

SENSOR MODULE AND METHOD OF USING A SENSOR MODULE

[0001] This application is being filed on 03 December 2013, as a PCT International patent application and claims priority to U.S. Provisional Application No. 61/732,783, filed December 3, 2012, which application is incorporated herein by reference in its entirety.

TECHNICAL FIELD

[0002] The present disclosure relates generally to sensors. More particularly, the present disclosure relates to sensors for measuring bio-analyte concentrations in blood samples.

BACKGROUND

[0003] Electrochemical bio-sensors have been developed for sensing (e.g., detecting or measuring) bio-analyte concentrations in fluid samples. For example, United States Patent Nos. 5,264,105; 5,356,786; 5,262,035; 5,320,725; and 6,464,849, which are hereby incorporated by reference in their entireties, disclose wired enzyme sensors for sensing analytes, such as lactate or glucose. Wired enzyme sensors have been widely used in blood glucose monitoring systems adapted for home use by diabetics to allow blood glucose levels to be closely monitored. Other example types of blood glucose monitoring systems are disclosed by United States Patent Nos. 5,575,403; 6,379,317; and 6,893,545.

SUMMARY

[0004] In general terms, this disclosure is directed to a sensor and a method of using the same.

[0005] One aspect of the present disclosure relates to a sensor module that includes a carrier and a skin piecing member carried by the carrier. The skin piecing member has a skin piecing end positioned opposite from a base end. The skin piecing member defines a lumen that extends along the central longitudinal axis from the skin piecing end toward the base end where the lumen has a lumen axis. The sensor module includes a blood sample analysis zone located entirely within the lumen of the skin piecing member and a capillary flow stop for stopping capillary flow at a predetermined location within the lumen of the skin piecing member. The sensor

module further includes an elongated working electrode positioned within the lumen. The working electrode has a length that extends along the lumen axis where at least a section of the working electrode is positioned within the analysis zone and the working electrode includes sensing chemistry.

[0006] Another aspect of the present disclosure relates to a sensor module including a carrier and a skin piecing member carried by the carrier. The skin piecing member has a skin piecing end positioned opposite from a base end. The skin piecing member defines a lumen that extends along the central longitudinal axis from the skin piecing end toward the base end and the lumen defines a lumen axis. The sensor module includes a blood sample analysis zone located within the lumen of the skin piecing member and an elongated working electrode positioned within the lumen. The working electrode having a length that extends along the lumen axis, at least a section of the working electrode is positioned within the analysis zone and the working electrode includes sensing chemistry. The working electrode has an end within 0.5 millimeters of a tip of the skin piecing member.

[0007] A further aspect of the present disclosure relates to a sensor module including a carrier movable relative to a base between a first position and second position and a skin piecing member carried by the carrier. The skin piecing member has a skin piecing end positioned opposite from a base end and the skin piecing member defines a lumen that extends along the central longitudinal axis from the skin piecing end toward the base end. The lumen defines a lumen axis. The sensor module includes a blood sample analysis zone located entirely within the lumen of the skin piecing member and an elongated working electrode positioned within the lumen. The working electrode has a length that extends along the lumen axis where at least a section of the working electrode is positioned within the analysis zone and the working electrode has sensing chemistry.

[0008] A further aspect of the present disclosure relates to a sensor module that includes a carrier and a skin piecing member carried by the carrier. The skin piecing member has a skin piecing end positioned opposite from a base end and the skin piecing member defines a lumen that extends along the central longitudinal axis from the skin piecing end toward the base end. The lumen defines a lumen axis. The sensor module includes a blood sample analysis zone located within the lumen of the skin piecing member and an elongated working electrode positioned within the lumen. The working electrode has a length that extends along the lumen axis where at

least a section of the working electrode is positioned within the analysis zone. The working electrode includes sensing chemistry and is formed by a single fiber or wire.

[0009] A further aspect of the present disclosure relates to a method for taking a blood analyte reading that includes puncturing skin with a skin piercing member having a lumen and positioning a tip of the skin piercing member in a capillary blood field less than 3 millimeters beneath the skin. The method includes initiating blood flow into the lumen by a combination of vascular blood pressure and capillary action to passively bring a blood sample to an analysis zone entirely within the lumen and sensing the blood analyte in the analysis zone.

[0010] Still another aspect of the present disclosure relates to a device for sensing an analyte in a blood sample. The device includes an elongated working electrode having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing. The device is configured for a one time use in which one analyte reading is taken.

[0011] Another aspect of the present disclosure relates to a device for sensing an analyte in a blood sample. The device includes a working electrode formed by a single conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing. The working electrode includes a layer of sensing chemistry on the first and second portions of the conductive fiber or wire.

[0012] A further aspect of the present disclosure relates to a device for sensing an analyte in a blood sample. The device includes a working electrode formed by a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing. The working electrode includes a layer of sensing chemistry on the first and second portions of the conductive fiber or wire. The device further includes a skin piercing member having a lumen in which the working electrode is positioned. The conductive fiber or wire has an outer diameter that is at least 10 percent as large as an outer diameter of the skin piercing member.

[0013] A further aspect of the present disclosure relates to a device for sensing an analyte in a blood sample. The device includes a working electrode formed by a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing. The working electrode includes a layer of sensing chemistry on the first and second portions of the

conductive fiber or wire. The device includes a skin piercing member having a lumen in which the working electrode is positioned. The conductive fiber or wire has an outer diameter that is greater than .001 inches.

[0014] Another aspect of the present disclosure relates to a device for sensing an analyte in a blood sample. The device includes an elongated working electrode including a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing. The conductive fiber or wire has a cross-dimension greater than .001 inches.

[0015] Still another aspect of the present disclosure relates to a device for sensing an analyte in a blood sample. The device includes a working electrode formed by a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing. The working electrode includes a layer of sensing chemistry on the first and second portions of the conductive fiber or wire. The device further includes a skin piercing member having a lumen in which the working electrode is positioned. The device is configured such that a blood analysis zone of the device fills passively.

[0016] A variety of additional aspects will be set forth within the description that follows. The aspects can relate to individual features and to combinations of features. It is to be understood that both the foregoing general description and the following detailed description are exemplary and explanatory only and are not restrictive of the broad concepts upon which the embodiments disclosed herein are based.

BRIEF DESCRIPTION OF THE DRAWINGS

[0017] FIG. 1 is perspective view of a sensor module in accordance with the principles of the present disclosure;

[0018] FIG. 2 is a top view of the sensor module of FIG. 1 with a skin piercing member of the sensor module in an extended position inserted in a vascular plexus;

[0019] FIG 3. is a plan view of the sensor module of FIG. 2 with various cross-section lines depicted;

[0020] FIG. 4 is a cross-sectional view taken along line 4-4 of FIG. 2;

[0021] FIG. 5 is a perspective view of another embodiment of a sensor module with a three electrode profile in accordance with the principles of the present disclosure;

- [0022] FIG. 6 is an alternate configuration of FIG. 5 with a different extrusion profile;
- [0023] FIG. 7 is an end view of another embodiment of a sensor module with a two electrode profile;
- [0024] FIG. 8 is a perspective view of the two electrode profile of FIG. 7 with a mask;
- [0025] FIG. 9 is a perspective view of the two electrode profile of FIG. 7 with an electrode guide;
- [0026] FIG. 10 is an end view of a portion of FIG. 7;
- [0027] FIG. 11 is an end view of the two electrode profile of FIG. 7 with contact points depicted;
- [0028] FIG. 12 is a schematic view of a sensing unit in accordance with the principles of the present disclosure that incorporates a plurality of the sensing modules of FIG. 1; and
- [0029] Figure 13 is a schematic view of a skin piercing member, reference electrode and working electrode configuration of the sensor module of Figures 1-3.

DETAILED DESCRIPTION

- [0030] Various embodiments will be described in detail with reference to the drawings, wherein like reference numerals represent like parts and assemblies throughout the several views. Reference to various embodiments does not limit the scope of the claims attached hereto. Additionally, any examples set forth in this specification are not intended to be limiting and merely set forth some of the many possible embodiments for the appended claims.
- [0031] The following definitions are provided for terms used herein:
- [0032] A "working electrode" is an electrode at which the analyte (or a second compound whose level depends on the level of the analyte) is electrooxidized or electroreduced with or without the agency of an electron transfer agent.
- [0033] A "reference electrode" is an electrode used in measuring the potential of the working electrode. The reference electrode should have a generally constant electrochemical potential as long as no current flows through it. As used herein, the term "reference electrode" includes pseudo-reference electrodes. In the context of the disclosure, the term "reference electrode" can include reference electrodes which also function as counter electrodes (i.e., a counter/reference electrode).

[0034] A "counter electrode" refers to an electrode paired with a working electrode to form an electrochemical cell. In use, electrical current passes through the working and counter electrodes. The electrical current passing through the counter electrode is equal in magnitude and opposite in sign to the current passing through the working electrode. In the context of the disclosure, the term "counter electrode" can include counter electrodes which also function as reference electrodes (i.e., a counter/reference electrode).

[0035] A "counter/reference electrode" is an electrode that functions as both a counter electrode and a reference electrode.

[0036] An "electrochemical sensing system" is a system configured to detect the presence and/or measure the level of an analyte in a sample via electrochemical oxidation and reduction reactions on the sensor. These reactions are converted (e.g., transduced) to an electrical signal that can be correlated to an amount, concentration, or level of an analyte in the sample. Further details about electrochemical sensing systems, working electrodes, counter electrodes and reference electrodes can be found at U.S. Patent No. 6,560,471, the disclosure of which is hereby incorporated herein by reference in its entirety.

[0037] "Electrolysis" is the electrooxidation or electroreduction of a compound either directly at an electrode or via one or more electron transfer agents.

[0038] An "electron transfer agent" is a compound that carries electrons between the analyte and the working electrode either directly or in cooperation with other electron transfer agents. One example of an electron transfer agent is a redox mediator.

[0039] A "sensing layer" is a component of the sensor which includes constituents that facilitate the electrolysis of the analyte. The sensing layer may include constituents such as an electron transfer agent, a catalyst which catalyzes a reaction of the analyte to produce a response at the electrode, or both.

[0040] FIGS. 1-3 are perspective views of an example of a sensor module 100. In this example, the sensor module 100 includes a carrier 102, a skin piercing member 110, a base 104 and two electrical contacts 112, 114. In FIG. 1, the skin piercing member 110 is shown in a retracted position.

[0041] The carrier member 102 is arranged and configured to slideably move along the base 104 within opposite channels 108 defined by the base 104. The channels 108 of the base 104 limit upward or downward movement of the carrier 102

relative to the base 104. The carrier 102 defines a first cavity 116 to mount the skin piercing member 110. In one example, the piercing member 40 is fixed relative to the carrier 102 such that the piercing member 40 is carried by the carrier 102 as the carrier 102 slides relative to the base 104. The carrier 102 is illustrated and described in more detail with reference to FIGS. 2-3.

[0042] The skin piercing member 110 extends along a length L of the carrier 102. The skin piercing member 110 is movable with the carrier 102 between a retracted position and an extended position (see FIG. 2) relative to the base. The skin piercing member 110 is illustrated and described in more detail with reference to FIGS. 2-4.

[0043] The electrical contacts 112, 114 mount on the carrier 102. The contacts 112, 114 respectively have contact tabs 120, 122. Tab 120 can be used to electrically connect the contact 112 to a reference electrode provided on an exterior surface of the member 110. Tab 122 can be used to electrically connect the contact 114 to a working electrode having a portion that extends into the piercing member 110 and a portion that extends axially outwardly from a base end of the piercing member 110. The contacts 112, 114 can include structures for electrically connecting the sensor module 100 to a sensor control system. In one example, the sensor control system applies a voltage across the working and reference electrodes and through a blood sample contained within a lumen of the skin piercing member 110. The reference electrode can be formed as a conductive layer provided on the outer surface of the skin piercing member 110. The electrical contacts 112, 114 are illustrated and described in more detail with reference to FIGS. 2-3.

[0044] FIGS. 2-3 illustrate features of the carrier 102.

[0045] In one example, the skin piercing member 110 is hollow and defines an interior lumen in which a working electrode is positioned. The working electrode can be formed by a conductive fiber or wire. A sensing layer can cover the conductive fiber or wire. A portion of the conductive fiber or wire covered with the sensing layer can form a sensing region located within a sample analysis zone contained entirely within the skin piercing member. The sample analysis zone 130 can provide for specific control of interrelated parameters such as active electrode area, response time, sensitivity, and drift to be engineered in as byproducts of static component features.

[0046] Figure 13 is a schematic view showing the skin piercing member 110 having the base end 140 and a tip end 138 (e.g., an insertion end). The skin piercing

member 110 defines a lumen 144 that extends through the entire length of the skin piercing member 110 along an axis 10 (e.g., a skin piercing member axis or a lumen axis). A capillary stop can be provide adjacent the base end 140. The elongated working electrode 142 can include a conductive fiber or wire coated or otherwise covered with a sensing layer. The working electrode 142 is positioned within the lumen 144 and can have a lower end within .5 millimeters of the tip 138 and an upper end portion that extends out of the lumen 144. The conductive fiber or wire of the working electrode can be covered with a sensing layer suitable for sensing an analyte such as glucose. The skin piercing member 110 can include a conductive core material encapsulated within a dielectric layer that prevents electrical connections between the working electrode and the skin piercing member. A reference electrode can be supported on the dielectric outer surface of the skin piercing member. A control unit can be electrically connected to the reference and working electrodes.

[0047] In use, the skin piercing member is inserted into the skin to a depth less than 3 millimeters such that the tip 138 resides in the capillary bed. As so positioned, a first portion of the working electrode is subcutaneous and a second portion extends outside the body beyond the skin. Upon insertion, the combination of vascular pressure and capillary action causes blood to rapidly fill the lumen 144 and surround the portion of the working electrode within the skin piercing member 110. The blood flows up the lumen to the capillary stop. The volume of space defined within the skin piercing member from the tip to the capillary stop forms an analysis zone having a length that corresponds to a length of a wetted surface area of the sensing chemistry supported on the conductive fiber or wire of the working electrode. The capillary stop combined ensures that the wetted surface area is precisely controlled (i.e., the wetted length of sensor chemistry corresponds to the length of the working electrode that extends below the capillary stop). Applying a voltage between reference and working electrodes causes the oxidation/reduction of glucose in the analysis zone thereby generating a current at the working electrode which can be measured to sense a concentration of glucose in the blood sample. Control circuitry can apply the voltage, measure the current, and provide a display showing a reading indicating the glucose level.

[0048] The sensor module 100 is relatively compact and disposable. For example, the sensor module 100 is generally rectangular in shape and has a length that

is less than 1 inch. The sensor module 100 includes opposite major sides and opposite minor sides that extend along the length of the sensor module 100.

[0049] The skin piercing member 110 of the sensor module 100 includes a skin piercing end 136 having a sharp tip 138 and a base end 140. The tip 138 of the skin piercing member 100 penetrates the skin of a patient and can be configured to provide a cutting action that generates a wound that self-closes upon removal of the piercing member 110 from the skin. The skin piercing member 110 can be a cannula, needle, or other similar structure preferably having a hollow interior. In this example, the sensor is configured to allow the analysis of the fluid sample to take place entirely within the skin piercing member 110. The skin piercing member 110 provides a volume or reservoir 107 for collecting blood received from a skin puncture site caused by the skin piercing member 110.

[0050] In one example, the skin piercing member 110 has a non-conductive construction. An example of a non-conductive construction includes a conductive base metal body (for strength) encased within a dielectric layer. The skin piercing member 110 includes a metal body 109 (FIG.4) that can include a conductive metal, such as, but not limited to, stainless steel, and can be covered by a insulation layer 111 (FIG. 4), such as, but not limited to, parylene, acting to keep the conductive metal from creating a direct electrical pathway from a reference electrode to a working electrode of a two electrode sensor (e.g., a glucose sensor). The skin piercing member 110 can be about 30 gauge or less to allow for an insertion into a patient's skin tissue without creating either a blood producing wound or noticeable pain or discomfort upon insertion. The skin piercing member 110 can have a length of about 12 to 13 mm. In one example, only a relatively short length of the piercing member 100 extends beyond the base 102 when the carrier is slid to an extended position. In one example, the module 100 is configured such that the insertion depth of the skin piercing member 110 will not exceed 2 millimeters. In another example, the skin insertion depth of the skin piercing member 110 is in the range of about 1.5 to 2 mm. This depth of piercing allows for the sensor in the sensor module 100 to communicate with the vascular plexus (VP) dermal layer of tissue. At this depth, the sensor encounters capillary blood that is representative of cellular glucose.

[0051] In use of the sensor module 100, a contact end 124 of the base 104 is placed against a patient's skin at a sampling site where it is desired to take a fluid (e.g., blood) sample. Once the contact end 124 is in contact with the skin, the skin

piercing member 110 is moved from the retracted position to the extended position (e.g., by sliding the carrier 102 relative to the base 104) thereby causing the tip 136 of the skin piercing member 110 to pierce the patient's skin. Upon insertion of the skin piercing member 110, blood from the capillary field fills the skin piercing member 110. Blood flow is caused at least in part by vascular pressure within the capillary bed. Capillary action also moves blood upwardly within the piercing member 110 to fill a sample analysis zone 130 within the piercing member 110. At the sample analysis zone 130, an analyte level (e.g., blood glucose level) in the blood sample is sensed through the use of a wired enzyme sensor arrangement including an elongated working electrode (WE) 142 (FIG. 4) positioned inside the piercing member 110. In certain embodiments, the electrode can be a fiber, wire, or other elongated member. In other embodiments, separate working, reference and counter electrodes can be provided in fluid communication with the sample analysis zone 130.

[0052] In some embodiments, a test is initiated by pressing an actuator button (not shown) on top of a meter (not shown) while holding the sensor module 100 on the test site (i.e., forearm or fingertip). This action causes a sequence of motions moving the sensor module 100 from a position within the sensor module 100 to an opening in the bottom of the meter. The meter can be placed on the approved testing site, (i.e., forearm or finger). The actuator button can be pressed again following a prompt causing the carrier 102 of the sensor module 100 carrying the skin piercing member 110 to move rapidly forward inserting the skin piercing member 110 to a prescribed depth. The skin piercing member 110 of the sensor module 100 enters a depth in tissue where a capillary blood field is encountered. The skin piercing member 110 stops at a capillary depth of about less than 3 mm below the skin surface and can reside for about less than 3 seconds to acquire a blood sample. The sample can be presented to the sensor module 100 by a rapid microfluidic flow initiated automatically by a combination of vascular blood pressure and capillary action. The sensor module 100 requires no other active mechanism to obtain a blood glucose value resulting in a passive system. Once the test is performed or completed, the carrier can be disposed by the user.

[0053] In one embodiment, the working electrode can include an elongated member that is coated or otherwise covered with a sensing layer and the reference/counter electrode can include any elongated member, such as a wire or fiber that is coated or otherwise covered with a layer, such as silver chloride. Preferably, at

least a portion of each elongated member is electrically conductive. In certain embodiments, each elongated member can include a metal wire or a glassy carbon fiber. In still other embodiments, each elongated member can each have a composite structure and can include a fiber having a dielectric core surrounded by a conductive layer suitable for forming an electrode. The core can be made of medical grade polyetheretherketone.

[0054] An example composite fiber is sold under the name Resistat® by Shakespeare Conductive Fibers LLC. This composite fiber includes a composite nylon, monofilament, conductive thread material made conductive by the suffusion of about a 1 micron layer of carbonized nylon isomer onto a dielectric nylon core material. The Resistat® material is comprised of isomers of nylon to create the basic 2 layer composite thread. However, many other polymers are available for the construction, such as: polyethylene terephthalate, nylon 6, nylon 6,6, cellulose, polypropylene cellulose acetate, polyacrylonitrile and copolymers of polyacrylonitrile for a first component and polymers such as of polyethylene terephthalate, nylon 6, nylon 6,6, cellulose, polypropylene cellulose acetate, polyacrylonitrile and copolymers of polyacrylonitrile as constituents of a second component. Inherently conductive polymers (ICP) such as doped polyaniline or polypyrrole can be incorporated into the conductive layer along with the carbon to complete the formulation. In certain embodiments, the ICP can be used as the electrode surface alone or in conjunction with carbon. The Resistat® fiber is availability in diameters of .0025 to .016 inches, which is suitable for sensor electrodes configured in accordance with the principles of the present disclosure. Example patents disclosing composite fibers suitable for use in practicing sensor modules configured in accordance with the principles of the present disclosure include U.S. Patent Nos. 3,823,035; 4,255,487; 4,545,835 and 4,704,311, which are hereby incorporated herein by reference in their entireties.

[0055] The sensing layers provided at working electrodes of sensor modules configured in accordance with the principles of the present disclosure can include a sensing chemistry, such as a redox compound or mediator. The term redox compound is used herein to mean a compound that can be oxidized or reduced. Example redox compounds include transition metal complexes with organic ligands. Preferred redox compounds/mediators include osmium transition metal complexes with one or more ligands having a nitrogen containing heterocycle such as 2, 2'-bipyridine. The sensing

material also can include a redox enzyme. A redox enzyme is an enzyme that catalyzes an oxidation or reduction of an analyte. For example, a glucose oxidase or glucose dehydrogenase can be used when the analyte is glucose. Also, a lactate oxidase or lactate dehydrogenase fills this role when the analyte is lactate. In sensor systems, such as the one being described, these enzymes catalyze the electrolysis of an analyte by transferring electrons between the analyte and the electrode via the redox compound. Further information regarding sensing chemistry can be found at U.S. Patent Nos. 5,264,105; 5,356,786; 5,262,035; and 5,320,725, which were previously incorporated by reference in their entireties.

[0056] In one embodiment, the skin piercing member 110 defines a lumen 144 that extends along an elongated axis 10 from the skin piercing end 136 of the skin piercing member 110 toward the base end 140. The elongated working electrode 142 is positioned within the lumen 144. The elongated working electrode 142 has a length that extends along the lumen axis 20 and at least a section of the elongated working electrode 142 is positioned within the sample analysis zone 130. The elongated working electrode 142 includes the sensing chemistry.

[0057] The interaction of the skin piercing member 110 in concert with microfluidic forces (e.g., surface tension) within the lumen 144 promotes capillary flow of blood. Flow is initiated by ambient capillary pressure at the proximal lumen of the skin piercing member 110 when the piercing member is inserted into the papillary dermis to a depth of between 1-2 mm below the skin. Flow may also be promoted by the treatment of the lumen 144 with a surfactant compound. When so prepared, the combined factors create a driving mechanism to enable a spontaneous flow of capillary blood to enter the proximal lumen 144 and fill the skin piercing member 110 throughout its length.

[0058] The capillary stop 154 is formed at the skin piercing member 110 to inhibit the spontaneous blood flow from exiting the skin piercing member 110 at the distal end of the lumen 144. The self-limiting action of the flow into the interior passage of the skin piercing member 110 facilitates the lumen 144 to function as both an analysis cell—defined by the volume of the skin piercing member 110 and the length of the wetted working electrode WE portion residing within the skin piercing member 110—and as a counter electrode component of a multi electrode electrochemical cell.

[0059] The lumen 144 of the piercing member 110 may be sized appropriately to the configuration of the electrode bundle within it so as to optimize the microfluidic

forces affecting the rate of transport thru the passage to the capillary stop 154. The lumen length must extend far enough above the tissue so as to provide sufficient surface area of the working electrode WE to produce a specified minimal output current. However, the lumen length may not be excessive or the time required to fill the lumen will drop rapidly with falling capillary pressure and fluid resistance slowing the transport rate.

[0060] The above described configuration of the electrode array within the piercing member 110 allows the major portion of the electrode surface to remain above the skin line presenting only the diameter of the piercing member 110 to the enervated tissue of the papillary dermis. This configuration allows the effective current produced by the electrode within the piercing member 110 to be two orders of magnitude larger than a traditional implanted sensor occupying the same footprint within tissue. In certain examples, the electrodes have an operational radius of less than 0.15 mm and a length of between 10 mm and 20 mm.

[0061] Referring to FIG. 4, a cross-sectional view of the skin piercing member 110 of the sensor module 100 is shown.

[0062] In this example, the elongated working electrode 142 is positioned within the lumen 144 of the skin piercing member 110 and includes a monofilament 146 (i.e., fiber) that can be coated with a conductive layer 148 (i.e., gold). The monofilament can be a polymeric material such as medical grade polyetheretherketone. The conductive monofilament 146 (fiber or wire) can have an outer diameter that is at least 10 percent as large as an outer diameter of the skin piercing member 110. The conductive layer 148 may be applied as a continuous Plasma Vacuum Deposition (PVD) process. The conductive layer 148 can be less than about 10 microns. The diameters of the multi-fiber composite can be between 0.001 inches and 0.004 inches. In this example, the elongated working electrode 142 may be coated in a secondary process step over the conductive layer 148 with glucose oxidase or other suitable enzyme chemistry. The elongated working electrode 142 may be inserted into the lumen 144 of the skin piercing member 110.

[0063] In this example, the skin piercing member 110 can include a conductive reference electrode (RE) layer 150 formed as a thick film of silver/silver chloride (Ag/Ag/Cl) on the insulated outer surface of the skin piercing member 110. The monofilament 146 (i.e., fiber) can be an aluminum conductive composite monofilament (CCM) with an enzyme sensing layer 152 which can serve as the

elongated working electrode 142. The sensor module 100 can become active when an ionic fluid such as blood simultaneously contacts both the outer surface of the conductive reference electrode layer 150 (RE) and the elongated working electrode 142 (WE) completing an electrical circuit through the fluid path. Once the circuit is established by a passive process of rapid capillary flow into the lumen 144 of the skin piercing member 110, blood continues up a defined open passage space (less than .004 inches circumferential clearance) surrounding the fiber until encountering a capillary stop 154 feature formed at the base end 140 of the piercing member 110. The lumen 144 can be kept partially open at the base end 140 to serve as an air vent to promote the capillary flow.

[0064] In this example, the insertion end of the lumen 144 should be free of tissue plugs and reside at or below the vascular plexus (VP) between about 1 to 2 mm deep in the dermal layer where capillary vascular pressure is sufficient (about 14 to 22 mm Hg) to promote initial blood flow into the flow passage 128 of the skin piercing member 110. Capillary flow can augment external vascular pressure to rapidly sweep up the interior of the flow passage 128 to the capillary stop 154. Rapid autonomous and complete filling of the sample analysis zone 130 – can be defined by the interior volume 107 of the skin piercing member 110 less the volume of the electrode within a space. This filling can be co-determinant of response time and is promoted by the addition of surfactants such as, but not limited to, Triton materials to either the skin piercing member 100 interior surface or to the detector chemistry or both.

[0065] In this example, the sample analysis zone 130 includes a controlled surface area of sensing chemistry that is wetted with blood during testing such that the entire controlled surface area is wetted when the analyte reading is taken. The controlled surface area is at least 10 times as large as a transverse cross-sectional area of the skin piercing member. In other embodiments, the controlled surface area can be at least 20 or 30 times larger than a transverse cross-sectional area of the working electrode.

[0066] Automation suitability can create a sensor configuration that will improve both quality of testing and the reliability of the test procedure for the consumer. The analysis zone method described can rely upon interdependent effects of defined part geometry, spatial relationships of components and specific transitional properties of the enzyme detector chemistry as it is hydrated by the incoming blood matrix. These factors in concert with the dynamic interaction of blood flowing into the cell in response to vascular pressure and capillary action function as the analysis zone

method for establishing a rapid and self-limiting amperometric assay cell formed along a defined section of a long fiber.

[0067] Flow up the lumen 144 of the skin piercing member 110 can be within the microfluidic domain of non-Newtonian laminar flow. This transport dynamic up the circumferential channel 128 defined within the lumen 144 between the working electrode 142 and the inner surface of the skin piercing member 110 can be optimized by promoting low surface energy properties for the WE to allow complete and rapid wetting of the enzyme sensing layer 152. This surface property in turn can act in concert with the laminar flow dynamics to sweep the entire cavity containing the WE, free of air pockets that could otherwise unpredictably affect the area of blood in contact with the electrode surface causing irreproducible sensor performance.

[0068] The capillary pressure, the viscosity of the blood media plus the surface energy interactions of the electrode coating and the skin piercing member 110 inner wall surface in concert with the distance separating the surfaces can all impact micro capillary flow characteristics.

[0069] The capillary stop 154 can be a mechanism that limits further fluid flow along the enzyme sensing layer 152 forming the WE and provides for venting of air displaced by the rapid filling of the capillary space by blood. In this example, one functional characteristic of the WE is that the dry enzyme detector chemistry can be an effective insulator and can transition in phases from insulator to semiconductor to conductor as it becomes hydrated. This property prevents errant signal contributions to any portion of the CCM fiber kept dry during the time of the glucose assay. By defining the hydrated area of the WE through the combined use of the capillary stop 154 feature with mechanical control of the length of CCM fiber extending down into the skin piercing member passage 134. This method of defining electrode surface area provides for both manufacturing and functional advantages.

[0070] Referring again to FIGS. 2-3, the electrical contacts 112, 114 can be made of an electrically conductive material, such as, but not limited to, metals (i.e. copper, silver, aluminum, gold, bronze, and magnesium). During sample analysis at the sample analysis zone, a voltage can be applied between the working and reference electrodes. When the potential is applied, an electrical current will flow through the fluid sample to the working electrode. The current is a result of the oxidation or reduction of an analyte, such as glucose, in the volume of fluid sample located within the sample analysis zone. This electrochemical reaction occurs via the electron

transfer agent in the enzyme sensing layer 152 and an optional electron transfer catalyst/enzyme in the enzyme sensing layer 152. By measuring the current flow generated at a given potential (e.g., with a controller described herein), the concentration of a given analyte (e.g., glucose) in the fluid sample can be determined. Those skilled in the art will recognize that current measurements can be obtained by a variety of techniques including, among other things, coulometric, potentiometric, perometric, voltometric, and other electrochemical techniques.

[0071] In this example, within a few hundredths of a second the defined sample analysis zone 130 is filled and the hydrating WE initiates an exchange electrons with the Ag/AgCl RE pattern on the non-conductive surface of the skin piercing member 110. A rising current appears at the data acquisition input of the sensor module 100 causing the software to start a countdown before initiating a data acquisition sequence for a prescribed number of discrete points (currently 500) taken at intervals over a set time window. The data set can be grouped by taking a mean of the discrete points. An area under the curve analysis can be applied to predict the plateau current for the sensor module 100. The correlation equates to a calibrated number representing a known glucose concentration at that current. The software then stores the value and can display it to the user on the meter LCD. The entire sequence from initiating actuator button to displayed blood glucose value requires less than 5 seconds. The result of the above testing sequence can be considered to be one reading. In certain examples, the modules 100 are single use and each can be used to provide one glucose reading.

[0072] In certain embodiments, the data can be acquired using wireless device or portable electronic device (PED) such as, but not limited to, cellular phones. The PED can be used to act as a control unit for the sensor module 100. The sensor module 100 can be configured to interface with the PED which can store and display the glucose concentration to the user. In other embodiments, a separate test unit may be utilized to interface with a wireless device or PED (i.e., cellular phone). A chipset or similar component can be used in a glucose module to link to a PED via a broadband connection. The glucose test module can be connected automatically to the PED to initiate an application that would perform and display all the data management tasks. The glucose test module can be configured to have wide area network (WAN) capability to link to therapeutic software resident on other servers, such as, but not limited to, Cloud, that would completely automate the diabetics

provisioning and treatment as well as link to a patient's physician or caregiver in real time. The glucose test module can be about 2.5 inch wide, about 3 inches long and about 1/4 inch high.

[0073] Referring to FIG. 5, a perspective view of another embodiment including three CCM electrodes 200.

[0074] In this example, the three CCM electrodes 200 are arranged and configured such that both RE and WE electrodes 202, 204 may be simultaneously inserted into a lumen of a skin piercing member. The skin piercing member can have a size between approximately 28 gauge to 30 gauge, preferably a 30 gauge size in the example of home blood glucose testing.

[0075] FIGS. 5-6 show the three CCM electrodes 200 or multi-electrode configurations that are assembled with two different profile extrusions, 208, 210 capable of insertion into a conductive needle, cannula or tube. In this example, the multi-electrode configuration does not need an insulated coating or a non-conductive piercing member as an RE electrode substrate; but may be only a piercing member, such as, but not limited to, a cannula, acting as the mechanical needle component. The piercing member can be made of a standard stainless steel material or similar material. The multi-electrode configuration may also be used as a catheter insert for continuous or remote monitoring applications.

[0076] The configuration of the multi-electrode may employ slightly different micro extrusion profiles as carriers for the three CCM electrodes 200, WE and RE fiber electrodes 202, 204. The micro extruded carrier profile is so designed in cross section as to maintain electrical isolation of the RE and WE 202,204 - both from each other, and from the stainless steel piercing member (i.e., cannula). In some embodiments, the multi-electrode may have two working electrodes and one RE. The specific properties and profile extrusions 208, 210 (FIG. 6) combined with dedicated features incorporated into the extruded profile helps provide for unrestricted capillary flow up a flow passage of the skin piercing member, electrical isolation of the conductive components without the co-extrusion of the three CCM electrodes 200 that may be incompatible with such extrusion process, and combining four extrusion components into a single reel structure capable of insertion into the lumen of the skin piercing member. In this example, the profile may be configured to allow three or more electrodes to reside within a skin piercing member and operate as a three electrode enzymatic detection system.

[0077] The multi-electrode configuration allows for the ability to eliminate an Ag/Ag/Cl film coating for the RE 202 and substitute a composite Ag substrate fiber that can be fabricated in the same substrate materials and PVD process as the WE. The multi-electrode configuration can include a third electrode 206 (e.g., a supplemental electrode) that has no need for enzyme or detector chemistry. The third electrode 206 may be employed as a comparison means to determine what portion of the raw sensor current comes from interferent compounds such as vitamin C rather than from glucose. The comparison means can be a result of oxidizing those compounds at the bare Au surface and subtracting the apparent signal from the total sensor output. This function eliminates a developmental step in the electrode design and can help to resolve a common background noise problem for sensor accuracy. In other embodiments, additional electrodes beyond the three electrode profile may be added to the structure for the purpose of multi-parameter assays. Multi-parameter sensors providing more than two analyte targets for use in continuous monitoring can be fabricated. The supplemental electrode can also be used to detect other analytes (e.g., lactate) or to detect oxygen concentration.

[0078] Turning to FIG. 6, a top view of the three CCM electrodes 200 is shown. The fabrication of the profile extrusion 210 can be made by means of micro extrusion technology combined with continuous PVD technology as can be used in the gold coating process for CCM electrodes. The profile extrusions 208, 210 can be used to insulate the three CCM electrodes 200 from one another as well as from the interior wall of a skin piercing member. In this example, the extrusion profiles 208, 210 can be about 0.001 inches thick. The profiles can be configured to maximize fluid transport fully along the interior passage of the skin piercing member.

[0079] Referring to FIG. 7, a perspective view of a two electrode 400 configuration for a sensor module is shown.

[0080] In this example, the two electrodes 400 configuration includes a monofilament 402 (i.e., fiber). The monofilament 402 can be insulated using an insulating substrate layer 404 similar to the embodiments described above. The monofilament 402 is capable of insertion into a lumen 406 of a skin piercing member 408. As shown, an AgAg/Cl layer 410 can be applied directly onto the insulating substrate layer 404.

[0081] Referring to FIG. 8, a perspective top view of the two electrodes 400 configuration is shown.

[0082] The AgAg/Cl layer 410 can be applied using fixtures and/or masks 412 to direct a plasma deposition of silver (Ag) travelling perpendicular to the plane P of the mask 412 onto a single surface of the profile for forming the RE. FIG. 9 shows an electrode guide 414 that can be used to help control the position of the profile in transit through the PVD process. The profile having the AgAg/Cl layer 410 deposition surface can act as the RE and the CCM monofilament 402 (i.e., fiber) coated with enzyme chemistry provides the WE function. FIG. 10 depicts arrows 416 that show features that can guide the profile and prevent both the RE and WE from contacting the conductive surface of the skin piercing member 408. FIG. 11 show contact points 418 that can be opposing where output contact features can impinge a sensor module and acquire a signal.

[0083] Referring to FIG. 12, a schematic of an analyte monitoring unit 300 is shown. The unit 300 where the modules 100 may be arrayed within a cartridge designed to provide a supply of multiple sensors that may be directly positioned on the skin of a patient's forearm or fingertip in order to obtain a blood glucose concentration. It will be appreciated that one or more sensor modules 100 can be incorporated as sub-components into an analyte monitoring unit 300. The unit 300 includes a controller 302 that couples to a module holder 304. The module holder 304 is configured to hold one or more sensor modules 100. Each sensor module 100 is configured to obtain one or more fluid samples, to measure a concentration level for one or more analytes (e.g., glucose, lactate, etc.), and to generate a signal (e.g., an electrical signal) indicating the concentration level. For example, the module holder 304 shown in FIG. 7 contains five sensor modules 100. In one embodiment, each sensor module 100 is configured to analyze a single fluid sample. In such an embodiment, the sensor module 100 can be removed from the module holder 304 after one use. In other embodiments, each sensor module 100 can be configured to analyze a greater number of fluid samples.

[0084] In general, the unit 300 includes a controller 302, an actuator 306, and input lines 308. The controller 302 controls the actuator 306 for driving the skin piercing members 110 of each sensor module 100 between the extended and retracted positions to obtain a fluid sample. The controller 302 can include a microcontroller, a mechanical controller, software driven controller, a hardware driven controller, a firmware driven controller, etc. The controller can include a microprocessor that interfaces with memory.

[0085] The controller 302 instructs the actuator 306 when to operate the sensor module 100 to obtain a fluid sample for analysis. The controller 302 also can instruct the module holder 304 and/or the actuator 306 to eject the used sensor module 100.

[0086] The input lines 308 carry the data/signals/readings (e.g., voltage values) generated at the elongated working electrode 142 of the sensor module 100 during analysis of a fluid sample to the controller 302 for analysis. The controller 302 converts the signals to an analyte concentration level (e.g., a blood glucose reading) or other desired information. The controller 302 causes the display 310 to indicate the processed information to the user. Other information also can be presented on the display 310. In one embodiment, the display 310 is a visual display. In other embodiments, an audio display also can be used. Additional information can be provided to the controller 302 via a user interface 312 (e.g., buttons, switches, etc.).

[0087] From the forgoing detailed description, it will be evident that modifications and variations can be made without departing from the spirit and scope of the disclosure.

WHAT IS CLAIMED IS:

1. A sensor module comprising:
 - a carrier;
 - a skin piercing member carried by the carrier, the skin piercing member having a skin piercing end positioned opposite from a base end, the skin piercing member defining a lumen that extends along the central longitudinal axis from the skin piercing end toward the base end, the lumen having a lumen axis;
 - a blood sample analysis zone located entirely within the lumen of the skin piercing member;
 - a capillary flow stop for stopping capillary flow at a predetermined location within the lumen of the skin piercing member; and
 - an elongated working electrode positioned within the lumen, the working electrode having a length that extends along the lumen axis, at least a section of the working electrode being positioned within the analysis zone, and the working electrode including sensing chemistry.
2. The sensor module of claim 1, further comprising a base to which the carrier is mounted, wherein the carrier can slide relative to the base between a first position where the skin piercing end of the skin piercing member is exposed and a second position wherein the skin piercing member is retracted.
3. The sensor module of claim 2, wherein the skin piercing member extends no more than 3 millimeters beyond the base when in the first position.
4. The sensor module of claim 3, wherein the skin piercing member extends 1-2 millimeters beyond the base when in the extended position.
5. The sensor module of claim 1, wherein the sensor module does not include an active means for drawing blood into the lumen.
6. The sensor module of claim 1, wherein the lumen extends completely through the skin piercing member from the skin piercing end to the base end.

7. The sensor module of claim 6, wherein the working electrode extends from the lumen outwardly beyond the base end of the skin piercing member.
8. The sensor module of claim 6, wherein the capillary flow stop is located at the base end of the skin piercing member.
9. The sensor module of claim 1, wherein a first end of the working electrode is within 2 millimeters of a tip of the skin piercing end of the skin piercing member.
10. The sensor module of claim 1, wherein a first end of the working electrode is within 1 millimeter of a tip of the skin piercing end of the skin piercing member.
11. The sensor module of claim 1, wherein a first end of the working electrode is within .5 millimeters of a tip of the skin piercing end of the skin piercing member.
12. The sensor module of claim 11, wherein the working electrode extends from the first end to a second end, and wherein the working electrode exits the skin piercing member at the base end of the skin piercing member such that the second end of the working electrode is positioned outside the skin piercing member.
13. The sensor module of claim 1, wherein the skin piercing member includes a skin penetrating portion adapted to be subdermal when a blood sample is taken, wherein the skin penetrating portion is less than 3 millimeters in length, wherein the working electrode includes a first end portion that extends along at least a majority of the skin penetrating portion of the skin piercing member.
14. The sensor module of claim 13, wherein the first end portion of the working electrode terminates within .5 millimeters of a tip of the skin piercing member.
15. The sensor module of claim 14, wherein a second end portion of the working electrode extends outwardly from the base end of the skin piercing member and is supported on the carrier.

16. The sensor module of claim 1, wherein the working electrode includes a fiber or a wire.
17. The sensor module of claim 1, wherein the working electrode includes a single fiber or a single wire having a diameter in the range of .001 to .004 inches.
18. The sensor module of claim 1, wherein the working electrode includes a fiber having a polymeric core surrounded by a conductive layer covered by the sensing chemistry.
19. The sensor module of claim 1, wherein the working electrode includes a single fiber having a polymeric core surrounded by a conductive layer covered by the sensing chemistry.
20. The sensor module of claim 1, further comprising a reference electrode, the sensor module also including first and second electrical contacts carried by the carrier, the first contact being electrically connected to the working electrode and the second electrical contact being electrically connected to the reference electrode.
21. The sensor module of claim 20, wherein the skin piercing member includes at least an outer layer that is dielectric, and wherein the reference electrode is supported on the outer dielectric layer of the skin piercing member.
22. The sensor module of claim 21, wherein the skin piercing member includes a metal main body encased within a dielectric material.
23. The sensor module of claim 1, wherein the sensing chemistry includes glucose oxidase or glucose dehydrogenase.
24. The sensor module of claim 1, further comprising a reference electrode positioned within the lumen, the working electrode and the reference electrode being separated by a dielectric spacer.

25. The sensor module of claim 24, wherein the dielectric spacer also separates the reference electrode and the working electrode from an interior surface of the skin piercing member.
26. The sensor module of claim 25, wherein the dielectric spacer is an extruded spacer that extends at least a majority of the length of the skin piercing member.
27. The sensor module of claim 25, wherein the reference electrode and the working electrode include fibers.
28. The sensor module of claim 27, wherein the fibers include polymeric cores surrounded by conductive layers.
29. The sensor module of claim 1, further comprising a reference electrode and an interferent detection electrode positioned within the lumen.
30. The sensor module of claim 29, wherein the working electrode, the interferent electrode and the reference electrode include fibers having polymeric cores surrounded by conductive layers, and wherein the working electrode, the interferent electrode and the reference electrode are separated by a spacer.
31. The sensor module of claim 1, further comprising a reference electrode and a supplemental electrode positioned within the lumen.
32. The sensor module of claim 31, wherein the working electrode is adapted to sense glucose and wherein the supplemental electrode is adapted to sense a chemical other than glucose.
33. The sensor module of claim 32, wherein the working electrode, the reference electrode and the supplemental electrode each include an optical fiber having a polymeric core surrounded by a conductive layer.
34. The sensor module of claim 33, wherein the working electrode, the reference electrode and the supplemental electrode are separated by an extruded dielectric

spacer that extends at least a majority of a length of the skin piercing member and that spaces the electrodes from an interior surface of the skin piercing member.

35. A sensor module comprising:

a carrier;

a skin piercing member carried by the carrier, the skin piercing member having a skin piercing end positioned opposite from a base end, the skin piercing member defining a lumen that extends along the central longitudinal axis from the skin piercing end toward the base end, the lumen defining a lumen axis;

a blood sample analysis zone located within the lumen of the skin piercing member; and

an elongated working electrode positioned within the lumen, the working electrode having a length that extends along the lumen axis, at least a section of the working electrode being positioned within the analysis zone, the working electrode including sensing chemistry, and the working electrode having an end within .5 millimeters of a tip of the skin piercing member.

36. The sensor module of claim 35, wherein the working electrode includes a fiber having a polymeric core surrounded by a conductive layer.

37. The sensor module of claim 25, wherein the working electrode is formed by a single fiber.

38. The sensor module of claim 1, wherein a surfactant is provided on the working electrode and/or on an interior surface of the skin piercing member.

39. A sensor module comprising:

a carrier movable relative to a base between a first position and second position;

a skin piercing member carried by the carrier, the skin piercing member having a skin piercing end positioned opposite from a base end, the skin piercing member defining a lumen that extends along the central longitudinal axis from the skin piercing end toward the base end, the lumen defining a lumen axis;

a blood sample analysis zone located entirely within the lumen of the skin piercing member; and

an elongated working electrode positioned within the lumen, the working electrode having a length that extends along the lumen axis, at least a section of the working electrode being positioned within the analysis zone, the working electrode including sensing chemistry.

40. A sensor module comprising:

a carrier;

a skin piercing member carried by the carrier, the skin piercing member having a skin piercing end positioned opposite from a base end, the skin piercing member defining a lumen that extends along the central longitudinal axis from the skin piercing end toward the base end, the lumen defining a lumen axis;

a blood sample analysis zone located within the lumen of the skin piercing member; and

an elongated working electrode positioned within the lumen, the working electrode having a length that extends along the lumen axis, at least a section of the working electrode being positioned within the analysis zone, the working electrode including sensing chemistry, and the working electrode being formed by a single fiber or wire.

41. A method for taking a blood analyte reading comprising:

puncturing skin with a skin piercing member having a lumen;

positioning a tip of the skin piercing member in a capillary blood field less than 3 millimeters beneath the skin;

initiating blood flow into the lumen by a combination of vascular blood pressure and capillary action to passively bring a blood sample to an analysis zone entirely within the lumen; and

sensing the blood analyte in the analysis zone.

42. The method of claim 41, further comprising stopping blood flow in the lumen at a capillary flow stop.

43. The method of claim 42, wherein the capillary flow stop includes a vent.

44. The method of claim 41, wherein a working electrode is present at the analysis zone within the lumen.
45. The method of claim 44, wherein the working electrode includes a fiber that extends through the lumen, and wherein less than .004 inches of circumferential clearance exists between the fiber and an interior surface of the skin piercing member.
46. The method of claim 41, wherein the capillary blood field has a capillary vascular pressure in the range of 14-22 millimeters of mercury.
47. The method of claim 42, wherein the blood analyte is sensed after blood flow has stopped in the lumen.
48. The method of claim 45, wherein the fiber is a single fiber having polymeric core, a conductive layer surrounding the core and sensing chemistry surrounding the conductive layer, the single fiber having an outer diameter in the range of .001 to .004 inches.
49. The method of claim 48, wherein the sensing chemistry is a dry sensing chemistry that is hydrated as the blood flows along the lumen.
50. The method of claim 49, wherein the sensing chemistry is dielectric when dry, and transitions from semi-conductive to conductive as the sensing chemistry hydrates.
51. A device for sensing an analyte in a blood sample, the device comprising:
an elongated working electrode having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing, the device being configured for a one time use in which one analyte reading is taken.
52. The device of claim 51, wherein the elongated working electrode includes a conductive fiber or wire, and wherein the first and second portions of the working electrode include sensing chemistry that cover the conductive fiber or wire.

53. The device of claim 51, wherein the elongated working electrode includes a single conductive fiber or a single wire, and wherein the first and second portions of the working electrode include sensing chemistry that cover the conductive fiber or wire.

54. The device of claim 51, wherein the working electrode includes a conductive wire or fiber, wherein sensing chemistry extends along at least a portion of a length of the working electrode, wherein the sensing chemistry is provided at the first and second portions of the working electrode, wherein the device includes an analysis zone including a controlled surface area of sensing chemistry that is wetted with blood during testing, the entire controlled surface area being wetted when the analyte reading is taken.

55. The device of claim 54, wherein the controlled surface area is at least 10 times larger than a transverse cross-sectional area of the working electrode.

56. The device of claim 54, wherein the controlled surface area is at least 20 times larger than a transverse cross-sectional area of the working electrode.

57. The device of claim 54, wherein the controlled surface area is at least 30 times larger than a transverse cross-sectional area of the working electrode.

58. The device of claim 51, wherein the first portion extends no more than 3 millimeters beneath the skin.

59. The device of claim 58, wherein the first portion extends 1-2 millimeters beneath the skin.

60. The device of claim 51, wherein the working electrode is positioned within a lumen of a skin piercing member.

61. The device of claim 60, wherein the skin piercing member is configured to generate a wound that shuts off blood flow when the skin piercing member is removed from the skin.

62. The device of claim 61, wherein the skin piercing member extends no more than 3 millimeters beneath the skin.
63. The device of claim 60, wherein the working electrode includes a conductive fiber or wire having an outer diameter that is at least 10 percent as large as an outer diameter of the skin piercing member.
64. The device of claim 60, wherein the working electrode is a conductive fiber or a wire, wherein sensing chemistry extends along at least a portion of a length of the working electrode, wherein the sensing chemistry is provided at the first and second portions of the working electrode, wherein the device includes an analysis zone including a controlled surface area of sensing chemistry that is wetted with blood during testing, wherein the entire controlled surface area is wetted when the analyte reading is taken, and wherein the controlled surface area is at least 10 times as large as a transverse cross-sectional area of the skin piercing member.
65. The device of claim 64, wherein the controlled surface area is at least 20 times as large as a transverse cross-sectional area of the skin piercing member.
66. The device of claim 64, wherein the transverse cross-sectional area of the skin piercing member corresponds to a transverse cross-sectional area of a puncture site created by the skin piercing member during testing.
67. The device of claim 61, wherein blood is fully contained within the device during testing.
68. The device of claim 60, wherein the working electrode includes a conductive fiber or wire having an outer diameter that is at least 20 percent as large as an outer diameter of the skin piercing member.
69. The device of claim 68, wherein the skin piercing member is 29 gauge or smaller.

70. A device for sensing an analyte in a blood sample, the device comprising:
a working electrode formed by a single conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing, the working electrode including a layer of sensing chemistry on the first and second portions of the conductive fiber or wire.
71. A device for sensing an analyte in a blood sample, the device comprising:
a working electrode formed by a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing, the working electrode including a layer of sensing chemistry on the first and second portions of the conductive fiber or wire, the device including a skin piercing member having a lumen in which the working electrode is positioned, the conductive fiber or wire having an outer diameter that is at least 10 percent as large as an outer diameter of the skin piercing member.
72. A device for sensing an analyte in a blood sample, the device comprising:
a working electrode formed by a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing, the working electrode including a layer of sensing chemistry on the first and second portions of the conductive fiber or wire, the device including a skin piercing member having a lumen in which the working electrode is positioned, the conductive fiber or wire having an outer diameter that is greater than .001 inches.
73. The device of claim 72, wherein the outer diameter of the conductive fiber or wire is greater than .002 inches.
74. The device of claim 72, wherein the outer diameter of the conductive fiber or wire is in the range of .0025 to .004 inches.
75. A device for sensing an analyte in a blood sample, the device comprising:
an elongated working electrode including a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing, conductive fiber or wire having a cross-dimension greater than .001 inches.

76. A device for sensing an analyte in a blood sample, the device comprising:
a working electrode formed by a conductive fiber or wire having a first portion that is subcutaneous during testing and a second portion that extends outside the body during testing, the working electrode including a layer of sensing chemistry on the first and second portions of the conductive fiber or wire, the device including a skin piercing member having a lumen in which the working electrode is positioned, the device being configured such that a blood analysis zone of the device fills passively.

FIG. 1

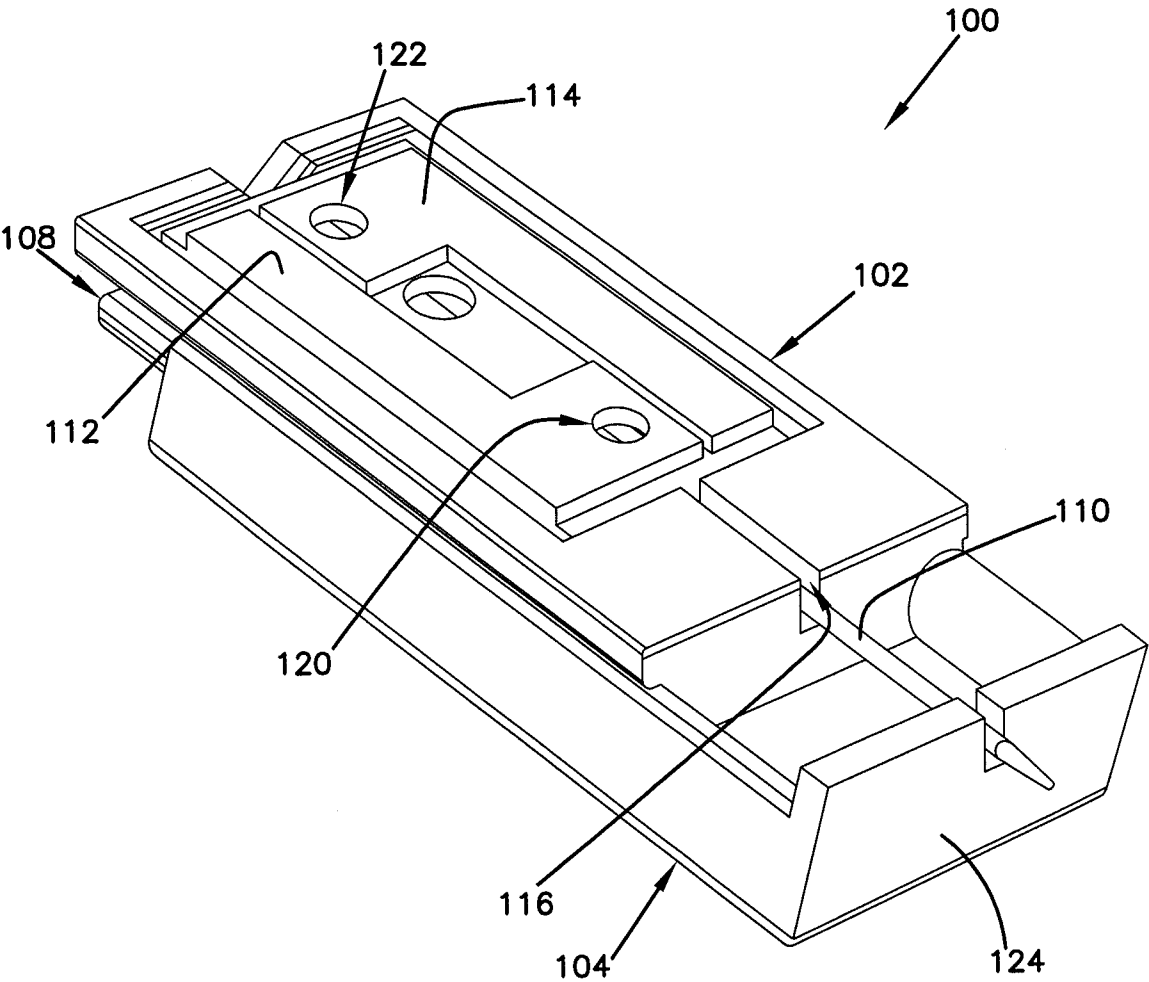
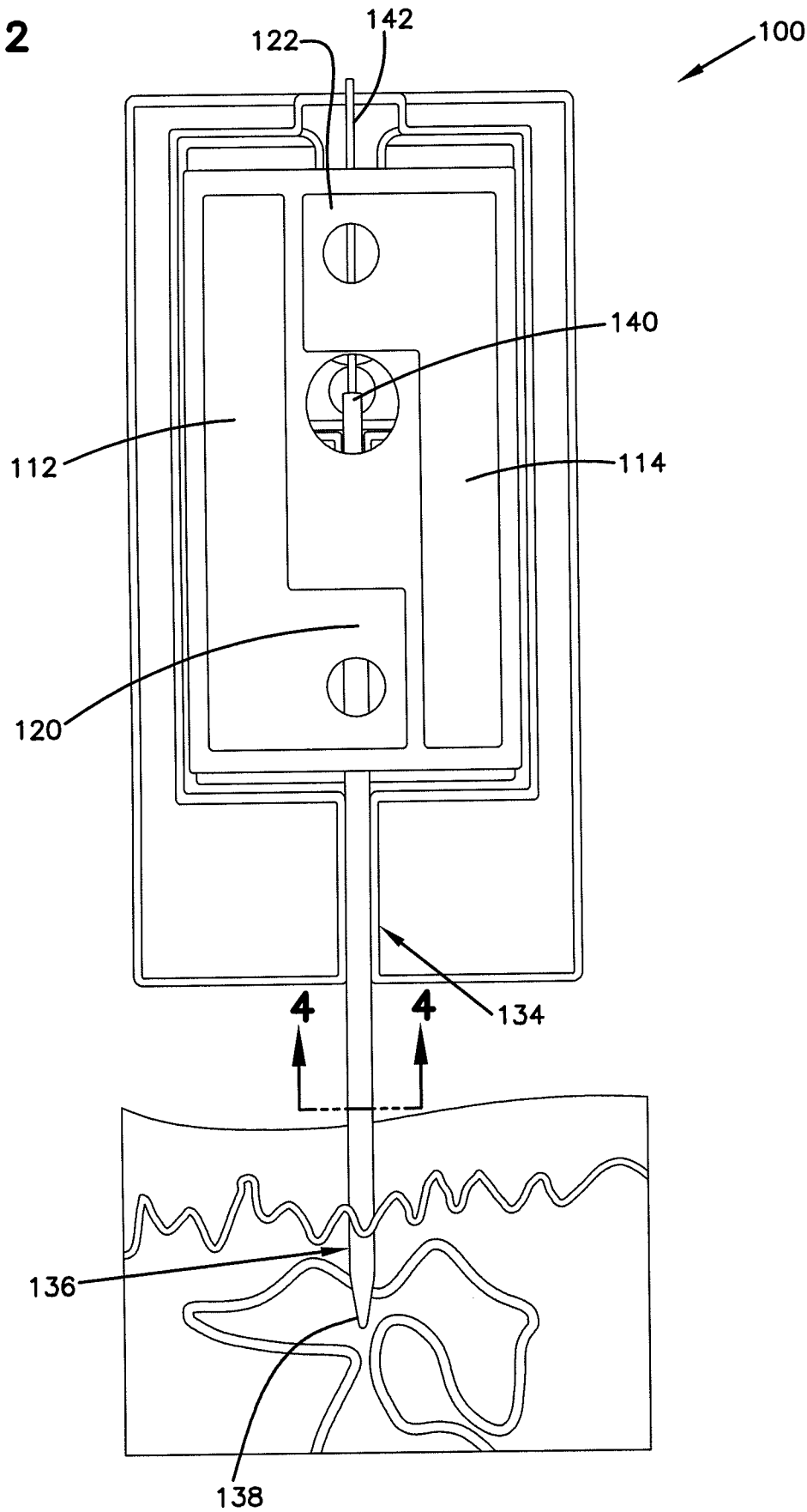


FIG. 2



3/13

FIG. 3

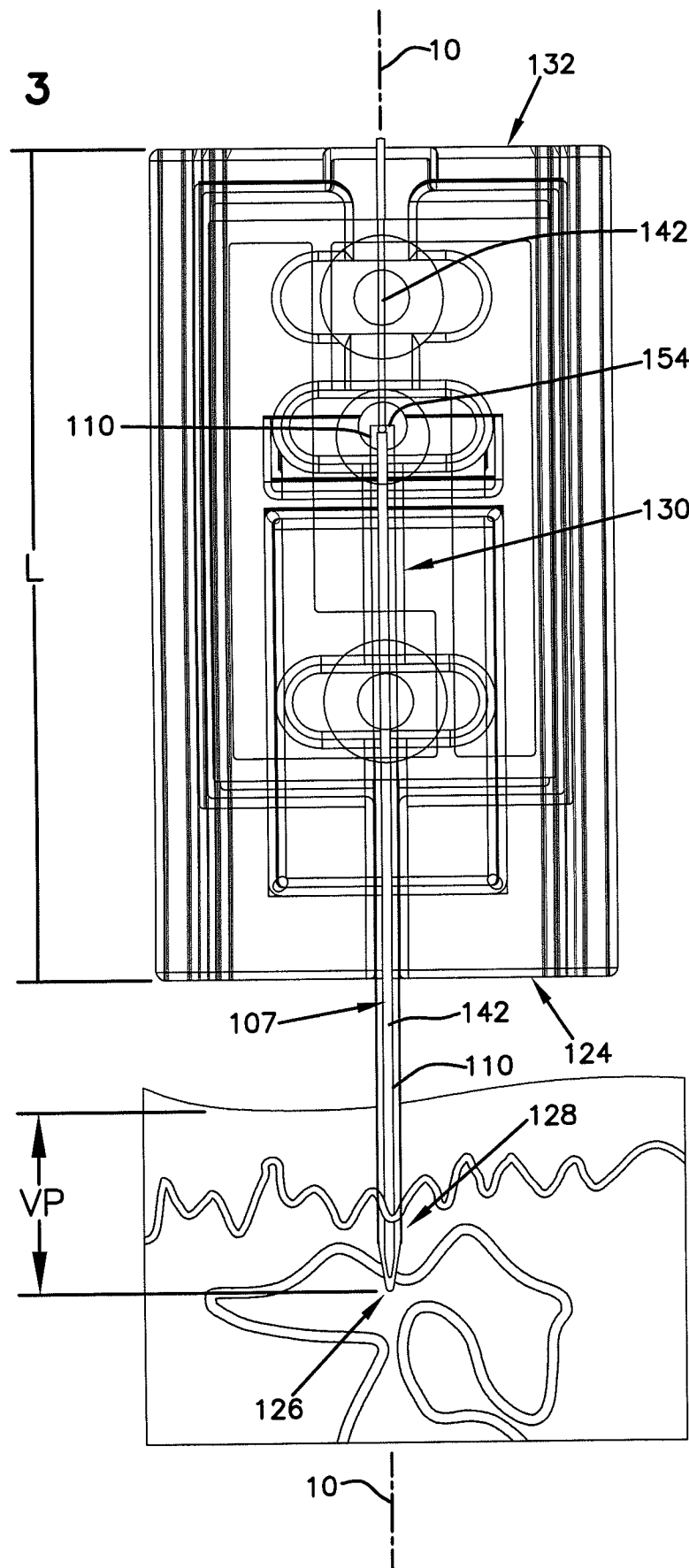


FIG. 4

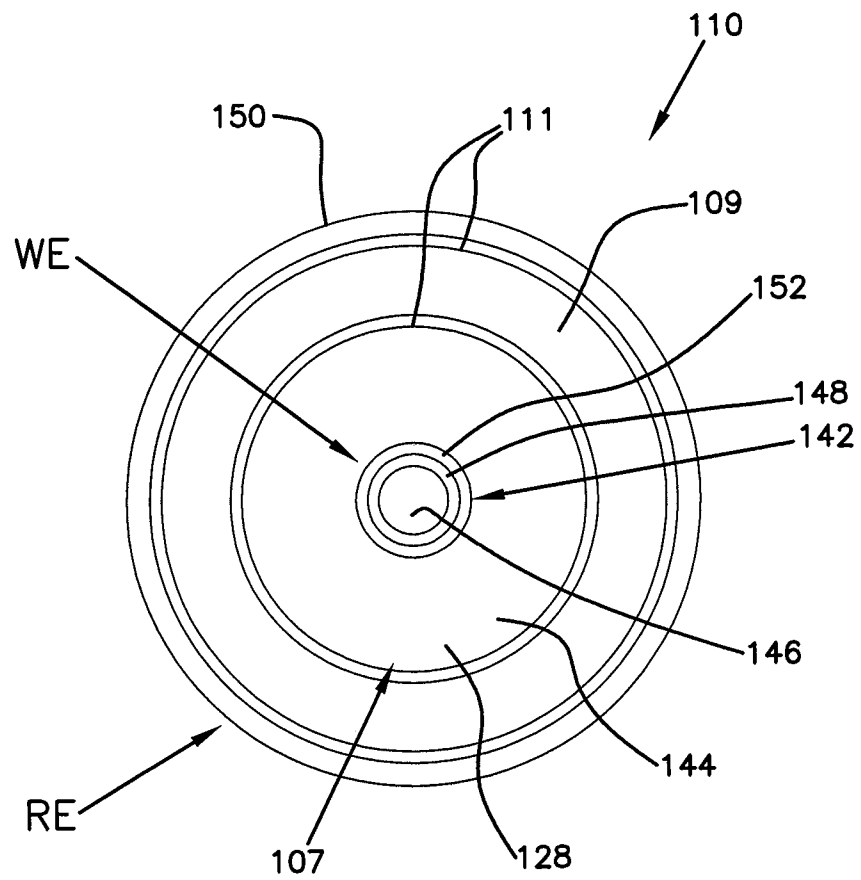


FIG. 5

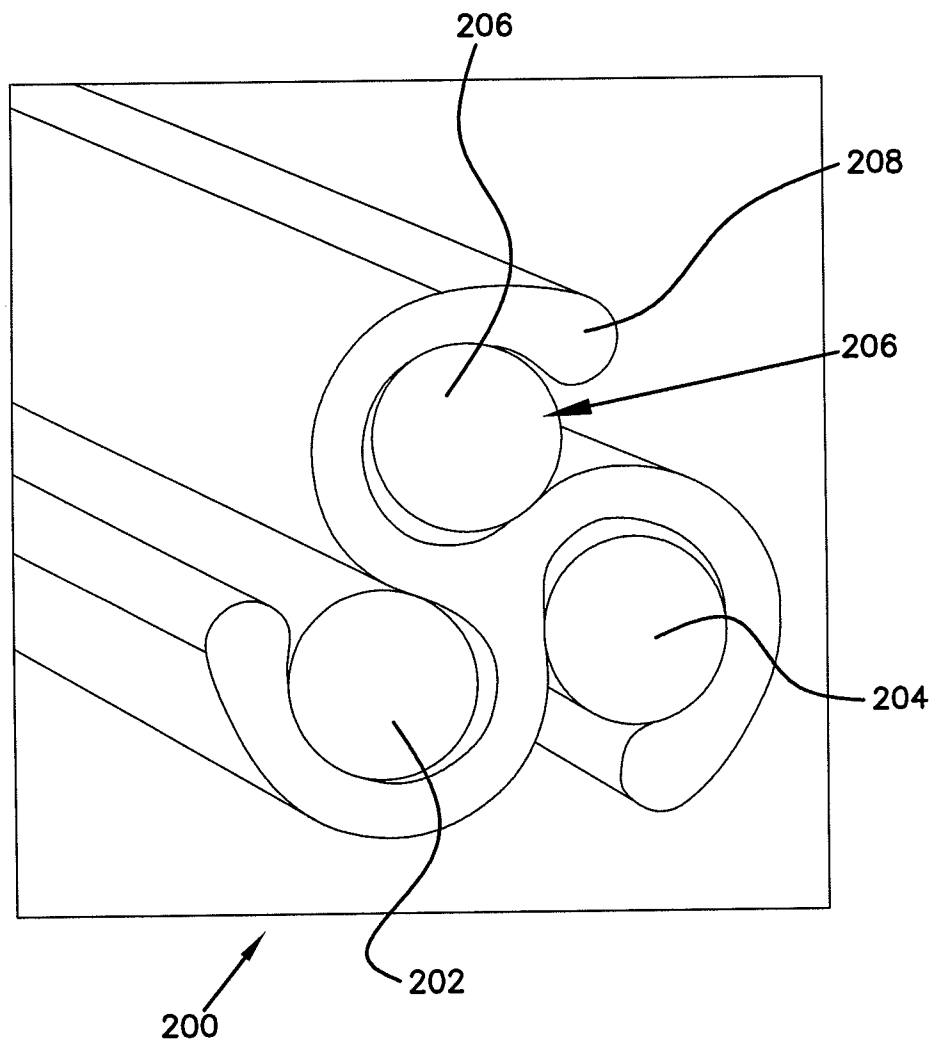


FIG. 6

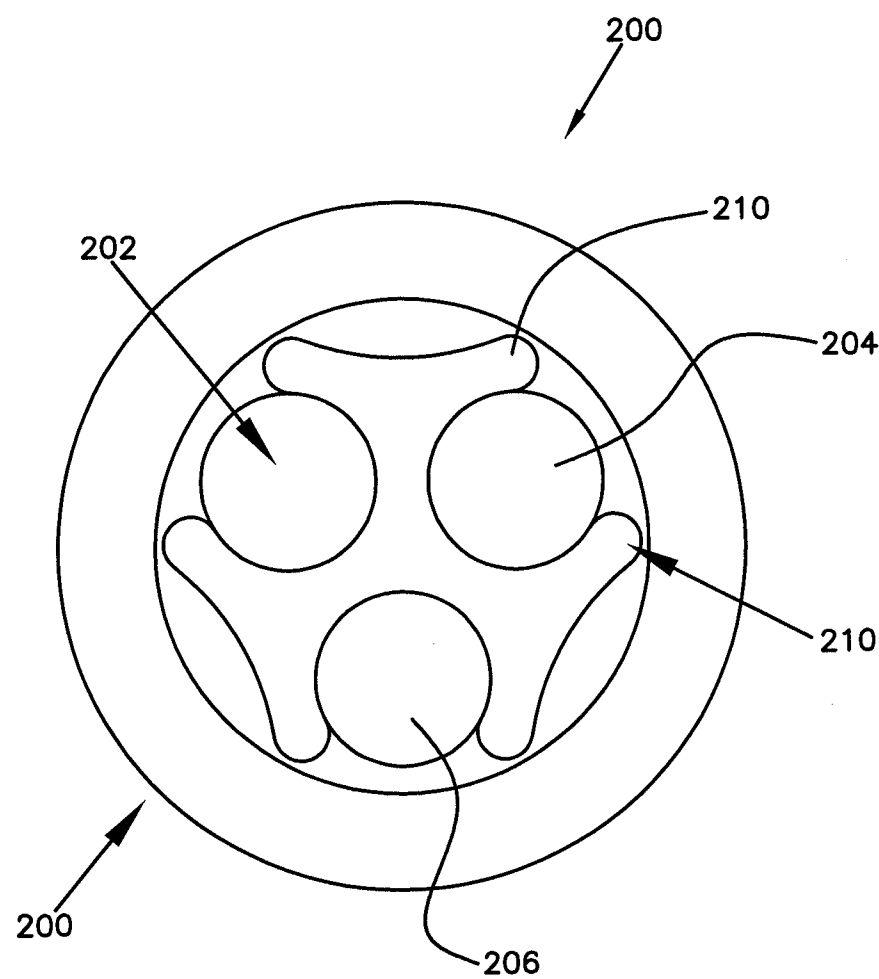


FIG. 7

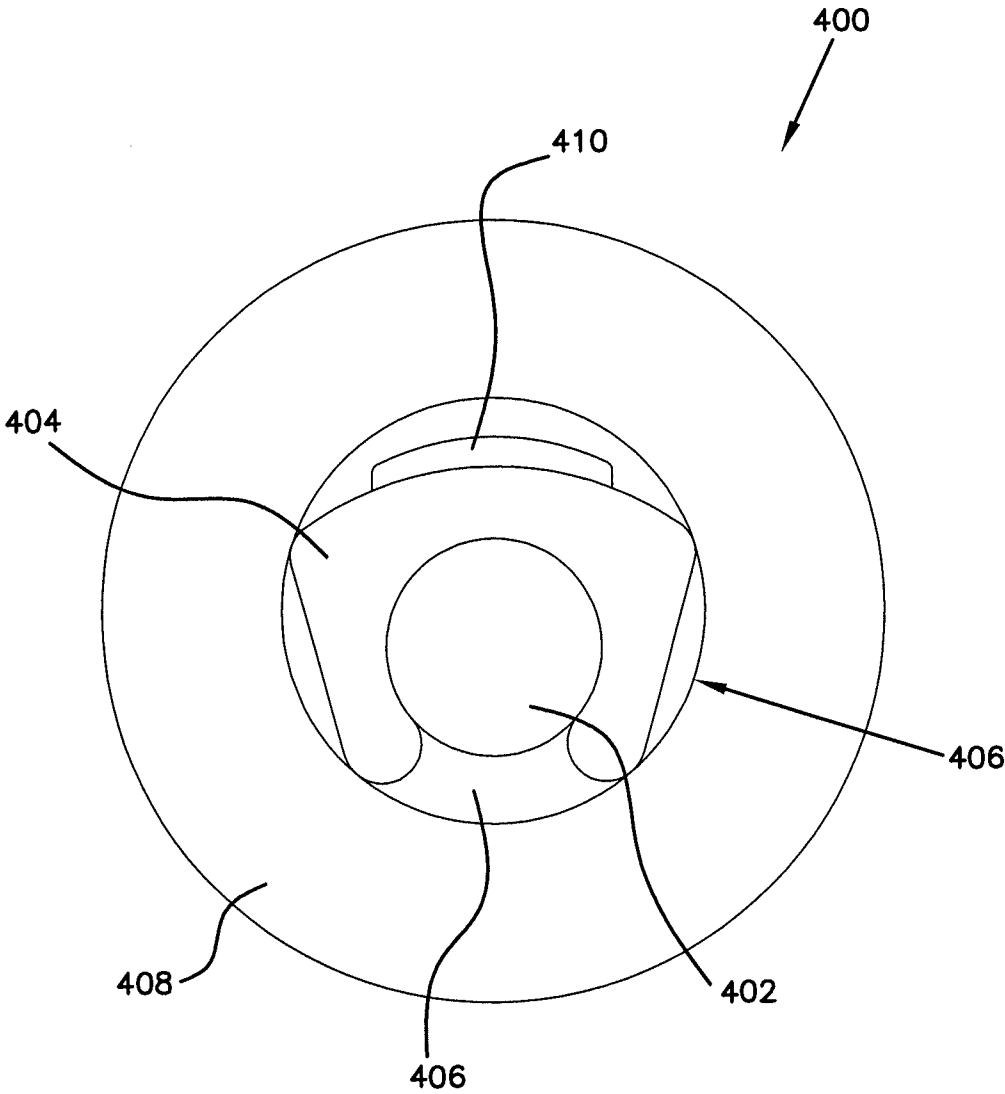


FIG. 8

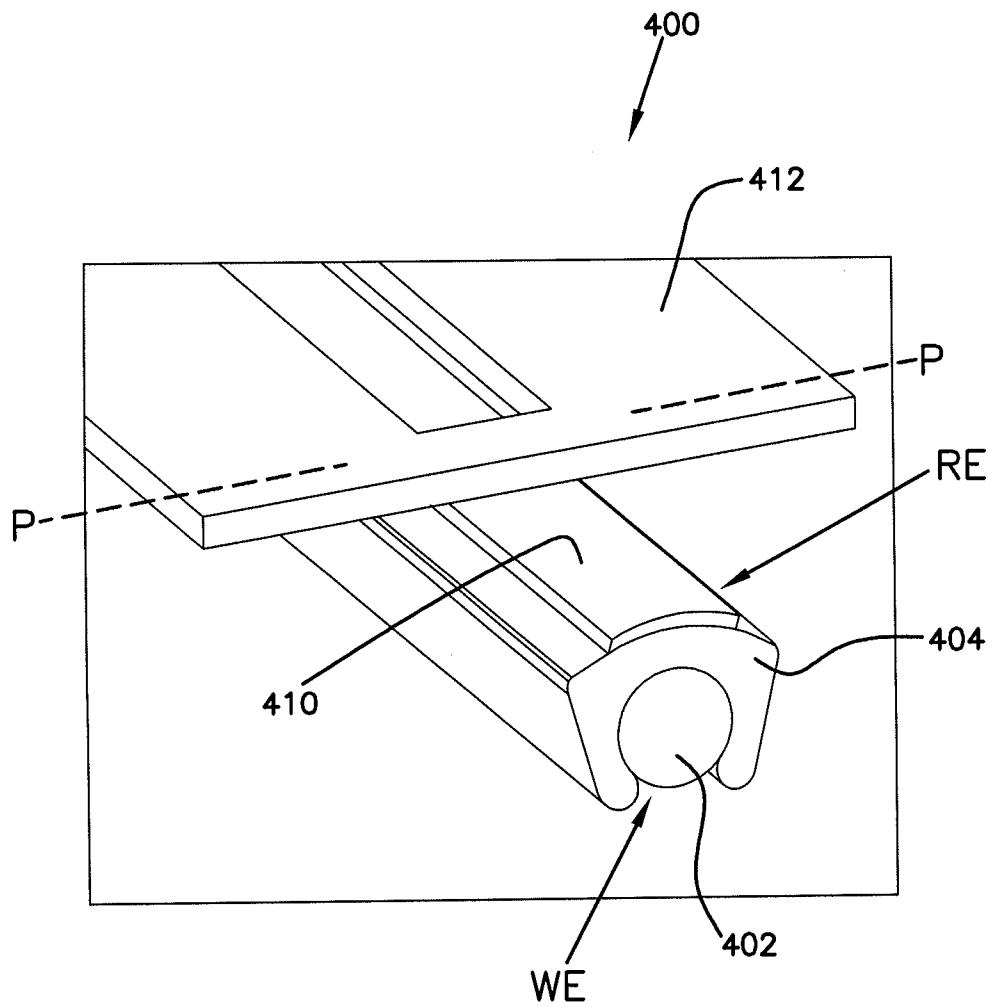


FIG. 9

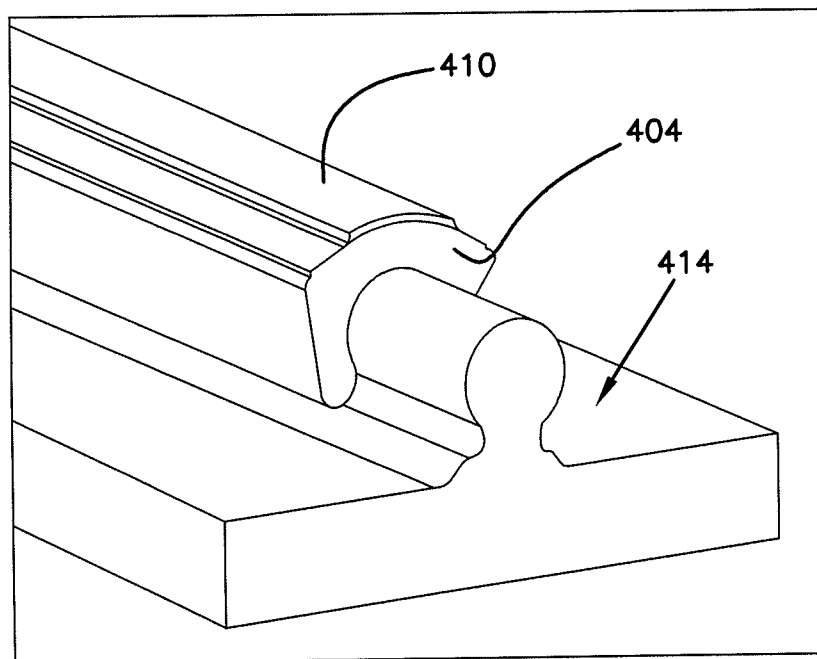


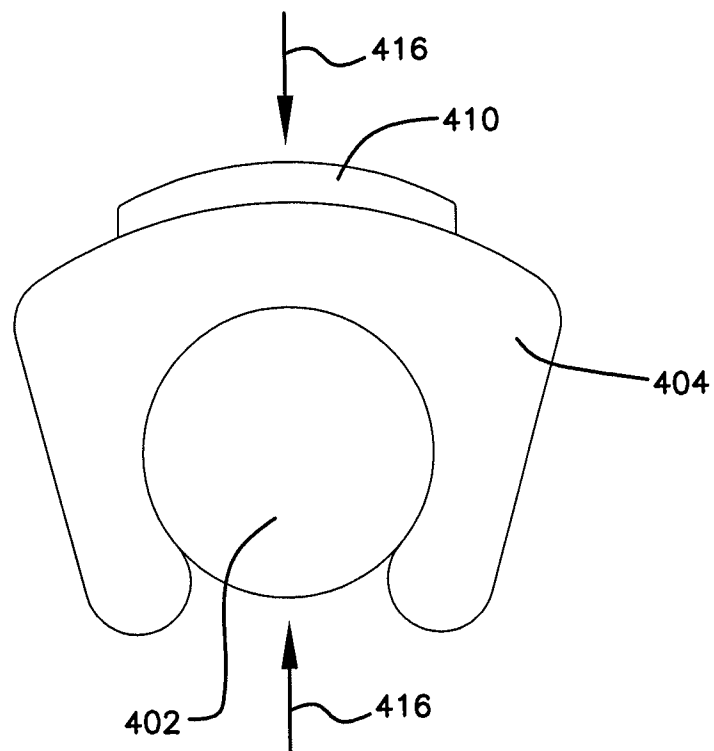
FIG. 10

FIG. 11

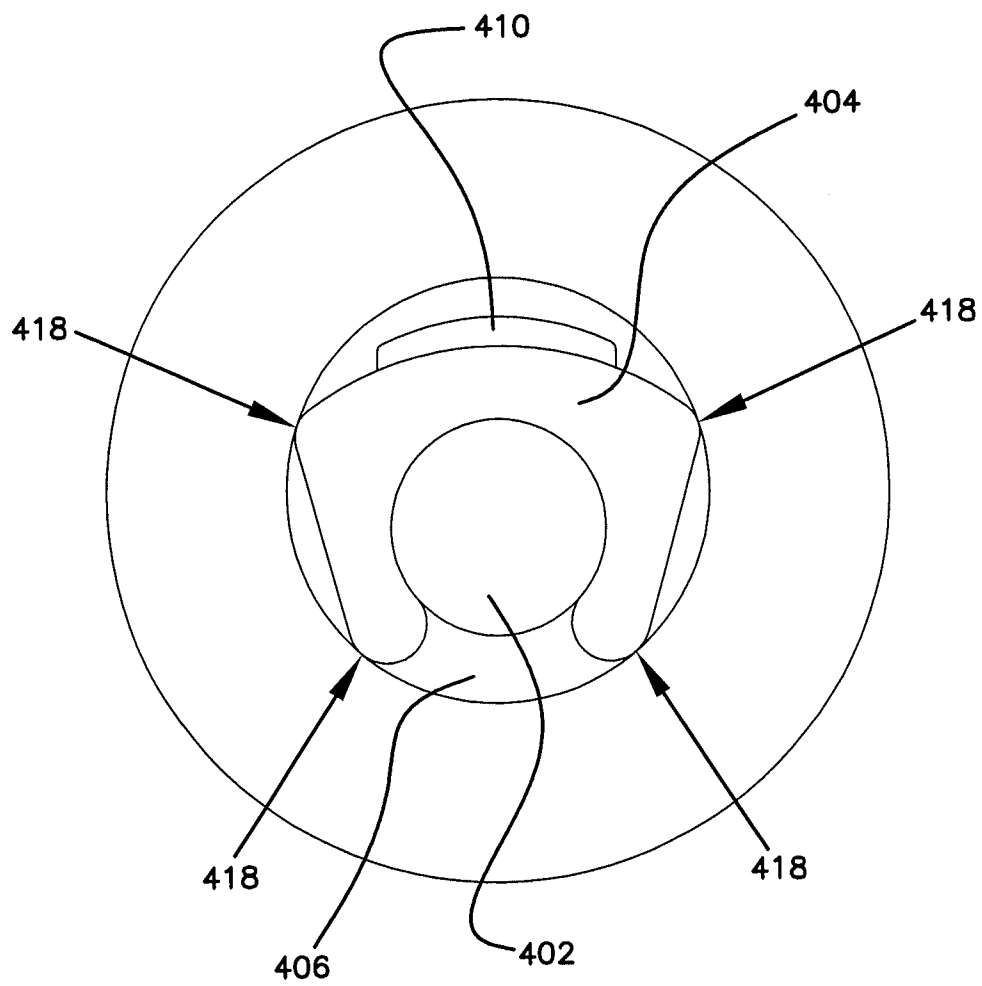
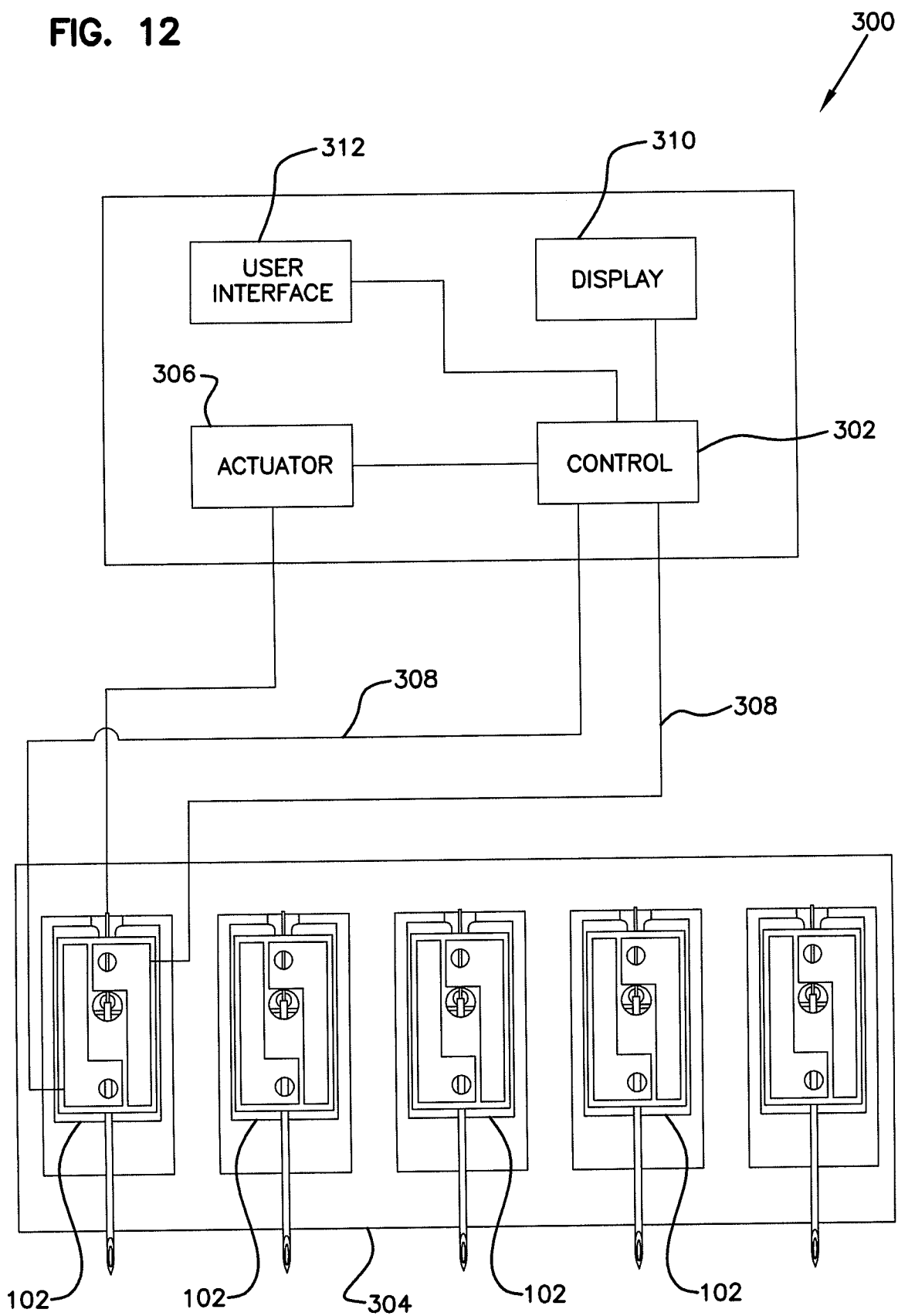
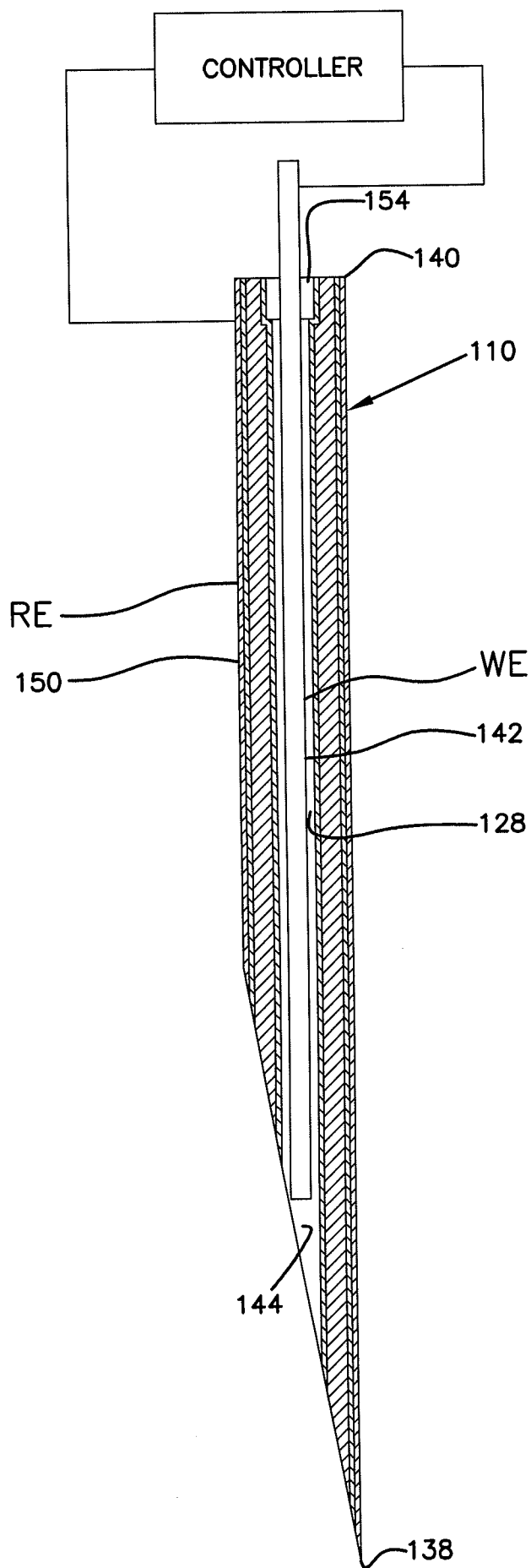


FIG. 12



13/13

FIG. 13



INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2013/072846

A. CLASSIFICATION OF SUBJECT MATTER

A61B 5/151(2006.01)i, A61B 5/157(2006.01)i

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

A61B 5/151; A61B 5/05; A61M 25/092; G01N 33/48; A61B 5/055; A61B 17/32; G01N 27/28; A61B 5/00; A61B 5/157

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Korean utility models and applications for utility models

Japanese utility models and applications for utility models

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

eKOMPASS(KIPO internal) & Keywords: sensor, blood, analysis, piercing, electrode

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 2011-0266149 A1 (JAMES L. SAY) 03 November 2011 See abstract, paragraphs [0034]-[0048],[0050], claim 1 and figures 3-5.	1-40,51-76
Y	WO 2008-118919 A1 (DEXCOM INC.) 02 October 2008 See abstract, paragraphs [0352]-[0371],[0540], claim 1 and figures 1A-1E.	1-40,51-76
A	US 2008-0319314 A1 (IRMA P. HILL et al.) 25 December 2008 See abstract, paragraphs [0030]-[0037],[0047]-[0050], claim 1 and figures 1-7.	1-40,51-76
A	US 5386828 A (ROBERT C. OWENS et al.) 07 February 1995 See abstract, column 5, line 42 - column 6, line 11 and figure 1.	1-40,51-76
A	US 2003-083686 A1 (DOMINIQUE M. FREEMAN et al.) 01 May 2003 See abstract, paragraphs [0120]-[0127] and figures 1-5,21.	1-40,51-76

☐ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

25 March 2014 (25.03.2014)

Date of mailing of the international search report

26 March 2014 (26.03.2014)

Name and mailing address of the ISA/KR



International Application Division
Korean Intellectual Property Office
189 Cheongsu-ro, Seo-gu, Daejeon Metropolitan City, 302-701,
Republic of Korea

Facsimile No. +82-42-472-7140

Authorized officer

KIM, Tae Hoon

Telephone No. +82-42-481-8407



INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2013/072846

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 41-50
because they relate to subject matter not required to be searched by this Authority, namely:
Claims 41-50 pertain to a method for treatment of the human body by surgery or by therapy and thus relate to a subject matter which this International Searching Authority is not required, under Article 17(2)(a)(i) of the PCT and Rule 39.1 (iv) of the Regulations under the PCT, to search.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of any additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2011-0266149 A1	03/11/2011	AU 2009-314069 A1 CN 102362172 A EP 2350629 A2 EP 2350629 A4 EP 2350629 B1 WO 2010-056869 A2 WO 2010-056869 A3	20/05/2010 22/02/2012 03/08/2011 07/11/2012 18/12/2013 20/05/2010 08/07/2010
WO 2008-118919 A1	02/10/2008	AU 2001-80886 A1 AU 2004-292229 A1 AU 2004-292229 B2 AU 2006-236319 A1 AU 2007-303239 A1 AU 2007-305002 A1 AU 2008-230832 A1 AU 2008-266162 A1 AU 2008-316630 A1 CA 2546072 A1 CA 2546072 C CA 2606770 A1 CA 2664426 A1 CA 2664528 A1 CA 2681412 A1 CA 2687980 A1 CA 2702799 A1 CN 100589003 C CN 101163998 A0 CN 101547633 A CN 101923195 A CN 1922525 A CN 1922525 C0 EP 1011425 A1 EP 1011425 B1 EP 1624907 A2 EP 1624908 A2 EP 1624908 B1 EP 1624908 B8 EP 1648293 A1 EP 1648298 A2 EP 1692556 A2 EP 1711789 A2 EP 1711790 A2 EP 1711790 B1 EP 1711791 A2 EP 1711802 A2 EP 1711802 B1 EP 1718350 A1 EP 1718350 B1 EP 1742568 A2	25/02/2002 02/06/2005 17/04/2008 26/10/2006 10/04/2008 10/04/2008 02/10/2008 24/12/2008 30/04/2009 02/06/2005 15/01/2013 26/10/2006 10/04/2008 10/04/2008 02/10/2008 24/12/2008 30/04/2009 10/02/2010 16/04/2008 30/09/2009 22/12/2010 28/02/2007 28/02/2007 28/06/2000 02/05/2007 15/02/2006 15/02/2006 05/01/2011 15/06/2011 26/04/2006 26/04/2006 23/08/2006 18/10/2006 18/10/2006 08/09/2010 18/10/2006 18/10/2006 14/07/2010 08/11/2006 01/05/2013 17/01/2007

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		EP 1776036 A1	25/04/2007
		EP 1776036 B1	01/05/2013
		EP 1804650 A1	11/07/2007
		EP 1804650 B1	14/03/2012
		EP 1875285 A2	09/01/2008
		EP 1893084 A2	05/03/2008
		EP 1914578 A2	23/04/2008
		EP 1914578 A3	30/04/2008
		EP 1991110 A2	19/11/2008
		EP 2004796 A2	24/12/2008
		EP 2069772 A2	17/06/2009
		EP 2091409 A2	26/08/2009
		EP 2129285 A1	09/12/2009
		EP 2155045 A1	24/02/2010
		EP 2203741 A1	07/07/2010
		EP 2223710 A1	01/09/2010
		EP 2226086 A1	08/09/2010
		EP 2228642 A1	15/09/2010
		EP 2239566 A2	13/10/2010
		EP 2239566 A3	17/11/2010
		EP 2239567 A2	13/10/2010
		EP 2239567 A3	17/11/2010
		EP 2256493 A1	01/12/2010
		EP 2264499 A2	22/12/2010
		EP 2264499 A3	05/01/2011
		EP 2264500 A2	22/12/2010
		EP 2264500 A3	05/01/2011
		EP 2264501 A2	22/12/2010
		EP 2264501 A3	05/01/2011
		EP 2264501 B1	25/07/2012
		EP 2301428 A1	30/03/2011
		EP 2316331 A1	04/05/2011
		EP 2322094 A1	18/05/2011
		EP 2327362 A1	01/06/2011
		EP 2327362 B1	13/11/2013
		EP 2327984 A2	01/06/2011
		EP 2327984 A3	07/09/2011
		EP 2329763 A1	08/06/2011
		EP 2329770 A1	08/06/2011
		EP 2329771 A2	08/06/2011
		EP 2329771 A3	22/06/2011
		EP 2332466 A1	15/06/2011
		EP 2335581 A1	22/06/2011
		EP 2335582 A1	22/06/2011
		EP 2335583 A2	22/06/2011
		EP 2335583 A3	31/08/2011
		EP 2335584 A2	22/06/2011
		EP 2335584 A3	31/08/2011
		EP 2335585 A2	22/06/2011
		EP 2335585 A3	31/08/2011

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		EP 2335586 A1	22/06/2011
		EP 2335587 A2	22/06/2011
		EP 2335587 A3	07/09/2011
		EP 2433563 A2	28/03/2012
		EP 2433563 A3	15/05/2013
		EP 2448485 A2	09/05/2012
		EP 2448486 A2	09/05/2012
		EP 2494921 A2	05/09/2012
		EP 2494921 A3	05/12/2012
		EP 2494922 A2	05/09/2012
		EP 2494922 A3	31/10/2012
		EP 2497415 A1	12/09/2012
		EP 2497420 A1	12/09/2012
		EP 2497421 A1	12/09/2012
		EP 2499969 A1	19/09/2012
		EP 2508129 A1	10/10/2012
		EP 2517623 A1	31/10/2012
		EP 2532302 A1	12/12/2012
		JP 2001-510382 A	31/07/2001
		JP 2006-525853 A	16/11/2006
		JP 2007-501028 A	25/01/2007
		JP 2007-501684 A	01/02/2007
		JP 2007-511737 A	10/05/2007
		JP 2007-514964 A	07/06/2007
		JP 2007-525276 A	06/09/2007
		JP 2007-535991 A	13/12/2007
		JP 2008-096448 A	24/04/2008
		JP 2008-506468 A	06/03/2008
		JP 2008-506469 A	06/03/2008
		JP 2008-538424 A	23/10/2008
		JP 2010-505534 A	25/02/2010
		JP 2011-224381 A	10/11/2011
		JP 2012-016597 A	26/01/2012
		JP 2012-110730 A	14/06/2012
		JP 2012-213654 A	08/11/2012
		JP 2013-099553 A	23/05/2013
		JP 4124827 B2	23/07/2008
		JP 4708342 B2	22/06/2011
		JP 4728249 B2	20/07/2011
		JP 4786653 B2	05/10/2011
		JP 4870075 B2	08/02/2012
		JP 5037128 B2	26/09/2012
		JP 5138819 B2	06/02/2013
		JP 5161341 B2	13/03/2013
		JP 5306521 B2	02/10/2013
		KR 10-1205756 B1	29/11/2012
		KR 10-1249602 B1	01/04/2013
		KR 10-2008-0005432 A	11/01/2008
		US 06001067 A	14/12/1999
		US 6435708 B1	20/08/2002

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		US 6558321 B1	06/05/2003
		US 6741877 B1	25/05/2004
		US 6862465 B2	01/03/2005
		US 6931327 B2	16/08/2005
		US 6983095 B2	03/01/2006
		US 7088899 B2	08/08/2006
		US 7103255 B2	05/09/2006
		US 7110803 B2	19/09/2006
		US 7136689 B2	14/11/2006
		US 7146089 B2	05/12/2006
		US 7171102 B2	30/01/2007
		US 7192450 B2	20/03/2007
		US 7200317 B2	03/04/2007
		US 7276029 B2	02/10/2007
		US 7310544 B2	18/12/2007
		US 7366556 B2	29/04/2008
		US 7369741 B2	06/05/2008
		US 7379765 B2	27/05/2008
		US 7400816 B2	15/07/2008
		US 7424318 B2	09/09/2008
		US 7460898 B2	02/12/2008
		US 7467003 B2	16/12/2008
		US 7471869 B2	30/12/2008
		US 7494465 B2	24/02/2009
		US 7497827 B2	03/03/2009
		US 7519408 B2	14/04/2009
		US 7583990 B2	01/09/2009
		US 7591801 B2	22/09/2009
		US 7599726 B2	06/10/2009
		US 7613491 B2	03/11/2009
		US 7615007 B2	10/11/2009
		US 7640048 B2	29/12/2009
		US 7646958 B1	12/01/2010
		US 7651596 B2	26/01/2010
		US 7654956 B2	02/02/2010
		US 7657297 B2	02/02/2010
		US 7711402 B2	04/05/2010
		US 7713574 B2	11/05/2010
		US 7715893 B2	11/05/2010
		US 7761130 B2	20/07/2010
		US 7771352 B2	10/08/2010
		US 7774145 B2	10/08/2010
		US 7775975 B2	17/08/2010
		US 7778680 B2	17/08/2010
		US 7783333 B2	24/08/2010
		US 7792562 B2	07/09/2010
		US 7797028 B2	14/09/2010
		US 7809232 B2	05/10/2010
		US 7809235 B2	05/10/2010
		US 7826981 B2	02/11/2010

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		US 7828728 B2	09/11/2010
		US 7831287 B2	09/11/2010
		US 7835777 B2	16/11/2010
		US 7844161 B2	30/11/2010
		US 7857760 B2	28/12/2010
		US 7860545 B2	28/12/2010
		US 7873255 B2	18/01/2011
		US 7875293 B2	25/01/2011
		US 7885697 B2	08/02/2011
		US 7896809 B2	01/03/2011
		US 7899511 B2	01/03/2011
		US 7901354 B2	08/03/2011
		US 7905833 B2	15/03/2011
		US 7914450 B2	29/03/2011
		US 7917186 B2	29/03/2011
		US 7925321 B2	12/04/2011
		US 7927274 B2	19/04/2011
		US 7933639 B2	26/04/2011
		US 7935057 B2	03/05/2011
		US 7946984 B2	24/05/2011
		US 7949381 B2	24/05/2011
		US 7955261 B2	07/06/2011
		US 7959569 B2	14/06/2011
		US 7970448 B2	28/06/2011
		US 7974672 B2	05/07/2011
		US 7976492 B2	12/07/2011
		US 7979104 B2	12/07/2011
		US 7986986 B2	26/07/2011
		US 7998071 B2	16/08/2011
		US 8000901 B2	16/08/2011
		US 8005335 B2	23/08/2011
		US 8005524 B2	23/08/2011
		US 8005525 B2	23/08/2011
		US 8010174 B2	30/08/2011
		US 8060173 B2	15/11/2011
		US 8064977 B2	22/11/2011
		US 8073519 B2	06/12/2011
		US 8073520 B2	06/12/2011
		US 8118877 B2	21/02/2012
		US 8128562 B2	06/03/2012
		US 8150488 B2	03/04/2012
		US 8155723 B2	10/04/2012
		US 8160669 B2	17/04/2012
		US 8160671 B2	17/04/2012
		US 8167801 B2	01/05/2012
		US 8170803 B2	01/05/2012
		US 8206297 B2	26/06/2012
		US 8216139 B2	10/07/2012
		US 8229534 B2	24/07/2012
		US 8229536 B2	24/07/2012

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		US 8231531 B2	31/07/2012
		US 8233958 B2	31/07/2012
		US 8233959 B2	31/07/2012
		US 8249684 B2	21/08/2012
		US 8251906 B2	28/08/2012
		US 8255030 B2	28/08/2012
		US 8255032 B2	28/08/2012
		US 8255033 B2	28/08/2012
		US 8260393 B2	04/09/2012
		US 8265725 B2	11/09/2012
		US 8275437 B2	25/09/2012
		US 8275438 B2	25/09/2012
		US 8280475 B2	02/10/2012
		US 8282549 B2	09/10/2012
		US 8285103 B2	09/10/2012
		US 8285354 B2	09/10/2012
		US 8287453 B2	16/10/2012
		US 8290559 B2	16/10/2012
US 2008-0319314 A1	25/12/2008	CA 2273467 A1	04/12/1999
		CA 2273467 C	17/02/2009
		DE 69936014 D1	21/06/2007
		DE 69936014 T2	10/01/2008
		EP 0962191 A1	08/12/1999
		EP 0962191 B1	09/05/2007
		JP 2000-024120 A	25/01/2000
		JP 2008-119522 A	29/05/2008
		JP 4312301 B2	12/08/2009
		JP 4541424 B2	08/09/2010
US 05386828 A	07/02/1995	AT 177612 T	15/04/1999
		AU 1993-21893 A	28/07/1993
		DE 69228680 D1	22/04/1999
		EP 0575580 A1	29/12/1993
		EP 0575580 B1	17/03/1999
		WO 93-12718 A1	08/07/1993
US 2003-0083686 A1	01/05/2003	AU 2002-312521 A8	23/12/2002
		AU 2002-315177 A8	23/12/2002
		AU 2002-315180 A8	23/12/2002
		AU 2002-320094 A8	23/12/2002
		AU 2002-330855 A8	23/12/2002
		AU 2002-344767 A1	11/11/2003
		AU 2002-344825 A8	23/12/2002
		AU 2002-348683 A8	23/12/2002
		AU 2002-367876 A1	03/11/2003
		AU 2003-231749 A1	03/11/2003
		AU 2003-234167 A1	03/11/2003
		AU 2003-239160 A1	03/11/2003
		AU 2003-268547 A1	29/03/2004

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		AU 2003-290589 A1	07/06/2004
		AU 2003-297205 A1	09/07/2004
		CA 2448681 A1	19/12/2002
		CA 2448790 A1	19/12/2002
		CA 2448790 C	07/09/2010
		CA 2448902 A1	19/12/2002
		CA 2448902 C	07/09/2010
		CA 2448905 A1	19/12/2002
		CA 2448905 C	07/09/2010
		CN 1662174 C0	31/08/2005
		CN 1735370 A	15/02/2006
		CN 1735370 C0	16/01/2008
		DK 1501427 T3	04/11/2013
		EP 1395185 A2	10/03/2004
		EP 1395185 B1	27/10/2010
		EP 1404218 A2	07/04/2004
		EP 1404232 A2	07/04/2004
		EP 1404232 B1	02/12/2009
		EP 1404233 A2	07/04/2004
		EP 1404233 B1	02/12/2009
		EP 1404234 A2	07/04/2004
		EP 1404234 B1	09/02/2011
		EP 1404235 A2	07/04/2004
		EP 1406537 A2	14/04/2004
		EP 1406537 B1	12/01/2011
		EP 1499247 A1	26/01/2005
		EP 1501402 A2	02/02/2005
		EP 1501409 A1	02/02/2005
		EP 1501410 A2	02/02/2005
		EP 1501427 A1	02/02/2005
		EP 1501427 A4	08/10/2008
		EP 1501427 B1	24/07/2013
		EP 1562470 A1	17/08/2005
		EP 1578286 A1	28/09/2005
		EP 1585558 A2	19/10/2005
		EP 1638454 A2	29/03/2006
		EP 1945093 A2	23/07/2008
		EP 1968461 A2	17/09/2008
		EP 1978865 A2	15/10/2008
		EP 2301438 A2	30/03/2011
		EP 2301438 A3	20/07/2011
		JP 2004-528936 A	24/09/2004
		JP 2005-504564 A	17/02/2005
		JP 2005-506857 A	10/03/2005
		JP 2005-517463 A	16/06/2005
		JP 2005-523065 A	04/08/2005
		JP 2006-504506 A	09/02/2006
		JP 2008-289939 A	04/12/2008
		JP 2009-106784 A	21/05/2009
		JP 2012-161627 A	30/08/2012

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		JP 2012-223612 A	15/11/2012
		JP 4149911 B2	17/09/2008
		JP 4150071 B2	01/10/2008
		JP 4209767 B2	14/01/2009
		JP 4272051 B2	03/06/2009
		JP 4522364 B2	11/08/2010
		JP 4996811 B2	08/08/2012
		JP 5134475 B2	30/01/2013
		JP 5232664 B2	10/07/2013
		US 2003-083685 A1	01/05/2003
		US 2003-199789 A1	23/10/2003
		US 2003-199790 A1	23/10/2003
		US 2003-199791 A1	23/10/2003
		US 2003-199893 A1	23/10/2003
		US 2003-199894 A1	23/10/2003
		US 2003-199895 A1	23/10/2003
		US 2003-199896 A1	23/10/2003
		US 2003-199897 A1	23/10/2003
		US 2003-199898 A1	23/10/2003
		US 2003-199899 A1	23/10/2003
		US 2003-199900 A1	23/10/2003
		US 2003-199901 A1	23/10/2003
		US 2003-199902 A1	23/10/2003
		US 2003-199903 A1	23/10/2003
		US 2003-199904 A1	23/10/2003
		US 2003-199905 A1	23/10/2003
		US 2003-199906 A1	23/10/2003
		US 2003-199907 A1	23/10/2003
		US 2003-199908 A1	23/10/2003
		US 2003-199909 A1	23/10/2003
		US 2003-199910 A1	23/10/2003
		US 2003-199911 A1	23/10/2003
		US 2003-212424 A1	13/11/2003
		US 2003-233112 A1	18/12/2003
		US 2003-233113 A1	18/12/2003
		US 2004-010279 A1	15/01/2004
		US 2004-049219 A1	11/03/2004
		US 2004-049220 A1	11/03/2004
		US 2004-067481 A1	08/04/2004
		US 2004-087990 A1	06/05/2004
		US 2004-092842 A1	13/05/2004
		US 2004-092994 A1	13/05/2004
		US 2004-092995 A1	13/05/2004
		US 2004-098009 A1	20/05/2004
		US 2004-102803 A1	27/05/2004
		US 2005-101981 A1	12/05/2005
		US 2005-256534 A1	17/11/2005
		US 2006-052810 A1	09/03/2006
		US 2006-085020 A1	20/04/2006
		US 2006-175216 A1	10/08/2006

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		US 2006-178687 A1	10/08/2006
		US 2006-178688 A1	10/08/2006
		US 2006-178689 A1	10/08/2006
		US 2006-178690 A1	10/08/2006
		US 2006-195047 A1	31/08/2006
		US 2006-195129 A1	31/08/2006
		US 2006-195130 A1	31/08/2006
		US 2006-195131 A1	31/08/2006
		US 2006-195132 A1	31/08/2006
		US 2006-195133 A1	31/08/2006
		US 2006-200044 A1	07/09/2006
		US 2006-241667 A1	26/10/2006
		US 2006-271083 A1	30/11/2006
		US 2007-038235 A1	15/02/2007
		US 2007-043305 A1	22/02/2007
		US 2007-043386 A1	22/02/2007
		US 2007-055174 A1	08/03/2007
		US 2007-064516 A1	22/03/2007
		US 2007-073188 A1	29/03/2007
		US 2007-073189 A1	29/03/2007
		US 2007-100255 A1	03/05/2007
		US 2007-123802 A1	31/05/2007
		US 2007-142747 A1	21/06/2007
		US 2007-142748 A1	21/06/2007
		US 2007-167870 A1	19/07/2007
		US 2007-167871 A1	19/07/2007
		US 2007-167872 A1	19/07/2007
		US 2007-167873 A1	19/07/2007
		US 2007-167874 A1	19/07/2007
		US 2007-167875 A1	19/07/2007
		US 2007-173741 A1	26/07/2007
		US 2007-173742 A1	26/07/2007
		US 2007-173743 A1	26/07/2007
		US 2007-185412 A1	09/08/2007
		US 2007-191736 A1	16/08/2007
		US 2007-191737 A1	16/08/2007
		US 2007-213601 A1	13/09/2007
		US 2007-213756 A1	13/09/2007
		US 2007-219462 A1	20/09/2007
		US 2007-219463 A1	20/09/2007
		US 2007-219573 A1	20/09/2007
		US 2007-219574 A1	20/09/2007
		US 2007-239189 A1	11/10/2007
		US 2007-239190 A1	11/10/2007
		US 2007-244499 A1	18/10/2007
		US 7025774 B2	11/04/2006
		US 7033371 B2	25/04/2006
		US 7041068 B2	09/05/2006
		US 7141058 B2	28/11/2006
		US 7175642 B2	13/02/2007

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		US 7198606 B2	03/04/2007
		US 7226461 B2	05/06/2007
		US 7229458 B2	12/06/2007
		US 7232451 B2	19/06/2007
		US 7244265 B2	17/07/2007
		US 7258693 B2	21/08/2007
		US 7291117 B2	06/11/2007
		US 7297122 B2	20/11/2007
		US 7297151 B2	20/11/2007
		US 7316700 B2	08/01/2008
		US 7331931 B2	19/02/2008
		US 7344507 B2	18/03/2008
		US 7371247 B2	13/05/2008
		US 7374544 B2	20/05/2008
		US 7410468 B2	12/08/2008
		US 7481776 B2	27/01/2009
		US 7485128 B2	03/02/2009
		US 7491178 B2	17/02/2009
		US 7524293 B2	28/04/2009
		US 7537571 B2	26/05/2009
		US 7547287 B2	16/06/2009
		US 7563232 B2	21/07/2009
		US 7582099 B2	01/09/2009
		US 7648468 B2	19/01/2010
		US 7648469 B2	19/01/2010
		US 7674232 B2	09/03/2010
		US 7708701 B2	04/05/2010
		US 7713214 B2	11/05/2010
		US 7717863 B2	18/05/2010
		US 7731729 B2	08/06/2010
		US 7749174 B2	06/07/2010
		US 7798979 B2	21/09/2010
		US 7833171 B2	16/11/2010
		US 7841992 B2	30/11/2010
		US 7850622 B2	14/12/2010
		US 7862520 B2	04/01/2011
		US 7874994 B2	25/01/2011
		US 7875047 B2	25/01/2011
		US 7892183 B2	22/02/2011
		US 7892185 B2	22/02/2011
		US 7901362 B2	08/03/2011
		US 7901365 B2	08/03/2011
		US 7909774 B2	22/03/2011
		US 7909775 B2	22/03/2011
		US 7909777 B2	22/03/2011
		US 7909778 B2	22/03/2011
		US 7914465 B2	29/03/2011
		US 7938787 B2	10/05/2011
		US 7959582 B2	14/06/2011
		US 7976476 B2	12/07/2011

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2013/072846

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		US 7981055 B2	19/07/2011
		US 7981056 B2	19/07/2011
		US 7988644 B2	02/08/2011
		US 7988645 B2	02/08/2011
		US 8007446 B2	30/08/2011
		US 8016774 B2	13/09/2011
		US 8062231 B2	22/11/2011
		US 8079960 B2	20/12/2011
		US 8123700 B2	28/02/2012
		US 8157748 B2	17/04/2012
		US 8162853 B2	24/04/2012
		US 8197421 B2	12/06/2012
		US 8197423 B2	12/06/2012
		US 8202231 B2	19/06/2012
		US 8206317 B2	26/06/2012
		US 8206319 B2	26/06/2012
		US 8211037 B2	03/07/2012
		US 8216154 B2	10/07/2012
		US 8221334 B2	17/07/2012
		US 8235915 B2	07/08/2012
		US 8267870 B2	18/09/2012
		US 8282577 B2	09/10/2012
		US 8333710 B2	18/12/2012
		US 8337419 B2	25/12/2012
		US 8337420 B2	25/12/2012
		US 8337421 B2	25/12/2012
		US 8343075 B2	01/01/2013
		US 8360991 B2	29/01/2013
		US 8382682 B2	26/02/2013
		US 8382683 B2	26/02/2013
		US 8403864 B2	26/03/2013
		US 8414503 B2	09/04/2013
		US 8430828 B2	30/04/2013
		US 8435190 B2	07/05/2013
		US 8491500 B2	23/07/2013
		US 8496601 B2	30/07/2013
		US 8556829 B2	15/10/2013
		US 8562545 B2	22/10/2013
		US 8574168 B2	05/11/2013
		US 8579831 B2	12/11/2013

专利名称(译)	传感器模块和使用传感器模块的方法		
公开(公告)号	EP2925229A4	公开(公告)日	2017-01-25
申请号	EP2013860026	申请日	2013-12-03
申请(专利权)人(译)	PEPEX生物医学，INC.		
当前申请(专利权)人(译)	Pepex生物医药公司		
[标]发明人	SAY JAMES L		
发明人	SAY, JAMES, L.		
IPC分类号	A61B5/151 A61B5/157 A61B5/00 A61B5/145 A61B5/1486 A61B5/15		
CPC分类号	A61B5/150396 A61B5/14532 A61B5/14865 A61B5/150022 A61B5/150213 A61B5/150389 A61B5/150755 A61B5/15105 A61B5/15142 A61B5/157 A61B5/6848 A61B2560/0468		
优先权	61/732783 2012-12-03 US		
其他公开文献	EP2925229A1		
外部链接	Espacenet		

摘要(译)

本文公开了一种传感器模块。传感器模块包括由载体承载的皮肤刺穿构件。皮肤刺穿构件具有与基端相对定位的皮肤刺穿端。皮肤刺穿构件限定沿着中心纵向轴线从皮肤穿透端朝向基端延伸的内腔，并且内腔具有内腔轴线。传感器模块还包括完全位于皮肤刺穿构件的内腔内的血液样品分析区和用于停止在皮肤刺穿构件的内腔内的预定位置处的毛细流动的毛细流动停止器。传感器模块包括定位在内腔内的细长工作电极。工作电极具有沿着内腔轴延伸的长度，其中工作电极的至少一部分位于分析区内。工作电极包括感测化学。