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(54) **PHYSIOLOGICAL SENSOR COMBINATION**

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(52) U.S. Cl. .... **600/324**; 600/344; 600/372; 600/483

(58) Field of Search ..... 600/310, 322, 600/323, 324, 340, 344, 372, 383, 384, 393, 481, 483

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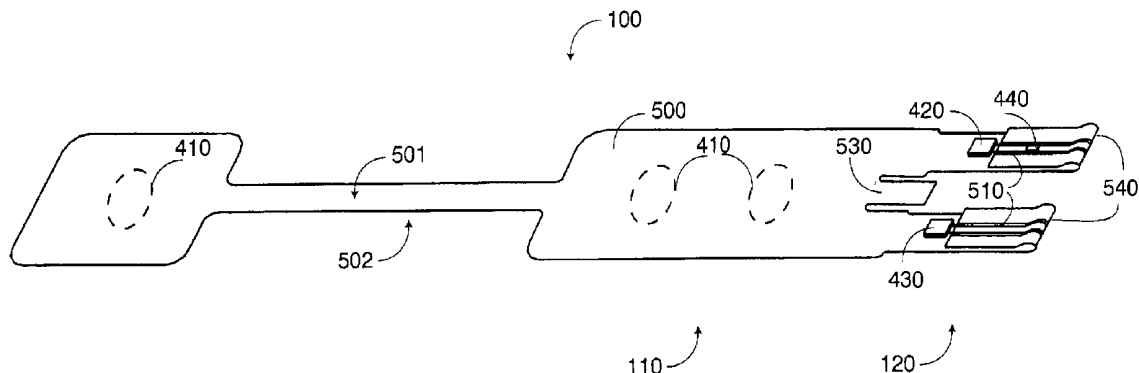
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(57) **ABSTRACT**

A physiological sensor combination has a flexible substrate configured to attach to a tissue site. Multiple sensors are disposed on the substrate, which generate physiological signals. Each of the signals is responsive to a different physiological parameter. Conductors are carried on the substrate and routed between the sensors and at least one connector. The connector is configured to communicate the physiological signals to at least one monitor, which derives measurements of the parameters.

**10 Claims, 7 Drawing Sheets**



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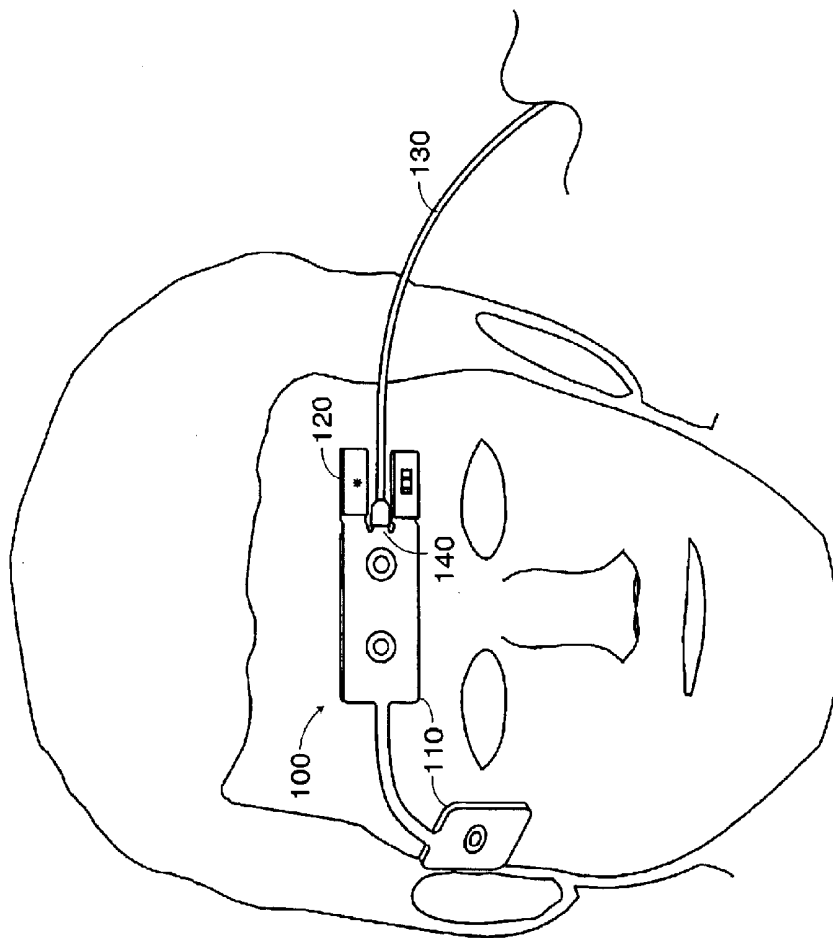


FIG. 1

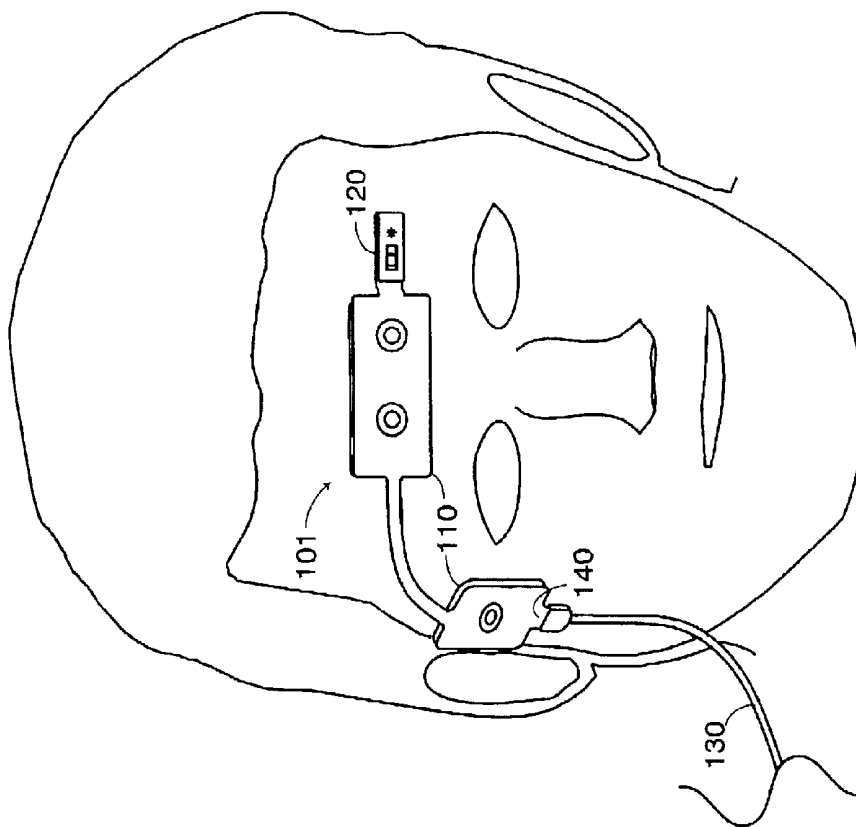


FIG. 2

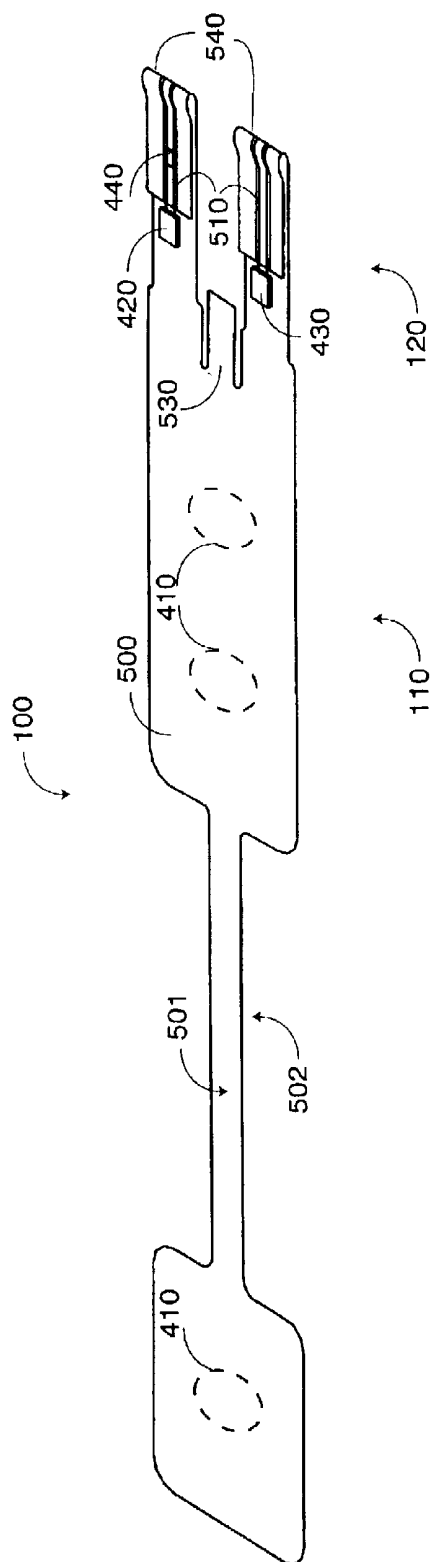


FIG. 3A

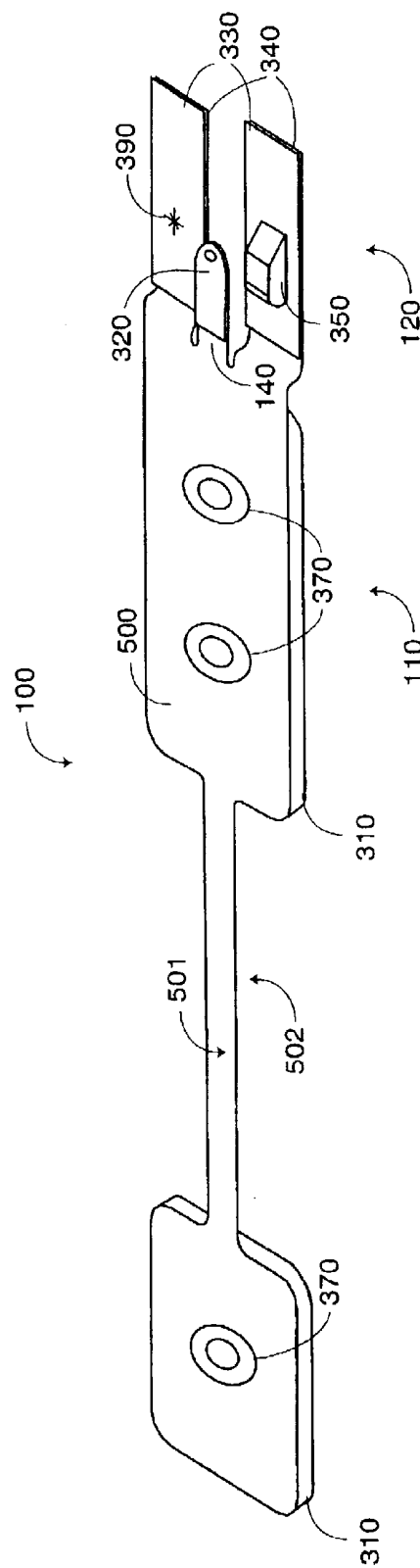


FIG. 3B

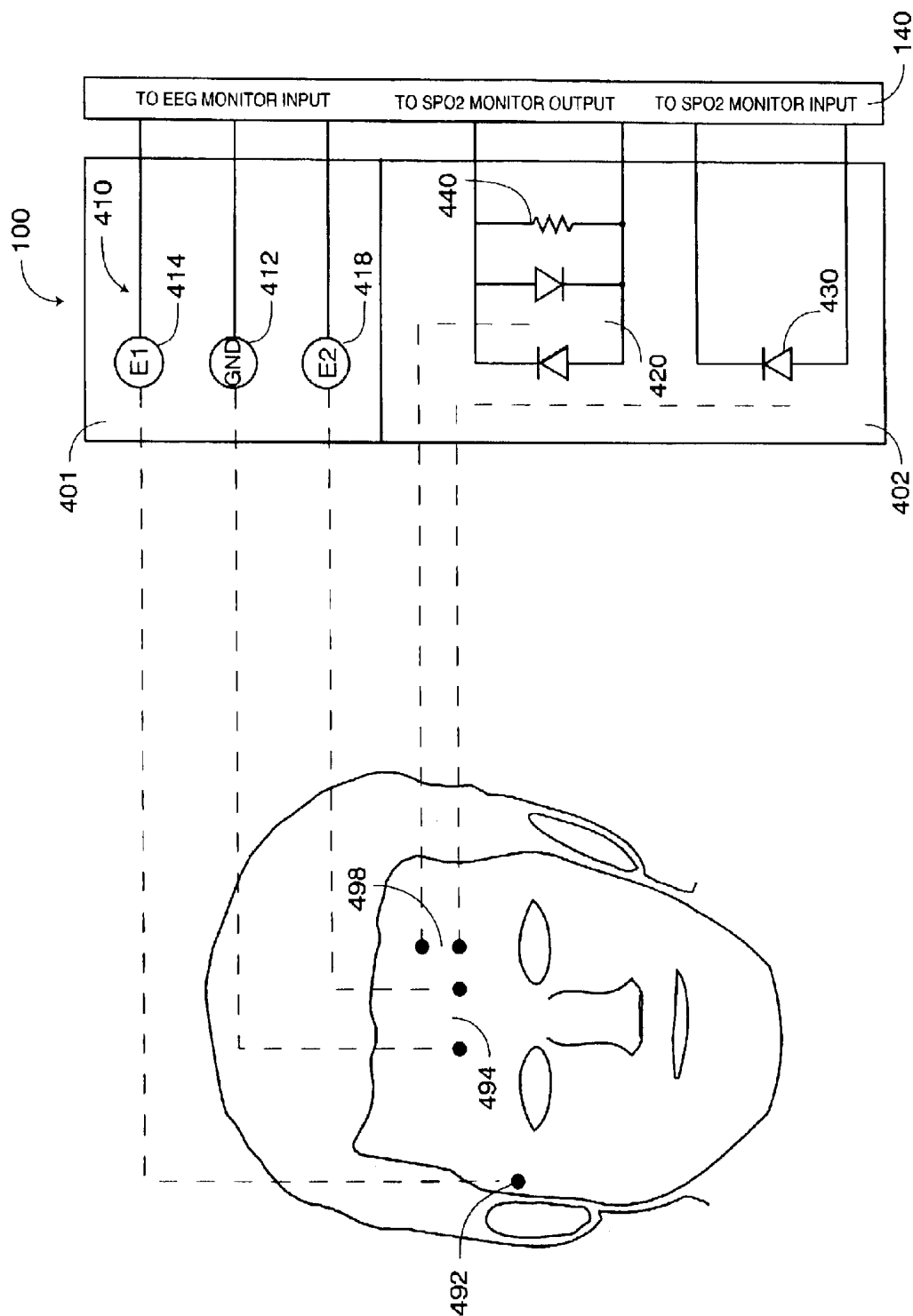


FIG. 4

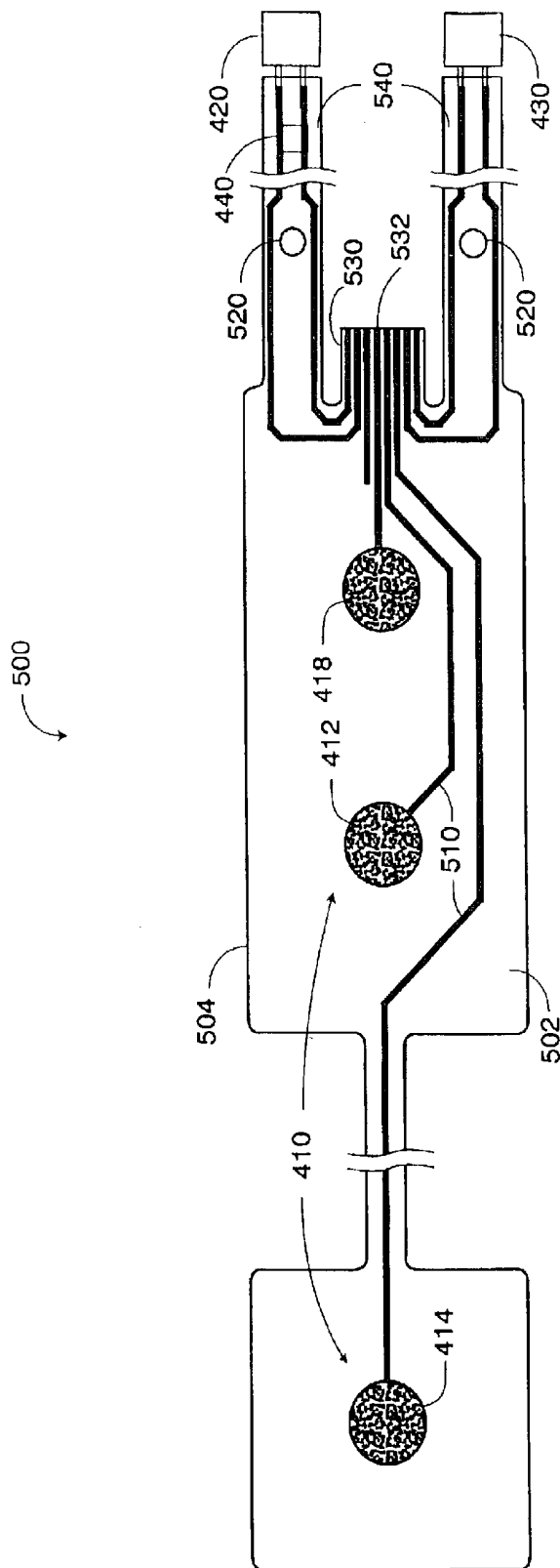


FIG. 5

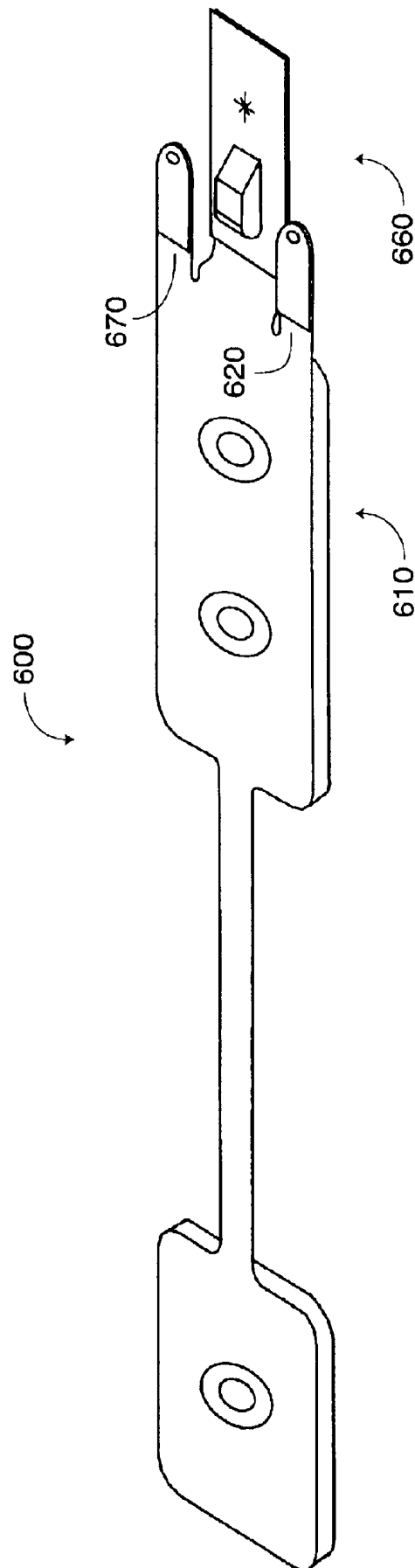


FIG. 6



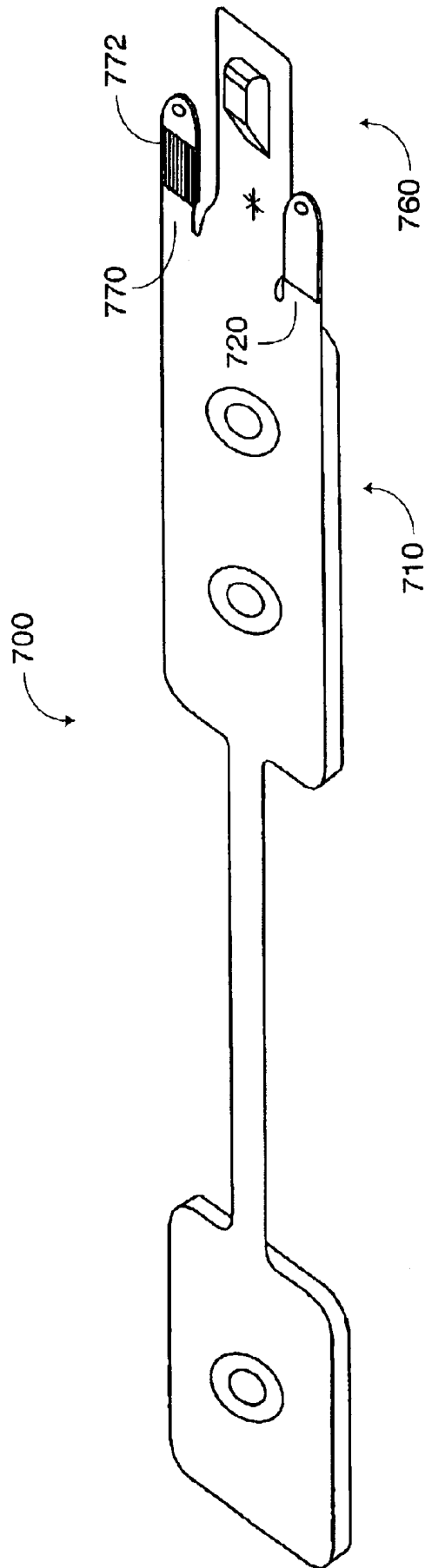


FIG. 7

**PHYSIOLOGICAL SENSOR COMBINATION****CROSS-REFERENCE TO RELATED APPLICATIONS**

This application relates to and claims the benefit of prior U.S. Provisional Patent Application No. 60/347,047 entitled Physiological Sensor Combination, filed Jan. 08, 2002, which is incorporated by reference herein.

**BACKGROUND OF THE INVENTION**

Pulse oximetry is a widely accepted noninvasive procedure for measuring the oxygen saturation level of arterial blood, an indicator of a person's oxygen supply. Early detection of low blood oxygen level is important in the medical field, for example in critical care and surgical applications, because an insufficient supply of oxygen can result in brain damage and death in a matter of minutes. A pulse oximetry system consists of a sensor applied to a patient, a pulse oximeter, and a patient cable connecting the sensor and the pulse oximeter. The pulse oximeter typically provides a numerical readout of the patient's oxygen saturation, a numerical readout of pulse rate, and an audible indication of each pulse. In addition, the pulse oximeter may display the patient's plethysmograph, which provides a visual indication of the patient's pulse contour and pulse rate.

Measuring a biopotential signal, such as an electroencephalogram (EEG) is also a widely accepted procedure for patient monitoring and diagnostic tests. An EEG measures cortical activity of the brain, which can reflect changes in cortical or subcortical cellular function due to insufficient oxygen or drugs, to name a few. For example, changes in EEG bandwidth and power can provide a measure of the effects of anesthetics on the brain. A biopotential measurement system consists of a biopotential sensor, a monitor and a patient cable connecting the sensor to the monitor. For example, an EEG monitor measures the potential difference between at least two well-spaced electrodes, using a separate ground electrode, and displays the resulting signal.

**SUMMARY OF THE INVENTION**

A physiological sensor combination has a flexible substrate configured to attach to a tissue site. Multiple sensors are disposed on the substrate, which generate physiological signals. Each of the signals is responsive to a different physiological parameter. Conductors are carried on the substrate and routed between the sensors and at least one connector. The connector is configured to communicate the physiological signals to at least one monitor, which derives measurements of the parameters. In one embodiment, the sensors comprise multiple electrodes disposed on the substrate. Each of the electrodes are adapted to be in electrical communication with the tissue site and electrically connect to at least one of the conductors. Further, an emitter and a detector are mounted to the substrate and electrically connected to at least one of the conductors. The emitter is adapted to transmit light into the tissue site, and the detector is adapted to receive reflected light from the tissue site.

In a particular embodiment, the substrate has a first side adapted to face toward the tissue site and a second side adapted to face away from the tissue site, where the conductors and the electrodes are disposed on the first side and the emitter and the detector are mounted to the first side. The substrate may comprise a fold-over portion having a circuit side corresponding to the first side, where the fold-over

portion is adapted to fold so that the circuit side is proximate the second side. Further, the emitter and the detector may be mounted to the a fold-over portion. The substrate may define at least one aperture configured so that the emitter and the detector each align with a corresponding aperture when the fold-over is in a folded position.

In another particular embodiment, the physiological sensor combination comprises a plurality of biopotential sensor pinouts corresponding to the electrodes, a plurality of optical sensor pinouts corresponding to the emitter and the detector, and a common connector extending from the substrate. The biopotential sensor pinouts and said optical sensor pinouts are each disposed on the common connector.

Another aspect of a physiological sensor combination is a substrate means for combining a first sensor and a second sensor, a connector means for communicating signals from the first sensor and the second sensor to at least one monitor, and an identifying means of conveying information about each of the first sensor and the second sensor to the monitor. The physiological sensor combination may further comprise a fold-over means for positioning sensor components so as to extend away from a tissue site. The physiological sensor combination may additionally comprise an aperture means for providing light communications between sensor components and the tissue site.

**BRIEF DESCRIPTION OF THE DRAWINGS**

FIG. 1 is an illustration of a physiological sensor combination applied to a patient and having a patient cable connected near the patient's forehead;

FIG. 2 is an illustration of a physiological sensor combination applied to a patient and having a patient cable connected near the patient's temple;

FIGS. 3A–B are perspective views of a circuit substrate and an assembled sensor, respectively, for a physiological sensor combination having a single-sided circuit substrate and a shared connector;

FIG. 4 is a schematic diagram of a physiological sensor combination showing the location of applied sensor components;

FIG. 5 is a layout diagram of a single-sided circuit for a physiological sensor combination;

FIG. 6 is a perspective view of a physiological sensor combination having a single-sided circuit substrate and dual connectors; and

FIG. 7 is a perspective view of a physiological sensor combination having a double-sided circuit substrate and dual connectors.

**DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS**

FIGS. 1–2 show a physiological sensor combination applied to a patient. FIGS. 3–5 illustrate a physiological sensor combination having a biopotential sensor and an optical sensor configured on a single-sided flexible circuit substrate with a shared patient cable connector. FIG. 6 illustrates a physiological sensor combination also having a biopotential sensor and an optical sensor configured on a single-sided flexible circuit substrate. The biopotential sensor and the optical sensor, however, each have separate patient cable connectors. FIG. 7 illustrates a physiological sensor combination having a biopotential sensor and an optical sensor configured on a double-sided circuit substrate, each sensor also having separate patient cable connectors.

FIGS. 1–2 illustrate a physiological sensor combination applied to the forehead and temple areas of a patient. A

patient cable 130 connects the physiological sensor combination 100 (FIG. 1), 101 (FIG. 2) to one or more monitoring devices (not shown). As shown in FIG. 1, the patient cable 130 may connect near the patient's forehead. As shown in FIG. 2, the patient cable 130 may alternatively connect near the patient's temple. The biopotential sensor 110 and optical sensor 120 may share a common connector 140. Alternatively, the biopotential sensor 110 and optical sensor 120 may each have a dedicated patient cable connector, as described in further detail with respect to FIGS. 6–7, below. The biopotential sensor 110 may be an EEG sensor for depth of consciousness monitoring, as described above. The optical sensor 120 may be a pulse oximetry reflectance sensor for oxygen saturation monitoring, also described above.

FIGS. 3A–B illustrate a physiological sensor combination 100 having a biopotential sensor 110 and an optical sensor 120 configured on a flexible circuit substrate 500. As shown in FIG. 3A, the flexible circuit 500 is single-sided, having a blank side 501 and a circuit side 502 with printed conductive traces 510 on the circuit side 502. The biopotential sensor 110 has electrodes 410 (not visible and shown as dashed lines) printed on the circuit side 502. The electrodes 410 are configured so that one electrode is applied to the temple area and two electrodes are applied to the forehead, as further described with respect to FIGS. 4–5, below.

Further shown in FIG. 3A, the optical sensor 120 includes a fold-over 540, an emitter 420, a detector 430 and an information element 440. The emitter 420, detector 430 and information element 440 are each mounted to the circuit side 502 on the fold-over 540 and electrically connected to traces 510, as described in detail with respect to FIGS. 4–5, below. The optical sensor 120 is configured so that emitter 420 and a detector 430 are applied over the forehead, also described with respect to FIGS. 4–5, below. The fold-over 540 is such that each of the emitter 420 and detector 430 align with corresponding apertures 520 (FIG. 5) so that light transmitted from the emitter 420 passes through an aperture 520 (FIG. 5) and into a patient's skin and that reflected light passes out of a patient's skin, through an aperture 520 (FIG. 5) and is received by the detector 430. The substrate 500 has a stub 530 that contains pinouts 532 (FIG. 5), which connect to the electrodes 410 and also to the emitter 420, detector 430 and information element 440, also described in detail with respect to FIGS. 4–5, below. Emitters and a detector for a pulse oximetry sensor are described in detail in U.S. Pat. No. 6,256,523 entitled "Low Noise Optical Probe," which is assigned to Masimo Corporation and incorporated by reference herein. An information element for a pulse oximetry sensor is described in detail in U.S. Pat. No. 6,001,986 entitled "Manual And Automatic Probe Calibration," which is assigned to Masimo Corporation and incorporated by reference herein.

As shown in FIG. 3B, the biopotential sensor 110 has an adhesive foam layer 310 disposed around the electrodes 410 on the circuit side 502. The foam layer 310 has an adhesive for patient skin attachment and cushions the biopotential sensor 110 against the skin. Further, the foam layer 310 forms cavities around the electrodes 410 that are filled with a conductive gel for electrical communication between a tissue site and the electrodes 410. Printed electrode indicators 370 facilitate sensor application on a tissue site. Electrodes printed on a substrate, an associated foam layer, and gel-filled foam cavities are described in detail in U.S. Pat. No. 6,032,064 entitled "Electrode Array System For Measuring Electrophysiological Signals," assigned to Aspect Medical Systems, Inc. and incorporated by reference herein. One of ordinary skill in the art will recognize that various electrode configurations may be utilized as the biopotential sensor 110.

Also shown in FIG. 3B, the optical sensor 120 has a face tape 330 and a base tape 340 that envelop the fold-over 540 along with the fold-over mounted components 420–440. In one embodiment, the face tape 330 and base tape 340 attach together and to the fold-over 540 with PSA. Further, the base tape 340 has a backing (not shown) that is removed to expose an adhesive for skin attachment. The face tape 330 also secures the detector 430 within an optical cavity and cover 350. A printed emitter indicator 390 facilitates sensor application on a tissue site. Emitters, detectors, optical cavities and corresponding covers are described in detail in U.S. Pat. No. 6,256,523, referenced above.

Further shown in FIG. 3B, the physiological sensor combination 100 has a tab 320 that attaches to the stub 530 (FIG. 3A) to complete the connector 140. In one embodiment, the attachment is accomplished with pressure sensitive adhesive (PSA) between the tab 320 and stub 530. The tab 320 provides a stiffener for the pinouts 532 (FIG. 5) and an insertion and locking mechanism for a mating patient cable connector, as described in U.S. Pat. No. 6,152,754 entitled "Circuit Board Based Cable Connector" and U.S. Pat. No. 6,280,213 entitled "Patient Cable Connector," each assigned to Masimo Corporation and incorporated by reference herein.

The physiological sensor combination 100 is described above with respect to a fold-over that positions the optical sensor components 420–440 so that they extend away from the tissue site. This advantageously allows a smooth surface to be positioned against the tissue site for patient comfort. In another embodiment, however, there is no fold-over 540 and the components 420–440 extend from the substrate toward the tissue site. In yet another embodiment, there is no fold-over and the components 420 are mounted on the substrate side opposite the conductors and utilize substrate feed-throughs to connect with the flex circuit traces 510. Further, the fold-over 540 is described above as positioning the emitter 420 and detector 430 over substrate apertures 520 (FIG. 5). In an alternative embodiment, the fold-over 540 is skewed so that the emitter 420 and detector 430 are positioned away from the substrate so that no apertures are necessary.

FIG. 4 illustrates a circuit diagram for a physiological sensor combination 100 having a biopotential sensor circuit 401 and an optical sensor circuit 402. The biopotential sensor circuit 401 has an electrode array 410, which is placed on well-separated skin areas. In one embodiment, a first electrode 414 is placed on a temple area 492 and a second electrode 418 is placed on a forehead area 494. A ground electrode 412 is also placed on the forehead area 494 near the second electrode 418. Each electrode of the array 410 provides a pinout to a connector 140. The connector 140 provides sensor input to a monitor. The electrodes placed on the patient's head transmit EEG signals to a monitor, which may include a separate digitizer located near the patient to reduce electrical noise. The difference in potential between the first electrode 414 and second electrode 418 reflects primarily a far-field electrical source, i.e. the EEG from the distant brain cortex, and not a near-field electrical source, such as transdermal nervous stimulation of muscle. The monitor filters the EEG data, analyzes it for artifact and extracts characteristic features from the complex signal to provide pattern recognition of changes over time.

Also shown in FIG. 4, the optical sensor circuit 402 has an emitter 420, a detector 430 and an information element 440. The emitter 420 includes both a red LED (light emitting diode) and an infrared (IR) LED in a back-to-back arrangement. In alternative embodiments, the red and IR LEDs are

arranged in three-wire, common anode or common cathode configurations, as is well-known in the art. The detector **430** is a photodiode. The LEDs **420** and photodiode **430** are located on the skin in close proximity, such as on a forehead area **498**. In this manner, the LEDs emit light into the blood vessels and capillaries underneath the skin, and the photodiode **430** is positioned to detect the LED emitted light reflected from the skin tissues. The emitter **420** and detector **430** provide pinouts to the connector **140**, which provides a sensor input to a monitor. The monitor determines oxygen saturation by computing the differential absorption by arterial blood of the two wavelengths of light projected into the skin from the emitter **420**, as is well-known in the art. The monitor provides LED drive current, which alternately activates the red and IR LEDs. The detector **430** uses a single photodiode that responds to both the red and infrared emitted light and generates a time-division-multiplexed ("modulated") output signal to the monitor, corresponding to the red and infrared light energy attenuated by absorption and reflection from the patient's tissue. The monitor has front-end circuitry for amplification, filtering and digitization of the detector signal. The monitor also has a signal processor that calculates a ratio of detected red and infrared intensities, and an arterial oxygen saturation value is empirically determined based on that ratio.

Further shown in FIG. 4, the optical sensor circuit **402** may have an information element **440**, such as a resistor configured in parallel with the emitter **420** LEDs. The information element **440** can be read by the monitor and used to determine such things as LED wavelength, sensor type or manufacturer. Information elements and monitor reading of information elements are described in U.S. Pat. No. 6,011,986, referenced above. Advantageously, although associated with the optical sensor circuit **402**, the information element **440** can be used to designate information regarding the biopotential sensor portion of the physiological sensor combination **100**. For example, the information element **440** can specify the number of electrodes as well as the electrode locations on the head.

FIG. 5 illustrates a flexible circuit **500** for a physiological sensor combination **100**. The flexible circuit **500** has a substrate **504**, traces **510**, electrodes **410**, pinouts **530** and apertures **520**. Conductors are deposited and/or etched on a circuit side **502** of the substrate **504** in a pattern to form the traces **510**, electrodes **410** and pinouts **532**, as is well known in the art. In one embodiment, the substrate **504** is a flexible polyester film and the conductors are silver/silver-chloride. In another embodiment, the conductors are copper. The components **420-440** attach to the flexible circuit **500** and are electrically connected to the traces **510**, such as with solder. The fold-over **540** is configured so that the emitter **420** and detector **430** align with the corresponding apertures **520**.

FIG. 6 illustrates a physiological sensor combination **600** having a biopotential sensor **610** and an optical sensor **660**. The biopotential sensor **610** is configured as described with respect to FIGS. 3-5, above, except that the physiological sensor combination **600** has a connector **620** that is dedicated to the biopotential sensor **610** rather than being shared with the optical sensor **660**. The optical sensor **660** also is configured as described with respect to FIGS. 3-5, above, except that a connector **670** is dedicated to the optical sensor **660** rather than being shared with the biopotential sensor **610**. Further, the optical sensor **660** has a single fold-over (not visible) on which is mounted the emitter **420** (FIG. 4) and detector **430** (FIG. 4) rather than having a separate fold-over **540** (FIG. 3A) for each.

FIG. 7 illustrates a physiological sensor combination **700** having a biopotential sensor **710** and an optical sensor **760**. The biopotential sensor **710** is configured as described with respect to FIG. 6, above. The optical sensor **760** also is configured as described with respect to FIG. 6, above, except that the flexible circuit **500** (FIG. 5) is double-sided, i.e. the traces **510** (FIG. 5) associated with the biopotential sensor **710** are on the side facing the patient's skin when applied, and the traces **510** (FIG. 5) associated with the optical sensor **760** are on the side away from the patient's skin when applied. As a result, the connector **770** is dedicated to the optical sensor **760** and has pinouts **772** facing away from the patient's skin when applied. Further, the optical sensor **760** does not have a fold-over **540** (FIG. 3A). Rather, the optical sensor components **420-440** (FIG. 4) are mounted on the flexible circuit side away from the patient's skin.

A physiological sensor combination is described above with either a shared patient cable connector or a patient cable connector dedicated to each sensor. One of ordinary skill will recognize that either connector configuration will allow the sensor to communicate with a single monitor that analyzes and displays multiple physiological parameters or, alternatively, multiple monitors that are dedicated to analyzing only related physiological parameters, such as oxygen saturation and pulse rate.

The physiological sensor combination as described above can be cost effectively manufactured, advantageously allowing disposable use. One of ordinary skill in the art will recognize that, however, that the physiological sensor combination as disclosed herein can be similarly applied to construct a reusable sensor combination.

The physiological sensor combination was also described above with respect to a shared substrate. One of ordinary skill in the art will recognize that a physiological sensor combination can be constructed from, for example, a biopotential sensor configured on a first substrate and an optical sensor configured on a second substrate, where the first substrate and the second substrate are joined together during the manufacturing process to form a multilayer substrate or an otherwise integrated substrate incorporating multiple sensors.

Although a physiological sensor combination is described above with respect to a biopotential sensor combined with an optical sensor applied to a patient's head, one of ordinary skill in the art will recognize that a physiological sensor combination may be applied to other tissue sites and utilize other sensor combinations, where there is a need to combine two or more sensors in one to accommodate sensors competing for the same tissue site. For example, a physiological sensor combination may include a noninvasive blood pressure (NIBP) sensor and a pulse oximetry sensor or a NIBP sensor and a respiration rate sensor for monitoring on the forearm or the wrist. As another example, a physiological sensor combination may include two optical sensors and one biopotential sensor applied to the forehead and configured as a pulse oximetry sensor and a EEG sensor, as described above, in addition to a near infrared spectroscopy sensor for measuring cerebral tissue oxygenation.

A biopotential sensor as described above could be used in conjunction with a depth of anesthesia monitor that uses not just passive EEG, but also active EEG. That is an Evoked Potential EEG can be used, where some kind of sound is played and changes in EEG are observed as the patient goes into consciousness.

A physiological sensor combination 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 physiological sensor combination comprising:

- a flexible substrate configured to attach to a tissue site;
- a plurality of sensors disposed on said substrate and adapted to provide a corresponding plurality of physiological signals, each of said signals responsive to at least one of a plurality of physiological parameters;
- a plurality of conductors disposed on said substrate between said sensors and at least one connector, and said at least one connector configured to communicate said signals to at least one monitor so as to derive a plurality of measurements of said parameters,

wherein said sensors comprise:

- a plurality of electrodes disposed on said substrate, each of said electrodes adapted to be in electrical communication with said tissue site and electrically connected to at least one of said conductors; and
- an emitter and a detector mounted to said substrate and electrically connected to at least one of said conductors, said emitter adapted to transmit light into said tissue site and said detector adapted to receive reflected light from said tissue site,

wherein:

- said substrate has a first side adapted to face toward said tissue site and a second side adapted to face away from said tissue site;
- said conductors and said electrodes disposed on said first side; and
- said emitter and said detector mounted to said first side wherein said substrate comprises a fold-over portion having a circuit side corresponding to said first side, and said fold-over portion adapted to fold so that said circuit side is proximate said second side.

**2.** The physiological sensor combination according to claim **1** wherein said emitter and said detector are mounted to said a fold-over portion.

**3.** The physiological sensor combination according to claim **2** wherein said substrate defines at least one aperture configured so that said emitter and said detector each align with a corresponding one of said at least one aperture when said fold-over is in a folded position.

**4.** The physiological sensor combination according to claim **3** further comprising:

- a plurality of biopotential sensor pinouts corresponding to said electrodes;
- a plurality of optical sensor pinouts corresponding to said emitter and said detector; and

a common connector extending from said substrate, said biopotential sensor pinouts and said optical sensor pinouts each disposed on said common connector.

**5.** The physiological sensor combination comprising:

- a substrate means for combining a first sensor and a second sensor;
- a connector means for communicating signals from said first sensor and said second sensor to at least one monitor;
- an identifying means of conveying information about each of said first sensor and said second sensor to said monitor; and
- fold-over means for positioning sensor components so as to extend away from a tissue site.

**6.** The physiological sensor combination according to claim **5** further comprising an aperture means for providing light communications between sensor components and said tissue site.

**7.** A sensor capable of outputting a signal indicative of one or more physiological parameters of body tissue of a patient, the sensor comprising:

- a substrate including a fold-over portion;
- a first sensor positioned on said substrate and including a detector capable of detecting light attenuated by body tissue, said detector also capable of outputting a detector signal indicative of one or more physiological parameters of said body tissue;
- a second sensor positioned on said substrate;
- conductors capable of communicating signals between said first sensor and at least one monitor, said signals including said detector signal; and
- conductors capable of communicating signals between said second sensor and said at least one monitor, wherein the fold-over portion of the substrate positions sensor components of said first sensor away from said body tissue to provide a substantially smooth sensor surface at said body tissue.

**8.** The sensor according to claim **7**, wherein said substrate further comprises apertures, and wherein said sensor components are aligned with said apertures when said substrate is folded over, thereby allowing said sensor components to emit and detect energy through said apertures.

**9.** The sensor according to claim **7**, further comprising an information element identifiable by said at least one monitor to convey information about at least one of said first sensor and said second sensor.

**10.** The sensor according to claim **7**, wherein said first sensor comprises a pulse oximetry sensor including a plurality of emitters and said detector, and wherein said second sensor comprises a biopotential sensor including electrodes.

\* \* \* \* \*

UNITED STATES PATENT AND TRADEMARK OFFICE  
**CERTIFICATE OF CORRECTION**

PATENT NO. : 6,934,570 B2  
APPLICATION NO. : 10/325699  
DATED : August 23, 2005  
INVENTOR(S) : Kiani et al.

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Title Page, column 1 Item [75] (Inventors), line 4, delete "La Mirada" and insert -- La Habra Heights --, therefore.

Title Page, column 1 Item [73] (Assignee), line 1, before "Irvine" insert -- 40 parker --.

Title Page, column 1 Item [73] (Assignee), line 1, after "CA" insert -- 92618 --.

At column 2, line 3, after "the" delete "a".

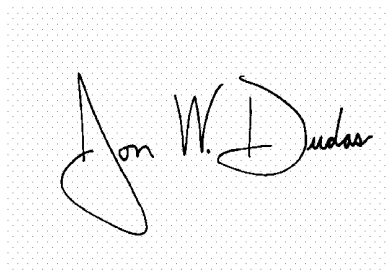
At column 2, line 53, delete "show" and insert -- shows --, therefore.

At column 3, line 13 (Approx.), after "above" insert -- . --.

At column 8, line 13, in Claim 5, before "fold-over" insert -- a --.

Signed and Sealed this

Twenty-sixth Day of June, 2007

A handwritten signature in black ink on a light gray dotted background. The signature is written in a cursive style and reads "Jon W. Dudas".

JON W. DUDAS  
*Director of the United States Patent and Trademark Office*

专利名称(译)	生理传感器组合		
公开(公告)号	<a href="#">US6934570</a>	公开(公告)日	2005-08-23
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外部链接	<a href="#">Espacenet</a> <a href="#">USPTO</a>		

#### 摘要(译)

生理传感器组合具有柔性基板，其配置成附接到组织部位。多个传感器设置在基板上，其产生生理信号。每个信号响应于不同的生理参数。导体承载在基板上并在传感器和至少一个连接器之间布线。连接器被配置为将生理信号传送到至少一个监视器，该监视器导出参数的测量值。

