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(54) **TELEMETERED CHARACTERISTIC
MONITOR SYSTEM AND METHOD OF
USING THE SAME**

(75) Inventors: **Alfred E. Mann**, Beverly Hills, CA
(US); **Richard E. Purvis**, Pasadena, CA
(US); **John J. Mastrototaro**, Los
Angeles, CA (US); **James D. Causey**,
Simi Valley, CA (US); **James Henke**,
Simi Valley, CA (US); **Peter Hong**,
Santa Clarita, CA (US); **John H.**
Livingston, Newport Beach, CA (US);
Clifford W. Hague, Sherman Oaks, CA
(US); **Brad T. Hite**, Valencia, CA (US)

(73) Assignee: **Medtronic Minimed, Inc.**, Northridge,
CA (US)

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patent is extended or adjusted under 35
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continuation of application No. 09/465,715, filed on
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continuation of application No. 09/377,472, filed on
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600/301, 331, 309, 323, 322; 340/870.07;
73/1.06

See application file for complete search history.

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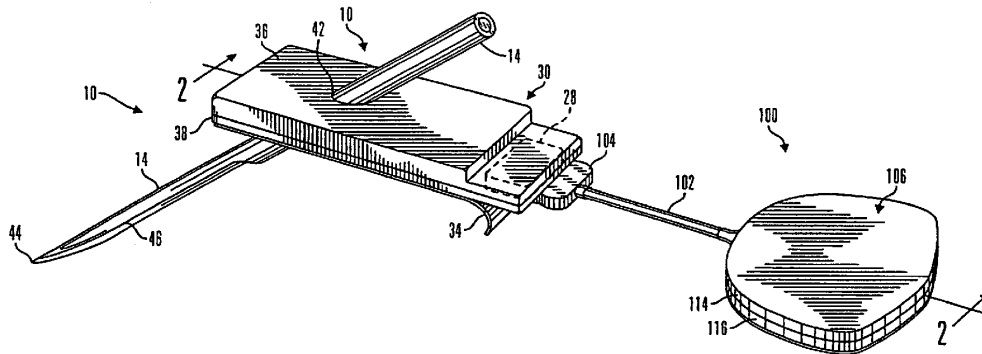
Assistant Examiner—Hung Q Dang

(74) *Attorney, Agent, or Firm*—Pillsbury Winthrop Shaw
Pittman LLP

(57) **ABSTRACT**

A telemetered characteristic monitor system includes a
remotely located data receiving device, a sensor for produc-
ing signal indicative of a characteristic of a user, and a
transmitter device. The transmitter device includes a hous-
ing, a sensor connector, a processor, and a transmitter. The
transmitter receives the signals from the sensor and wire-
lessly transmits the processed signals to the remotely located
data receiving device. The processor coupled to the sensor
processes the signals from the sensor for transmission to the
remotely located data receiving device. The data receiving
device may be a characteristic monitor, a data receiver that
provides data to another device, an RF programmer for a
medical device, a medication delivery device (such as an
infusion pump), or the like.

11 Claims, 5 Drawing Sheets



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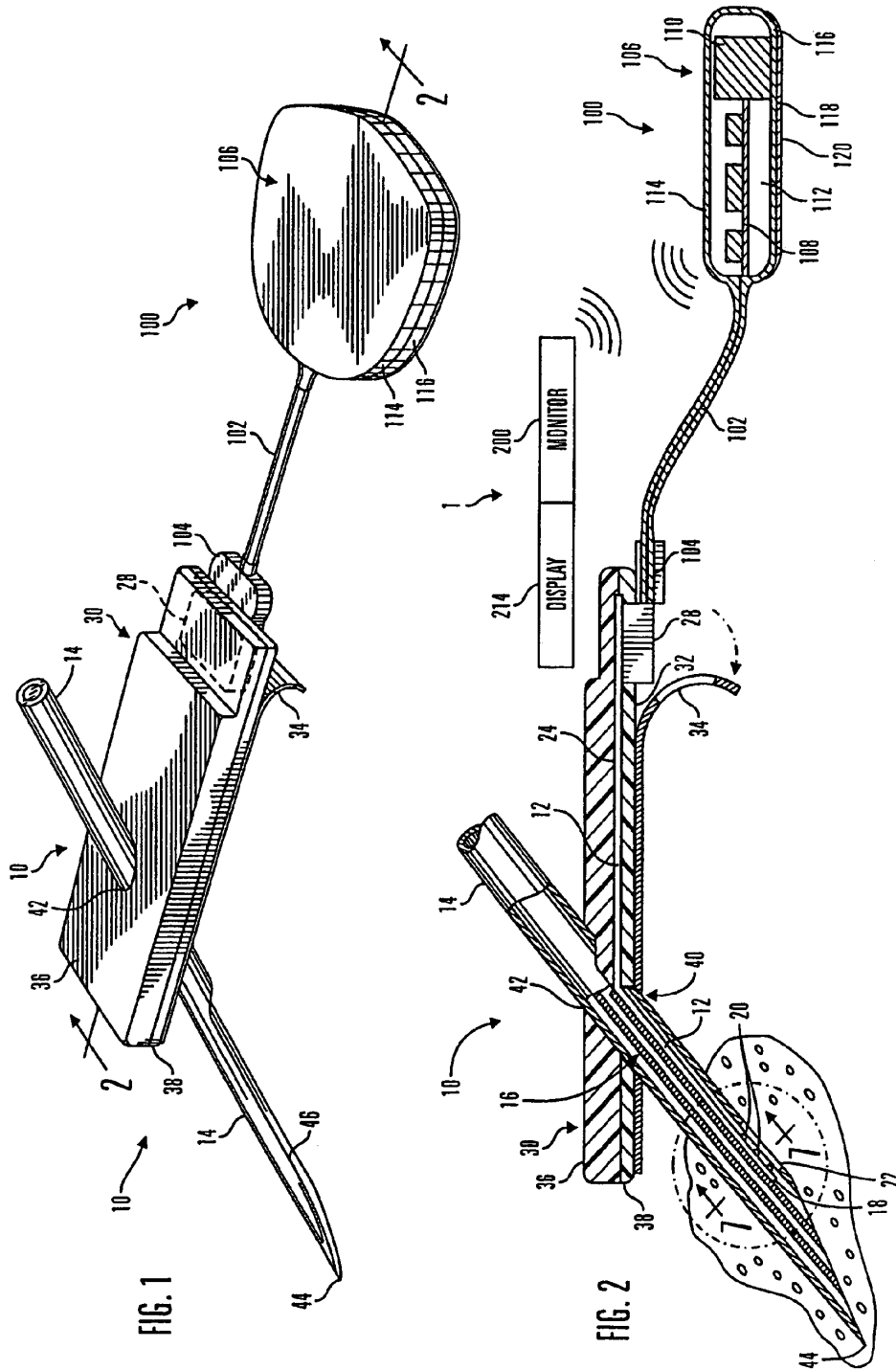
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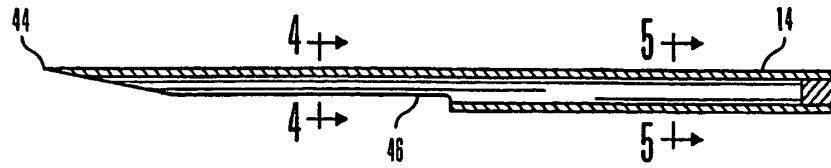


FIG. 3

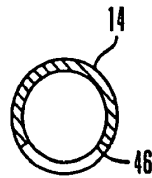


FIG. 4

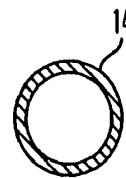


FIG. 5

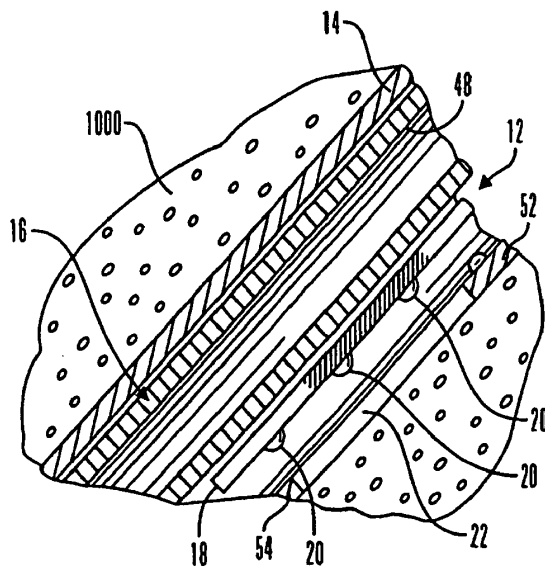


FIG. 6

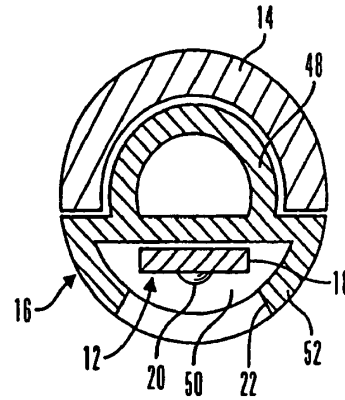


FIG. 7

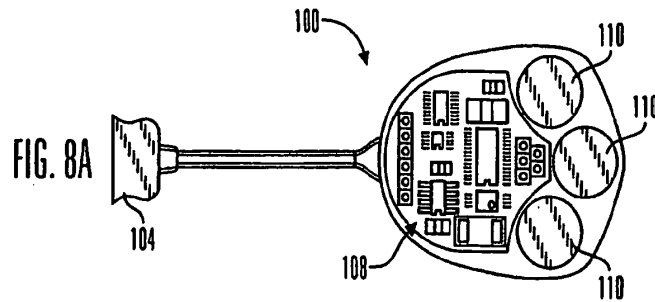
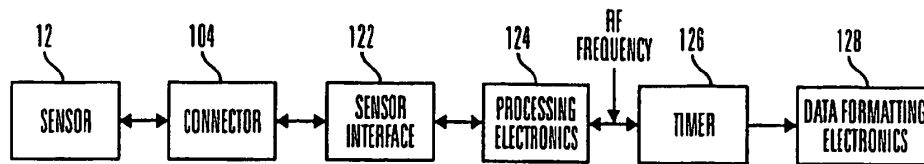


FIG. 8B

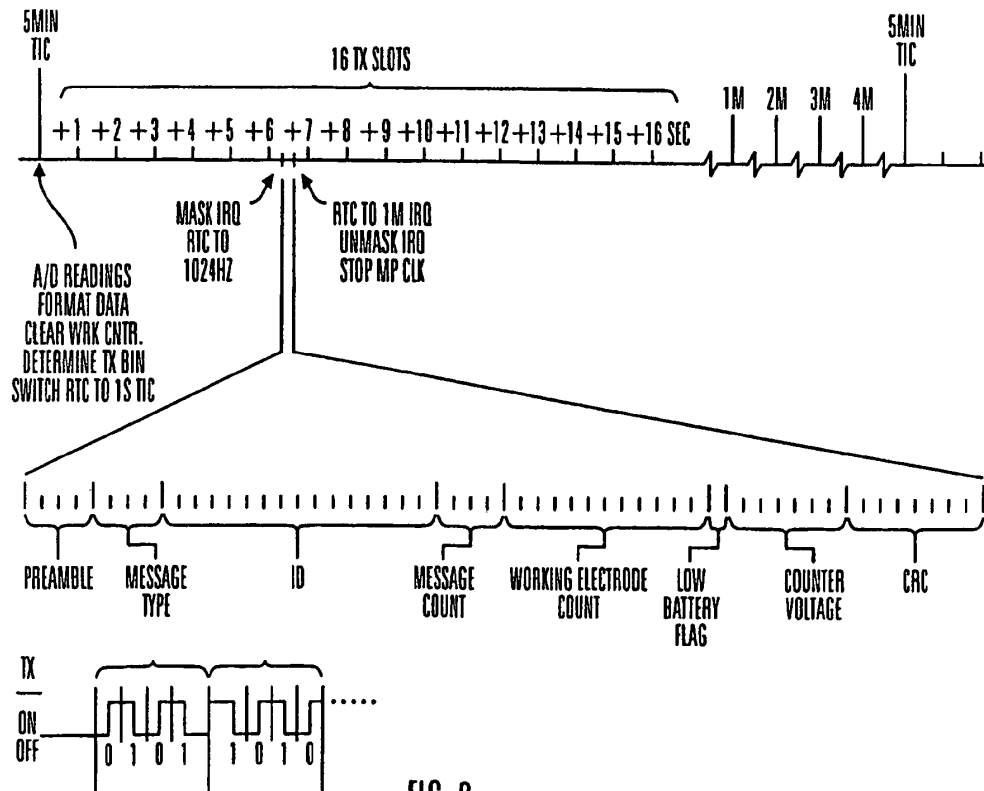
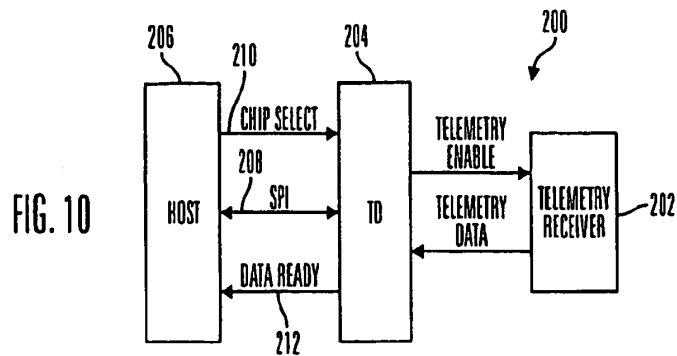
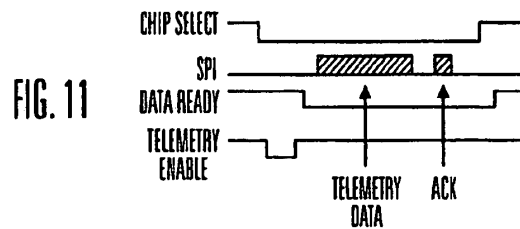


FIG. 9



HOST INITIATES RF SESSION

TD WAKES UP
THROUGH RBD / INT

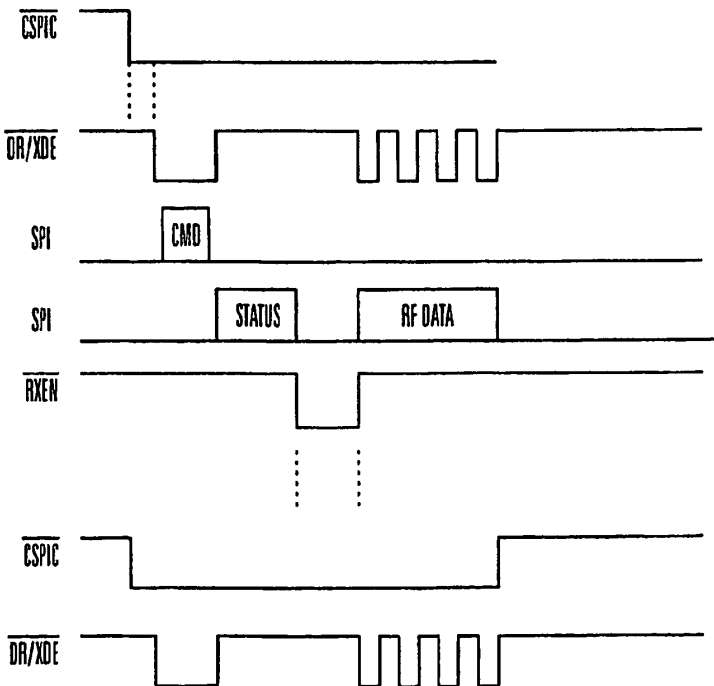
TD SIGNALS DATA READY

HOST SENDS COMMAND

TD SENDS STATUS
DATA TO HOST

TD ENABLES RF RECEIVER

TD COLLECTS RF DATA

HOST TERMINATES
RF SESSIONTD CLOSES OUT
RF SESSION AND
GOES TO SLEEP**FIG. 12**

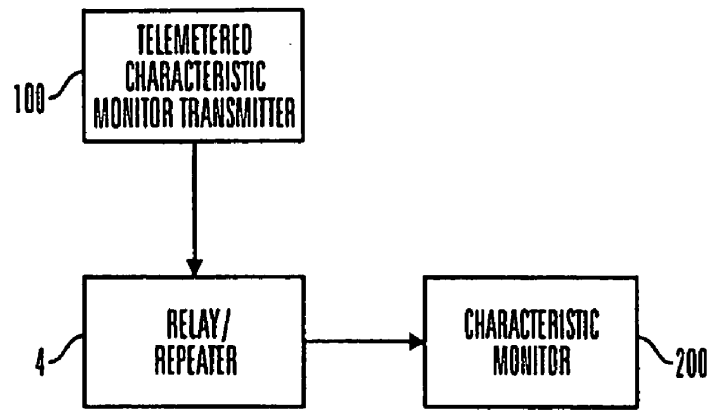


FIG. 13

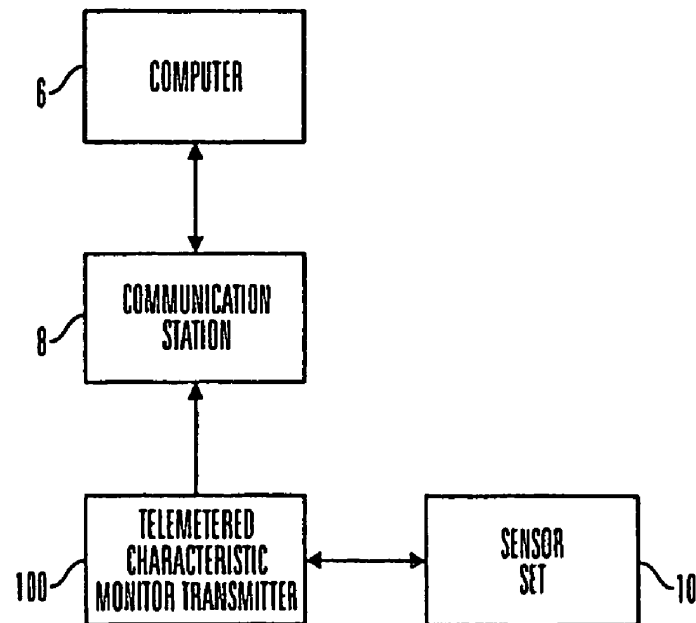


FIG. 14

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TELEMETERED CHARACTERISTIC MONITOR SYSTEM AND METHOD OF USING THE SAME

RELATED APPLICATIONS

This application is a continuation of U.S. patent application Ser. No. 10/898,589, filed on Jul. 23, 2004 and issued as U.S. Pat. No. 7,098,803 on Aug. 29, 2006, which is a continuation of U.S. patent application Ser. No. 09/465,715, filed on Dec. 17, 1999 and issued as U.S. Pat. No. 6,809,653 on Oct. 26, 2005, which is a continuation of U.S. patent application Ser. No. 09/377,472, filed on Aug. 19, 1999 and later abandoned, which claimed priority from U.S. Provisional Application Ser. No. 60/103,812, filed on Oct. 8, 1998, each of which is herein incorporated by reference.

FIELD OF THE INVENTION

This invention relates to telemetered subcutaneous sensor devices and, in particular embodiments, to devices and methods for wireless communication between an implantable subcutaneous sensor set at a selected insertion site within the body of a user and a remotely located characteristic monitor.

BACKGROUND OF THE INVENTION

Over the years, a variety of implantable electrochemical sensors have been developed for detecting and/or quantifying specific agents or compositions in a patient's blood. For instance, glucose sensors have been developed for use in obtaining an indication of blood glucose levels in a diabetic patient. Such readings are useful in monitoring and/or adjusting a treatment regimen which typically includes the regular administration of insulin to the patient. Thus, blood glucose readings improve medical therapies with semi-automated medication infusion pumps of the external type, as generally described in U.S. Pat. Nos. 4,562,751; 4,678,408; and 4,685,903; or automated implantable medication infusion pumps, as generally described in U.S. Pat. No. 4,573,994, which are herein incorporated by reference.

Generally, small and flexible electrochemical sensors can be used to obtain periodic readings over an extended period of time. In one form, flexible subcutaneous sensors are constructed in accordance with thin film mask techniques in which an elongated sensor includes thin film conductive elements encased between flexible insulative layers of polyimide sheets or similar material. Such thin film sensors typically include a plurality of exposed electrodes at one end for subcutaneous placement with a user's interstitial fluid, blood, or the like, and a corresponding exposed plurality of conductive contacts at another end for convenient external electrical connection with a suitable monitoring device through a wire or cable. Typical thin film sensors are described in commonly assigned U.S. Pat. Nos. 5,390,671; 5,391,250; 5,482,473; and 5,586,553 which are incorporated by reference herein. See also U.S. Pat. No. 5,299,571.

Drawbacks to the use of implantable sensors arise from the use of a wired connection between the implantable sensor set and the monitor. The use of the wire or cable is an additional inconvenience to users that already utilize an external infusion pump that includes an infusion insertion set and tube to infuse the medication. Also, the preferred site for some sensing device may be inconvenient for connection by wire to a characteristic monitor. For implantable pumps, the wire or cable negates the very benefit of having an

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internal device without external wires or cables. For Type 2 diabetics, who do not necessarily need or use an infusion pump, the use of a cable is seen as an inconvenience that may inhibit use of the device. In addition, the use of a wire or cable limits a user's ability to position the monitor, since it can be placed no further away than the ultimate length of the wire or cable. Thus, the user must normally wear the monitor, which can be problematic. For example, removal of the monitor for sleeping can be difficult, since a user would tend to become "tangled" in the wire or cable, between the sensor and the monitor, during the normal tossing and turning that occurs during sleep. Furthermore, the more connections the user must deal with (e.g., infusion pump and catheter and/or monitor with wire to sensor), the more complicated it is to use the devices, and the less likely the user will maintain compliance with the medical regimen due to perceived and actual difficulties with all of the wires and cables.

SUMMARY OF THE DISCLOSURE

It is an object of an embodiment of the present invention to provide an improved telemetered implantable sensor set (such as a subcutaneous or percutaneous sensor) and monitor connection device, which obviates for practical purposes, the above mentioned limitations.

According to an embodiment of the invention, a telemetered characteristic monitor system includes a remotely located data receiving device, a sensor for producing signal indicative of a characteristic of a user, and a transmitter device. In preferred embodiments, the transmitter device includes a housing, a sensor connector, a processor, and a transmitter. A potentiostat within the transmitter device may be coupled to the sensor connector and applies power to the sensor. The sensor connector receives the produced signals from the sensor. The processor is coupled to the sensor connector and processes the signals from the sensor for delivery to the remotely located data receiving device. The transmitter is coupled to the processor for wirelessly transmitting the processed signals to the remotely located data receiving device. In preferred embodiments, the data receiving device is a characteristic monitor. However, in other embodiments, the data receiving device is a data receiver that provides data to another device, an RF programmer, a medication delivery device (such as an infusion pump), or the like.

In particular embodiments, the transmitter of the transmitter device transmits the processed signals by radio frequencies. In other embodiments, the sensor may be implanted in and/or through subcutaneous, dermal, subdermal, intra-peritoneal or peritoneal tissue, and the sensor connector of the transmitter device includes a cable that is connected to the sensor. Also, the implantable sensor can be configured for a wired connection to a characteristic monitor, and the sensor connector of the transmitter device is formed to connect to the configured implantable sensor. Still further embodiments of the transmitter device include a receiver to receive data and instructions from the characteristic monitor, or the like.

Embodiments of the transmitter device (when used with a subcutaneous or percutaneous sensor) may include a bio-compatible adhesive to secure the housing to a skin surface of the user. Preferably, the housing of the transmitter device is less than about 3.0 inches in diameter by 0.5 inches thick. In addition, the housing is resistant to fluids when immersed in a fluid, operable in a temperature range of 0° C. to 50.° C., and has an operable life of at least 3 months. If the sensor

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is fully implanted, the transmitter that is connected to the sensor may be secured by sutures, sewing rings, or the like.

Other features and advantages of the invention will become apparent from the following detailed description, taken in conjunction with the accompanying drawings which illustrate, by way of example, various features of embodiments of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

A detailed description of embodiments of the invention will be made with reference to the accompanying drawings, wherein like numerals designate corresponding parts in the several figures.

FIG. 1 is a perspective view illustrating a subcutaneous sensor insertion set and telemetered characteristic monitor transmitter device embodying the novel features of the invention;

FIG. 2 is an enlarged longitudinal vertical section taken generally on the line 2-2 of FIG. 1;

FIG. 3 is an enlarged longitudinal sectional of a slotted insertion needle used in the insertion set of FIGS. 1 and 2;

FIG. 4 is an enlarged transverse section taken generally on the line 4-4 of FIG. 3;

FIG. 5 is an enlarged transverse section taken generally on the line 5-5 of FIG. 3;

FIG. 6 is an enlarged fragmented sectional view corresponding generally with the encircled region 6 of FIG. 2; and

FIG. 7 is an enlarged transverse section taken generally on the line 7-7 of FIG. 2.

FIG. 8(a) is a top plan and partial cut-away view of the telemetered characteristic monitor transmitter device in accordance with the embodiment shown in FIG. 1.

FIG. 8(b) is a simplified block diagram of the printed circuit board of the telemetered characteristic monitor transmitter device in accordance with the embodiments shown in FIG. 1.

FIG. 9 is a timing diagram illustrating an embodiment of a message and timing format used by the telemetered characteristic monitor transmitter device shown in FIG. 1.

FIG. 10 is a simplified block diagram of a characteristic monitor used in accordance with an embodiment of the present invention.

FIG. 11 is a timing diagram for the characteristic monitor shown in FIG. 10.

FIG. 12 is another timing diagram for the characteristic monitor shown in FIG. 10.

FIG. 13 is a simplified block diagram of a telemetered characteristic monitor transmitter and sensor set system in accordance with another embodiment of the present invention.

FIG. 14 is a simplified block diagram of a telemetered characteristic monitor transmitter and characteristic monitor system in accordance with still another embodiment of the present invention.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

As shown in the drawings for purposes of illustration, the invention is embodied in a telemetered characteristic monitor transmitter coupled to a sensor set, that may be implanted in and/or through subcutaneous, dermal, sub-dermal, interperitoneal or peritoneal tissue, that transmits data from the sensor set to the characteristic monitor for determining body characteristics. In preferred embodiments of the present

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invention, the sensor set and monitor are for determining glucose levels in the blood and/or body fluids of the user without the use of, or necessity of, a wire or cable connection between the transmitter and the monitor. However, it will be recognized that further embodiments of the invention may be used to determine the levels of other agents, characteristics or compositions, such as hormones, cholesterol, medication concentrations, pH, oxygen saturation, viral loads (e.g., HIV), or the like. In other embodiments, the sensor set may also include the capability to be programmed or calibrated using data received by the telemetered characteristic monitor transmitter device, or may be calibrated at the monitor device (or receiver). The telemetered characteristic monitor system is primarily adapted for use in subcutaneous human tissue. However, still further embodiments may be placed in other types of tissue, such as muscle, lymph, organ tissue, veins, arteries or the like, and used in animal tissue. Embodiments may provide sensor readings on an intermittent or continuous basis.

The telemetered characteristic monitor system 1, in accordance with a preferred embodiment of the present invention include a percutaneous sensor set 10, a telemetered characteristic monitor transmitter device 100 and a characteristic monitor 200. The percutaneous sensor set 10 utilizes an electrode-type sensor, as described in more detail below. However, in alternative embodiments, the system may use other types of sensors, such as chemical based, optical based or the like. In further alternative embodiments, the sensors may be of a type that is used on the external surface of the skin or placed below the skin layer of the user. Preferred embodiments of a surface mounted sensor would utilize interstitial fluid harvested from underneath the skin. The telemetered characteristic monitor transmitter 100 generally includes the capability to transmit data. However, in alternative embodiments, the telemetered characteristic monitor transmitter 100 may include a receiver, or the like, to facilitate two-way communication between the sensor set 10 and the characteristic monitor 200. The characteristic monitor 200 utilizes the transmitted data to determine the characteristic reading. However, in alternative embodiments, the characteristic monitor 200 may be replaced with a data receiver, storage and/or transmitting device for later processing of the transmitted data or programming of the telemetered characteristic monitor transmitter 100.

In addition, a relay or repeater 4 may be used with a telemetered characteristic monitor transmitter 100 and a characteristic monitor 200 to increase the distance that the telemetered characteristic monitor transmitter 100 can be used with the characteristic monitor 200, as shown in FIG. 13. For example, the relay 4 could be used to provide information to parents of children using the telemetered characteristic monitor transmitter 100 and the sensor set 10 from a distance. The information could be used when children are in another room during sleep or doing activities in a location remote from the parents. In further embodiments, the relay 4 can include the capability to sound an alarm. In addition, the relay 4 may be capable of providing telemetered characteristic monitor transmitter 100 data from the sensor set 10, as well as other data, to a remotely located individual via a modem connected to the relay 4 for display on a monitor, pager or the like. The data may also be downloaded through a Communication-Station 8 to a remotely located computer 6 such as a PC, lap top, or the like, over communication lines, by modem or wireless connection, as shown in FIG. 14. Also, some embodiments may omit the Communication Station 8 and uses a direct modem or wireless connection to the computer 6. In further

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embodiments, the telemetered characteristic monitor transmitter **100** transmits to an RF programmer, which acts as a relay, or shuttle, for data transmission between the sensor set **10** and a PC, laptop, Communication-station, a data processor, or the like. In further alternatives, the telemetered characteristic monitor transmitter **100** may transmit an alarm to a remotely located device, such as a communication-station, modem or the like to summon help. In addition, further embodiments may include the capability for simultaneous monitoring of multiple sensors and/or include a sensor for multiple measurements.

Still further embodiments of the telemetered characteristic monitor transmitter **100** may have and use an input port for direct (e.g., wired) connection to a programming or data readout device and/or be used for calibration of the sensor set **10**. Preferably, any port would be water proof (or water resistant) or include a water proof, or water resistant, removable cover.

The purpose of the telemetered characteristic monitor system **1** (see FIG. 2) is to provide for better treatment and control in an outpatient or a home use environment. For example, the monitor system **1** can provide indications of glucose levels, a hypoglycemia/hyperglycemia alert and outpatient diagnostics. It is also useful as an evaluation tool under a physician's supervision.

The monitor system **1** also removes inconvenience by separating the monitor electronics into two separate devices; a telemetered characteristic monitor transmitter **100**, which attaches to the implantable sensor set **10**; and a characteristic monitor **200** (or other receiver), which is carried like a pager. This provides several advantages over wire connected devices. For instance, the user can more easily conceal the presence of the monitor system **1**, since a wire will not be visible (or cumbersome), within clothing. Such remote communication also provides greater convenience and flexibility in the placement of the sensor. It also makes it is easier to protect the characteristic monitor **200**, which can be removed from the user's body during showers, exercise, sleep or the like. In addition, the use of multiple components (e.g., transmitter **100** and characteristic monitor **200**) facilitates upgrades or replacements, since one module or the other can be modified or replaced without requiring complete replacement of the monitor system **1**. Further, the use of multiple components can improve the economics of manufacturing, since some components may require replacement on a more frequent basis, sizing requirements may be different for each module, there may be different assembly environment requirements, and modifications can be made without affecting the other components.

The telemetered characteristic monitor transmitter **100** takes characteristic information, such as glucose data or the like, from the percutaneous sensor set **10** and transmits it via wireless telemetry to the characteristic monitor **200**, which displays and logs the received glucose readings. Logged data can be downloaded from the characteristic monitor **200** to a personal computer, laptop, or the like, for detailed data analysis. In further embodiments, the telemetered characteristic monitor system **1** may be used in a hospital environment or the like. Still further embodiments of the present invention may include one or more buttons (on the telemetered characteristic monitor transmitter **100** or characteristic monitor **200**) to record data and events for later analysis, correlation, or the like. In addition, the telemetered characteristic monitor transmitter **100** may include a transmit on/off button for compliance with safety standards and regulations to temporarily suspend transmissions. Further buttons can include a sensor on/off button to conserve power

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and to assist in initializing the sensor set **10**. The telemetered characteristic monitor transmitter **100** and characteristic monitor **200** may also be combined with other medical devices to combine other patient data through a common data network and telemetry system.

Further embodiments of the percutaneous sensor set **10** would monitor the temperature of the sensor set **10**, which can then be used to improve the calibration of the sensor. For instance, for a glucose sensor, the enzyme reaction activity may have a known temperature coefficient. The relationship between temperature and enzyme activity can be used to adjust the sensor values to more accurately reflect the actual characteristic levels. In addition to temperature measurements, the oxygen saturation level can be determined by measuring signals from the various electrodes of the sensor set **10**. Once obtained, the oxygen saturation level may be used in calibration of the sensor set **10** due to changes in the oxygen saturation levels, and its effects on the chemical reactions in the sensor set **10**. For instance, as the oxygen level goes lower the sensor sensitivity may be lowered. The oxygen level can be utilized in calibration of the sensor set **10** by adjusting for the changing oxygen saturation. In alternative embodiments, temperature measurements may be used in conjunction with other readings to determine the required sensor calibration.

As shown in FIGS. 1-7, a percutaneous sensor set **10** is provided for subcutaneous placement of an active portion of a flexible sensor **12** (see FIG. 2), or the like, at a selected site in the body of a user. The subcutaneous or percutaneous portion of the sensor set **10** includes a hollow, slotted insertion needle **14**, and a cannula **16**. The needle **14** is used to facilitate quick and easy subcutaneous placement of the cannula **16** at the subcutaneous insertion site. Inside the cannula **16** is a sensing portion **18** of the sensor **12** to expose one or more sensor electrodes **20** to the user's bodily fluids through a window **22** formed in the cannula **16**. After insertion, the insertion needle **14** is withdrawn to leave the cannula **16** with the sensing portion **18** and the sensor electrodes **20** in place at the selected insertion site.

In preferred embodiments, the percutaneous sensor set **10** facilitates accurate placement of a flexible thin film electrochemical sensor **12** of the type used for monitoring specific blood parameters representative of a user's condition. Preferably, the sensor **12** monitors glucose levels in the body, and may be used in conjunction with automated or semi-automated medication infusion pumps of the external or implantable type as described in U.S. Pat. No. 4,562,751; 4,678,408; 4,685,903 or 4,573,994, to control delivery of insulin to a diabetic patient.

Preferred embodiments of the flexible electrochemical sensor **12** are constructed in accordance with thin film mask techniques to include elongated thin film conductors embedded or encased between layers of a selected insulative material such as polyimide film or sheet, and membranes. The sensor electrodes **20** at a tip end of the sensing portion **18** are exposed through one of the insulative layers for direct contact with patient blood or other body fluids, when the sensing portion **18** (or active portion) of the sensor **12** is subcutaneously placed at an insertion site. The sensing portion **18** is joined to a connection portion **24** (see FIG. 2) that terminates in conductive contact pads, or the like, which are also exposed through one of the insulative layers. In alternative embodiments, other types of implantable sensors, such as chemical based, optical based, or the like, may be used.

As is known in the art, and illustrated schematically in FIG. 2, the connection portion **24** and the contact pads are

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generally adapted for a direct wired electrical connection to a suitable monitor **200** for monitoring a user's condition in response to signals derived from the sensor electrodes **20**. Further description of flexible thin film sensors of this general type are be found in U.S. Pat. No. 5,391,250, entitled METHOD OF FABRICATING THIN FILM SENSORS, which is herein incorporated by reference. The connection portion **24** may be conveniently connected electrically to the monitor **200** or a telemetered characteristic monitor transmitter **100** by a connector block **28** (or the like) as shown and described in U.S. Pat. No. 5,482,473, entitled FLEX CIRCUIT CONNECTOR, which is also herein incorporated by reference. Thus, in accordance with embodiments of the present invention, subcutaneous sensor sets **10** are configured or formed to work with either a wired or a wireless characteristic monitor system.

The proximal part of the sensor **12** is mounted in a mounting base **30** adapted for placement onto the skin of a user. As shown, the mounting base **30** is a pad having an underside surface coated with a suitable pressure sensitive adhesive layer **32**, with a peel-off paper strip **34** normally provided to cover and protect the adhesive layer **32**, until the sensor set **10** is ready for use. As shown in FIGS. 1 and 2, the mounting base **30** includes upper and lower layers **36** and **38**, with the connection portion **24** of the flexible sensor **12** being sandwiched between the layers **36** and **38**. The connection portion **24** has a forward section joined to the active sensing portion **18** of the sensor **12**, which is folded angularly to extend downwardly through a bore **40** formed in the lower base layer **38**. In preferred embodiments, the adhesive layer **32** includes an anti-bacterial agent to reduce the chance of infection; however, alternative embodiments may omit the agent. In the illustrated embodiment, the mounting base is generally rectangular, but alternative embodiments may be other shapes, such as circular, oval, hour-glass, butterfly, irregular, or the like.

The insertion needle **14** is adapted for slide-fit reception through a needle port **42** formed in the upper base layer **36** and further through the lower bore **40** in the lower base layer **38**. As shown, the insertion needle **14** has a sharpened tip **44** and an open slot **46** which extends longitudinally from the tip **44** at the underside of the needle **14** to a position at least within the bore **40** in the lower base layer **36**. Above the mounting base **30**, the insertion needle **14** may have a full round cross-sectional shape, and may be closed off at a rear end of the needle **14**. Further description of the needle **14** and the sensor set **10** are found in U.S. Pat. No. 5,586,553, entitled "TRANSCUTANEOUS SENSOR INSERTION SET" and co-pending U.S. patent application Ser. No. 08/871,831, entitled "DISPOSABLE SENSOR INSERTION ASSEMBLY," which are herein incorporated by reference.

The cannula **16** is best shown in FIGS. 6 and 7, and includes a first portion **48** having partly-circular cross-section to fit within the insertion needle **14** that extends downwardly from the mounting base **30**. In alternative embodiments, the first portion **48** may be formed with a solid core; rather than a hollow core. In preferred embodiments, the cannula **16** is constructed from a suitable medical grade plastic or elastomer, such as polytetrafluoroethylene, silicone, or the like. The cannula **16** also defines an open lumen **50** in a second portion **52** for receiving, protecting and guideably supporting the sensing portion **18** of the sensor **12**. The cannula **16** has one end fitted into the bore **40** formed in the lower layer **38** of the mounting base **30**, and the cannula **16** is secured to the mounting base **30** by a suitable adhesive, ultrasonic welding, snap fit or other

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selected attachment method. From the mounting base **30**, the cannula **16** extends angularly downwardly with the first portion **48** nested within the insertion needle **14**, and terminates before the needle tip **44**. At least one window **22** is formed in the lumen **50** near the implanted end **54**, in general alignment with the sensor electrodes **20**, to permit direct electrode exposure to the user's bodily fluid when the sensor **12** is subcutaneously placed. Alternatively, a membrane can cover this area with a porosity that controls rapid diffusion of glucose through the membrane.

As shown in FIGS. 1, 2 and 8(a), the telemetered characteristic monitor transmitter **100** is coupled to a sensor set **10** by a cable **102** through a connector **104** that is electrically coupled to the connector block **28** of the connector portion **24** of the sensor set **10**. In alternative embodiments, the cable **102** may be omitted, and the telemetered characteristic monitor transmitter **100** may include an appropriate connector (not shown) for direct connection to the connector portion **24** of the sensor set **10** or the sensor set **10** may be modified to have the connector portion **24** positioned at a different location, such as for example, on the top of the sensor set **10** to facilitate placement of the telemetered characteristic monitor transmitter over the subcutaneous sensor set **10**. This would minimize the amount of skin surface covered or contacted by medical devices, and tend to minimize movement of the sensor set **10** relative to the telemetered characteristic monitor transmitter **100**. In further alternative embodiments, the cable **102** and the connector **104** may be formed as add-on adapters to fit different types of connectors on different types or kinds of sensor sets. The use of adapters would facilitate adaptation of the telemetered characteristic monitor transmitter **100** to work with a wide variety of sensor systems. In further embodiments, the telemetered characteristic monitor transmitter **100** may omit the cable **102** and connector **104** and is instead optically couple with an implanted sensor, in the subcutaneous, dermal, sub-dermal, inter-peritoneal or peritoneal tissue, to interrogate the implanted sensor using visible, and/or IR frequencies, either transmitting to and receiving a signal from the implanted sensor or receiving a signal from the implanted sensor.

The telemetered characteristic monitor **100** (also known as Potentiostat Transmitter Device) includes a housing **106** that supports a printed circuit board **108**, batteries **110**, antenna **112**, and the cable **102** with the connector **104**. In preferred embodiments, the housing **106** is formed from an upper case **114** and a lower case **116** that are sealed with an ultrasonic weld to form a waterproof (or resistant) seal to permit cleaning by immersion (or swabbing) with water, cleaners, alcohol or the like. In preferred embodiments, the upper and lower case **114** and **116** are formed from a medical grade plastic. However, in alternative embodiments, the upper case **114** and lower case **116** may be connected together by other methods, such as snap fits, sealing rings, RTV (silicone sealant) and bonded together, or the like, or formed from other materials, such as metal, composites, ceramics, or the like. In other embodiments, the separate case can be eliminated and the assembly is simply potted in epoxy or other moldable materials that is compatible with the electronics and reasonably moisture resistant. In preferred embodiments, the housing **106** is disk or oval shaped. However, in alternative embodiments, other shapes, such as hour glass, rectangular or the like, may be used. Preferred embodiments of the housing **106** are sized in the range of 2.0 square inches by 0.35 inches thick to minimize weight, discomfort and the noticeability of the telemetered characteristic monitor transmitter **100** on the body of the user.

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However, larger or smaller sizes, such as 1.0 square inches and 0.25 inches thick or less, and 3.0 square inches and 0.5 inches thick or more, may be used. Also, the housing may simply be formed from potted epoxy, or other material, especially if the battery life relative to the device cost is long enough, or if the device is rechargeable.

As shown, the lower case **116** may have an underside surface coated with a suitable pressure sensitive adhesive layer **118**, with a peel-off paper strip **120** normally provided to cover and protect the adhesive layer **118**, until the sensor set telemetered characteristic monitor transmitter **100** is ready for use. In preferred embodiments, the adhesive layer **118** includes an anti-bacterial agent to reduce the chance of infection; however, alternative embodiments may omit the agent. In further alternative embodiments, the adhesive layer **118** may be omitted and the telemetered characteristic monitor transmitter **100** is secured to the body by other methods, such as an adhesive overdressing, straps, belts, clips or the like.

In preferred embodiments, the cable **102** and connector **104** are similar to (but not necessarily identical to) shortened versions of a cable and connector that are used to provide a standard wired connection between the sensor set **10** and the characteristic monitor **200**. This allows the telemetered characteristic monitor transmitter **100** to be used with existing sensor sets **10**, and avoids the necessity to re-certify the connector portion **24** of the sensor set **10** for use with a wireless connection. The cable **102** should also include a flexible strain relief portion (not shown) to minimize strain on the sensor set **10** and prevent movement of the inserted sensor **12**, which can lead to discomfort or dislodging of the sensor set **10**. The flexible strain relief portion is intended to minimize sensor artifacts generated by user movements that might cause the sensing area of the sensor set **10** to move relative to the body tissues in contact with the sensing area of the sensor set **10**.

The printed circuit board **108** of the telemetered characteristic monitor transmitter **100** includes a sensor interface **122**, processing electronics **124**, timers **126**, and data formatting electronics **128**, as shown in FIG. 8(b). In preferred embodiments, the sensor interface **122**, processing electronics **124**, timers **126**, and data formatting electronics **128** are formed as separate semiconductor chips; however, alternative embodiments may combine the various semiconductor chips into a single customized semiconductor chip. The sensor interface **122** connects with the cable **102** that is connected with the sensor set **10**. In preferred embodiments, the sensor interface is permanently connected to the cable **102**. However, in alternative embodiments, the sensor interface **122** may be configured in the form of a jack to accept different types of cables that provide adaptability of the telemetered characteristic monitor transmitter **100** to work with different types of sensors and/or sensors placed in different locations of the user's body. In preferred embodiments, the printed circuit board **108**, and associated electronics, are capable of operating in a temperature range of 0° C. and 50° C. However, larger or smaller temperature ranges may be used.

Preferably, the battery assembly will use a weld tab design to connect power to the system. For example, it can use three series silver oxide 357 battery cells **110**, or the like. However, it is understood that different battery chemistries may be used, such as lithium based chemistries, alkaline batteries, nickel metalhydride, or the like, and different numbers

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of batteries can be used. In further embodiments, the sensor interface **122** will include circuitry and/or a mechanism for detecting connection to the sensor set **10**. This would provide the capability to save power and to more quickly and efficiently start initialization of the sensor set **10**. In preferred embodiments, the batteries **110** have a life in the range of 3 months to 2 years, and provide a low battery warning alarm. Alternative embodiments may provide longer or shorter battery lifetimes, or include a power port, solar cells or an inductive coil to permit recharging of rechargeable batteries in the telemetered characteristic monitor transmitter **100**.

In preferred embodiments, the telemetered characteristic monitor transmitter **100** provides power, through the cable **102** and cable connector **104** to the sensor set **10**. The power is used to monitor and drive the sensor set **10**. The power connection is also used to speed the initialization of the sensor **12**, when it is first placed under the skin. The use of an initialization process can reduce the time for sensor **12** stabilization from several hours to an hour or less. The preferred initialization procedure uses a two step process. First, a high voltage (preferably between 1.0-1.2 volts—although other voltages may be used) is applied to the sensor **12** for 1 to 2 minutes (although different time periods may be used) to allow the sensor **12** to stabilize. Then, a lower voltage (preferably between 0.5-0.6 volts—although other voltages may be used) is applied for the remainder of the initialization process (typically 58 minutes or less). Other stabilization/initialization procedures using differing currents, currents and voltages, different numbers of steps, or the like, may be used. Other embodiments may omit the initialization/stabilization process, if not required by the sensor or if timing is not a factor.

At the completion of the stabilizing process, a reading may be transmitted from the sensor set **10** and the telemetered characteristic monitor transmitter **100** to the characteristic monitor **200**, and then the user will input a calibrating glucose reading into characteristic monitor **200**. In alternative embodiments, a fluid containing a known value of glucose may be injected into the site around the sensor set **10**, and then the reading is sent to the characteristic monitor **200** and the user inputs the known concentration value, presses a button (not shown) or otherwise instructs the monitor to calibrate using the known value. During the calibration process, the telemetered characteristic monitor transmitter **100** checks to determine if the sensor set **10** is still connected. If the sensor set **10** is no longer connected, the telemetered characteristic monitor transmitter **100** will abort the stabilization process and sound an alarm (or send a signal to the characteristic monitor **200** to sound an alarm).

Preferably, the transmissions (or telemetry) of the telemetered characteristic monitor transmitter **100** will contain at least the following information: a unique ID code that uniquely identifies each telemetered characteristic monitor transmitter **100**, a sensor characteristic data signal representative of the measured characteristic value (e.g., glucose or the like) from the sensor **18** of the subcutaneous sensor set **10**, a counter electrode voltage, a low battery flag, and error detection bits (such as CRC). FIG. 9 and Table 1 illustrate a preferred message format for the telemetry of the telemetered characteristic monitor transmitter **100**. However, it will be understood that different message protocols and structures may be used.

TABLE 1

Message format	
1. Encoding method: OOK MANCHESTER (1 = 1/0, 0 = 0/1 sequence, where 1 = transmitter (TX) on)	
2. Clock rate 1024 Hz (512 Hz symbol/bit rate).	
3. Message format:	
Preamble:	4 bits (0101, want only one transition per bit)
Message Type:	4 bits (1010 = transmitter 100, 15 others for pump/ppc protocol)
Unique ID #:	16 bits (65536 unique numbers)
Message count #:	4 bits (also determines TX time slot)
Working	12 bits (9 MSBs + 3 magnitude bits =
Electrode:	16 bit range - converted by the characteristic monitor 200 into a value representative of characteristic level, such as glucose level)
Low battery flag:	1 bit (0 = ok, 1 = low)
Counter Voltage:	7 bits (0-1.2 V typ., 8 bit a/d 3.2 V FS)
CRC:	8 bits
Total	56 bits (times, 1/512 hz = 109 mS)
4. Message TX interval: 300 seconds (5 min) + 1-16 seconds pseudo- random delay (TX time slot)	
5. TX duty cycle: 56 bits*1/512 Hz* 1/300 S*1/2 = 1.823e-4	

Preferred embodiments utilize a time-slicing transmission protocol. Use of the time slicing protocol facilitates the use of multiple signals on the same frequency bands or to the same receiver from multiple transmitters. The time-slicing may also be used to obviate the need for a receiver in the telemetered characteristic monitor transmitter 100. For instance, the use of intermittent transmission reduces the amount of power required to operate the transmitter 100 and to extend the life of the device. It also saves power in the characteristic monitor 200 by reducing the amount of time the characteristic monitor 200 must spend in the receive mode.

In preferred embodiments, when the telemetered characteristic monitor transmitter 100 is connected to the sensor set 10, it detects the connection and is activated. Next, if desired or necessary, the telemetered characteristic monitor transmitter 100 initializes the sensor 12 of the sensor set 10. After (or in some cases during) initialization, the telemetered characteristic monitor transmitter 100 sends out a message of between 100-150 ms length every 5 minutes. Although other timing intervals ranging from 1 second to 30 minutes may be used.

Preferably, the message is transmitted in a pseudo-randomly selected time window within the 128 seconds following the 5 minute interval. In preferred embodiments, the telemetered characteristic transmitter 100 utilizes its own unique ID as a random-seed to set up a table of transmission time windows that defines the order in which the telemetered characteristic monitor 100 will transmit a message following the 5 minute interval. The order is repeated after the table is set-up. Included in the message sent will be the message count number, which indicates where in the sequence of time windows the telemetered characteristic monitor transmitter is currently transmitting. The characteristic monitor 200 uses the unique ID code of the telemetered characteristic monitor transmitter 100 to set up a corresponding table in the characteristic monitor 200 and the received message count to synchronize the characteristic monitor 200 with the current position in the table being used by the telemetered characteristic monitor transmitter 100 to predict the next time window to be used. The use of pseudo-random time windows prevents multiple transmitters from continuously interfering with other transmitting devices that are tempo-

rarily, or inadvertently, synchronized with the telemetered characteristic monitor transmitter 100. The characteristic monitor 200 acquires the transmitted message, and determines the time window in which the characteristic monitor 200 must be in a receive mode to acquire the next message. The characteristic monitor 200 then places itself in the receive mode every 5 minutes (although other timing intervals from 1 second to 30 minutes may be used) to receive the next message and data from the telemetered characteristic monitor transmitter 100 at the next predicted time window. Thus, the characteristic monitor 200 needs only be in the receive mode for 1 second (i.e., 1 time window); rather than 128 seconds (128 time windows). In alternative embodiments, the characteristic monitor 200 may not use the unique ID and the message count and may remain in the receive mode during the entire period (e.g., for 128 time windows) during which a transmission is possible. In addition, other embodiments may cause the characteristic monitor 200 to enter the receive mode 1 time window ahead and stay on for 1 time window longer to maximize the likelihood of receiving the next transmission. In further alternative embodiments, the telemetered characteristic monitor transmitter 100 and/or characteristic monitor 200 may utilize other methods or numbers to determine when transmission time windows are selected. Alternative message time-slicing transmission parameters, such as message length, number of time windows, frequency of transmissions, or the like, that are larger or smaller than those describe above, may also be used. Preferred embodiments transmit the data and/or information at a data rate between 1000 Hz to 4000 Hz modulated onto a high frequency carrier wave. However, alternative embodiments may use smaller or larger transmission rates, with the rate being selected based on user environment, power requirements, interference issues, redundancy criteria, or the like.

If a transmitted message is not received by the characteristic monitor 200 after a predetermined period of time, an alarm will be sounded or provided. In addition, the characteristic monitor 200 may continue to attempt to receive the next message by entering the receive mode at the next anticipated transmission time or may expand to enter the receive mode to cover all time windows until the next message is received.

In another alternative embodiment, if there is little or no likelihood of interference from other telemetered characteristic monitor transmitter 100, such as by message length, frequency selections or the like, the telemetered characteristic monitor transmitter 100 may transmit at one time window for all cases (typically the choice of window may be randomly selected at connection of a sensor set 10 or set at the factory). This permits the characteristic monitor 200 to be in the receive mode for even shorter periods of time (i.e., approximately 200 ms to bracket the telemetered characteristic monitor transmitter 100 transmission instead of the 128 seconds (or 1 second if able to predict the next time window) needed to bracket 128 windows) to conserve power in the characteristic monitor 200. For instance, in this scenario, the characteristic monitor 200 will be in a non-receive mode for 299.8 seconds and in a receive mode for 200 ms. In particular embodiments the non-receive mode and receive mode periods will be determined by the message length and expected frequency of transmission. It is also noted that in a system where the receiver must cover a range of time windows, the receiver may lock on to a particular range of time windows to permit the receiver being in the receive mode for shorter periods of time.

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The use of these transmission protocols obviates the need for a transmitter and receiver in both the telemetered characteristic monitor transmitter **100** and characteristic monitor **200**, which reduces costs, simplifies the system design, reduces power consumption and the like. However, alternative embodiments may include the capability for two-way communication, if desirable. In further embodiments, the telemetered characteristic monitor transmitter **100** transmits continuously and the characteristic monitor **200** enters the receive mode when desired or required to determine a characteristic value, such as a glucose level or the like.

In preferred embodiments, the telemetered characteristic monitor transmitter **100** will have the ability to uniquely identify itself to the characteristic monitor **200**. The telemetered characteristic monitor transmitter **100** will have an operating range to the characteristic monitor **200** of at least 10 feet. In alternative embodiments, larger or smaller ranges may be used, with the selection being dependent on the environment in which the telemetered characteristic monitor transmitter **100** will be used, the size and needs of the user, power requirements, and the like.

In further alternative embodiments, the telemetered characteristic monitor transmitter **100** can be combined with a sensor set **10** as a single unit. This would be particularly well adapted where batteries and the transmitter can be made cheaply enough to facilitate changing the transmitter **100** with each new sensor set **10**.

As shown in FIG. **10**, the characteristic monitor **200** includes a telemetry receiver **202**, a Telemetry Decoder (TD) **204** and a host micro-controller (Host) **206** for communication with the telemetered characteristic monitor transmitter **100**. The TD **204** is used to decode a received telemetry signal from the transmitter device and forward the decoded signal to the Host **206**. The Host **206** is a microprocessor for data reduction, data storage, user interface, or the like. The telemetry receiver **202** receives the characteristic data (e.g., glucose data) from the telemetered characteristic monitor transmitter, and passes it to the TD **204** for decoding and formatting. After complete receipt of the data by the TD **204**, the data is transferred to the Host **206** for processing, where calibration information, based upon user entered characteristic readings (e.g., blood glucose readings), is performed to determine the corresponding characteristic level (e.g., glucose level) from measurement in the characteristic data (e.g., glucose data). The Host **206** also provides for storage of historical characteristic data, and can download the data to a personal computer, lap-top, or the like, via a com-station, wireless connection, modem or the like. For example, in preferred embodiments, the counter electrode voltage is included in the message from the telemetered characteristic monitor transmitter **100** and is used as a diagnostic signal. The raw current signal values generally range from 0 to 999, which represents sensor electrode current in the range between 0.0 to 99.9 nanoAmperes, and is converted to characteristic values, such as glucose values in the range of 40 to 400 mg/dl. However, in alternative embodiments, larger or smaller ranges may be used. The values are then displayed on the characteristic monitor **200** or stored in data memory for later recall.

The characteristic monitor **200** also includes circuitry in the TD **204** to uniquely mate it to an identified telemetered characteristic monitor transmitter **100**. In preferred embodiments, the identification number of the telemetered characteristic monitor transmitter **100** is entered manually by the user using keys located on the characteristic monitor **200**. In alternative embodiments, the characteristic monitor **200** includes a "learn ID" mode. Generally, the "learn ID" mode

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is best suited for the home environment, since multiple telemetered characteristic monitor transmitters **100**, typically encountered in a hospital setting, are less likely to cause confusion in the characteristic monitor **200** when it attempts to learn an ID code. In addition, the characteristic monitor **200** will include the ability to learn or be reprogrammed to work with a different (or replacement) telemetered characteristic monitor transmitter **100**. The preferred operating distance is at least 10 feet. In alternative embodiments, larger or smaller ranges may be used, with the selection being dependent on the environment in which the telemetered characteristic monitor transmitter **100** will be used, the size and needs of the user, power requirements, and the like. Furthermore, if the characteristic monitor **200** does not receive a transmission from the identified telemetered characteristic monitor transmitter **100** after a certain period of time (e.g., one or more missed transmissions), an alarm will be sounded.

In preferred embodiments, the characteristic monitor **200** utilizes a two processor system, in which the Host **206** is the master processor and the TD **204** is a slave processor dedicated to telemetry processing. A first communication protocol between the Host **206** and the TD **204** is shown in FIG. **11**. The first protocol uses a serial peripheral interface (SPI) **208** and two control lines **210** and **212**; one control line (chip select—CSPIC) **210** is used by Host **206** to wake up the TD **204** to initiate telemetry receiving task; and the other control line (data ready—DR) **212** is used by the TD **204** to indicate to the Host **206** that the telemetry data from the telemetered characteristic monitor transmitter has been received and is ready to be transferred to the HCO **206**. Upon receiving data through the SPI **208**, the Host **206** sends an acknowledgment through the SPI **208** to the TD **204**. In preferred embodiments, fixed length data blocks are used. However, in alternative embodiments, variable length data blocks may be used. In preferred embodiments, the Host **206** may pull the Chip Select (CSPIC) **210** high at any time to abort the telemetry data transfer from the TD **204**. Alternatively, an additional line (not shown) may be used to reset the TD **204**. FIG. **12** shows a second, more complex, alternative protocol that is used by the Host **206** and the TD **204**.

In alternative embodiments, the TD **204** and Host **206** may be combined together in a single semiconductor chip to obviate the need for dual processors and to reduce the space needed for the electronics. In further embodiments, the functions of the TD **204** and Host **206** may be allocated differently between one or more processors.

As shown in FIG. **2**, the characteristic monitor may include a display **214** that is used to display the results of the measurement received from the sensor **18** in the sensor set **10** via the telemetered characteristic monitor transmitter **100**. The results and information displayed includes, but is not limited to, trending information of the characteristic (e.g., rate of change of glucose), graphs of historical data, average characteristic levels (e.g., glucose), or the like. Alternative embodiments include the ability to scroll through the data. The display **214** may also be used with buttons (not shown) on the characteristic monitor to program or update data in the characteristic monitor **200**. It is noted that the typical user can be expected to have somewhat diminished visual and tactile abilities due to complications from diabetes or other conditions. Thus, the display **214** and buttons should be configured and adapted to the needs of a user with diminished visual and tactile abilities. In alternative embodiments, the value can be conveyed to the user by audio signals, such as beeps, speech or the like. Still further

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embodiments may use a touch screen instead of (or in some cases addition to) buttons to facilitate water proofing and to ease changes in the characteristic monitor 200 hardware to accommodate improvements or upgrades.

Preferably, the characteristic monitor uses batteries (not shown) to provide power to the characteristic monitor. For example, a plurality of silver oxide batteries may be used. However, it is understood that different battery chemistries may be used, such as lithium based, alkaline based, nickel metalhydride, or the like, and different numbers of batteries can be used. In preferred embodiments, the batteries have a life in the range of 1 month to 2 years, and provide a low battery warning alarm. Alternative embodiments may provide longer or shorter battery lifetimes, or include a power port, solar cells or an induction coil to permit recharging of rechargeable batteries in the characteristic monitor 200. In preferred embodiments, the batteries are not replaceable to facilitate waterproofing the housing 106.

In further embodiments of the present invention, the characteristic monitor 200 may be replaced by a different device. For example, in one embodiment, the telemetered characteristic monitor transmitter 100 communicates with an RF programmer (not shown) that is also used to program and obtain data from an infusion pump or the like. The RF programmer may also be used to update and program the transmitter 100, if the transmitter 100 includes a receiver for remote programming, calibration or data receipt. The RF programmer can be used to store data obtained from the sensor 18 and then provide it to either an infusion pump, characteristic monitor, computer or the like for analysis. In further embodiments, the transmitter 100 may transmit the data to a medication delivery device, such as an infusion pump or the like, as part of a closed loop system. This would allow the medication delivery device to compare sensor results with medication delivery data and either sound alarms when appropriate or suggest corrections to the medication delivery regimen. In preferred embodiments, the transmitter 100 would include a transmitter to receive updates or requests for additional sensor data. An example of one type of RF programmer can be found in U.S. patent application Ser. No. 60/096,994 filed Aug. 18, 1998 and is entitled "INFUSION DEVICE WITH REMOTE PROGRAMMING, CARBOHYDRATE CALCULATOR AND/OR VIBRATION ALARM CAPABILITIES," or U.S. patent application Ser. No. 09/334,858 filed Jun. 17, 1999 and is entitled "EXTERNAL INFUSION DEVICE WITH REMOTE PROGRAMMING, BOLUS ESTIMATOR AND/OR VIBRATION ALARM CAPABILITIES," both of which are herein incorporated by reference.

In further embodiments, the telemetered characteristic monitor transmitter can include a modem, or the like, to transfer data to and from a healthcare professional. Further embodiments, can receive updated programming or instructions via a modem connection.

In use, the sensor set 10 permits quick and easy subcutaneous placement of the active sensing portion 18 at a selected site within the body of the user. More specifically, the peel-off strip 34 (see FIG. 1) is removed from the mounting base 30, at which time the mounting base 30 can be pressed onto and seated upon the patient's skin. During this step, the insertion needle 14 pierces the user's skin and carries the protective cannula 16 with the sensing portion 18 to the appropriate subcutaneous placement site. During insertion, the cannula 16 provides a stable support and guide structure to carry the flexible sensor 12 to the desired placement site. When the sensor 12 is subcutaneously placed, with the mounting base 30 seated upon the user's

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skin, the insertion needle 14 can be slidably withdrawn from the user. During this withdrawal step, the insertion needle 14 slides over the first portion 48 of the protective cannula 16, leaving the sensing portion 18 with electrodes 20 directly exposed to the user's body fluids via the window 22. Further description of the needle 14 and the sensor set 10 are found in U.S. Pat. No. 5,586,553, entitled "TRANSCUTANEOUS SENSOR INSERTION SET"; co-pending U.S. patent application Ser. No. 08/871,831, entitled "DISPOSABLE SENSOR INSERTION ASSEMBLY"; and co-pending U.S. patent application Ser. No. 09/161,128, filed Sep. 25, 1998, entitled "A SUBCUTANEOUS IMPLANTABLE SENSOR SET HAVING THE CAPABILITY TO REMOVE OR DELIVER FLUIDS TO AN INSERTION SITE," which are herein incorporated by reference.

Next, the user connects the connection portion 24 of the sensor set 10 to the cable 102 of the telemetered characteristic monitor transmitter 100, so that the sensor 12 can then be used over a prolonged period of time for taking blood chemistry or other characteristic readings, such as blood glucose readings in a diabetic patient. Preferred embodiments of the telemetered characteristic monitor transmitter 100 detect the connection of the sensor 12 to activate the telemetered characteristic monitor transmitter 100. For instance, connection of the sensor 12 may activate a switch or close a circuit to turn the telemetered characteristic monitor transmitter 100 on. The use of a connection detection provides the capability to maximize the battery and shelf life of the telemetered characteristic monitor transmitter prior to use, such as during manufacturing, test and storage. Alternative embodiments of the present invention may utilize an on/off switch (or button) on the telemetered characteristic monitor transmitter 100.

The transmitter 100 is then affixed to the user's body with an adhesive overdressing. Alternatively, the peel-off strip 34 (see FIG. 1) is removed from the lower case 116, at which time the lower case 116 can be pressed onto and seated upon the patient's skin. The user then activates the transmitter 100, or the transmitter is activated by detection of the connection to the sensor 12 of the sensor set 10. Generally, the act of connecting (and disconnecting) the sensor 12 activates (and deactivates) the telemetered characteristic monitor 100, and no other interface is required. In alternative steps, the sensor set 10 is connected to the transmitter 100 prior to placement of the sensor 12 to avoid possible movement or dislodging of the sensor 12 during attachment of the transmitter 100. Also, the transmitter may be attached to the user prior to attaching the sensor set 10 to the transmitter 100.

The user then programs the characteristic monitor (or it learns) the identification of the transmitter 100 and verifies proper operation and calibration of the transmitter 100. The characteristic monitor 200 and transmitter 100 then work to transmit and receive sensor data to determine characteristic levels. Thus, once a user attaches a transmitter 100 to a sensor set 10, the sensor 12 is automatically initialized and readings are periodically transmitted, together with other information, to the characteristic monitor 200.

After a sensor set 10 has been used for a period of time, it is replaced. The user will disconnect the sensor set 10 from the cable 102 of the telemetered characteristic monitor transmitter 100. In preferred embodiments, the telemetered characteristic monitor transmitter 100 is removed and positioned adjacent the new site for a new sensor set 10. In alternative embodiments, the user does not need to remove the transmitter 100. A new sensor set 10 and sensor 12 are attached to the transmitter 100 and connected to the user's

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body. Monitoring then continues, as with the previous sensor 12. If the user must replace the telemetered characteristic monitor transmitter 100, the user disconnects the transmitter 100 from the sensor set 10 and the user's body. The user then connects a new transmitter 100, and reprograms the characteristic monitor (or learns) to work with the new transmitter 100. Monitoring then continues, as with the previous sensor 12.

Additional embodiments of the present invention may include a vibrator alarm (or optional indicator such as an L.E.D.) in either or both the telemetered characteristic monitor transmitter 100 and the characteristic monitor 200 to provide a tactile (vibration) alarm to the user, such as sensor set malfunction, improper connection, low battery, missed message, bad data, transmitter interference, or the like. The use of a vibration alarm provides additional reminders to an audio alarm, which could be important with someone suffering an acute reaction, or to have non-audio alarms to preserve and conceal the presence of the telemetered characteristic monitor system 1.

While the description above refers to particular embodiments of the present invention, it will be understood that many modifications may be made without departing from the spirit thereof. The accompanying claims are intended to cover such modifications as would fall within the true scope and spirit of the present invention.

The presently disclosed embodiments are therefore to be considered in all respects as illustrative and not restrictive, the scope of the invention being indicated by the appended claims, rather than the foregoing description, and all changes which come within the meaning and range of equivalency of the claims are therefore intended to be embraced therein.

What is claimed is:

1. A telemetered characteristic monitor system for monitoring a characteristic of a user, the system comprising:
 - a remotely located data receiving device;
 - a sensor for producing a first signal indicative of a characteristic of a user and a second signal indicative of the oxygen saturation level of the sensor; and
 - a transmitter device including:
 - a housing;
 - a sensor connector coupled to the housing and that is connectable to the sensor to receive signals from the sensor;

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a processor within the housing coupled to the sensor connector to process the signals from the sensor for delivery to the remotely located data receiving device; and

a transmitter for wirelessly transmitting the processed signals to the remotely located data receiving device.

2. The telemetered characteristic monitor system of claim 1, wherein the sensor is calibrated based on the oxygen level.

3. The telemetered characteristic monitor system of claim 1, further comprising a sensor set including the sensor and a connector coupled to the sensor, wherein the sensor connector of the transmitter device is directly connected to the connector of the sensor set without any intermediate cable between the two.

4. The telemetered characteristic monitor system of claim 1 wherein the size of the housing is not greater than 2.0 square inches by 0.35 inches thick.

5. The telemetered characteristic monitor system of claim 1 wherein the size of the housing is not greater than 1.0 square inches by 0.25 inches thick.

6. The telemetered characteristic monitor system of claim 1, wherein the sensor is implantable in tissue selected from the group consisting of subcutaneous, dermal, sub-dermal, intra-peritoneal, and peritoneal tissue.

7. The telemetered characteristic monitor system of claim 1, wherein the sensor is a percutaneous sensor.

8. The telemetered characteristic monitor system of claim 1, wherein the transmitter device further includes a wireless receiver to receive data and instructions from the data receiving device.

9. The telemetered characteristic monitor system of claim 1, wherein the data receiving device is an insulin pump.

10. The telemetered characteristic monitor system of claim 1, wherein the transmitter transmits the processed signals by radio frequencies.

11. The telemetered characteristic monitor system of claim 1, wherein the characteristic of the user is a glucose level in the body of the user.

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