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(54) **SEGMENTED SENSOR**

(71) Applicant: **Nonin Medical, Inc.**, Plymouth, MN
(US)

(72) Inventors: **Stefan Skorich**, St. Louis Park, MN (US); **Brian Joseph Kane**, Inver Grove Heights, MN (US); **Matthew Prior**, Plymouth, MN (US)

(73) Assignee: **Nonin Medical, Inc.**, Plymouth, MN
(US)

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2562/04 (2013.01); *A61B 2562/146* (2013.01);
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See application file for complete search history.

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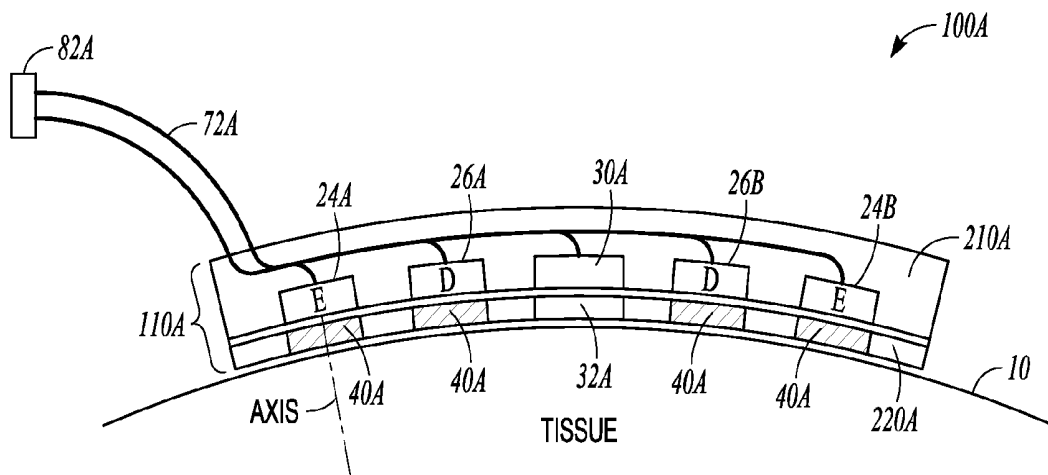
Primary Examiner — Eric F Winakur

(74) *Attorney, Agent, or Firm* — Schwegman Lundberg & Woessner, P.A.

(57) **ABSTRACT**

A device includes a first portion, an information element, a second portion, and an encoder element. The first portion includes a planar substrate and at least one optical element coupled to the substrate. The optical element has an axis and the axis is substantially normal to the substrate. The information element sensor is coupled to the substrate. The information element sensor is configured to provide an information signal. The second portion includes a membrane configured to couple to the planar substrate. The membrane has a contact surface configured to couple with tissue. The encoder element is coupled to the membrane. The information signal corresponds to proximity of the encoder element.

16 Claims, 5 Drawing Sheets



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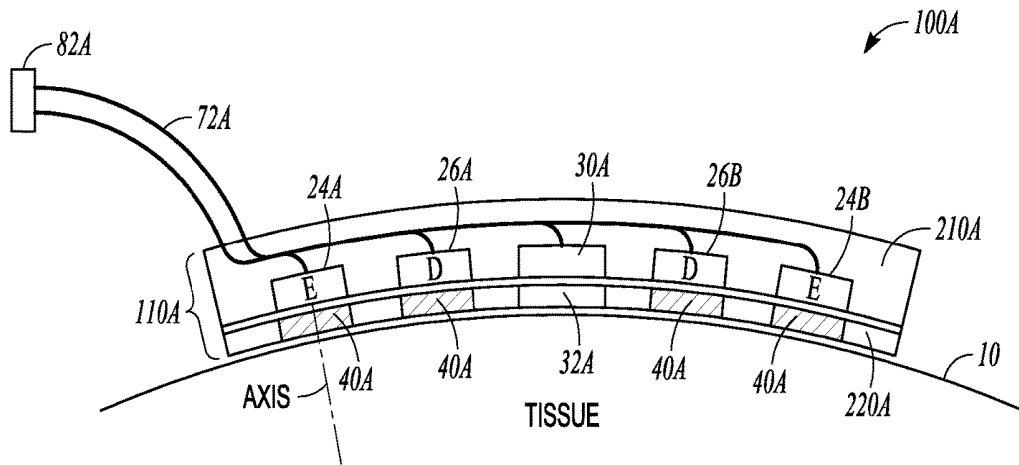


FIG. 1

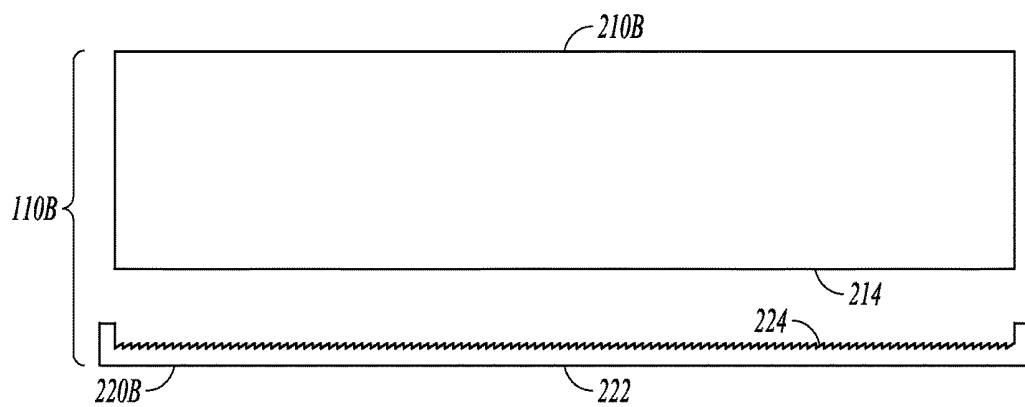


FIG. 2A

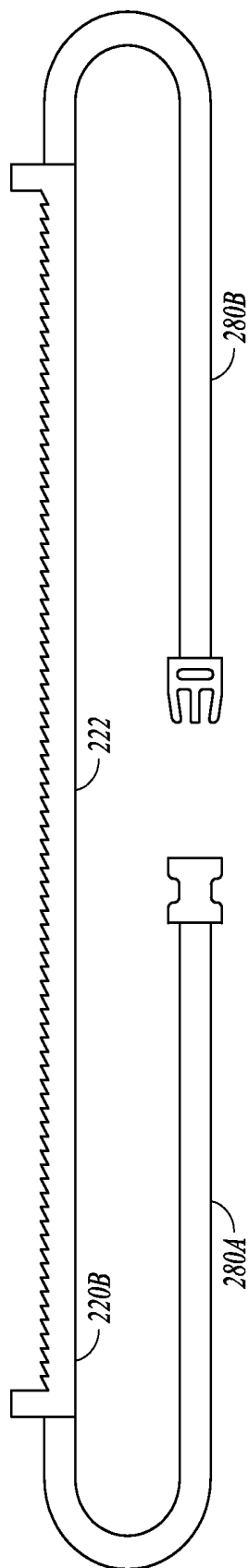


FIG. 2B

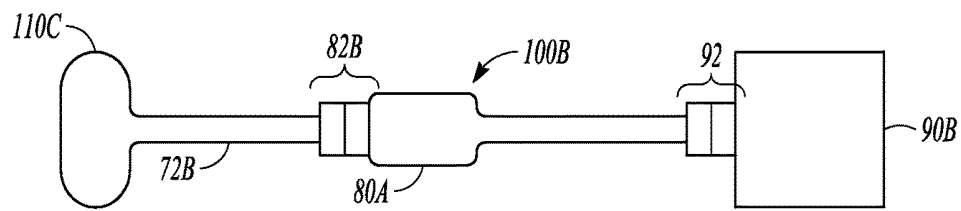


FIG. 3A

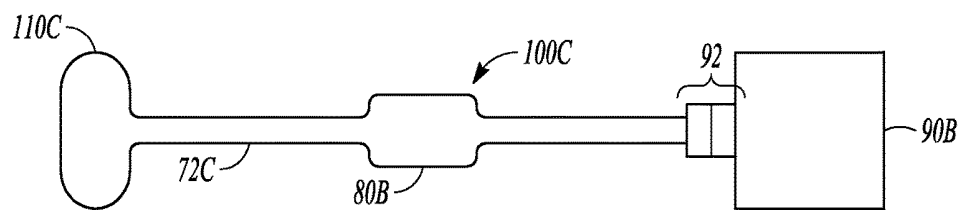


FIG. 3B

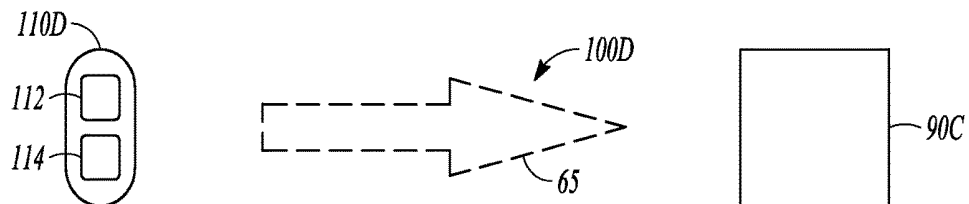


FIG. 3C

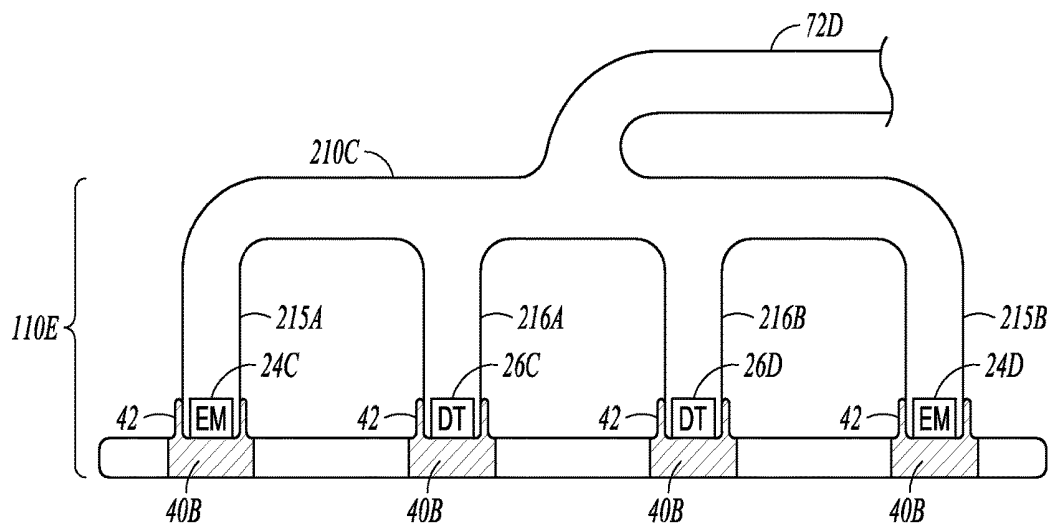


FIG. 4

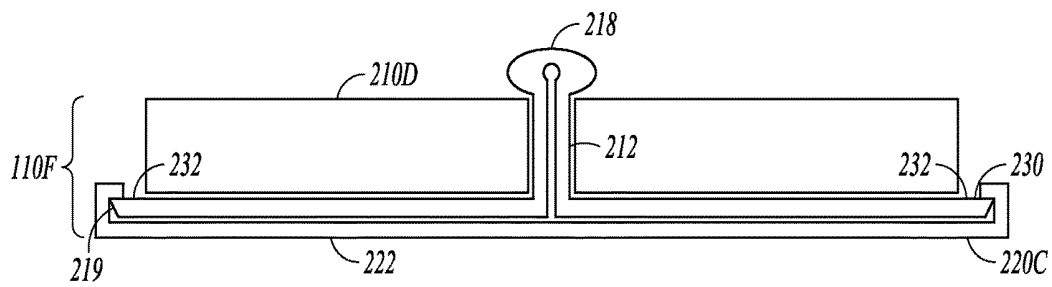


FIG. 5

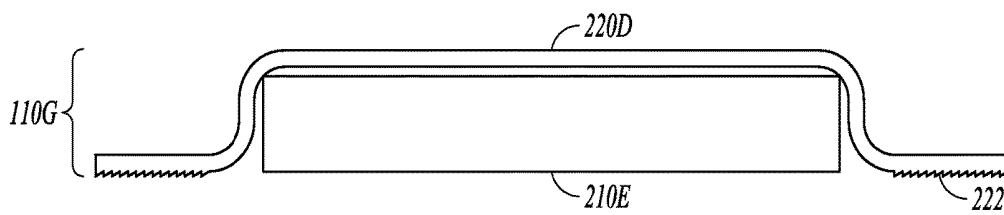


FIG. 6

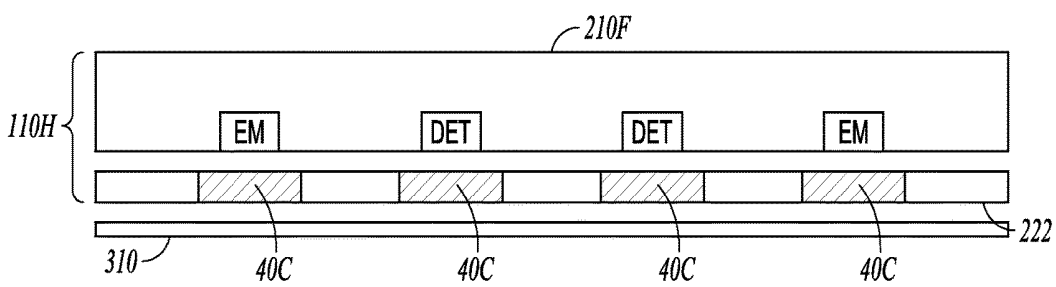
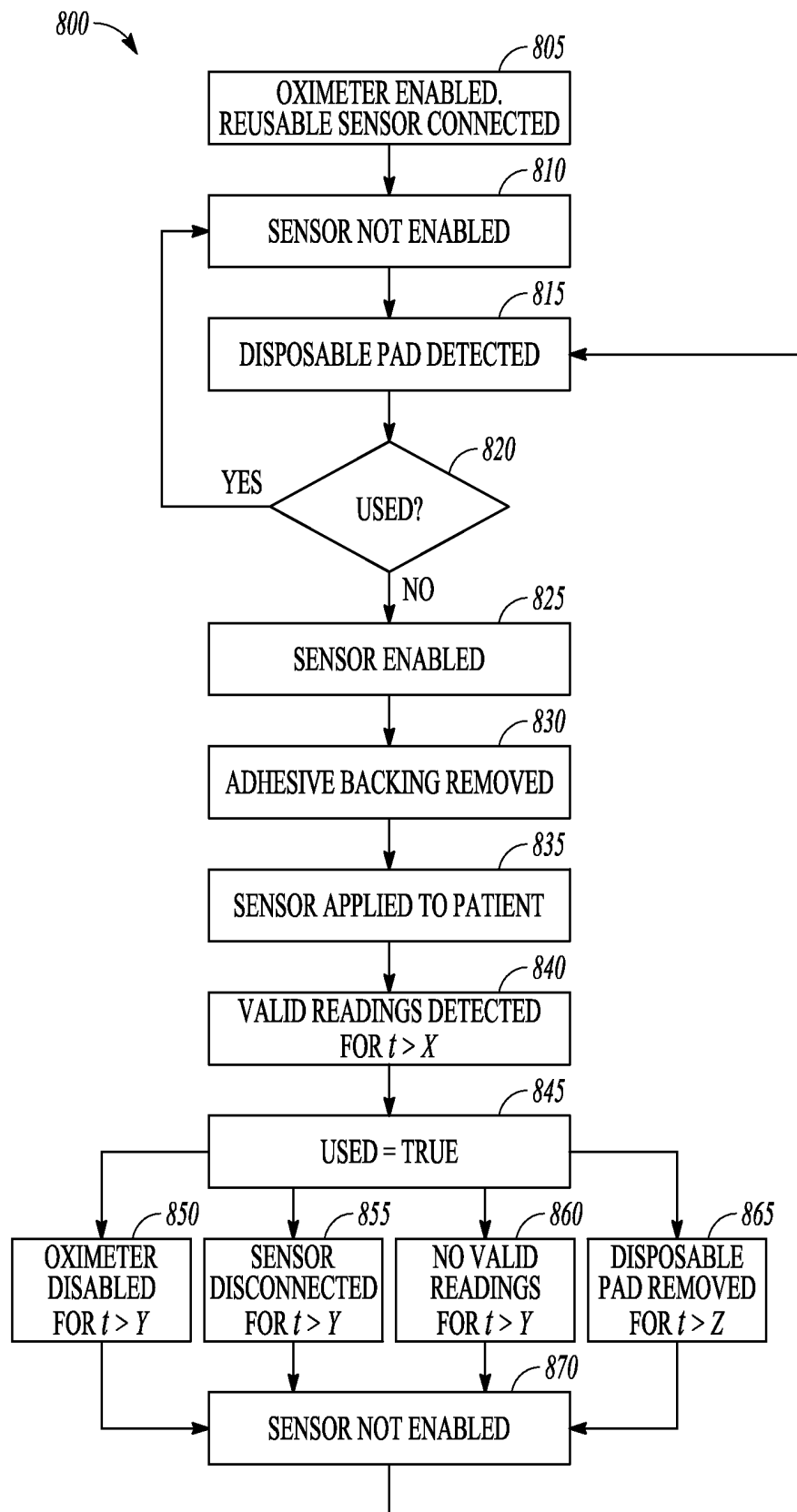


FIG. 7

**FIG. 8**

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SEGMENTED SENSOR

BACKGROUND

Presently available technology for physiological sensors is inadequate. For example, a typical sensor includes high value components that are often discarded after a single use.

OVERVIEW

The present inventors have recognized, among other things, that a problem to be solved can include providing a lower cost sensor system. The present subject matter can help provide a solution to this problem, such as by providing a segmented sensor having a first portion including high value components and a second portion having low value components.

According to one example, the present subject matter includes a sensor module configured to noninvasively measure a physiological parameter. The sensor module can include a first portion and can include a second portion. The first portion can be viewed as a reusable portion that can be used multiple times. This can include multiple usages associated with a single patient or multiple usages with multiple patients. The second portion can be of relatively low cost and, can be construed as disposable.

The second portion of the sensor module includes at least one information element and includes a substantially planar substrate having opposing surfaces. The opposing surfaces include a proximal surface configured to couple with the tissue of the patient and includes a distal surface configured to couple with the first portion of the sensor.

In one example, the first portion of the sensor module includes optical elements such as a radiation emitter and a radiation detector. In addition, the first portion can include an information element sensor. The first portion can include a housing of a protective material such as plastic, silicone, or metal.

The second portion of the sensor module can include an encoder element. The encoder element is configured to communicate wirelessly with the information element sensor in the first portion. The first portion is configured to wirelessly receive information from the second portion, and in this manner, the second portion communicate wirelessly with the first portion of the sensor module. The first portion is electrically isolated from the second portion.

In one example, the encoder element and the information element sensor can communicate using radio frequency communication. For example, the encoder element can include a radio frequency identification (RFID) tag and the information element sensor can include an RFID reader.

In one example, the encoder element and the information element sensor can communicate using magnetic coupling. For example, the encoder can include a magnetic element and the information element sensor can include a Hall Effect sensor.

In one example, the encoder element and the information element sensor can communicate using capacitive coupling. For example, the encoder and information element sensor can each include conductive elements in which case an oscillator circuit is included in the first portion.

In one example, the encoder element and the information element sensor can communicate using inductive sensing, and include a magnet, a metal object, or a conductive element in the second portion, and include a coil in the first portion.

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The information element sensor allows the first portion to detect proximity or detect the presence of the second portion. The information element associated with the second portion can be encoded with information pertaining to characteristics of the sensor, such as sensor type. The information element can include encrypted data or security data to prevent reuse or excessive use of the second portion. In addition, the information element can be encoded to indicate authenticity of the second portion.

The first portion and the second portion can have mutually complementary features to allow one to couple with the other through mechanical means. This can include mechanical features in both portions. One example includes an interlocking mechanism by which a deformable feature of one portion engages with a feature of another portion. In one example, an adhesive is disposed on a mating surface of either or both of the first portion and the second portion.

The optical elements of the first portion can include an emitter and a detector. The optical elements of the first portion can be electrically connected with circuitry to allow measurement of a physiological parameter of a patient. This circuitry can be housed in an external (or remote) device such as an oximeter, which connects to the sensor via an electrical connector. The circuitry can be housed in the first portion in which case the measurement data is transmitted wirelessly to an external device such as a computer or a display monitor.

This overview is intended to provide an overview of subject matter of the present patent application. It is not intended to provide an exclusive or exhaustive explanation of the invention. The detailed description is included to provide further information about the present patent application.

BRIEF DESCRIPTION OF THE DRAWINGS

In the drawings, which are not necessarily drawn to scale, like numerals may describe similar components in different views. Like numerals having different letter suffixes may represent different instances of similar components. The drawings illustrate generally, by way of example, but not by way of limitation, various embodiments discussed in the present document.

FIG. 1 illustrates a segmented sensor, according to one example.

FIG. 2A illustrates an interface for a segmented sensor, according to one example.

FIG. 2B illustrates a second portion having a mechanical fastener, according to one example.

FIG. 3A illustrates a wired system including an interconnected segmented sensor, according to one example.

FIG. 3B illustrates a wired system including a segmented sensor, according to one example.

FIG. 3C illustrates a wireless system including a segmented sensor, according to one example.

FIG. 4 illustrates a segmented sensor, according to one example.

FIG. 5 illustrates a segmented sensor having a user separable linkage, according to one example.

FIG. 6 illustrates a segmented sensor having an overlay, according to one example.

FIG. 7 illustrates a segmented sensor having a protective layer, according to one example.

FIG. 8 shows a validation scheme for a segmented sensor, according to one example.

DETAILED DESCRIPTION

FIG. 1 illustrates system 100A according to one example. System 100A includes segmented sensor 110A having a first

portion **210A** and a second portion **220A**, according to one example. First portion **210A** can be regarded as reusable and second portion **220A** can be regarded as disposable however, this view considers only the costs associated with the particular portions.

In the example illustrated, system **100A** includes line **72A** and connector **82A**. Line **72A** can include a multiple conductor electric cord or fiber optic link. Connector **82A** is configured to engage with a complementary connector of a remote device or an external device. Line **72A** and connector **82A** can provide electric power for operation of segmented sensor **110A**.

In the example illustrated, first portion **210A** includes optical elements. The optical elements include emitter **24A** and emitter **24B**. Emitter **24A** and emitter **24B** can include an infrared emitter, a near-infrared light emitter, or a visible light emitter. The optical elements of first portion **210A** include photodetectors **26A** and photodetector **26B**. The optical elements are arranged to provide spacing and alignment to enable optical detection of a physiological parameter such as oxygen saturation.

The optical elements, including emitter **24A**, emitter **24B**, detector **26A**, and detector **26B** are each connected to line **72A** and are each aligned such that an axis is substantially normal to a planar substrate of first portion **210A**. In the example, the optical elements of first portion **210A** can be viewed as coupled to a planar substrate disposed along a bottom surface.

First portion **210A** includes information element sensor **30A**, which is connected to line **72A**. Information element sensor **30A** is configured to read data or detect proximity associated with information encoder **32A** of second portion **220A**.

Second portion **220A** includes a membrane. In one example, the membrane is coupled to tissue **10** by an adhesive. In the example shown, tissue **10** has a curvature to represent, for example, a scalp. In other figures, the curvature is omitted for clarity and the second portion is shown having a flat contour. The second portion can be rigid or flexible and conformable to accommodate a tissue contour.

Second portion **220A** is also coupled to first portion **210A** via an adhesive or via a mechanical feature.

Second portion **220A** includes a plurality of openings or windows, each of which are here denoted as aperture **40A**. Aperture **40** can include an optically transparent window or a cutout portion of a membrane.

FIG. **2A** illustrates an interface for segmented sensor **110B**, according to one example. First portion **210B** includes mating surface **214**. Second portion **220B** includes tissue surface **222** and mating surface **224**. Mating surface **214** and mating surface **224** can be bonded by an adhesive (here denoted by a jagged surface) affixed to either one or both.

In one example, tissue surface **222** is coupled to patient tissue by an adhesive bond. In other examples, tissue surface **222** of second portion **220B** is coupled, or held securely, to the patient by a strap, a belt, a headband, an elastic wrap, a bandage wrap, or a sleeve.

FIG. **2B** illustrates second portion **220B** having a mechanical fastener. In the figure, mechanical fastener is represented by a strap having portion **280A** and portion **280B**, configured to encircle a patient body. Portions **280A** and **280B** terminate in buckle ends or other fastener. In addition to a strap, the mechanical fastener can include a belt, a headband, an elastic wrap, a bandage wrap, or a sleeve. The mechanical fastener is configured to couple the second portion to the tissue of the patient. The coupling

between the patient and the second surface is configured to occlude ambient light and provide good optical coupling to the tissue.

FIGS. **3A**, **3B**, and **3C** illustrate configurations of systems according to various examples.

FIG. **3A** illustrates system **100B** having wired interconnections. In the figure, segmented sensor **110C** is coupled to device **80A** by link **72B** and by connector **82B**. Device **80A** can include an oximeter having measurement circuitry. Device **80A** can control operation and receive data from segmented sensor **110C**. Device **80A** is coupled to device **90B** by connector **92**. Device **90B** can include a display or a computer.

FIG. **3B** illustrates system **100C** including a sensor with an integrated oximeter. System **100C** includes wired interconnections as shown. In the figure, segmented sensor **110C** is coupled to device **80B** by link **72C**. Device **80B** can include measurement circuitry. Device **80B** can control operation and receive data from segmented sensor **110C**. Device **80B** is coupled to device **90B** by connector **92**. Device **90B** can include a display or a computer.

FIG. **3C** illustrates system **100D** having wireless interconnections. In the figure, segmented sensor **110D** is coupled to device **90C** by wireless link **65**. Device **90C** can include a display. Device **90C** can control operation and receive data from segmented sensor **110D**. Segmented sensor **110D** includes wireless communication module **112** and measurement module **114**. In one example, segmented sensor **110D** includes an oximeter and measurement module **114** includes measurement circuitry. Segmented sensor **110D** communicates with device **90C** the processed data by a radio frequency communication link or other non-wired communication link.

FIG. **4** illustrates segmented sensor **110E**, according to one example. Segmented sensor **110E** includes first portion **210C** and second portion **220C**. First portion **110E** includes an arrangement of optical elements disposed on flexible members. For example, emitter **24C** is disposed at an end of linear member **215A**, emitter **24D** is disposed at an end of linear member **215B**, detector **26C** is disposed at an end of linear member **216A** and detector **26D** is disposed at an end of linear member **216B**. Emitter **24C** is coupled to second portion **220C** at aperture **40B** by socket **42**. In addition, detector **26C** is coupled to second portion **220C** at aperture **40B** by socket **42**.

Linear members **215A**, **215B**, **216A**, and **216C** are flexible and can be manipulated to engage with sockets of second portion **220C** in the manner shown. The sockets of second portion **220C** are aligned with apertures and are configured to receive the limbs and establish suitable spacing. The spacing can be selected according to the patient age, site, health, or other parameter.

FIG. **5** illustrates segmented sensor **110F** having a user-separable linkage, according to one example. First portion **210D** includes an aperture to receive stem **212**. Stem **212**, affixed to coupler plate **230**, includes a split member that terminates with bulb **218** on one end and is coupled to second portion **220C** at the other end. Second portion **220C** includes tissue surface **222** and beveled feature **219**. Second portion **220C** and first portion **210D** are coupled together by forcing bulb **218** through an aperture of first portion **210D**. Beveled feature **219**, located at each end **232** of coupler plate **230**, is configured to receive an adhesively bonded membrane.

First portion **210D** and second portion **220C** are configured to allow single-time usage. After assembly in the manner shown in the figure, segmented sensor **110F** is

configured to resist separation without damage. As such, bulb **218** will separate from stem **212** if physically forced apart. In this manner, repeated usage of second portion **220C** is precluded.

FIG. **6** illustrates segmented sensor **110G** having second portion **220D** configured as an overlay, according to one example. In this example, first portion **210E** includes optical elements and an information element sensor. Second portion **220D** includes adhesively bonded tissue surfaces **222** and information encoder. Second portion **220D** engulfs first portion **210E**.

FIG. **7** illustrates segmented sensor **110H** having protective layer **310**, according to one example. First portion **210F** includes emitters and detectors as shown. Second portion **220E** includes apertures **40C** aligned with the optical elements of first portion **210F**. In addition, protective layer **310** is adhesively bonded to the tissue surface **222** of second portion **220E**. The configuration shown is suitable for storage or shipping.

FIG. **8** illustrates method **800** according to one example. Method **800** includes a validation scheme for a segmented sensor. Method **800**, at **805** includes enabling a remote device (such as an oximeter) and includes coupling the segmented sensor with the remote device.

At **810**, the method indicates the state in which the segmented sensor is not enabled.

At **815**, the method includes monitoring for the information element. This can include detecting presence or proximity.

At **820**, method **800** includes an inquiry to determine if the segmented sensor has been used previously. This can include checking a counter value, checking a date or time code, or checking status of a fusible element. If the inquiry indicates that the segmented sensor has been used previously, then processing continues at **810** where the flag or state is set to indicate that the sensor is not enabled. If the inquiry returns a negative answer, then, at **825**, processing continues by setting a flag to indicate that the sensor is enabled.

At **830**, method **800** includes removing a backing film from an adhesive layer on the second portion of the segmented sensor.

At **835**, method **800** includes applying the segmented sensor to the patient tissue.

At **840**, method **800** includes an inquiry to determine if the segmented sensor has returned a valid reading for a time greater than value X.

If the inquiry at **840** indicates that the sensor has been used, then, at **845**, method **800** includes setting a flag to indicate that the sensor has been used previously. Further inquiries are presented at **850**, **855**, **860**, and at **865**. At **850**, the inquiry considers if the oximeter has been disabled for time greater than a selected value Y. At **855**, method **800** can include an inquiry to consider if the sensor has been disconnected for a time greater than a selected value Y. At **860**, method **800** can include an inquiry to determine if the segmented sensor has provided invalid readings for a time greater than a selected value Y. At **865**, method **800** includes an inquiry to consider if the second portion has been removed for a time greater than selected value Z.

If any inquiry presented at **850**, **855**, **860**, or **865** provides a 'true' indication, then, at **870**, the segmented sensor is disabled and thereafter, processing returns to **815**.

Other validation routines are also contemplated, including different inquiries and different processing.

An example of the present subject matter utilizes a wireless communication link between a first portion and a

second portion. The wireless communication link can include a rolling security protocol in which each information or data exchange includes a different encoding. In one example, a segmented sensor system includes a counter that is incremented to monitor usage instances or includes a timer to monitor usage duration. On detecting a predetermined count or time, the segmented sensor is disabled.

The optical elements of a segmented sensor can be configured to measure and monitor regional oximetry, pulse oximetry, DC-based parameter (such as a signal amplitude), or other measured physiological parameter.

In various examples, the second portion can include zero, one, two or more optical elements. In various examples, the first portion can include zero, one, two or more optical elements.

The second portion can include a substrate having opposing surfaces in which one surface can be affixed to tissue and another surface can be coupled to the first portion.

The first portion can include a housing to which is coupled the optical elements. The optical elements are affixed to a bottom surface and are aligned to interact with the tissue. The housing can include a protective material such as plastic or silicone.

In one example, the first portion and the second portion are wirelessly coupled to enable functionality. Failure to wirelessly couple in the manner provided will preclude operability of the segmented sensor.

The segmented sensor can be connected to circuitry that enables the measurement of physiological parameters of the patient. This circuitry can be contained in an external device such as an oximeter, which connects to the sensor via an electrical connector in the reusable portion. The circuitry can be contained in the first portion of the sensor itself, with the measurement data being transmitted wirelessly to an external device such as a computer or a display monitor.

The first portion and the second portion of the segmented sensor can be removably attached to each other. The second portion of the sensor carries the sensor to the patient using an adhesive layer.

In one example, the segmented sensor includes a tissue contact surface on a bottom of the second portion and a mating surface configured to engage with a bottom surface of the first portion. The second portion includes windows aligned with the emitters and detectors of the first portion that allows the radiation to pass through unimpeded.

The first portion and the second portion are held in alignment by an alignment feature. The alignment feature can provide visual indication (such as a visible light), audible indication (an audible tone), a tactile (a snap fit), or a combination. One example includes an alignment feature at the mating surface between the first portion and the second portion. The alignment feature can include a pin, a tab, a hole, a ridge, or other element that ensures alignment of the optical element of the first portion and the aperture of the second portion. An alignment feature can include a raised perimeter, an example of which is shown in FIG. **2A** and in FIG. **2B**. In addition to aligning the optical element and the aperture, the alignment feature can also mechanically secure the first portion to the second portion. For example, FIG. **5** illustrates bulb **218** and beveled features **219** which engage corresponding features to provide alignment and also secures the first portion to the second portion.

In one example, the mating surface of the second portion includes an adhesive layer that adheres to the bottom surface of the first portion. To attach the second portion to the first portion, an adhesive liner is removed from the second portion, thus exposing the adhesive. The second portion is

aligned and coupled to the first portion. After using the segmented sensor, the second portion can be peeled off the first portion.

To aid in this removal, a tissue contact surface of the second portion contains an adhesive layer that bonds to the patient. The adhesive bond on the tissue is stronger than the mating surface bond between the first portion and the second portion, so that when the segmented sensor is removed from the patient, the second portion separates from the first portion and remains affixed to the patient.

In one example, the first portion or the second portion includes a tab and the user can grasp and pull the tab to separate the portions.

In one example, a region of the adhesive contains a non-stick surface (for example, PTFE), such that no adhesive residue remains when the second portion is removed.

In one example, the second portion and the first portion include mating features that provides a locking interface. To attach the second portion to the first portion, the locking features of the two portions are mated and a lock is engaged. According to various examples, this can include a retractable protrusion in the first portion that mates with a cavity in the second portion. Upon insertion of the first portion into the second portion, the protrusions are forced into a refracted position, similar to the action of a door latch, and once fully inserted, the protrusions snap into place with an audible click. A mechanism on the first portion, such as a button, knob, or lever, retracts the protrusions to allow for removal of the second portion.

In one example, a retractable protrusion in the first portion mates with a cavity in the second portion. A knob on the first portion adjusts the protrusions between a protruded position and a retracted position. Upon insertion of the first portion into the second portion, the knob holds the protrusions in the retracted position. After insertion, the knob is adjusted to the protruded position, locking the first portion in place. In one example, after the second portion is locked in place, retracting the protrusions results in the destruction of the cavities, such that a used second portion cannot be locked to the first portion again.

In one example, the second portion includes a laminated stack of adhesive layers, with the bottom-most layer contacting the patient tissue and sticking to the patient during use. After each use, the bottom-most adhesive layer is removed and thus exposing a fresh adhesive layer.

In one example, the second portion and the first portion includes mating features that create a friction-fit interface. To attach the second portion to the first portion, the two portions are pressed together, forcing the friction-fit mating features together. In one example, this includes a tongue-in-groove configuration. In one example, this includes a raised outer radius on the second portion that captures the perimeter of the first portion. In one example, a raised cylindrical node is configured to fit into a matching recess, similar to children's plastic bricks. In one example, the second portion and the first portion include mating features that snap together. To attach the second portion to the first portion, the two portions are pressed together, snapping the mating features together. For example, the interface can include a spline configured to engage a groove, a snap-fit element, or the second portion includes snap-fit hooking posts protruding from the mating surface, and the first portion includes recessed areas around the perimeter, such that when the first portion is pressed into the second portion, the perimeter of the first portion is captured in place by the hooking posts.

In one example, the second portion and the first portion mate together using a magnetic interface. One portion includes a magnet, such as high-pull rare earth magnet, and the other portion includes a piece of ferromagnetic material, such as iron.

In one example, the second portion has a larger footprint than the first portion and the second portion is applied over the top of the first portion, such that the first side of the second portion contacts both the patient tissue and a top side of the first portion. The surface of the second portion extends beyond the perimeter of the first portion and contains an adhesive layer that bonds to the patient and holds the segmented sensor in place. The second portion and the first portion can be attached together using a variety of couplings.

In one example, the second portion forms an envelope or sleeve which surrounds the first portion. The sleeve is sealed closed around the first portion. This sealing could be accomplished through a variety of features, including a hook-and-loop strip, adhesive, or a locking strip similar to those used in a zip top plastic bag. The second portion contains windows aligned with the emitters and detectors of the first portion that allow the radiation to pass through unimpeded. The first portion and the second portions can be coupled using a variety of coupling features.

The spacing and arrangement of optical elements (for example, the distance between the emitters and detectors) remains static and is set by the construction of the first portion. A variety of different sensor spacing can be provided, such as 20 mm spacing for an adult patient, and 15 mm spacing for a pediatric patient. In one example, the segmented sensor includes user-selectable spacing and arrangement of optical elements. In one example, the emitters and detectors are disposed at the end of flexible "fingers" which couple with receptacles in the second portion.

In one example, the second portion includes "pipes" which guide the radiation from the emitters and detectors, configured for a particular spacing, to emission and detection sites on the patient in a different spacing. The pipes can be fabricated of optical fibers or clear plastic.

The attachment configuration between the first and second portion can allow multiple shapes of the second portion, such that site-specific shapes are available to the end user. For example, a segmented sensor for the forehead may include a different adhesive shape than a segmented sensor for a calf muscle.

In one example, the second portion includes at least one information element, and the first portion includes at least one sensing element. The sensing element in the first portion detects and/or communicates with the information element in the second portion without requiring electrical contact between the two elements. In one example, the coupling includes radio frequency signaling. An RFID tag in the second portion communicates with an RFID reader in the first portion. In one example, a magnetic sensor is provided. A magnetic element in the second portion and a Hall Effect sensor in the first portion can provide a wireless coupling. One example includes a capacitive sensing configuration, such as conductive elements in the first and second portions, coupled with an oscillator circuit. One example includes an inductive sensing configuration, such as a conductive element in the first portion, and a coil in the second portion. One example includes an optical signaling configuration, such as an LED coupled with a microcontroller and an energy source in the second portion, and a photodiode in the first portion.

The information element allows the first portion of the segmented sensor to detect the presence of the second

portion. The information element can include information pertaining to characteristics of the segmented sensor, such as sensor type. The information element can include security measures such as encryption to ensure quality of second portions.

Various Notes & Examples

Each of these non-limiting examples can stand on its own, or can be combined in various permutations or combinations with one or more of the other examples.

The above detailed description includes references to the accompanying drawings, which form a part of the detailed description. The drawings show, by way of illustration, specific embodiments in which the invention can be practiced. These embodiments are also referred to herein as “examples.” Such examples can include elements in addition to those shown or described. However, the present inventors also contemplate examples in which only those elements shown or described are provided. Moreover, the present inventors also contemplate examples using any combination or permutation of those elements shown or described (or one or more aspects thereof), either with respect to a particular example (or one or more aspects thereof), or with respect to other examples (or one or more aspects thereof) shown or described herein.

In the event of inconsistent usages between this document and any documents so incorporated by reference, the usage in this document controls.

In this document, the terms “a” or “an” are used, as is common in patent documents, to include one or more than one, independent of any other instances or usages of “at least one” or “one or more.” In this document, the term “or” is used to refer to a nonexclusive or, such that “A or B” includes “A but not B,” “B but not A,” and “A and B,” unless otherwise indicated. In this document, the terms “including” and “in which” are used as the plain-English equivalents of the respective terms “comprising” and “wherein.” Also, in the following claims, the terms “including” and “comprising” are open-ended, that is, a system, device, article, composition, formulation, or process that includes elements in addition to those listed after such a term in a claim are still deemed to fall within the scope of that claim. Moreover, in the following claims, the terms “first,” “second,” and “third,” etc. are used merely as labels, and are not intended to impose numerical requirements on their objects.

Method examples described herein can be machine or computer-implemented at least in part. Some examples can include a computer-readable medium or machine-readable medium encoded with instructions operable to configure an electronic device to perform methods as described in the above examples. An implementation of such methods can include code, such as microcode, assembly language code, a higher-level language code, or the like. Such code can include computer readable instructions for performing various methods. The code may form portions of computer program products. Further, in an example, the code can be tangibly stored on one or more volatile, non-transitory, or non-volatile tangible computer-readable media, such as during execution or at other times. Examples of these tangible computer-readable media can include, but are not limited to, hard disks, removable magnetic disks, removable optical disks (e.g., compact disks and digital video disks), magnetic cassettes, memory cards or sticks, random access memories (RAMs), read only memories (ROMs), and the like.

The above description is intended to be illustrative, and not restrictive. For example, the above-described examples

(or one or more aspects thereof) may be used in combination with each other. Other embodiments can be used, such as by one of ordinary skill in the art upon reviewing the above description. The Abstract is provided to allow the reader to quickly ascertain the nature of the technical disclosure. It is submitted with the understanding that it will not be used to interpret or limit the scope or meaning of the claims. Also, in the above Detailed Description, various features may be grouped together to streamline the disclosure. This should not be interpreted as intending that an unclaimed disclosed feature is essential to any claim. Rather, inventive subject matter may lie in less than all features of a particular disclosed embodiment. Thus, the following claims are hereby incorporated into the Detailed Description as examples or embodiments, with each claim standing on its own as a separate embodiment, and it is contemplated that such embodiments can be combined with each other in various combinations or permutations. The scope of the invention should be determined with reference to the appended claims, along with the full scope of equivalents to which such claims are entitled.

The claimed invention is:

1. A device comprising:

a first portion having a planar substrate and having at least one optical element coupled to the substrate, the optical element configured to emit or detect light aligned substantially normal to the substrate, the first portion having an electrical circuit;

an information element sensor coupled to the substrate; a second portion having a membrane configured to couple to the planar substrate, the membrane having a contact surface configured to couple with tissue wherein the membrane is configured to occlude ambient light and wherein the membrane includes at least one transparent aperture aligned with the at least one optical element and wherein the aperture is configured to pass optical radiation between the at least one optical element and the contact surface;

an encoder element coupled to the membrane; and wherein the circuit is configured to read data from the encoder element or configured to detect proximity of the encoder element relative to the information element sensor and wherein the circuit is configured to enable or disable the at least one optical element based on a wireless coupling between the first portion and the second portion and based on information encoded in the encoder element.

2. The device of claim 1 wherein the contact surface includes an adhesive.

3. The device of claim 1 wherein the optical element is configured to emit radiation.

4. The device of claim 1 wherein the optical element is configured to detect radiation.

5. The device of claim 1 wherein the optical element includes at least one radiation emitter and at least one photodetector and wherein the photodetector is configured to provide an electrical signal corresponding to oximetry.

6. The device of claim 1 wherein the encoder element includes at least one of a magnet, a capacitor plate, a metal object, and a radio frequency identification device.

7. The device of claim 1 wherein the information element sensor includes at least one of a magnet sensor, a capacitor plate, an inductor, and a radio frequency identification device reader.

8. The device of claim 1 wherein the membrane includes a mechanical alignment feature configured to engage with the planar substrate.

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9. The device of claim 1 further including a wireless communication module coupled to the first portion, wherein the wireless communication module is configured to communicate wirelessly with a remote device.

10. A method comprising:

coupling an electrical circuit and an optical element to a first portion having a planar substrate, the optical element configured to emit or detect light aligned substantially normal to the substrate;

coupling an information element sensor to the substrate;

configuring a second portion having a membrane to couple with the planar substrate, the membrane having an adhesive surface configured to bond with tissue and wherein the membrane is configured to occlude ambient light and wherein the membrane includes a transparent aperture aligned with the optical element and wherein the aperture is configured to pass light between the optical element and the adhesive surface;

coupling an encoder element to the membrane; and wherein the circuit is configured to read data from the encoder element or configured to detect proximity of the encoder element relative to the information element sensor and wherein the circuit is configured to enable or disable the optical element based on a wireless cou-

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pling between the first portion and the second portion and based on information encoded in the encoder element.

11. The method of claim 10 further including configuring the optical element to emit radiation.

12. The method of claim 10 further including configuring the optical element configured to detect radiation.

13. The method of claim 10 further including configuring the optical element to provide a signal corresponding to at least one of regional oximetry, pulse oximetry, or a DC-based parameter.

14. The method of claim 10 wherein coupling the encoder element includes coupling at least one of a magnet, a capacitor plate, an inductor, and a radio frequency identification device.

15. The method of claim 10 wherein coupling the information element sensor to the substrate includes coupling at least one of a magnet sensor, a capacitance sensor, an inductance sensor, and a radio frequency identification device reader.

16. The method of claim 10 further including coupling a wireless communication module coupled to the first portion, wherein the wireless communication module is configured to communicate wirelessly with a remote device.

* * * * *

专利名称(译)	分段传感器		
公开(公告)号	US10342467	公开(公告)日	2019-07-09
申请号	US14/644972	申请日	2015-03-11
[标]申请(专利权)人(译)	NONIN医疗		
申请(专利权)人(译)	NONIN MEDICAL , INC.		
当前申请(专利权)人(译)	NONIN MEDICAL , INC.		
[标]发明人	SKORICH STEFAN KANE BRIAN JOSEPH PRIOR MATTHEW		
发明人	SKORICH, STEFAN KANE, BRIAN JOSEPH PRIOR, MATTHEW		
IPC分类号	A61B5/1455 A61B5/00 A61B5/024		
CPC分类号	A61B5/14552 A61B5/0002 A61B5/02416 A61B5/6831 A61B5/6832 A61B2562/164 A61B2562/0223 A61B2562/0238 A61B2562/04 A61B2562/146 A61B5/0024 A61B5/68335		
代理机构(译)	SCHWEGMAN和Lundberg & 沃斯纳 , P.A.		
其他公开文献	US20160262673A1		
外部链接	Espacenet		

摘要(译)

一种装置包括第一部分，信息元件，第二部分和编码器元件。第一部分包括平面基板和耦合到基板的至少一个光学元件。光学元件具有轴，并且轴基本垂直于基板。信息元件传感器耦合到基板。信息元素传感器被配置为提供信息信号。第二部分包括被配置为耦合到平面基板的膜。膜具有配置成与组织耦合的接触表面。编码器元件耦合到膜。信息信号对应于编码器元件的接近度。

