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(54) **SYSTEMS AND METHODS FOR OPERATING AN ALERT SYSTEM OF MEDICAL DEVICES**

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(51) **Int. Cl.**
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A61B 5/00 (2006.01)

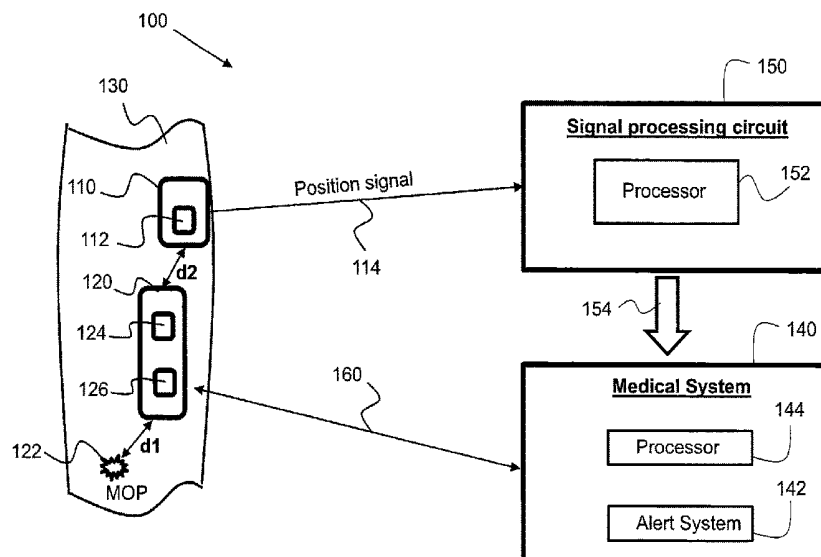
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(57) **ABSTRACT**

A system for a medical system includes a position sensing unit to sense a displacement of a medical device from a medically operational point on a subject, and a signal processing circuit to output, based on the sensed displacement, a signal to disable an alert system when the medical device is for sensing a physiological parameter of a subject during a medical procedure and the displacement is greater than a first threshold value, or to output a signal to enable the alert system when the medical device is for delivering treatment to the subject and the displacement is greater than a second threshold value.

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A61M 16/08 (2006.01)
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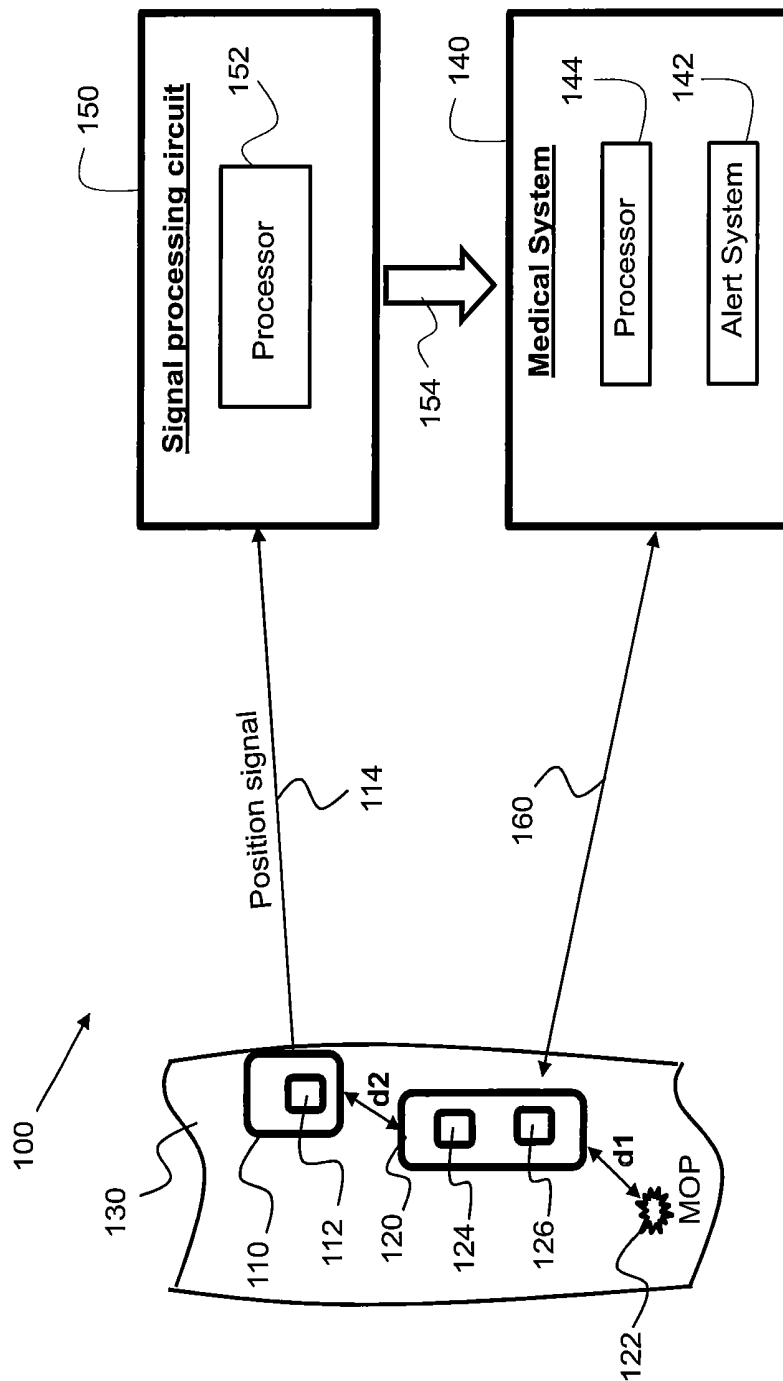


Fig. 1

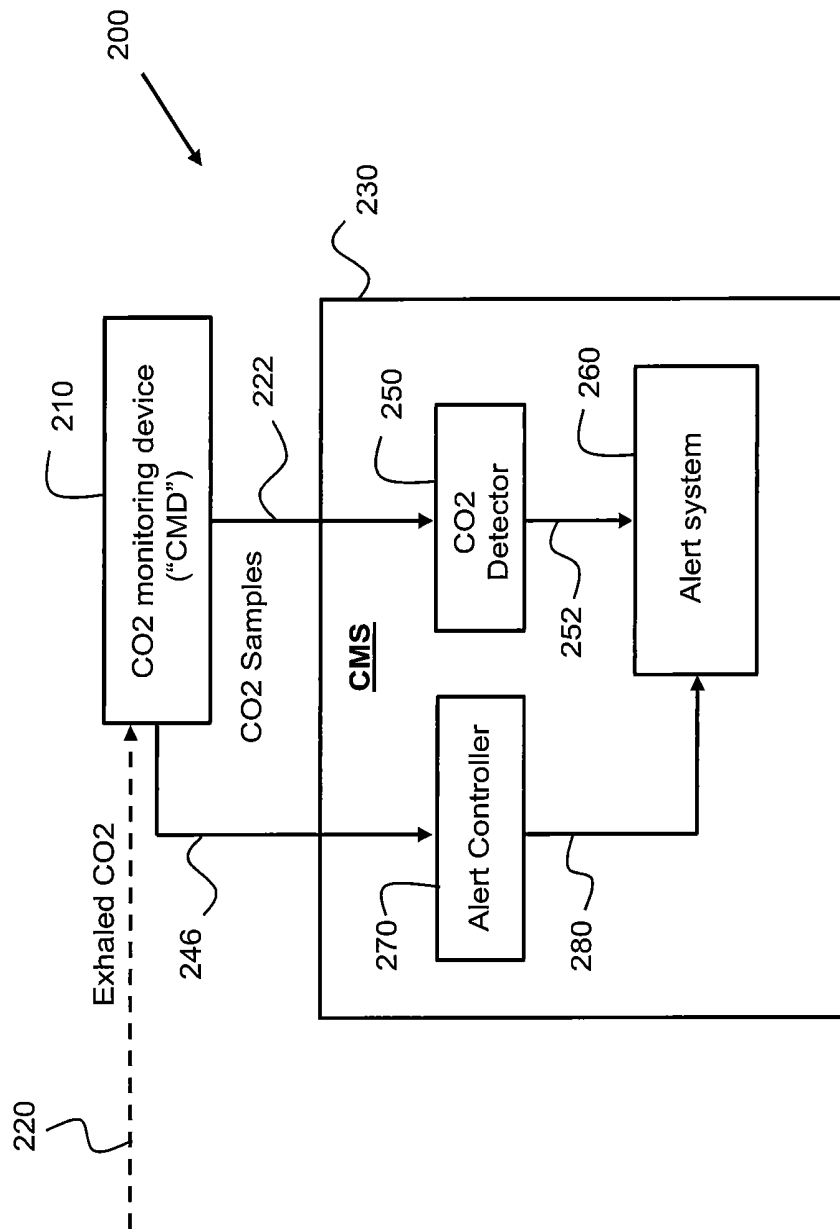


Fig. 2

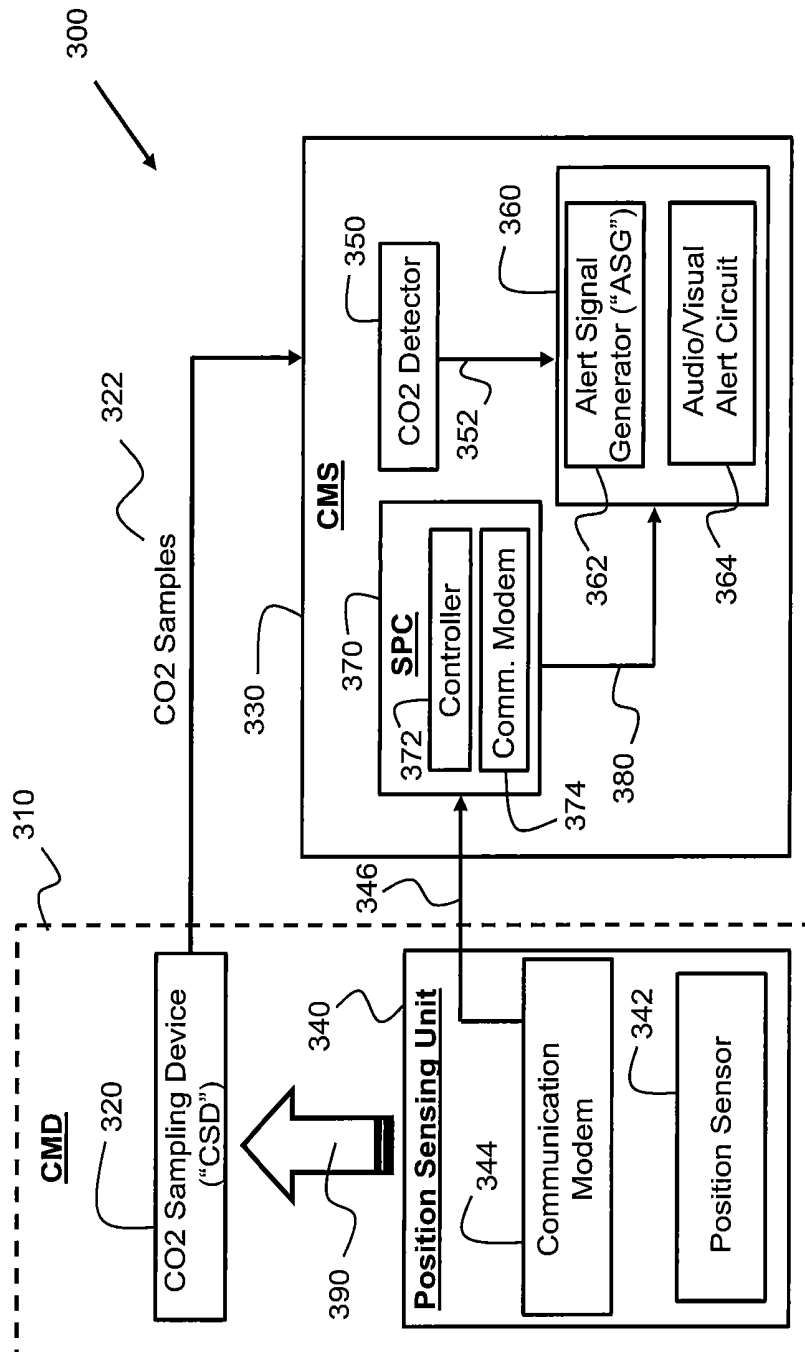


Fig 3

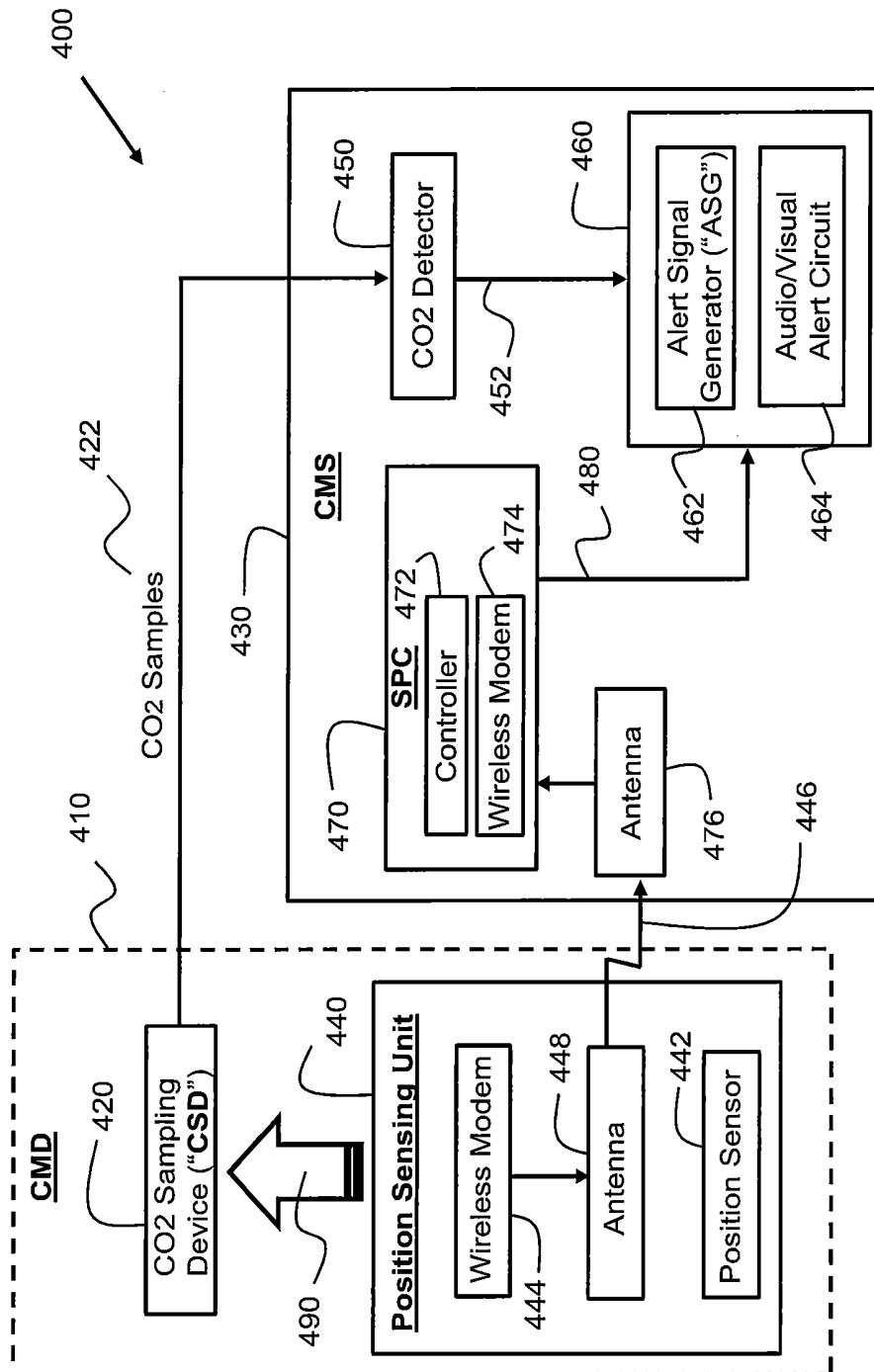


Fig. 4A

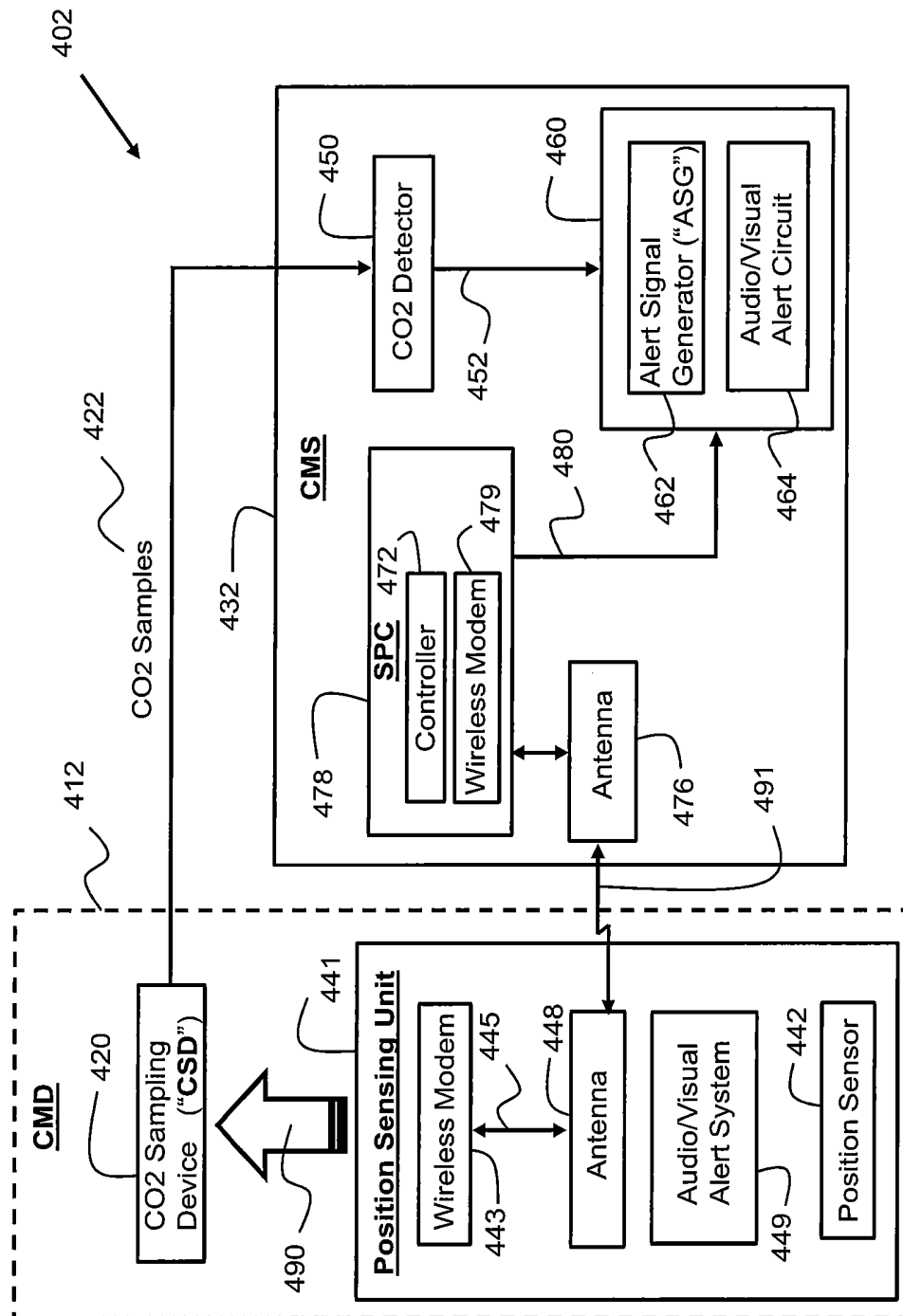


Fig. 4B

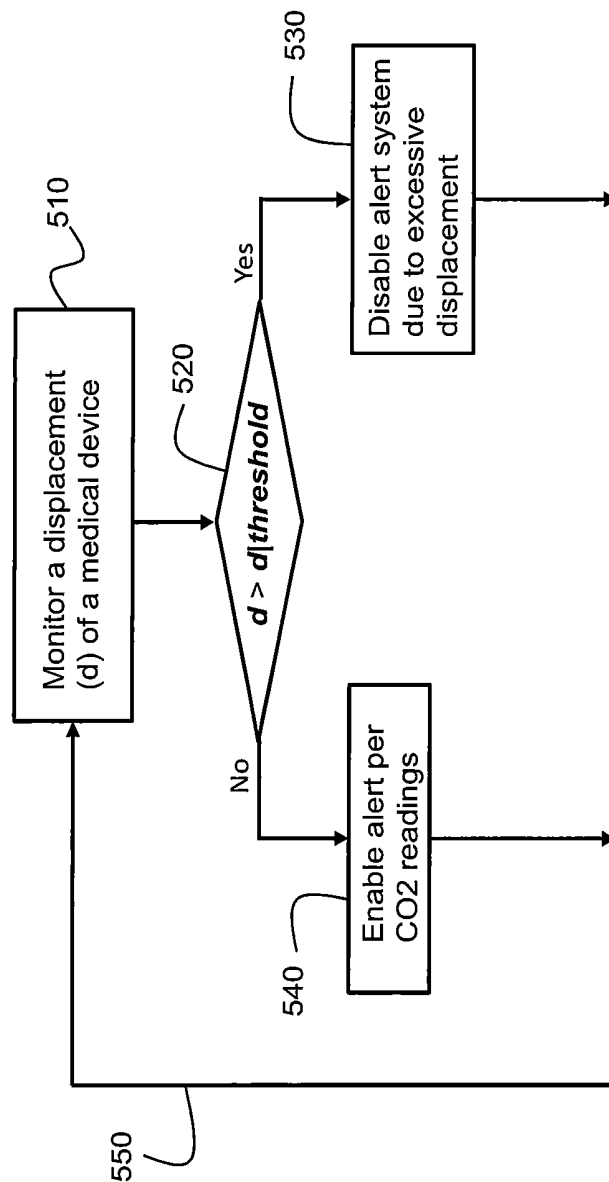


Fig. 5

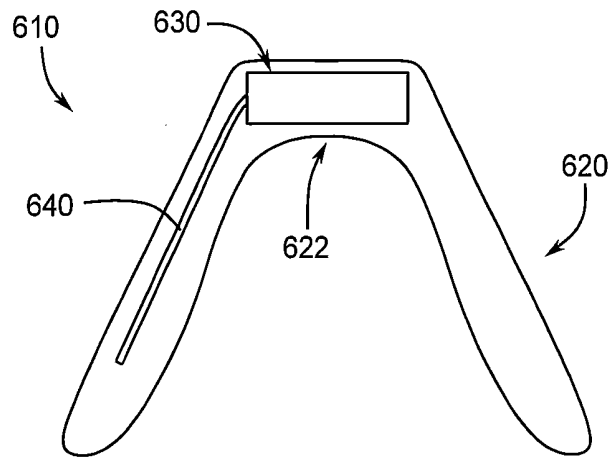


Fig. 6A

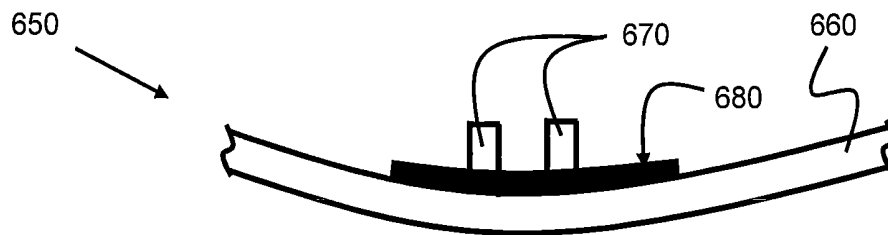


Fig. 6B

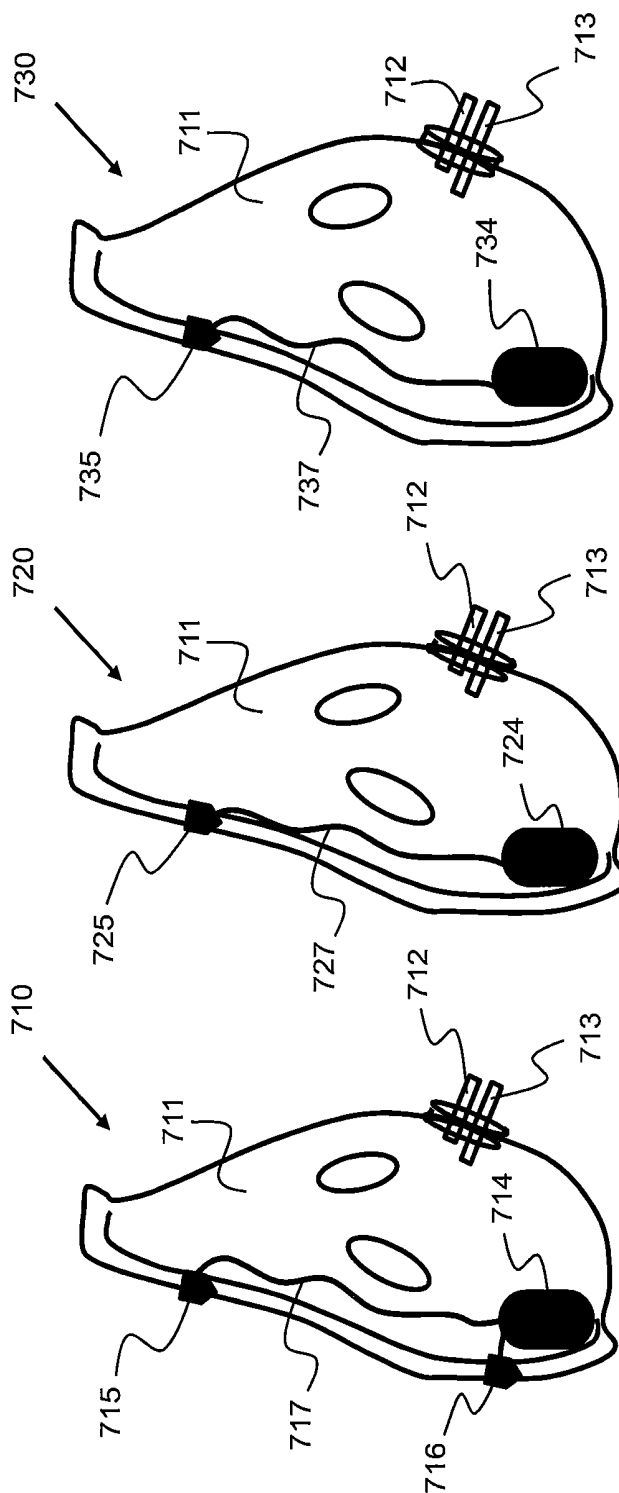


Fig. 7A

Fig. 7B

Fig. 7C

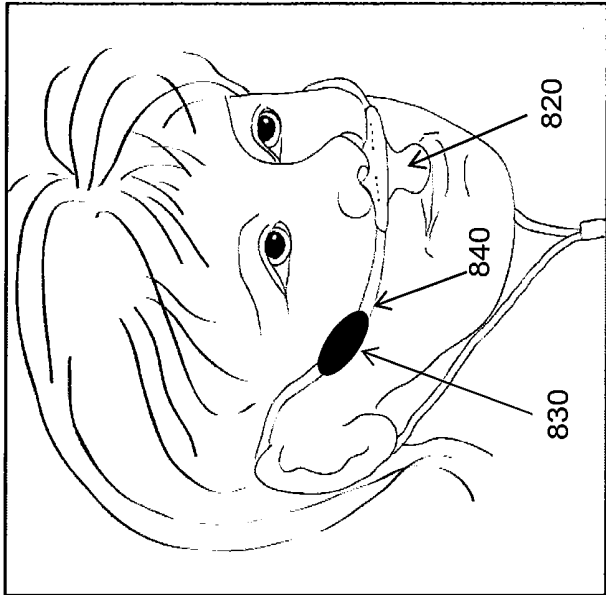


Fig. 8B

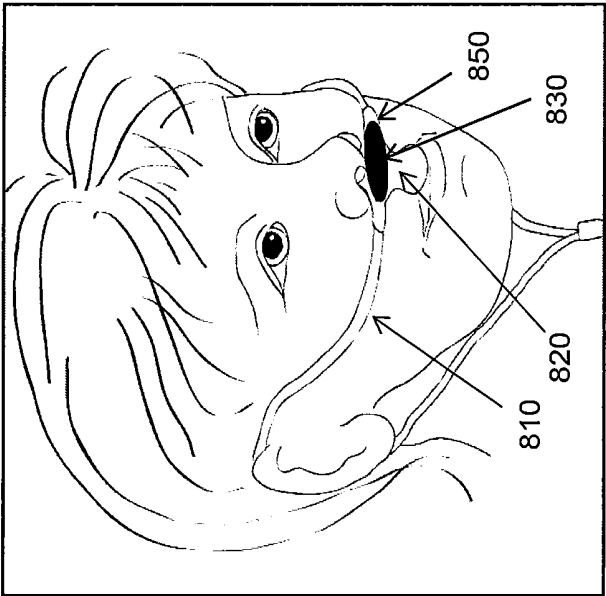


Fig. 8A

SYSTEMS AND METHODS FOR OPERATING AN ALERT SYSTEM OF MEDICAL DEVICES

CROSS-REFERENCE TO RELATED APPLICATION

The present application claims priority to and the benefit of U.S. Provisional Application No. 62/558,715, entitled "SYSTEMS AND METHODS FOR OPERATING AN ALERT SYSTEM OF MEDICAL DEVICES," filed Sep. 14, 2017, the content of which is incorporated herein by reference in its entirety for all purposes.

TECHNOLOGICAL FIELD

The present disclosure generally relates to medical systems and, more particularly, to systems and methods for disabling an alarm system in a medical system in order to prevent (e.g., reduce or block) false alarms during a medical procedure that is rendered to a subject.

BACKGROUND

This section is intended to introduce the reader to various aspects of art that may be related to various aspects of the present techniques, which are described and/or claimed below. This discussion is believed to be helpful in providing the reader with background information to facilitate a better understanding of the various aspects of the present disclosure. Accordingly, it should be understood that these statements are to be read in this light, and not as admissions of prior art.

Some medical systems include a medical device (for example, cannula, mask, nasal tubing, carbon dioxide [CO₂] sampling device, physiological sensor, etc.) that is attached to a patient (for example to a face, nose, index finger, wrist, etc.) and by which a physiological parameter of the patient is monitored. Sometimes, a patient may accidentally remove or move the medical device from its intended location, causing false measurements of the patient's physiological parameter. Often, movement of the medical device is intentional, as a patient wearing or using the medical device moves it aside in order to be able, for example, to talk or to eat more comfortably.

Some medical systems are designed to output an alert when a monitored physiological parameter indicates a medical problem, or when their operation is compromised (e.g., gets interrupted). However, some medical systems may also output nuisance alerts as a result of false measurements, for example, due to accidental removal or movement of the medical device. Nuisance alerts may be distracting to the medical staff and may result in inefficient medical care, for example.

SUMMARY

A summary of certain embodiments disclosed herein is set forth below. It should be understood that these aspects are presented merely to provide the reader with a brief summary of these certain embodiments and that these aspects are not intended to limit the scope of this disclosure. Indeed, this disclosure may encompass a variety of aspects that may not be set forth below.

It would, therefore, be beneficial to provide a system (e.g., a nuisance alert preventing system) that can prevent (e.g., reduce or block) nuisance alerts in medical systems. For example, the system may include components that can be

mounted in or on various medical devices to detect movement or removal of the medical devices from a medically operational point (MOP) on a body of a subject during a medical procedure, and components that are configured to prevent nuisance alerts by the related medical system whenever the system detects such movement or removal of the medical device. It would also be beneficial to provide a system that can trigger an alert when a medical device that delivers treatment to a patient ((for example oxygen or medication via a nebulizer, venturi mask, or the like) moves (e.g., unintentionally moves) or is removed from a MOP.

In some embodiments, a system to prevent nuisance alerts in a medical system, or to trigger an alert when this is justified, may include a position sensing unit configured to sense a displacement of a medical device from a medically operational point on a subject, and a signal processing circuit to output, based on the sensed displacement, a signal to enable an alert system of a medical system cooperating with the medical device when the displacement is less than a threshold value, and to output a signal to disable the alert system when the displacement is equal to or greater than the threshold value.

In some embodiments, an alert activation/deactivation system is provided for a medical system and may include a position sensing unit to sense a displacement of a medical device from a medically operational point on a subject, and a signal processing circuit to output, based on the sensed displacement, a signal to disable an alert system of a medical system cooperating with the medical device if the medical device is for sensing a physiological parameter of a subject during a medical procedure and the displacement is greater than a first threshold value, and to output a signal to enable the alert system when the medical device is for delivering treatment to the subject and the displacement is greater than a second threshold value. The medical device may be configured to deliver treatment (e.g., oxygen, medication, etc.) to the subject. The physiological parameter may be oxygen saturation level, heart pulse rate, blood pressure, electrocardiogram (ECG), temperature and impedance, to name a few.

BRIEF DESCRIPTION OF THE DRAWINGS

Various exemplary embodiments are illustrated in the accompanying figures with the intent that these examples not be restrictive. It will be appreciated that for simplicity and clarity of the illustration, elements shown in the figures referenced below are not necessarily drawn to scale. Also, where considered appropriate, reference numerals may be repeated among the figures to indicate like, corresponding or analogous elements. Of the accompanying figures:

FIG. 1 schematically illustrates a nuisance alert preventing system, according to an embodiment of the present disclosure;

FIG. 2 schematically illustrates a CO₂ monitoring system and a nuisance alert preventing system, according to an embodiment of the present disclosure;

FIG. 3 schematically illustrates a CO₂ monitoring system and a nuisance alert preventing system, according to an embodiment of the present disclosure;

FIG. 4A schematically illustrates a CO₂ monitoring system and a nuisance alert preventing system that wirelessly communicate with one another, according to an embodiment of the present disclosure;

FIG. 4B schematically illustrates a nuisance alert preventing system having an audio/visual alert system associated with a position sensing unit, according to an embodiment of the present disclosure;

FIG. 5 shows a method for preventing nuisance alerts in medical system, according to an embodiment of the present disclosure;

FIG. 6A shows a nose clip, according to an embodiment of the present disclosure;

FIG. 6B shows a cannula, according to an embodiment of the present disclosure;

FIG. 7A illustrates an oxygen/capnography mask, in accordance with an embodiment of the present disclosure;

FIG. 7B illustrates an oxygen/capnography mask, in accordance with an embodiment of the present disclosure;

FIG. 7C illustrates an oxygen/capnography mask, in accordance with an embodiment of the present disclosure;

FIG. 8A illustrates a cannula, in accordance with an embodiment of the present disclosure; and

FIG. 8B illustrates a cannula in accordance with an embodiment of the present disclosure.

DETAILED DESCRIPTION

One or more specific embodiments of the present disclosure will be described below. These described embodiments are only examples of the presently disclosed techniques. Additionally, in an effort to provide a concise description of these embodiments, all features of an actual implementation may not be described in the specification. It should be appreciated that in the development of any such actual implementation, as in any engineering or design project, numerous implementation-specific decisions must be made to achieve the developers' specific goals, such as compliance with system-related and business-related constraints, which may vary from one implementation to another. Moreover, it should be appreciated that such a development effort might be complex and time consuming, but may nevertheless be a routine undertaking of design, fabrication, and manufacture for those of ordinary skill having the benefit of this disclosure. Further, the current embodiments may be implemented by one or more computer-processors that implement one or more machine-readable instructions stored on a tangible, non-transitory, machine-readable medium and/or by specialized circuitry designed to implement the discussed features.

During a medical procedure or at various other times, a medical device may be positioned at an intended location on the body of the patient. The intended location may be the location at which the medical device is usually attached in order to obtain optimal operation (e.g., generally desirable or effective operation) of the medical system as a whole, for example, in order to obtain optimal measurements (e.g., reliable or accurate measurements) of a physiological parameter and/or in order to provide optimal medical care (e.g., appropriate or intended medical care). The intended location is referred to herein as a medically operational point (MOP).

As noted above, some medical systems are designed to output an alert when a monitored physiological parameter indicates a medical problem, or when their operation is compromised (e.g., gets interrupted). However, some medical systems may also output nuisance alerts as a result of false measurements, for example, due to removal or movement of the medical device from the MOP. For example, a medical system monitoring a physiological parameter of a patient (e.g., oxygen saturation level in a subject, concen-

tration level of exhaled CO₂, etc.) outputs an alert when the value of a measured, or sensed, physiological parameter is lower than a threshold. In another example, a capnography system is designed to normally output a CO₂ level alert when the concentration of CO₂ exhaled by the patient is lower than a threshold value. Continuing the examples, a patient may accidentally remove or move the cannula, or mask, through which the CO₂ monitoring system obtains CO₂ samples, from its medically operational point, thus causing a nuisance alert.

FIG. 1 schematically illustrates a nuisance alert preventing system 100 (e.g., nuisance alert blocking system) according to an example embodiment. Nuisance alert preventing system 100 may include a position sensing unit (PSU) 110. PSU 110 may include a position sensor 112 configured to output a signal 114 (e.g., a position signal) that represents a displacement d1 of a medical device 120 from a MOP 122 on a subject 130. Medical device 120 is configured to cooperate with a medical system 140 to perform a medical procedure. Nuisance alert preventing system 100 may also include a signal processing circuit (SPC) 150 that includes, among other things, a processor 152. Processor 152 may be configured to output 154, to medical system 140, based on position signal 114, a signal to enable an alert system 142 of medical system 140 when position signal 114 indicates a displacement d1 of medical device 120, which is smaller, less or lower than a threshold value, and to output a signal to disable alert system 142 when position signal 114 indicates an excess displacement, that is, a displacement d1 that is equal to or greater than the threshold value. In some embodiments, the value of displacement d1 may be set to a reference value (e.g., baseline, first, or initial value or position) that represents an optimal alignment (e.g., baseline, first, or initial alignment) of medical device 120 with MOP 122, and a deviation from the reference value of d1 may be interpreted (e.g., by medical system 140 or SPC 150) as misalignment, or displacement, of medical device 120 from MOP 122.

In some embodiments, a sensing facilitating means 124 (e.g., element detectable by position sensor 112) may be incorporated in or attached to medical device 120 and configured to facilitate sensing, by or in conjunction with position sensor 112, a displacement d1 of medical device 120 from MOP 122. For example, position sensor 112 may be or include a proximity sensor (e.g., switch), and sensing facilitating means 124 may be or include a metal body that is designed to change the state (on/off or open/closed) of the proximity sensor/switch as a function of the distance between the two devices. In some embodiments, sensing facilitating means 124 may be or include a miniature transponder (e.g., a radio frequency identification (RFID) tag/system) that may be designed to cooperate with position sensor 112 (that may be, in this example, a conjugated RFID reader) when the two devices are within a certain distance from one another. Briefly, an RFID system uses electromagnetic fields to automatically identify and track an RFID tag that is attached to an object. RFID tags contain electronically stored information. Passive RFID tags, for example, collect energy from interrogating radio waves transmitted from a nearby RFID reader. In some embodiments, position sensor 112 may be a proximity sensor/switch, and sensing facilitating means 124 may be, or include, in these embodiments, a metallic element as a target object to which the proximity sensor/switch reacts (e.g., by changing state from "close" to "open", and vice versa). It should be appreciated that in some embodiments, the position sensor 112 may be coupled to or incorporated into the medical device 120, while the

sensing facilitating means **124** is a separate component that may be coupled to the subject **130** at a location at or near the MOP **122**, for example.

Medical device **120** functionally cooperates **160** (e.g., via wireless or wired communication) with medical system **140** to perform a medical procedure. The medical procedure that medical system **140** may perform may be, for example, monitoring a physiological parameter which may be, for example, the concentration level CO₂ of the CO₂ exhaled by the subject, the oxygen saturation level in the subject's blood, the blood pressure of the subject, the heart pulse rate of the subject, electrocardiogram (ECG) monitoring by an ECG monitoring system, temperature, tissue impedance, etc. The medical procedure that medical system **140** may perform may be delivery of oxygen to the subject, administering medication to the subject, providing nutrition, etc. Cooperation **160** means that a processor **144** of medical system **140** may control transfer of oxygen and/or medication to the subject **130** via medical device **120**, or that processor **144** may receive sensory signal(s) from medical device **120**, which represent a physiological parameter(s) of the subject **130**. Processor **144** may receive and process the signal **114** or the output **154** that is indicative of the position of the medical device **120** relative to the MOP **122** and, based on the processing result, processor **144** may enable or disable alert system **142**.

Nuisance alert preventing system **100** may also include a signal processing circuit (SPC) **150**. Signal processing circuit **150** may include a processor **152**. Based on position signal **114**, processor **152** may output **154**, to or for medical system **140**, a first signal to enable an alert system **142** of medical system **140** when position signal **114** indicates that medical device **120** and MOP **122** are aligned (for example, when position signal **114** indicates a relative displacement that is less than a threshold value). Based on position signal **114**, processor **152** may also output **154** a second signal to medical system **140**, to disable alert system **142** when the position signal **114** indicates that medical device **120** is misaligned with (e.g., displaced from) MOP **122** (for example, when position signal **114** indicates an excess displacement, that is, displacement that is equal to or greater than a threshold value).

Medical device **120** may be or include one or more of: a cannula, a nasal tubing, a face mask, a CO₂ sampling device to collect exhaled CO₂ for the CO₂ monitoring system, an oxygen delivery device (e.g., mask, tube) to deliver oxygen to the subject, and a physiological parameter sensor (e.g., physiological parameter sensor **126**) to sense a physiological parameter of the subject. Physiological parameter sensor **126** may be, for example, an oxygen sensor, a heart pulse rate sensor, a blood pressure sensor, a temperature sensor, ECG and an impedance sensor, to name a few. It should be appreciated that additional sensors may be used to sense different physiological parameters of the subject. In some embodiments, the first signal and the second signal that processor **152** of SPC **150** may output **154** are, respectively, to enable and disable alert system **142** of medical system **140** with respect to sensed values of the physiological parameter. In some embodiments, if, for example, medical device **120** is an oxygen delivery device, the first signal and the second signal that processor **152** of SPC **150** output **154** may, respectively, enable and disable alert system **142** in the related oxygen monitoring system **140** with respect to the delivery of oxygen to the subject.

Medically operational point (MOP) **122** on the subject may be selected from the group consisting of, for example: the face of the subject, the nose of the subject, the mouth of

the subject, an index finger of the subject, a wrist of the subject, an elbow of the subject and the chest of the subject. In some embodiments, position sensor **112** may be positioned at a predetermined distance **d2** from medical device **120**, or position sensor **112** may be incorporated in or be attached to the medical device **120** (e.g., via a clip, fastener, tether, adhesive, or the like). In some embodiments, PSU **110**, or only position sensor **112**, is part of, incorporated in or attached to medical device **120**. In some embodiments, SPC **150** is incorporated in medical system **140**.

Position sensor **112** may be selected from the group consisting of, for example: an impedance sensor, a temperature sensor, a pressure sensor, a capacitive sensor, a proximity sensor, a motion sensor, an acceleration sensor and an optical sensor. Position signal **114** may be selected from the group of signals consisting of, for example: an electrical signal and a wireless signal. For example, PSU **110** may be wired to SPC **150**, and position signal **114** may be transferred to signal processing circuit **150** via an electrical wire or cable. In another example, PSU **110** may use a radio frequency ("RF") communication circuit, for example a Bluetooth circuit, to transfer position signal **114** to SPC **150** wirelessly.

In some embodiments each one of PSU **110** and SPC **150** may include a communication modem (e.g., **344, 374** FIG. **3**) to enable the transfer of position signal **114** to SPC **150**. As used herein, a modem may be a transmitter configured to wirelessly transmit signals, a receiver configured to wirelessly receive signals, or a transceiver configured to wirelessly transmit and receive signals to facilitate communication between components. In some embodiments, at least one of position sensor **112** and the communication modem of PSU **110** may be an add-on device that is attachable (e.g., removably attachable) to medical device **120**, and thus, may be attached to the medical device **120** after manufacturing of the medical device **120** (e.g., at a clinical site prior to the medical procedure) and/or may be detached from the medical device **120** and attached to another medical device to enable reuse with multiple different medical devices **120**. In some embodiments, at least one of position sensor **112** and the communication modem of PSU **110** may be embedded in medical device **120**. In some embodiments, each communication modem is a bi-directional modem (e.g., modems **544** and **574**, FIG. **5**). In some embodiments, the communication modems are wireless. In some embodiments, the wireless modems are Bluetooth modems.

In some embodiments medical system **140** may be or include a CO₂ monitoring system (e.g., system **200**, FIG. **2**), that is, a capnograph system, that is configured to measure CO₂ concentration level (or another CO₂-related parameter), or an oxygen delivery system to deliver oxygen to the subject, or an oximeter to measure oxygen saturation, or a heart pulse rate monitoring system to measure the subject's pulse rate, or an ECG system. "Capnography" is, in general, the monitoring of the concentration level or partial pressure of CO₂ in the respiratory gases.

In some embodiments, the first signal that processor **152** may output **154** to medical system **140** is configured (is an instruction or message) to enable normal operation of a CO₂ alert circuit (e.g., CO₂ alert circuit **260**, FIG. **2**) associated with the CO₂ monitoring system (e.g., CO₂ monitoring system **230**, FIG. **2**), and the second signal that processor **152** may output **154** to medical system **140** is configured (is an instruction or message) to disable the CO₂ alert circuit (e.g., CO₂ alert circuit **260**, FIG. **2**).

In some embodiments, the CO₂ monitoring system (e.g., CO₂ monitoring system **300**, FIG. **3**) may include a CO₂

monitoring device (e.g., CO₂ monitoring device [CMD] 310). The CMD (e.g., CMD 310) may include a CO₂ sampling device (e.g., CO₂ sampling device [CSD] 320, FIG. 3) as an example medical device. The CSD (e.g., CSD 320) may be designed, or it may include a device that is designed, to collect (e.g., 220, FIG. 2) CO₂ samples of CO₂ that is exhaled by the subject. For example, the device collecting the CO₂ samples may be designed as a scoop or as a funnel. The CMD (e.g., CMD 310) may also include or be coupled to the PSU (e.g., PSU 340), which may be configured to output the position signal (e.g., 246, 346) that represents the position of the CO₂ sampling device on the subject (e.g., position relative to the MOP and/or relative to a reference position). CO₂ monitoring system 230 may be configured to receive 222, detect and process, by CO₂ detector 250, the CO₂ samples. Concurrently to CO₂ detector 250 receiving 222 the CO₂ samples, CO₂ detector 250 receives, by alert controller 270, a position signal 246 from CO₂ monitoring device 210 in order to determine whether the CO₂ sampling device (e.g., CSD 320) was removed or moved from the designated MOP at the time when the related CO₂ samples were collected by the CSD.

Carbon dioxide monitoring system 230 may be further configured to conditionally output, by alert system 260, an alert signal when a value associated with the processed CO₂ samples exceeds a predetermined level or range, and the condition to output the alert signal is that the CO₂ sampling device was not removed or moved from the MOP (that is, it is still at the MOP). Alert controller 270 processes position signal 246 in order to determine whether the CO₂ sampling device has moved or been removed from the designated MOP, and outputs 280 to alert system 260 an “enable” or a “disable” signal according to the determination. In other words, CO₂ monitoring system 230 is allowed to output an alert only if alert controller 270 determines (based on position signal 246) that the CO₂ sampling device is still at (e.g., aligned with) the designated MOP; otherwise, alert controller 270 disables 280 alert system 260.

An example method of operating the system of FIG. 1 is described below. In some embodiments, PSU 110 may be attached to, or be positioned near or at, medical device 120. Medical device 120 may be positioned on, at or near medically operational point (MOP) 122 on subject 130. Once the medical device 120 and the PSU 110 are positioned in this manner, a reference value or a baseline position may be set. During patient monitoring, a change in the position of the medical device 120 relative to the MOP 122 may be detected via the PSU 110, and the change may be utilized to disable the alert system, for example.

Physiological parameter sensor 126 may sense a physiological parameter of subject 130, and transfer 160, to medical system 140, a parameter signal that represents the value of the physiological parameter. The parameter signal representing the value of the physiological parameter may be transferred 160 to medical system 140, for example, continuously, or according to a predetermined interval (for example, once every 10 seconds; however, other intervals may be used).

Processor 144 may process the parameter signal, or the information embodied in it, and, based on the value of the parameter signal (based on the information) which represents the physiological parameter, processor 144 may determine that alert system 142 should output or generate an alert due to the physiological parameter having abnormal value (s). However, concurrently to the monitoring of the physiological parameter, PSU 110, which uses position sensor 112, or position sensor 112 in conjunction with means 124,

transfers position signal 114 to SPC 150 for processing, for example, by processor 152. Processor 152 may continuously, intermittently, or according to a predetermined time interval, determine from position signal 114 whether medical device 120 is at designated MOP 122, or whether medical device 120 is displaced from designated MOP 122 by an extent that renders measurements of the physiological parameter unreliable, which means that such measurements should be ignored and the alert system 142 disabled in order to prevent nuisance alert(s). Depending on the determination result, regarding enabling or disabling alert system 142, processor 152 may output 154 a corresponding instruction, or message, to processor 144 of medical system 140. In response to the instruction, or message, that processor 144 receives from processor 152, processor 144 may enable alert system 142 (allowing alert system 142 to output an alert), or disable it to prevent nuisance alerts when medical device 120 is displaced from MOP 122. SPC 150 may be external to medical system 140 or be part of (e.g., embedded in) medical system 140.

FIG. 2 schematically illustrates a system 200 in accordance with an example embodiment. System 200 may include a CO₂ monitoring device (CMD) 210 and a CO₂ monitoring system (CMS) 230. CMD 210 is configured to receive CO₂ 220 that is exhaled by a subject, to sample CO₂ from the exhaled CO₂, and to transfer 222, or to enable the transfer 222 of, the CO₂ samples to CMS 230.

CMS 230 may include a CO₂ detector 250 to detect the concentration level of, or another parameter related to, the CO₂ samples. CO₂ detector 250 may include a processor (the processor is not shown in FIG. 2). CO₂ detector 250 (or the processor associated with it) may output an activation signal 252 to activate an alert system 260, which may be included in, for example, CMS 230, when the CO₂ concentration level (or another parameter of the CO₂ samples) has an abnormal value, for example, for a predetermined amount of time.

Concurrently with the transfer 222 of the CO₂ samples to CMS 230, CMD 210 may transfer 246 a position signal to CMS 230. The position signal may represent the position of CMD 210 relative to a MOP on the body of the subject undergoing a CO₂ monitoring procedure. In general, the location of a MOP depends on the medical procedure/monitoring being performed. If the medical procedure/monitoring being performed is CO₂ monitoring, the MOP may be the subject’s mouth and/or nose. That is, CMD 210 may be positioned at, or near, the subject’s mouth and/or nose in a way that reliable CO₂ measurements can be made.

Carbon dioxide monitoring system 230 may also include an alert controller 260 to which CO₂ detector 250 (or the processor associated with it) may output the activation signal 252, and an alert controller 270. Alert controller 270 may analyze the position signal it receives 246 from CMD 210, and, based on the analysis result, may determine whether CMD 210 is still at, or operationally near, the designated MOP when CO₂ samples are taken. If alert controller 270 detects (from the position signal) movement of CMD 210 away, or deviation, from the MOP, alert controller 270 outputs a signal 280 that disables alert system 260, so that if CO₂ measurements are abnormal, the abnormal measurements are attributed to movement or removal of CMD 210 from the MOP, rather than these measurements indicating a genuine medical problem, and alert system 260 is, therefore, disabled.

If alert controller 270 detects (from the position signal) that CMD 210 is still at or operationally near the MOP, alert controller 270 outputs a signal 280 that enables alert system

260, so that if CO2 measurements are abnormal, the abnormal measurements are attributed to a genuine medical problem, and alert system 260 is, therefore, enabled.

FIG. 3 schematically illustrates a system 300 in accordance with an example embodiment. System 300 may include a CO2 monitoring device (CMD) 310 and a CO2 monitoring system (CMS) 330. CO2 monitoring device 310 may include a CSD 320 as an example medical device. CSD 320 may be configured to obtain samples from CO2 that is exhaled by a subject, and to transfer 322, or to enable the transfer of, the CO2 samples to CMS 330. It should be appreciated that CMS 330 may include a gas sampling pump to draw CO2 samples from CSD 320.

CO2 monitoring device 310 may also include a position sensing unit (PSU) 340. PSU 340 may include a position sensor 342 to sense the position of CSD 320 relative to a MOP on a body of the subject, and a communication modem 344 to transmit 346 the position signal to CMS 330. For example, in some embodiments, the CSD 320 may be initially positioned on the subject at the MOP, and the PSU 340 may establish or set this as a reference position for the CSD 320 (e.g., in response to a user input indicating that the CSD 320 is at the MOP, upon initiation of monitoring or therapy, or the like). Then, any subsequent movement away from the reference position may be detected by the PSU 340, and the PSU 340 may generate the position signal that indicates that the CSD 320 has moved relative to the MOP. Position sensor 342 may be entirely included in PSU 340, or only some components of position sensor 342 (for example, a controller, a power source, a signal processing/shaping circuit, etc.) may be incorporated in PSU 340, while other components of position sensor 342 (for example, a sensing electrode(s)) may be external to PSU 340 but operationally connected (for example, by being wired) to PSU 340.

CMS 330 may include a CO2 detector 350 to detect the concentration level of, or another parameter related to, the CO2 samples 322. CO2 detector 350 may include a processor (the processor is not shown in FIG. 3). CO2 detector 350 (or the processor associated with it) may output an activation signal 352 to activate an alert system 360, which may be included in, for example CMS 330, when the CO2 concentration level (or another parameter of the CO2 samples) has abnormal values, for example for a predetermined amount of time. Alert system 360 may include an alert signal generator 362 to generate an alert signal when CO2 detector 350 detects CO2 samples with abnormal values, and an audio-visual alert circuit 364 to effect the alert. CMS 330 may also include an alert system 360 and a signal processing unit (SPC) 370. SPC 370 may include a communication modem 374 to receive 346 the position signal from communication modem 344 of PSU 340, and a controller 372. Communication modem 344 may include a controller to process the output signal of position sensor 342, and to manage the transfer of position signal 346 to CMS 330.

Controller 372 may process the received 346 position signal and, based on the processed position signal, determine whether CSD 320 is still at, or operationally near, the designated MOP, or not. If CSD 320 is still at the designated MOP, or it is operationally near, the MOP (e.g., the displacement between the reference position and a current position is within a threshold value), controller 372 enables 380, or sets on or "arms," alert system 360 so that alert system 360 can operate normally or in a default operational mode (e.g., can go off) based on the signal 352 that CO2 detector 350 outputs regarding whether the measured CO2 samples indicate normal readings or abnormal readings. However, if CSD 320 is not at the designated MOP (e.g., the

displacement between the reference position and a current position is not within a threshold value), controller 372 disables 380, or turns off, alert system 360 in order to prevent nuisance alerts. In some embodiments, PSU 340 may be releasably attached 390 to CSD 320, for example by using a snap fit connector or a snap fastener. SPC 370 may be external to medical system 330 or be part of (e.g., embedded in) medical system 330.

FIG. 4A schematically illustrates a system 400 in accordance with an example embodiment. System 400 may include a CO2 monitoring device (CMD) 410 and a CO2 monitoring system (CMS) 430. CO2 monitoring device 410 may include a CO2 sampling device (CSD) 420 as an example medical device. CSD 420 may be configured to obtain samples from CO2 that is exhaled by a subject, and to transfer 422, or to enable the transfer of, the CO2 samples to CMS 430. It should be appreciated that CMS 430 may include a gas sampling pump to draw CO2 samples from CSD 420. CO2 monitoring device 410 may also include a PSU 440. PSU 440 may include a position sensor 442 to sense the position of CSD 420 relative to a MOP on the body of a subject, and a wireless communication modem 444 connected to an antenna 448 (e.g., radio frequency (RF) antenna) to wirelessly transmit 446, via antenna 448, the position signal to CMS 430. Position sensor 442 may be entirely included in PSU 440, or only some components of position sensor 442 (for example, a controller, a power source, a signal processing/shaping circuit, etc.) may be incorporated in PSU 440 while other components of position sensor 442 (for example, a sensing electrode(s)) may be external to PSU 440 but operationally connected (for example, by being wired) to PSU 440.

CMS 430 may include a CO2 detector 450 to detect the concentration level of, or another parameter related to, the CO2 samples 422 that are transferred to CMS 430 from CSD 420. Carbon dioxide detector 450 may include a processor (the processor is not shown in FIG. 4A). Carbon dioxide detector 450 (or the processor associated with it) may output an activation signal 452 to activate an alert system 460, which may be included in, for example CMS 430, when the CO2 concentration level (or the other measured parameter of the CO2 samples) has abnormal values, for example for a predetermined amount of time (e.g., for 10 seconds; however, other time periods may be used). Alert system 460 may include an alert signal generator (ASG) 462 to generate an alert signal when CO2 detector 450 detects, for example, CO2 samples with abnormal values, and an audio/visual alert circuit 464 to effect the alert. CMS 430 may also include a signal processing unit (SPC) 470. SPC 470 may include a wireless communication modem 474 to receive 446, via an antenna 476, the position signal from wireless communication modem 444 of PSU 440. SPC 470 may also include a controller 472. Communication modem 444 may include a controller to process the output signal of position sensor 442, and to manage the transfer of signal 446 to CMS 430. (The controller of communication modem 444 is not shown in FIG. 4A.)

Controller 472 may process the received 446 position signal and, based on the processed position signal, determine whether CSD 420 is still at, or operationally near, the designated MOP, or not. If CSD 420 is still at the designated MOP, or it is operationally near, the MOP (e.g., the displacement between the reference position and a current position is within a threshold value), controller 472 enables 480, or sets on or "arms", alert system 460 so that alert system 460 can operate normally or in a default operational mode (e.g., can go off) based on the signal 452 that CO2

detector **450** outputs regarding whether the measured CO₂ samples indicate normal readings or abnormal readings. However, if CSD **420** is not at the designated MOP (e.g., the displacement between the reference position and a current position is not within a threshold value), or it has operationally displaced from, the MOP, controller **472** disables **480**, or turns off, alert system **460** in order to prevent nuisance alerts. PSU **440** may be releasably attached **490** to CSD **320**, for example by using a snap fit connector or a snap fastener.

In general, the location of a MOP, that may be, for example, a conspicuous point, area or feature on an organ, or an entire organ (for example a mouth, a nose, etc.) of, the subject, may be selected for initial or proper placement of the medical device by a medical professional depending, for example, on the medical procedure/monitoring being performed. For example, if the medical procedure/monitoring being performed is CO₂ monitoring, the MOP may be, for example, the subject's mouth and/or nose (where reliable CO₂ measurements can be made), and a deviation of the CSD (for example deviation of CSD **320** or CSD **420**) may be detected with respect to the mouth and/or nose.

In some embodiments, enabling and disabling the alert system (e.g., alert system **360**, alert system **460**) may be effected by the CMS controller (e.g., controller **372**, controller **472**) sending the enable/disable signal (signal **380**, signal **480**) directly to the CO₂ detector (e.g., CO₂ detector **330**, CO₂ detector **430**), for example to a controller that manages operation of the CO₂ detector. In these embodiments, the CO₂ detector (or its controller) may be allowed to output a "CO₂ level low" (for example) alert signal (e.g., signals **352**, **452**) to set the alert system (e.g., alert system **360**, alert system **460**) off when the enable/disable signal (e.g., signals **380**, **480**) indicates that the medical device (for example CSDs **320**, **420**) is at, or operationally near, the designated MOP. Conversely, the CO₂ detector (or its controller) may be inhibited from sending a "CO₂ level low" (for example) alert signal (e.g., signals **352**, **452**) to the alert system (e.g., alert system **360**, alert system **460**) when the enable/disable signal (for example signals **380**, **480**) indicates that the medical device (for example CSDs **320**, **420**) is displaced (operationally moved away) from the designated MOP.

In some embodiments, enabling and disabling the alert system (e.g., system **360** or **460**) may be effected by the communication modem of the PSU (e.g., modem **344** or **444**), or a controller managing this modem, sending an enable/disable signal (e.g., signals **346**, **446**) (for example via a modem in the CMS; e.g., modem **374** or **474**) to the CO₂ detector (e.g., directly to detector **350** or **450**), or directly to the alert system, to enable or to disable the alert system depending on whether the signal that the PSU (e.g., PSU **340** or **440**) transfers to the CMS indicates that the medical device (for example, CSD **320** or **420**) is operationally at or near the designated MOP. SPC **470** may be external to medical system **430** or be part of (e.g., embedded in) medical system **430**.

FIG. 4B schematically illustrates a system **402** in accordance with another example embodiment. Reference numerals are repeated among FIG. 4A and FIG. 4B to indicate like, corresponding or analogous elements, and, accordingly, the elements in FIGS. 4A and 4B that are referred to by the same reference numerals operate in a same way or in a similar way. Therefore, the way these elements operate is not described in connection with FIG. 4B.

System **402** may include a CMD **412** and a CMS **432**. CMD **412** may include a CSD **420** as an example medical

device, and a PSU **441**. PSU **441** may include a wireless modem **443** which, in this example, is bidirectional modem that is capable of bidirectional communication with CMS **432**. PSU **441** may also include an in-situ audio-visual alert system **449** to locally indicate (for example, to the subject wearing CSD **420**) a displacement of CSD **420** (or another medical device) from a designated MOP. PSU **441** may also include an antenna **448** and a position sensor **442**.

CMS **432** may include a CO₂ detector **450**, an alert system **460** and a signal processing unit (SPC) **478**. SPC **478** may include a bidirectional wireless modem **479** to bidirectionally communicate with bidirectional wireless modem **443** via bidirectional communication path **491**. SPC **478** may also include a controller **472** to control, among other things, the operation of communication modem **479**. SPC **478** may be external to medical system **432** or be part of (e.g., embedded in) medical system **432**.

PSU **441** and CMS **432** may exchange information with regard to activation, or deactivation, of an alert that is to be introduced (audibly and/or visually), or inhibited, in-situ (e.g., on CMD **412**) or remotely (e.g., on CMS **432**), as a result of a displacement of CSD **420**. For example, exchanging information between PSU **441** and CMS **432** may include sending a disabling signal (**491**) from PSU **441** to CMS **432** to disable the CMS's alert system (**460**) so that low CO₂ readings would not trigger an alert in CMS **432** when CSD **420** is not at the designated MOP, and sending CO₂ measurements from CMS **432** to PSU **441** in order to trigger an in-situ alert signal by audio/visual alert system **449**. In some embodiments, position sensor **442** may sense a displacement of CSD **420** from the designated MOP, and a controller managing the operation of PSU **441** may cause audio/visual alert system **449** to introduce (audibly or visually) an alert when such a displacement is sensed (e.g., displacement that exceeds a threshold value).

FIG. 5 shows a method of disabling an alert system in a medical system monitoring a physiological parameter of a subject according to an example embodiment. By way of example, the method is described in connection with a CO₂ monitoring system (CMS) as a medical system. However, a similar method may likewise be applied to other types of medical systems, for example it may be applied to an oxygen monitoring system, to an oxygen delivering system, to a heart pulse monitoring system, etc.

Initially, a medical device (e.g., CSD **420**) may be positioned at a designated MOP of a subject to monitor a physiological parameter and/or to carry out a medical procedure. Then, while a subject's exhaled CO₂ is sampled by a CO₂ sampling device and monitored by a CO₂ monitoring system, the following steps may be performed. At step **510**, a displacement, *d*, of a medical device (e.g., CSD **420**) used to sample exhaled CO₂ from the designated MOP may be monitored by a PSU (e.g., PSU **440**) continuously or once in a while (e.g., once every *n* seconds; *n*=1, 2, 3, . . .). The PSU may send a position signal to the CMS. The position signal may be or represent the displacement, *d*. The PSU may monitor the displacement via any of a variety of techniques. For example, the PSU may establish a reference displacement value or a reference position while the medical device is positioned at the MOP (e.g., upon input by an operator indicating the medical device is positioned at the MOP, upon initiation of monitoring or the medical procedure, upon detection of the means **124**, or the like). The PSU may then detect any subsequent movement or displacement of the medical device relative to the reference position, which may correspond to movement or displacement relative to the MOP. In some embodiments, the PSU may include one or

more position sensors (e.g., position sensor **112**) and one or more detectable elements (e.g., means **124**). In some such cases, the position sensor may be placed at a first position on the subject and the detectable element may be coupled to or incorporated into the medical device, which may be placed at the MOP of the subject. The position sensor may then monitor for a change in a distance between the position sensor and the detectable element (e.g., the position sensor may be a proximity sensor that changes state when a distance between the position sensor and the detectable element exceeds a threshold value), which may be indicative of movement or displacement of the medical device relative to the MOP.

At step **520**, a controller (e.g., a controller of the CMS) may compare the displacement d to a threshold value ($d/\text{threshold}$). If the displacement d is greater than the threshold value (if $d > d/\text{threshold}$, this condition is shown as “Yes” at step **520**), the CMS’s controller may, at step **530**, disable an alert system of the CMS such that abnormal CO₂ measurements would not set off the alert system of the CMS. However, if the displacement d is not greater than the threshold value (this condition is shown as “No” at step **520**), the CMS’s controller may, at step **540**, enable the alert system of the CMS such that abnormal CO₂ measurements would set off the alert system of the CMS. Monitoring the displacement of the medical device from the designated MOP may continue, as shown at step **550**, or be repeated, for example, for as long as the physiological parameter (in this example, exhaled CO₂) is monitored by the medical system (in this example, a CMS).

In some embodiments, the medical device is a cannula. The cannula may be configured to collect samples of CO₂ that is exhaled by a subject, and, in addition, it may be configured, in accordance with the present disclosure, to facilitate sensing of its own position with respect to a designated MOP. FIG. **6A** depicts an example nose clip **610** according to an example embodiment. Nose clip **610** may include a V-shaped member **620** that is releasably attachable to the nose of a subject, for example in a similar way as an eyeglass’s nose pad (by being lightly pressed against the nose). Member **620** may be an elongated member (e.g., clip) having a middle semi-straight segment (a bridge) **622** that bridges the two side segments of V-shaped member **620**. Bridge section **622** may contain an electronic circuit **630**. Electronic circuit **630** may be or include a PSU that may be identical or similar to, for example, PSU **420** (FIG. **4A**).

The PSU may include a position sensor, a wireless communication modem and a controller to manage transmission of a position signal (that the position sensor outputs), for example, to a remote medical system. V-shaped (or U-shaped) member **620** may include an antenna **640** via which the PSU (e.g., its controller) may transmit the position signal. In some embodiments, the position sensor may be a proximity sensor (e.g., a proximity switch). The proximity sensor may be magnetic (inductive), capacitive, etc. A proximity switch is a device that causes a switching action without physical contact. In general, proximity switches respond to ‘targets’ that enter the active range of their generated sensing fields. Inductive proximity sensors, for example, are used for non-contact detection of metallic objects. Their operating principle of inductive proximity sensors is based on a coil and oscillator that creates an electromagnetic field in the close surroundings of the sensing surface.

FIG. **6B** depicts an example cannula **650** according to an example embodiment. Cannula **650** is used as an example medical device. It should be appreciated that the medical

device subject of the present disclosure may be any medical device from a relatively simple device, for example, a tube, to a relatively complex device, for example, a face mask that enables both efficient delivery of oxygen to a subject via the mask, and concomitant extraction of exhaled CO₂ from the mask. Cannula **650** may conventionally include a cannula tube **660** and prongs **670** (one prong per nostril). Prongs **670** may be used to deliver oxygen enriched air to a subject during respiration and/or to collect samples of CO₂ that the subject exhales. Cannula **650** may also include an object **680** (e.g., a sensing target for a proximity sensor to sense or detect) in order for it to be sensed by the position sensor (e.g., proximity sensor coupled to or mounted on, for example, electric circuit **630**). Object **680** may be, for example, metallic or ferromagnetic. In some embodiments, object **680** may be an RFID tag, and electric circuit **630** may include a RFID reader. The RFID reader may be configured to output a very low power radio frequency (RF) signal (e.g., a sensing signal) that is in compliance with the relatively small operational distance (e.g., gap) between the position sensor (e.g., in electric circuit **630**) and, for example, cannula prongs **670** to enable the RFID reader to read the RFID tag when separated by the gap. When a distance between the RFID tag and the RFID reader exceeds the gap, the RFID reader may not read the RFID tag, thereby indicating that the RFID tag has moved away from the RFID reader and away from the MOP.

In some embodiments, the position sensor of nose clip **610** may be or include a light source (e.g., light emitting diode), a light guiding element (e.g., optical fiber) to guide light to a certain point on cannula **650**, and a light sensor (e.g., photodiode) to sense a reflected light. In these embodiments, object **680** of cannula **650** may be or include a light reflecting element. The light source of the position sensor and the light reflecting element may be configured such that they are aligned (light is reflected back to the light sensor) when the cannula, or prongs, are at the designated MOP, and are misaligned when the cannula, or prongs, are displaced from the designated MOP, thereby enabling the electronic circuit **630** to monitor the position of the cannula relative to the designated MOP.

FIGS. **7A-7C** schematically illustrate medical devices according to some embodiments. Referring to FIG. **7A**, medical device **710**, which is another example medical device, may include a face mask **711**, an oxygen port **712** for delivering oxygen to a subject wearing face mask **711**, and a CO₂ port **713** through which samples of CO₂ (which is exhaled by the subject) may be transferred from the interior space of mask **711** to a remote CO₂ monitoring system. Medical device **710** may include only oxygen port **712**, or only CO₂ port **713**, or both ports **712** and **713**. In operation, mask **711** is positioned on the face of a subject such that delivery of oxygen to the interior space of the mask is optimal (e.g., maximized, desired, or adequate for treatment), and the sampling of the exhaled CO₂ is also optimal (e.g., maximized, desired, or adequate for monitoring). Such optimal performance of the mask may be obtained when the mask is in an optimal position (e.g., desired, intended, or appropriate position) with respect to the subject’s face, the optimal position being referred to herein as the medically operational position (MOP).

Medical device **710** also includes a position sensing unit (PSU) **714**. PSU may **714** include an electric circuit that is connected to two electrodes (**715**, **716**). Electric circuit **714** may include a controller and a wireless communication modem or transceiver. Electrodes **715** and **716** may be used to sense, for example, an impedance of a tissue that is

contacted by mask **711**. When the mask is removed from the subject's face, the impedance sensed between the two electrodes is expected to be infinite, and when the mask is moved on the subject's face, the impedance sensed between the two electrodes is expected to change or be unstable. Each electrode may be wired to electric circuit **714**. In some embodiments, in operation, the PSU **714** may obtain a first impedance measurement (e.g., reference or baseline measurement) with the electrodes **715**, **716** (e.g., upon input from an operator that the medical device **710** is position at the MOP, upon initiation of monitoring or therapy, or the like), and then monitor for a change in the impedance, which is indicative of movement of the medical device **710** relative to the MOP. For example, electrode **715** is connected to electric circuit **714** via electric wire **717**. During operation, the PSU may use electrodes **715**, **716** to produce a position signal that represents the position of the mask on the face of the subject, or a position signal that indicates whether the mask is in the right position (e.g., at, or operationally near, the MOP) or not. The PSU's controller may use the wireless communication modem to transmit the position signal to the remote CO₂ monitoring system and/or to the remote oxygen delivery system to enable or to disable the alert system, for example.

FIG. 7B shows a medical device **720** which is somewhat different than medical device **710**, the difference being that the PSU (PSU **724**) is connected to a temperature sensor **725**, rather than to impedance electrodes. Temperature sensor **725** may sense the temperature of the subject's skin, which, under normal conditions, is expected to be higher than the room temperature. The electric signal that PSU **724** receives from temperature sensor **725** (e.g., via electrical wire **727**) may be indicative of a position of the medical device **720** relative to the subject, and thus, to the MOP. For example, in operation, the PSU **724** may obtain a first temperature (e.g., reference or baseline measurement) from the temperature sensor **725** (e.g., upon input from an operator that the medical device **720** is position at the MOP, upon initiation of monitoring or therapy, or the like), and then monitor for a change in the temperature (e.g., a decrease, such as by more than 1, 2, 3, 4, 5, 10, or 15 percent), which may be indicative of movement of the medical device **720** away from the subject's skin and relative to the MOP. PSU **724** may transmit the signal to a remote medical system that operates with medical device **720** to enable or to disable the alert system, for example.

FIG. 7C shows a medical device **730** which is somewhat different than medical devices **710** and **720**, the difference being that the PSU (PSU **734**) is connected to an optical scanner **735**. Optical scanner **735** may be designed and function in a similar way as an optical computer mouse. For example, the optical scanner **735** may include a light source, typically a light-emitting diode ("LED"), and a light detector, such as an array of photodiodes, that together are configured to detect movement of the optical scanner **735** relative to a surface, such as a surface of the subject's skin. The electric signal that PSU **734** receives from optical scanner **735** (e.g., via electrical wire **737**) may be indicative of a position of the medical device **730** relative to the subject, and thus, to the MOP. PSU **734** may transmit the signal to a remote medical system that operates with medical device **730** to enable or to disable the alert system, for example.

In some embodiments, an alert activation/deactivation system is provided for a medical system and may include a PSU (for example, a PSU similar to PSU **110**) to sense a displacement **d1** of a medical device (for example a medical

device similar to medical device **120**) from a MOP (e.g., MOP **122**) on a subject (e.g., subject **130**), and a signal processing circuit (for example, a signal processing circuit similar to signal processing circuit **150**) to output **154**, based on the sensed displacement **d1**, a signal to disable an alert system (for example, an alert system similar to alert system **142**) of the medical system (for example, a medical system similar to medical system **140**) cooperating with the medical device if the medical device is for sensing a physiological parameter of a subject during a medical procedure and the displacement **d1** is greater than a first threshold value, and, in other embodiments, to output a signal to enable an alert system when the medical device is for delivering a treatment to the subject and the displacement **d1** is greater than a second threshold value. In some embodiments, the second threshold value may be different than and/or less than the first threshold value. The medical device may be configured to deliver any treatment to the subject, for example oxygen, nutrition, medication, etc. The physiological parameter may be, for example, concentration level of exhaled CO₂, oxygen saturation level, heart pulse rate, blood pressure, electrocardiogram (ECG), body temperature and tissue impedance, to name a few.

Various aspects of the embodiments disclosed herein are combinable with various types of medical devices. Although the discussion herein relates to cannulas and to face masks, embodiments of the disclosure are not limited in this regard. For example, the embodiments may, similarly, be incorporated or built into, or attached to, or mounted in or on any medical device, such as (but not limited to) a medical device that senses oxygen saturation in a subject, or exhaled CO₂, or a body temperature, or a heart pulse, or blood pressure, signal, or a medical device that deliver treatment (e.g., oxygen, medication) to a subject. The embodiments may, similarly, be incorporated or built into, or attached to, or mounted in or on any medical device that delivers, or facilitates delivery of, for example, oxygen, nutrition or medication, to name a few medical procedures that can be performed by various medical devices.

FIGS. 8A-8B illustrate a cannula in accordance with some embodiments of the present disclosure. Referring to FIG. 8A, a cannula **810** includes a CO₂ sampling device (CSD, such as a scoop) **820** to collect samples of exhaled CO₂. Cannula **810** is mounted on a subject, and a position sensor unit (PSU) **830** is shown mounted on (e.g., releasably attached to) CSD **820**. PSU **820** may be mounted on, or attached to, CSD **820** by using, for example, a snap fit connector, a snap fastener, or a Velcro strip. Alternatively, the PSU may be mounted on a side section or face section **840** of cannula **810**, as shown in FIG. 8B at **830**, instead of mounting the PSU (e.g., PSU **820**) on the CSD (e.g., CSD **820**), or on the cannula's mouth section **850** (as shown in FIG. 8A).

While certain features of the disclosure have been illustrated and described herein, many modifications, substitutions, changes, and equivalents will now occur to those of ordinary skill in the art, and the appended claims are intended to cover all such modifications and changes. It should be appreciated that the various features disclosed and illustrated in FIGS. 1-8B may be combined with one another in any suitable manner.

The invention claimed is:

1. A system to block nuisance alerts in a medical system, comprising:
 - a position sensing unit configured to sense a displacement of a medical device from a medically operational point on a subject; and

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a signal processing circuit remote from the medical device and configured to output, based on the sensed displacement, a first signal to enable an alert system of a medical system cooperating with the medical device when the displacement is less than a threshold value, and to output a second signal to disable the alert system when the displacement is equal to or greater than the threshold value.

2. The system as in claim 1, wherein the position sensing unit comprises:

a position sensor configured to output a position signal representing the displacement of the medical device from the medically operational point on the subject, wherein the medical device is configured to cooperate with the medical system to perform a medical procedure, and wherein the signal processing circuit comprises a processor that is configured to output for the medical system, based on the position signal, the first signal to enable the alert system of the medical system, and to output the second signal to disable the alert system.

3. The system as in claim 2, further comprising a first element incorporated in or attached to the medical device to facilitate sensing the displacement of the medical device from the medically operational point.

4. The system as in claim 1, wherein the medical device comprises: a cannula, a nasal tubing, a face mask, a CO₂ sampling device to collect exhaled CO₂ for the CO₂ monitoring system, and a physiological parameter sensor to sense a physiological parameter of the subject.

5. The system as in claim 4, wherein the physiological parameter sensor is selected from the group consisting of: an oxygen sensor, a heart pulse sensor, a blood pressure sensor, an electrocardiogram (ECG), temperature sensor, and an impedance sensor.

6. The system as in claim 1, wherein the medically operational point on the subject is selected from the group consisting of: a face of the subject, a nose of the subject, a mouth of the subject, a finger of the subject, a wrist of the subject, an elbow of the subject, and the chest of the subject.

7. The system as in claim 2, wherein the position sensor is configured to be positioned at a predetermined distance from the medical device or is incorporated in or attached to the medical device.

8. The system as in claim 2, wherein the signal processing circuit is incorporated in the medical system.

9. The system as in claim 2, wherein the position sensor is selected from the group consisting of: an impedance sensor, a temperature sensor, a pressure sensor, a capacitive sensor, a proximity sensor, a motion sensor, an acceleration sensor, and an optical sensor.

10. The system as in claim 2, wherein each of the position sensing unit and the signal processing circuit comprises a communication modem configured to transfer the position signal from the position sensing unit to the signal processing circuit.

11. The system as in claim 10, wherein any of the position sensor and the communication modem of the position sensing unit is an add-on device attachable to, or embedded in, the medical device.

12. The system as in claim 11, wherein each communication modem is wireless.

13. The system as in claim 1, wherein the medical system is a CO₂ monitoring system configured to measure CO₂ concentration, an oximeter configured to measure oxygen saturation level, a heart pulse rate monitoring system configured to measure the subject's pulse rate, a blood pressure

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monitoring system configured to measure the subject's blood pressure, or an electrocardiogram (ECG) monitoring system configured to monitor electrical activity of the subject's heart.

14. The system as in claim 13, wherein the first signal enables a CO₂ alert circuit of the CO₂ monitoring system and the second signal disables the CO₂ alert circuit.

15. The system as in claim 13, wherein the medical device is a CO₂ monitoring device, the CO₂ monitoring device comprising:

a CO₂ sampling device configured to collect CO₂ samples exhaled by the subject;

the position sensing unit configured to output the position signal representing a position of the CO₂ sampling device on the subject;

the alert system configured to output an alert signal based on a detected CO₂ concentration; and

the signal processing circuit configured to activate or deactivate the alert system based on the position signal, wherein the CO₂ monitoring system is configured to receive and process the CO₂ samples and to receive the position signal from the CO₂ monitoring device, and to conditionally output an alert signal when a value associated with the processed CO₂ samples exceeds a predetermined level or range, the condition to output said alert signal being that the CO₂ monitoring system interprets said position signal as the CO₂ monitoring device being properly positioned on or relative to a face of the subject.

16. An alert activation/deactivation system for a medical system, comprising:

a position sensing unit configured to sense a displacement of a medical device from a medically operational point on a subject; and

a signal processing circuit remote from the medical device and configured to output, based on the sensed displacement, a first signal to disable an alert system of a medical system cooperating with the medical device when the medical device is for sensing a physiological parameter of a subject during a medical procedure and the displacement is greater than a first threshold value, or to output a second signal to enable the alert system when the medical device is for delivering treatment to the subject and the displacement is greater than a second threshold value.

17. The system as in claim 16, wherein the medical device is configured to deliver oxygen or medication to the subject.

18. The system as in claim 16, wherein the physiological parameter is selected from the group consisting of: concentration level of exhaled carbon dioxide, oxygen saturation level, heart pulse rate, blood pressure, electrocardiogram (ECG), body temperature and tissue impedance.

19. A method for monitoring carbon dioxide (CO₂), comprising:

generating a position signal representing a position of a CO₂ monitoring device relative to a medically operational point on a subject and measuring CO₂ exhaled by the subject by a CO₂ monitoring system; and

disabling, by an alert controller, an alert system of the CO₂ monitoring system in response to receipt of the position signal by a signal processing unit remote from the CO₂ monitoring device, the signal indicating an excess displacement of the CO₂ monitoring device with respect to the medically operational point.

20. The method as in claim 19, wherein the step of disabling the alert system comprises activating an indication

in or on the CO₂ monitoring device to indicate displacement of the CO₂ monitoring device from the medically operational point.

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专利名称(译)	用于操作医疗设备警报系统的系统和方法		
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摘要(译)

一种用于医疗系统的系统，包括：位置感测单元，用于感测医疗设备相对于受试者的医疗操作点的位移；以及信号处理电路，用于基于感测到的位移来输出信号，以在以下情况时禁用警报系统：该医疗设备用于在医疗过程中感测对象的生理参数，并且该位移大于第一阈值，或者当该医疗设备用于向对象和患者提供治疗时输出信号以启用警报系统。位移大于第二阈值。

