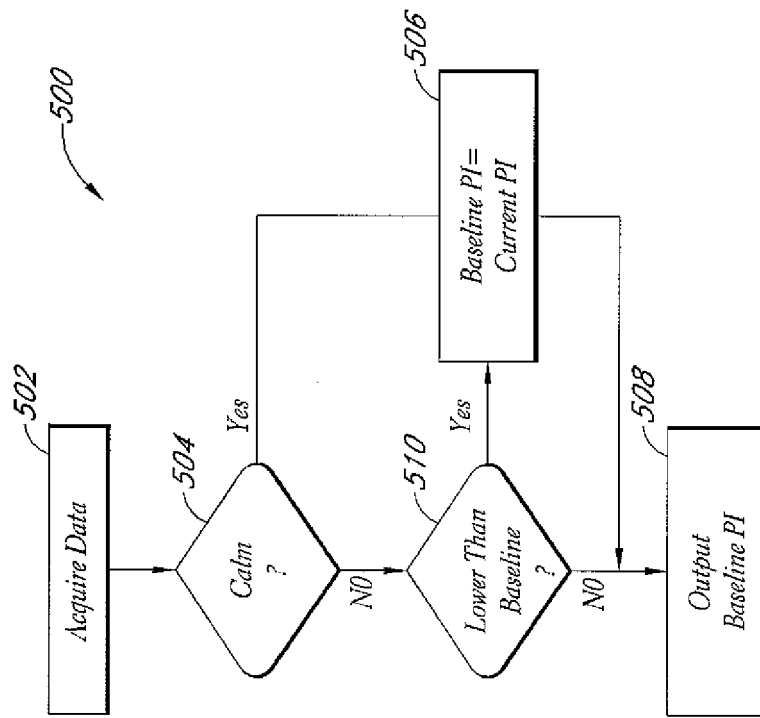


FIG. 5

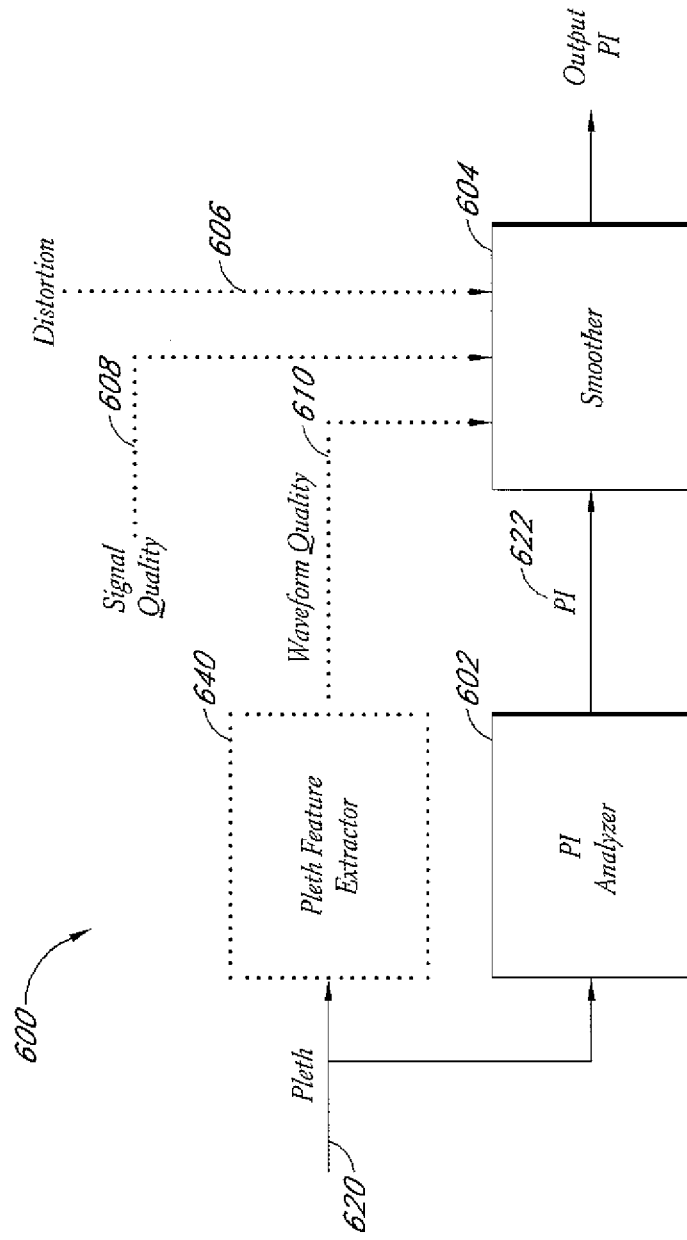


FIG. 6

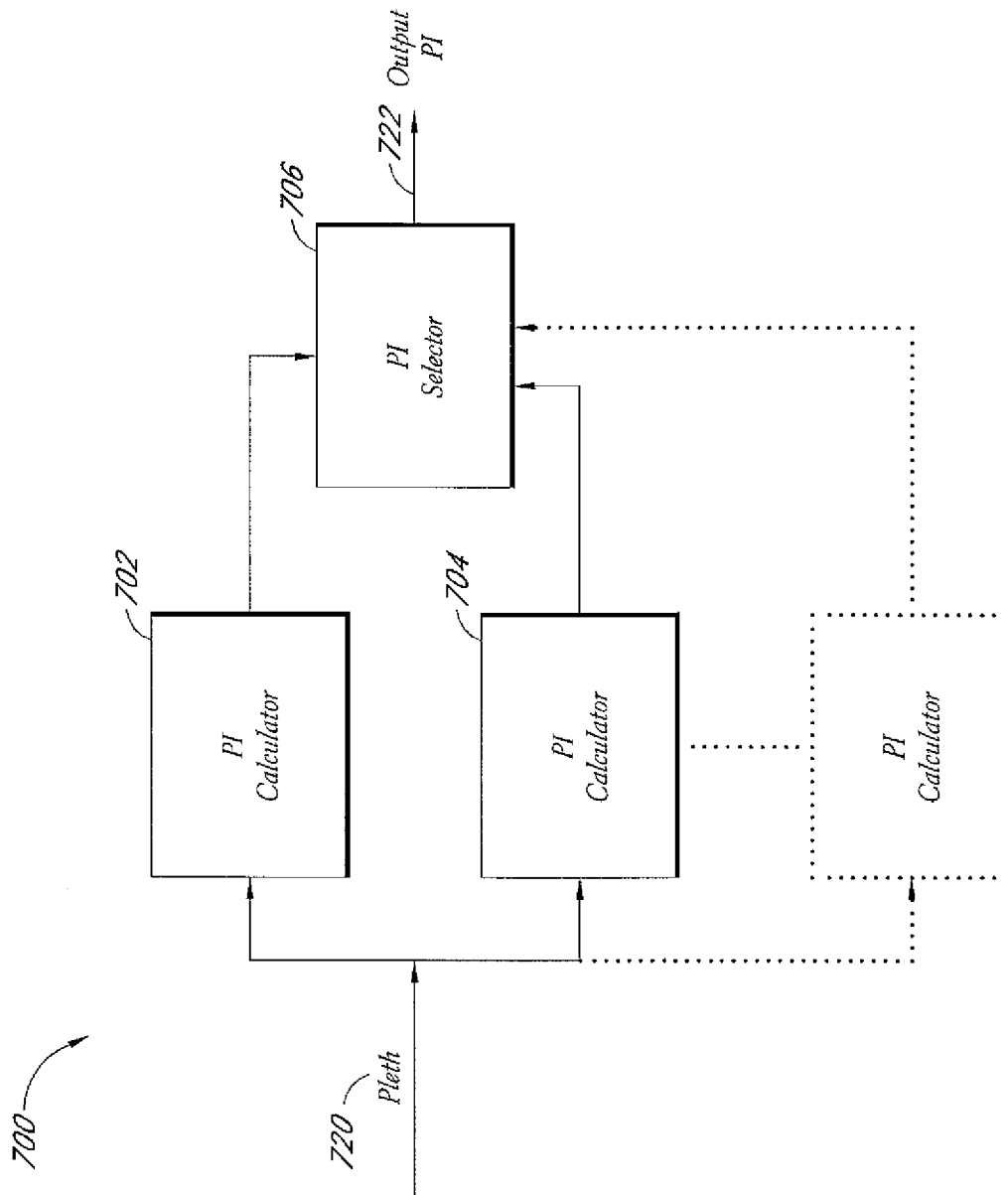


FIG. 7

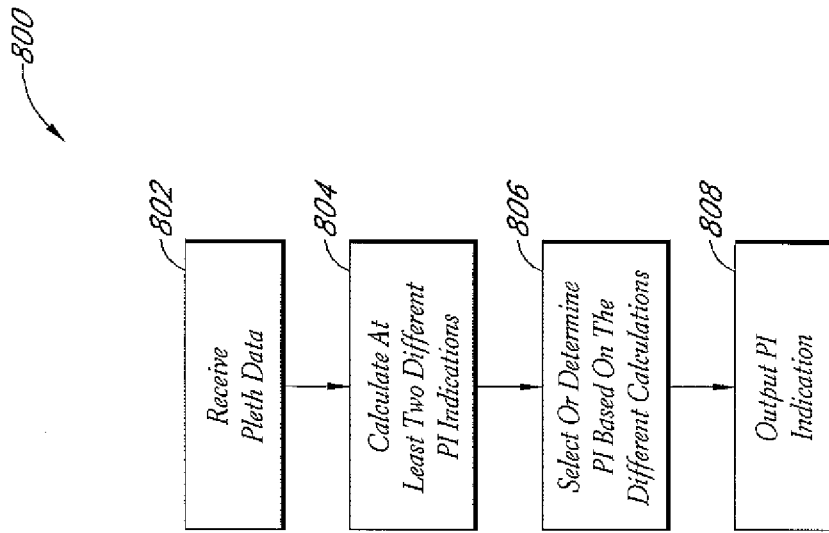


FIG. 8

REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- WO 2004034898 A2 [0007]
- US 6606511 B [0019]

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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		
发明人	AL-ALI, AMMAR		
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摘要(译)

本公开的实施例寻求通过使用基线灌注指数测量和/或通过多个PI计算来平滑灌注指数测量。基线灌注指数测量的组合减少了计算的PI测量值与实际条件之间的误差。

$$PI = \frac{PI_{max} - PI_{min}}{AOC}$$