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(54) **SMALL VOLUME IN VITRO ANALYTE SENSOR AND METHODS**

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(52) **U.S. Cl.** **205/792; 204/403.03**

(57) **ABSTRACT**

A small volume sensor, and methods of making, for determining the concentration of an analyte, such as glucose or lactate, in a biological fluid, such as blood or serum, using techniques such as coulometry, amperometry, and potentiometry. The sensor includes a working electrode and a counter electrode, and can include an insertion monitoring trace to determine correct positioning of the sensor in a connector. In one embodiment, the sensor determines the concentration of the analyte by discharging an amount of charge into the sample, determining the time needed to discharge the charge, and determining the current used to electrolyze a portion of the analyte using the amount of charge and the amount of time.

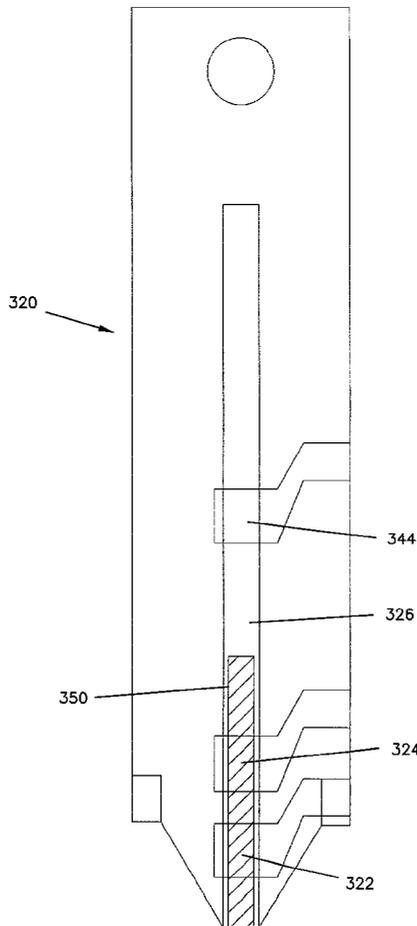
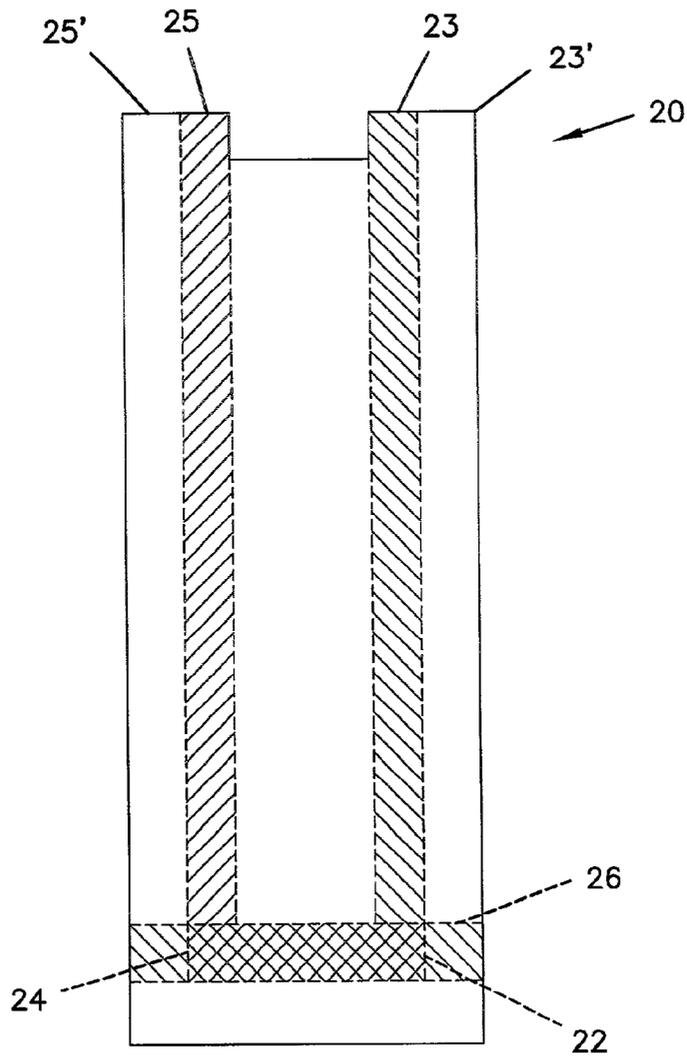


FIG. 1



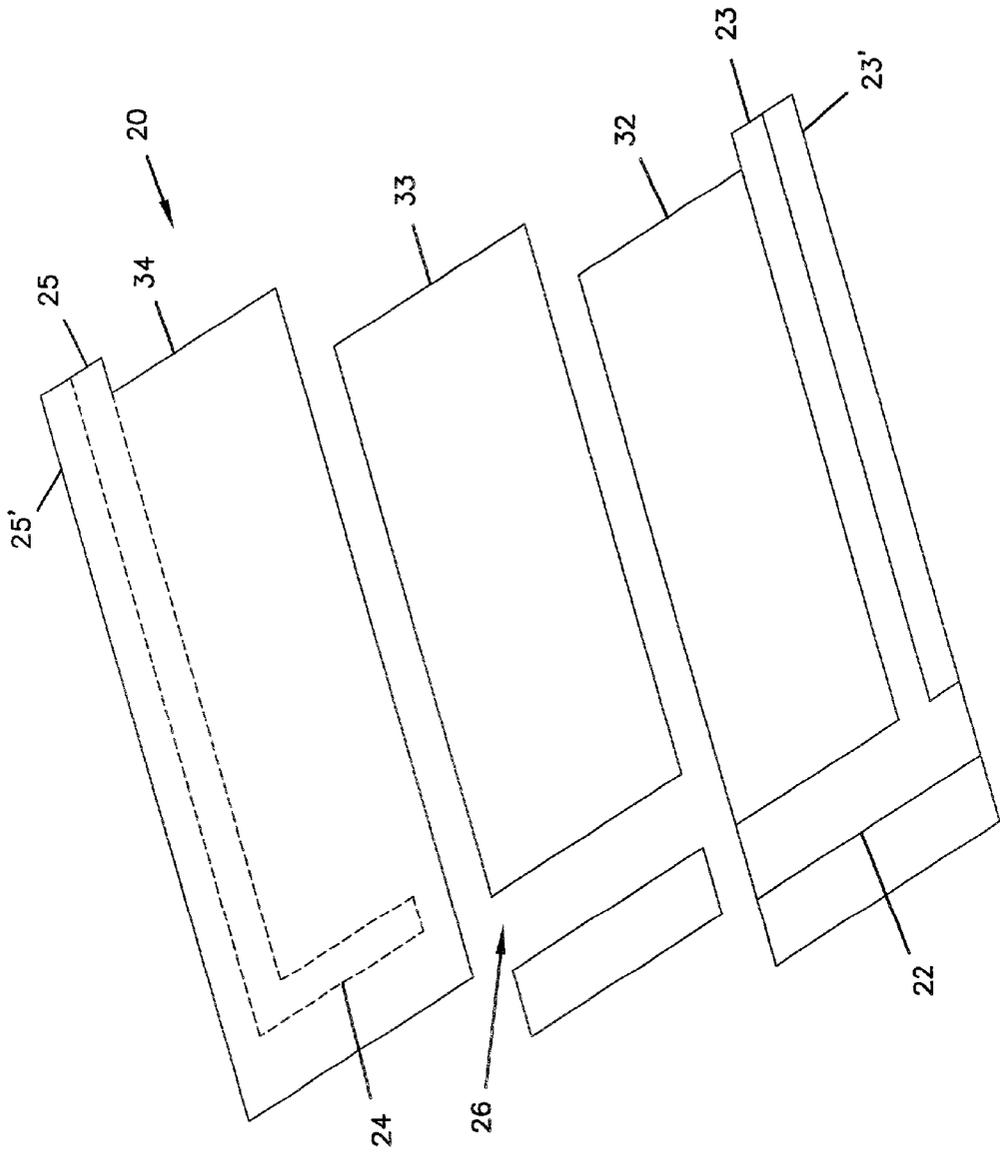


FIG. 2

FIG. 4

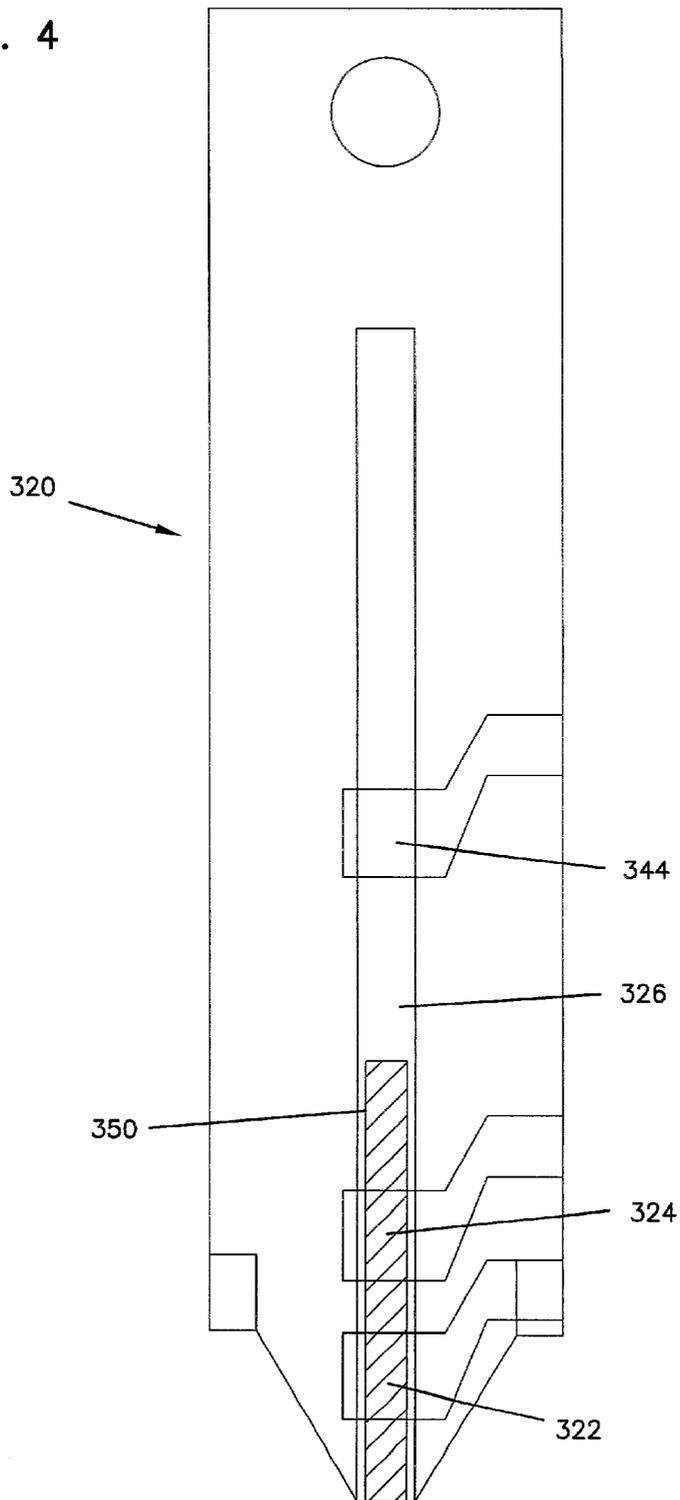


FIG. 5A

FIG. 5B

FIG. 5C

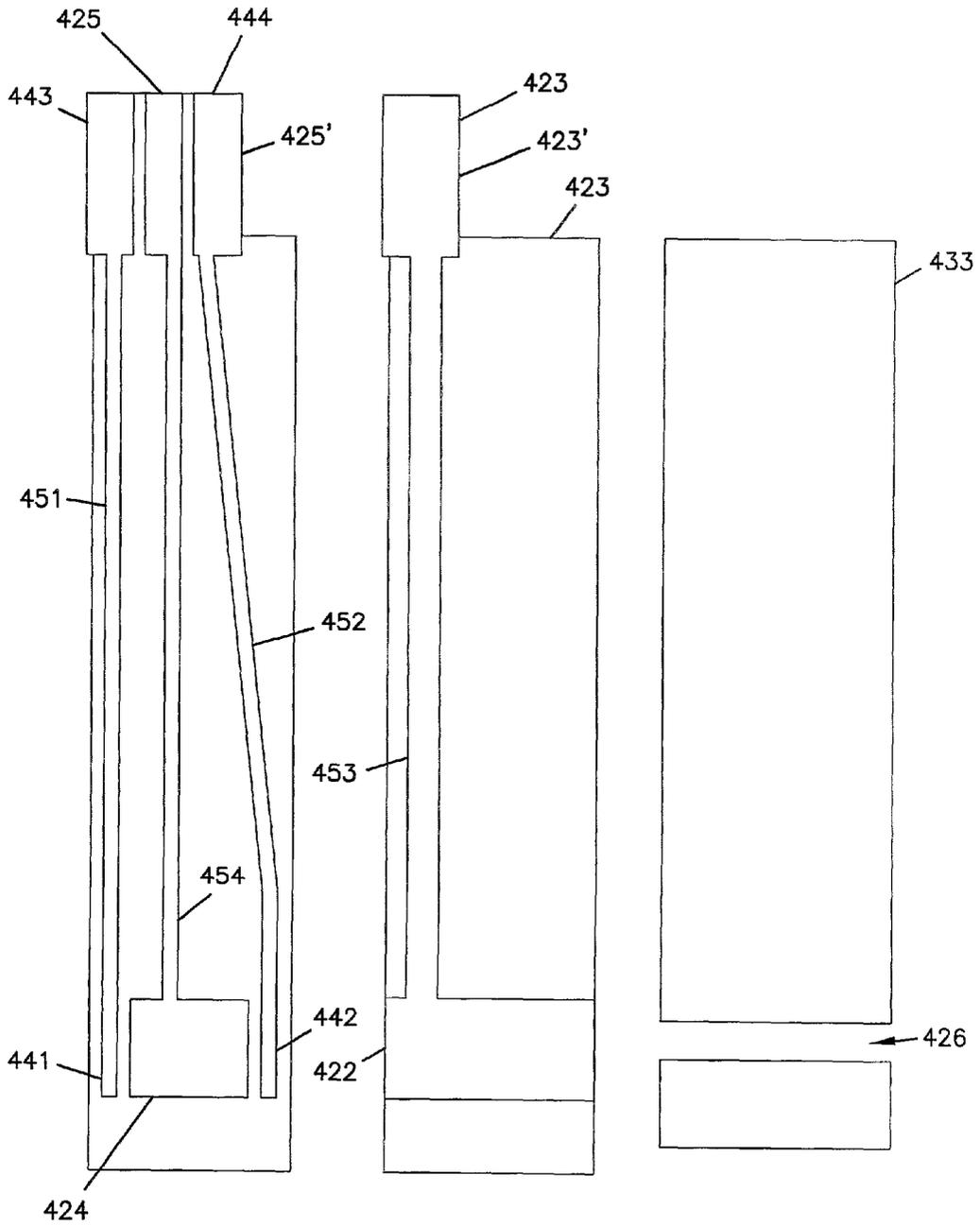
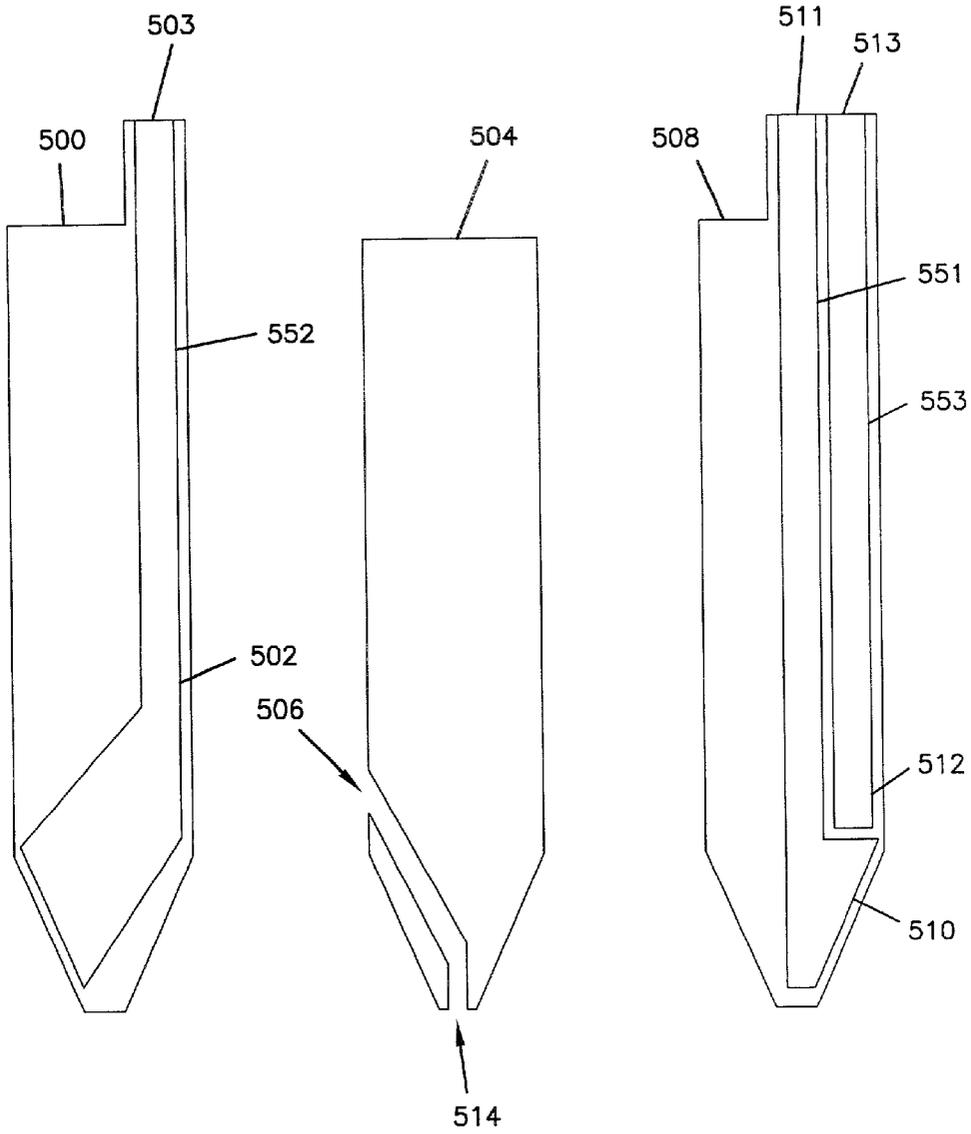
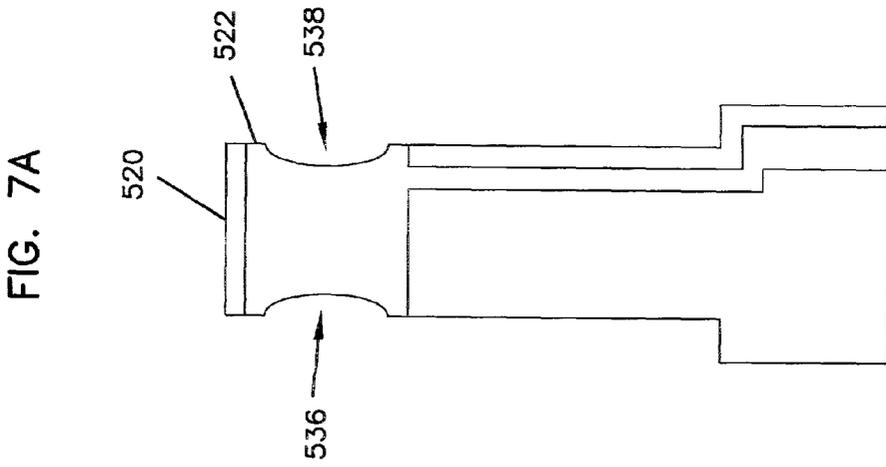
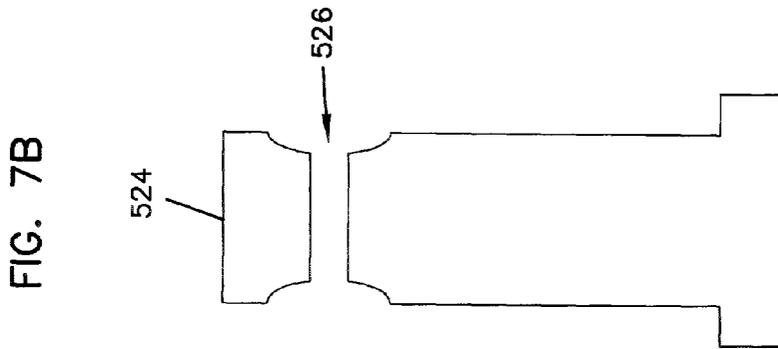
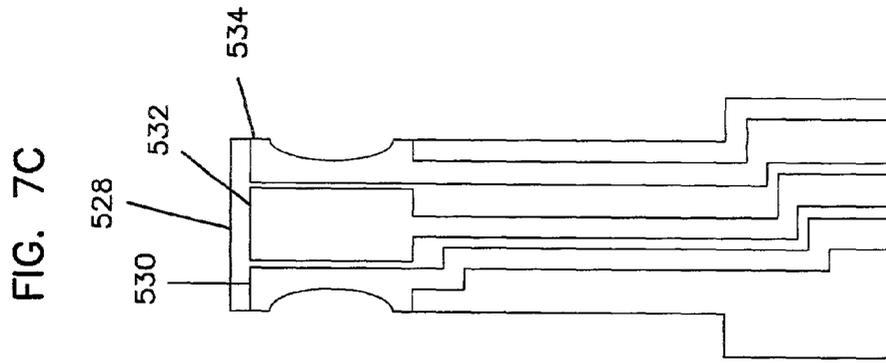


FIG. 6A

FIG. 6B

FIG. 6C





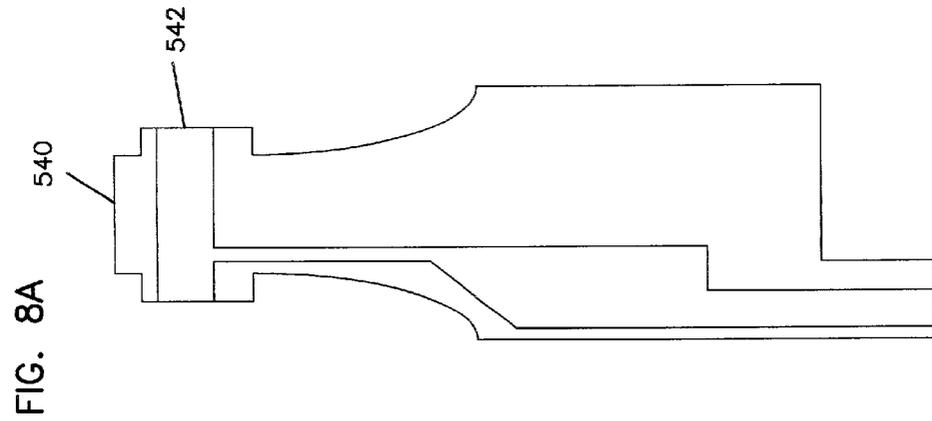
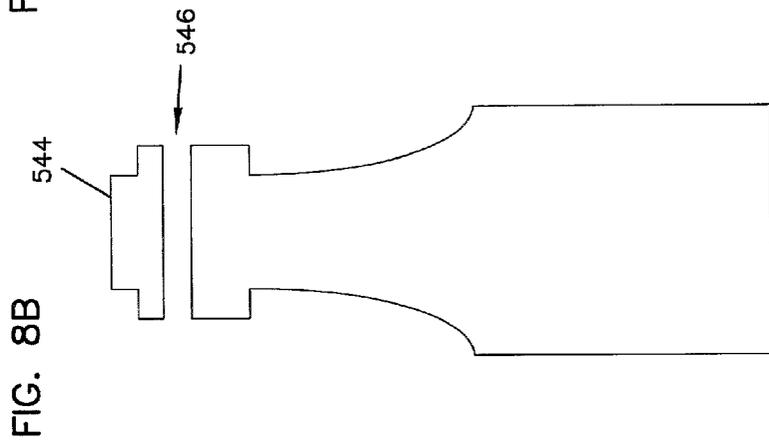
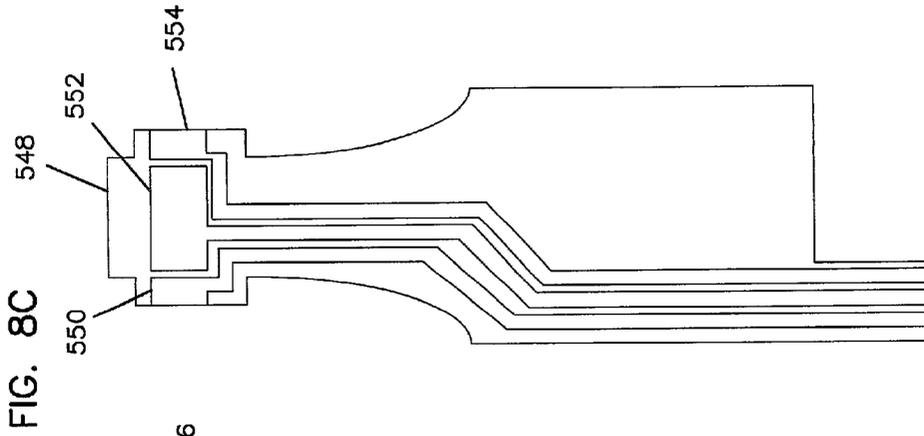


FIG. 9A

FIG. 9B

FIG. 9C

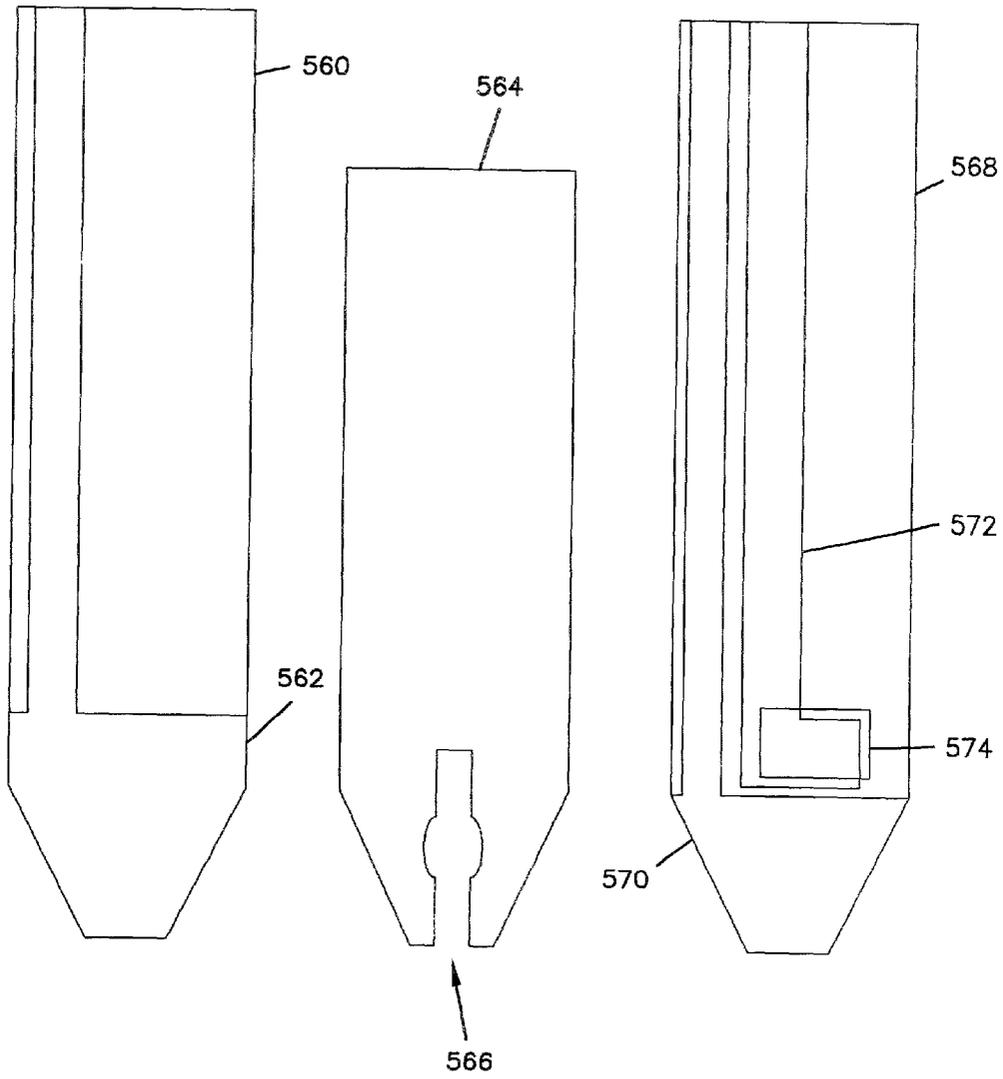


FIG. 10A

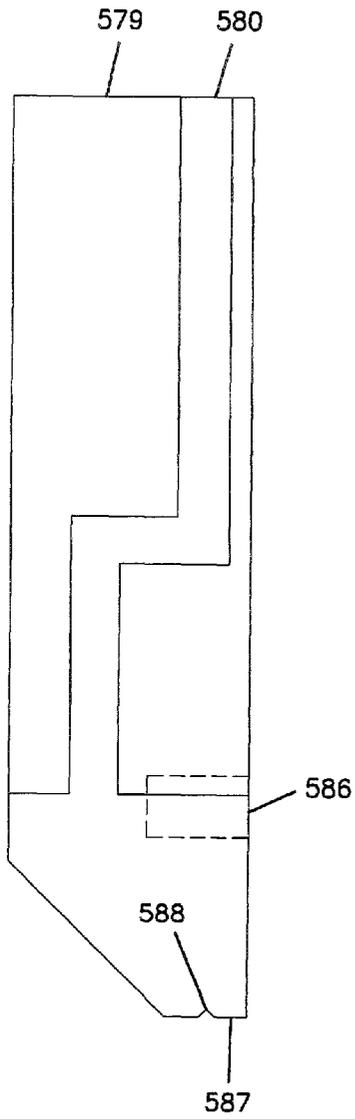


FIG. 10B

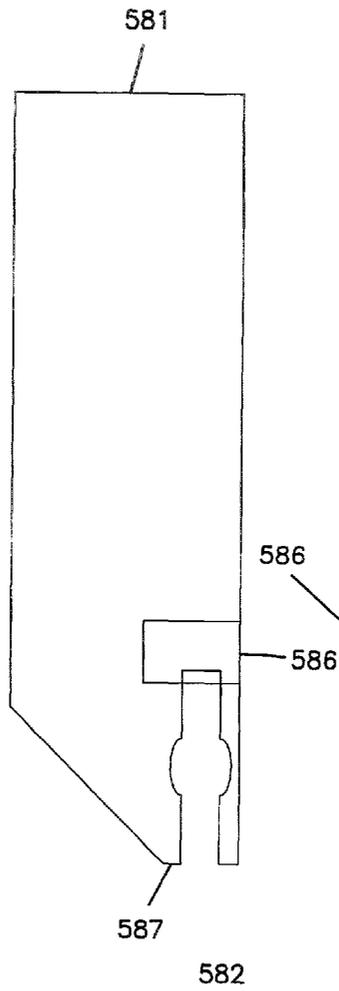


FIG. 10C

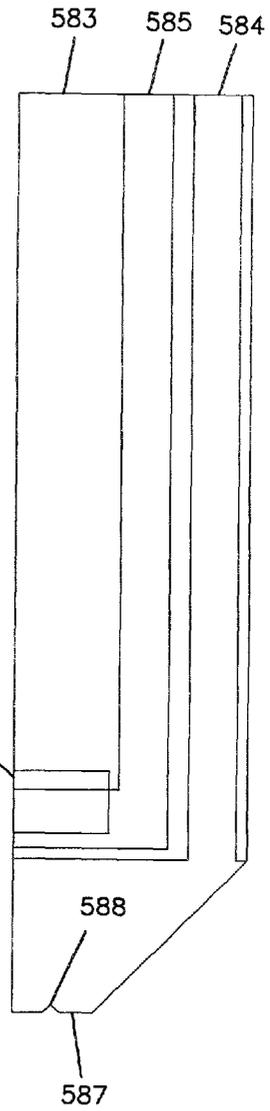


FIG. 11A

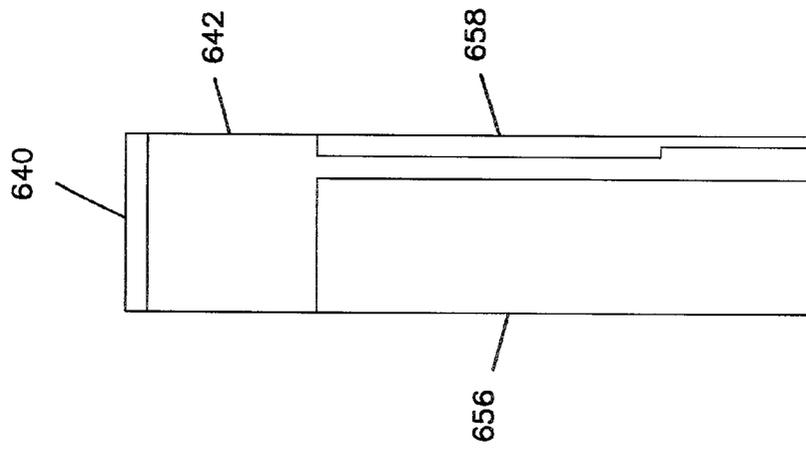


FIG. 11B

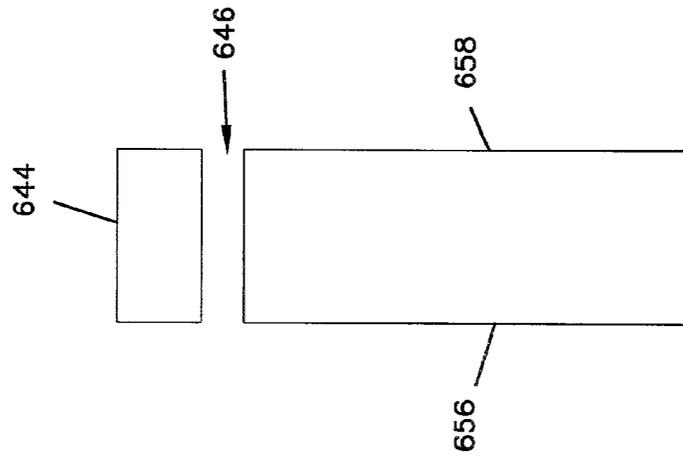


FIG. 11C

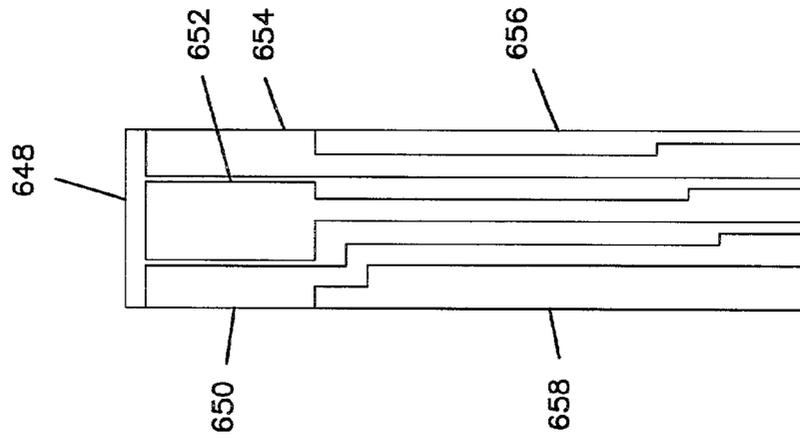


FIG. 12C

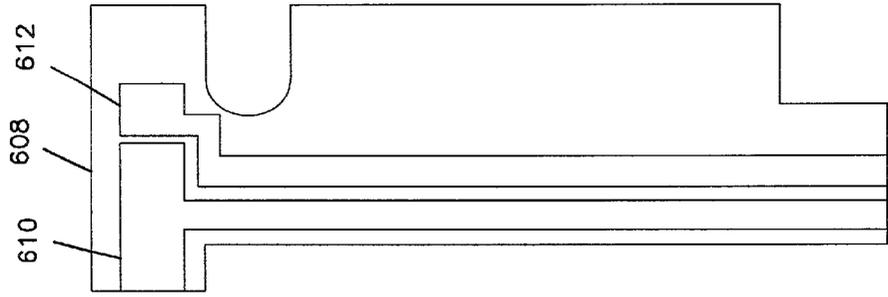


FIG. 12B

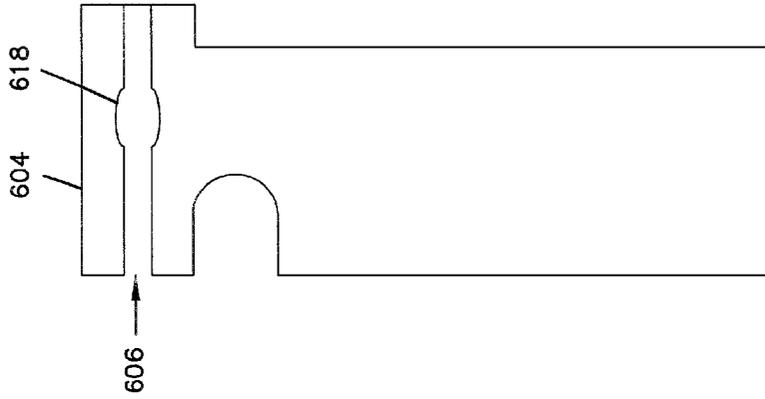


FIG. 12A

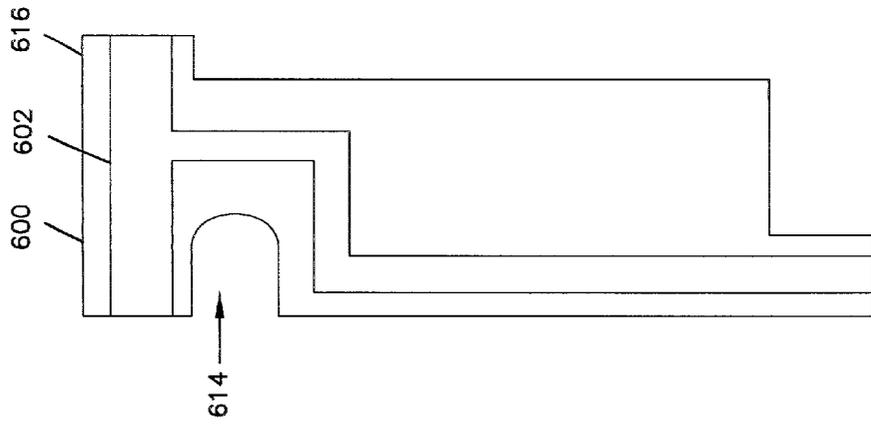


FIG. 13A

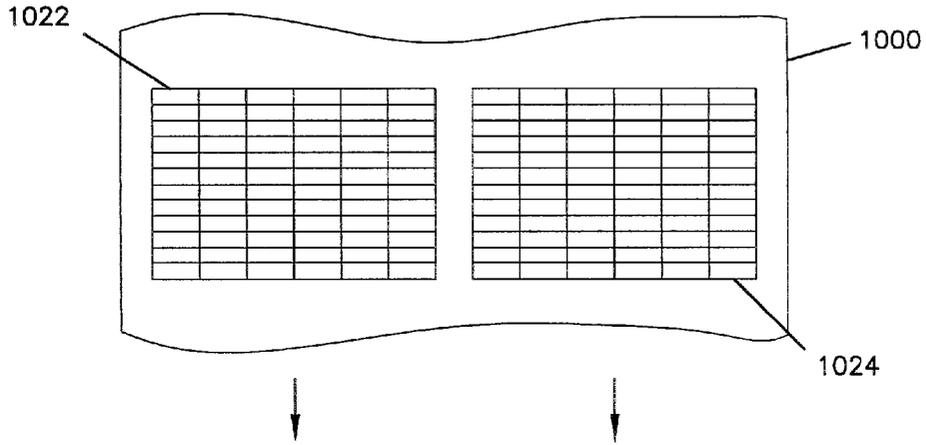


FIG. 13B

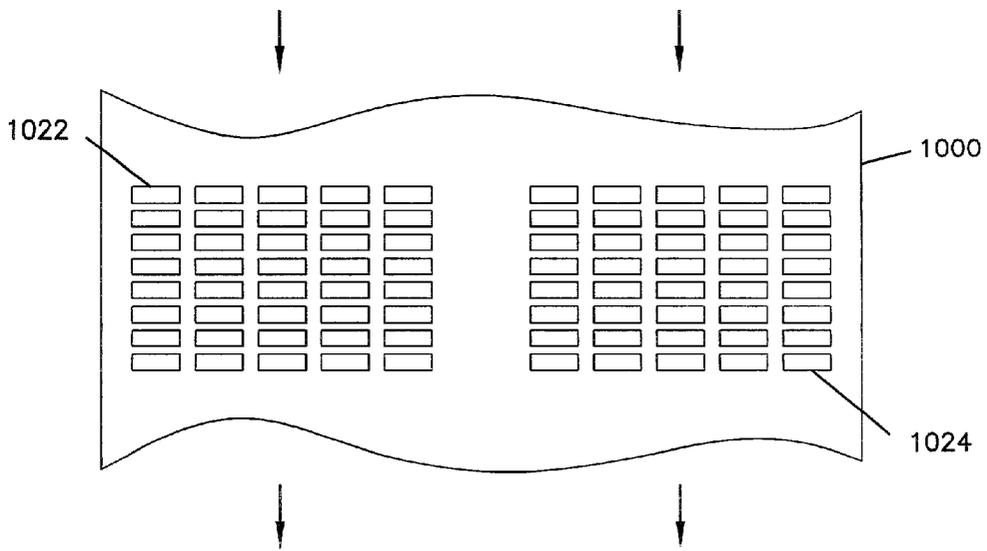


FIG. 14A

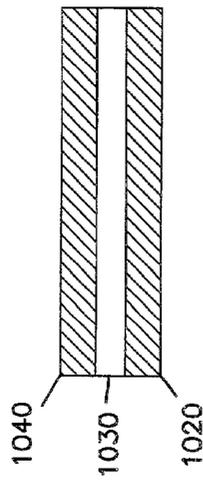


FIG. 14B

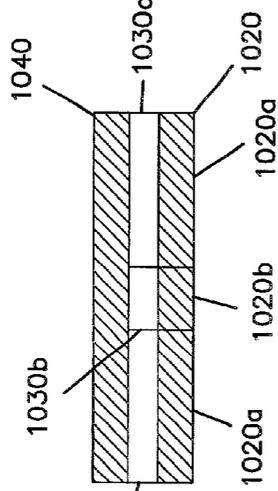


FIG. 14C

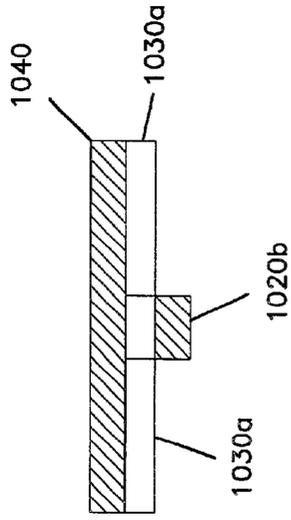


FIG. 14D

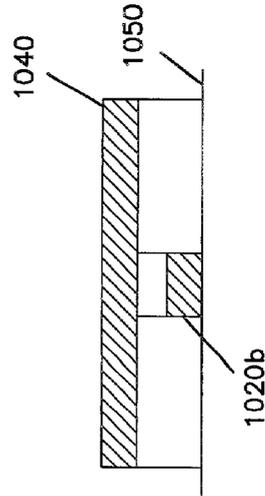


FIG. 14E

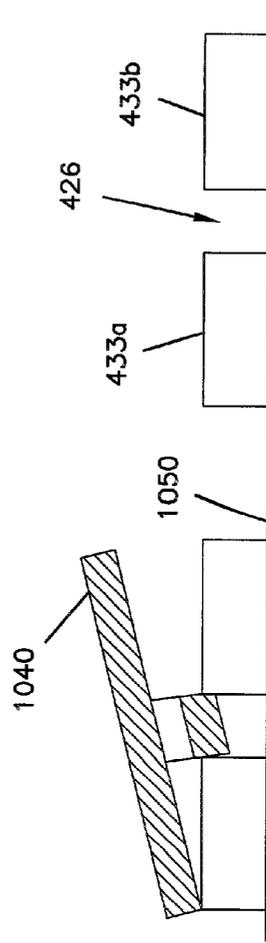
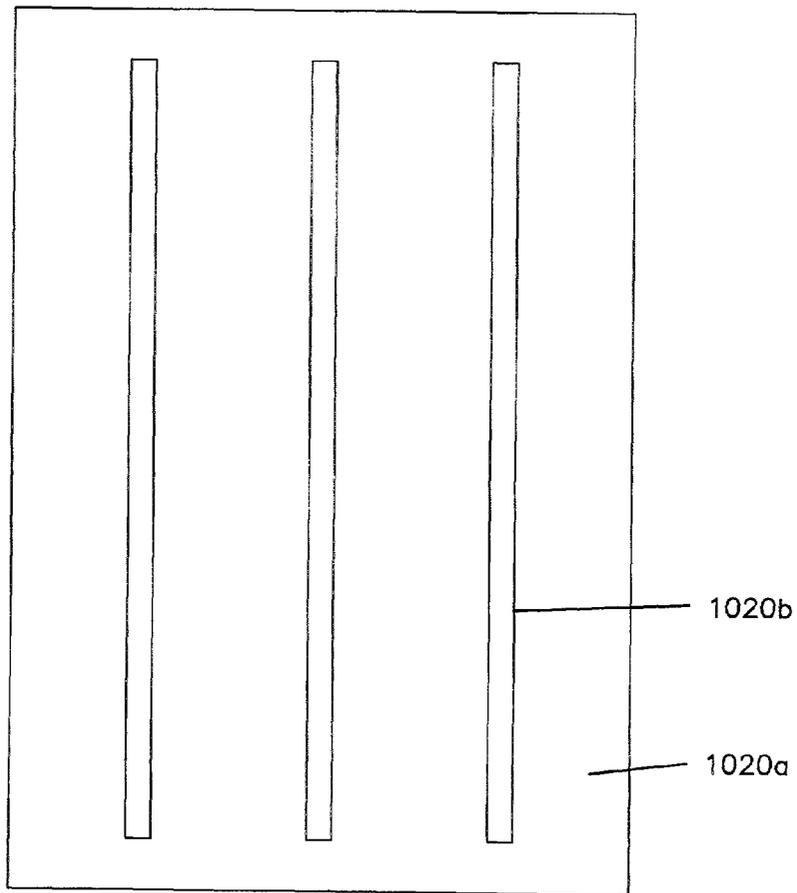
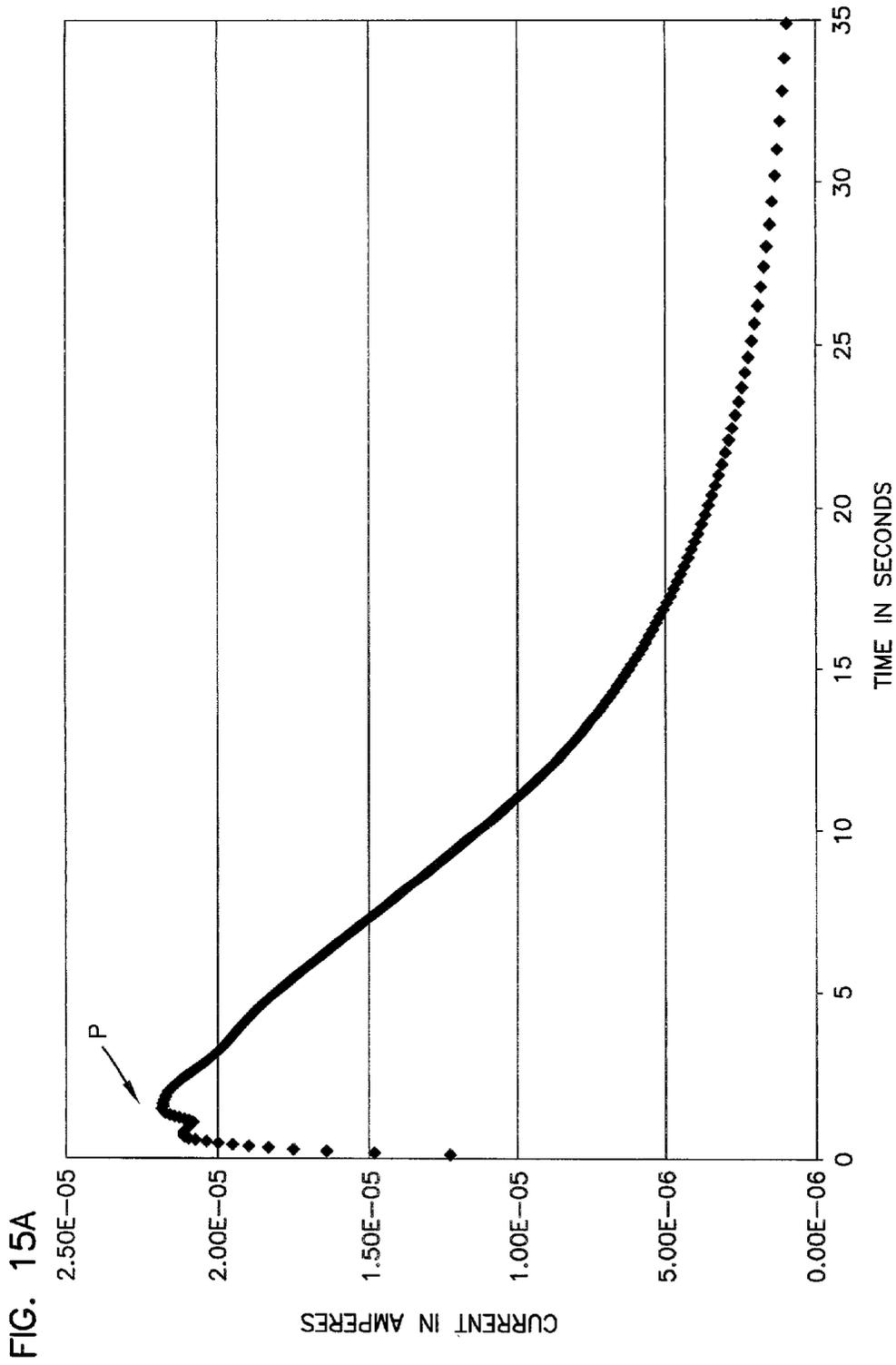


FIG. 14F

FIG. 14G





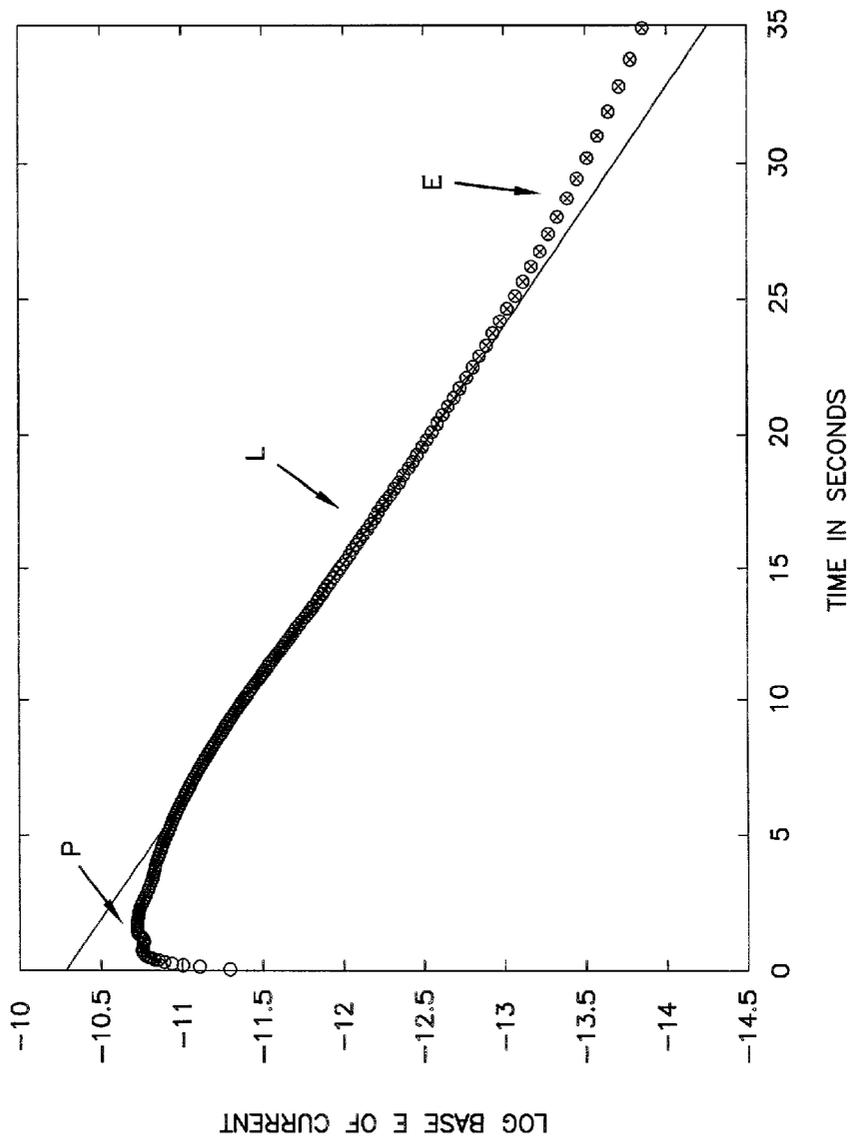


FIG. 15B

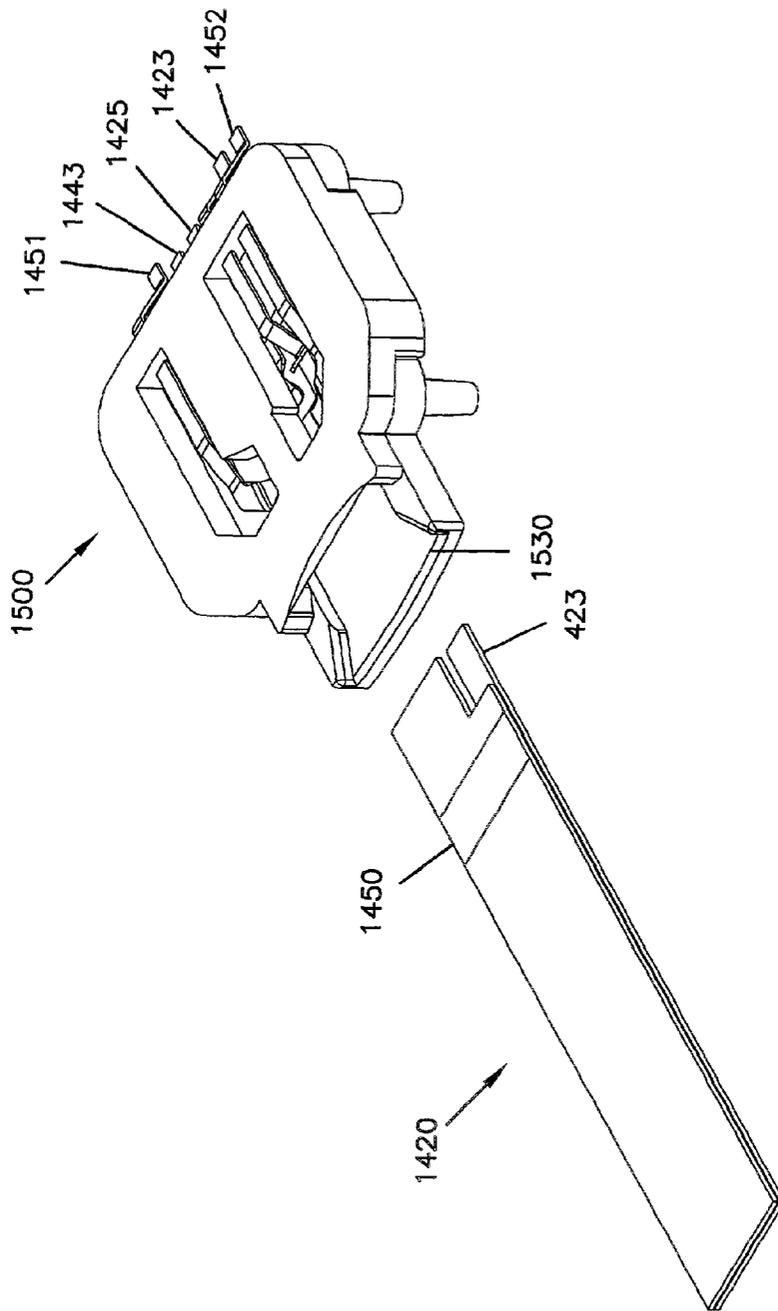


FIG. 16A

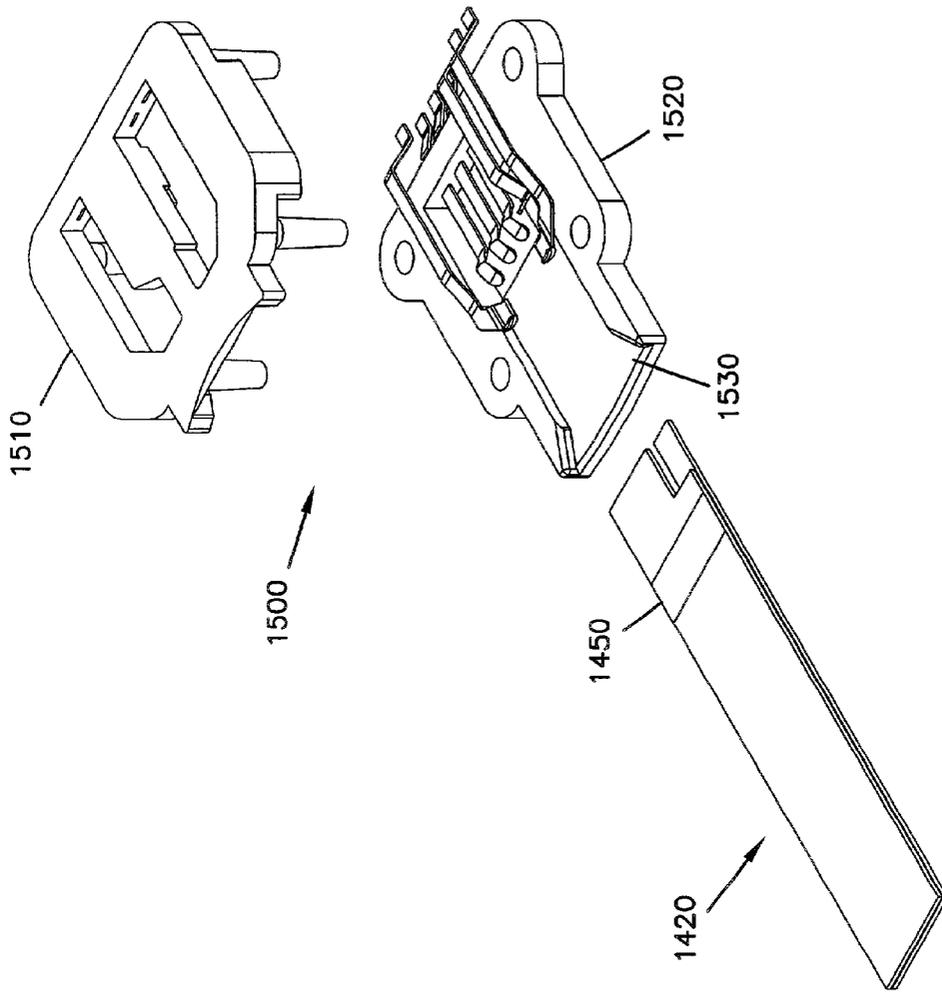


FIG. 16B

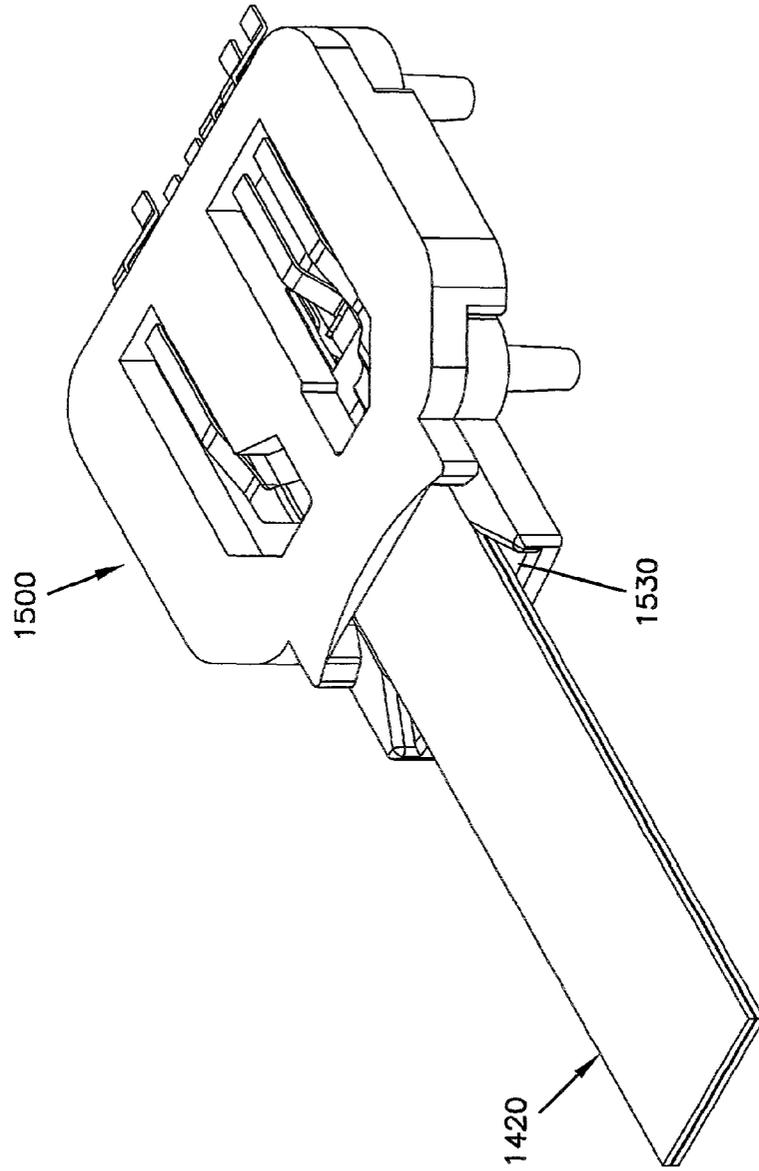
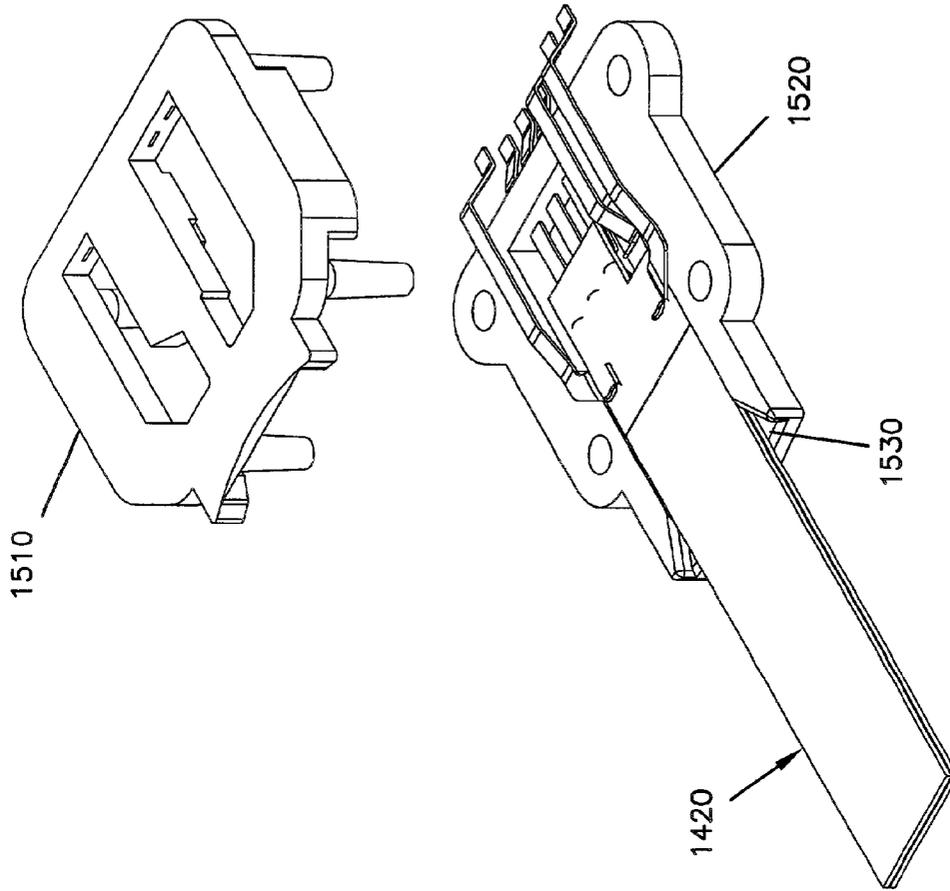


FIG. 17A

FIG. 17B



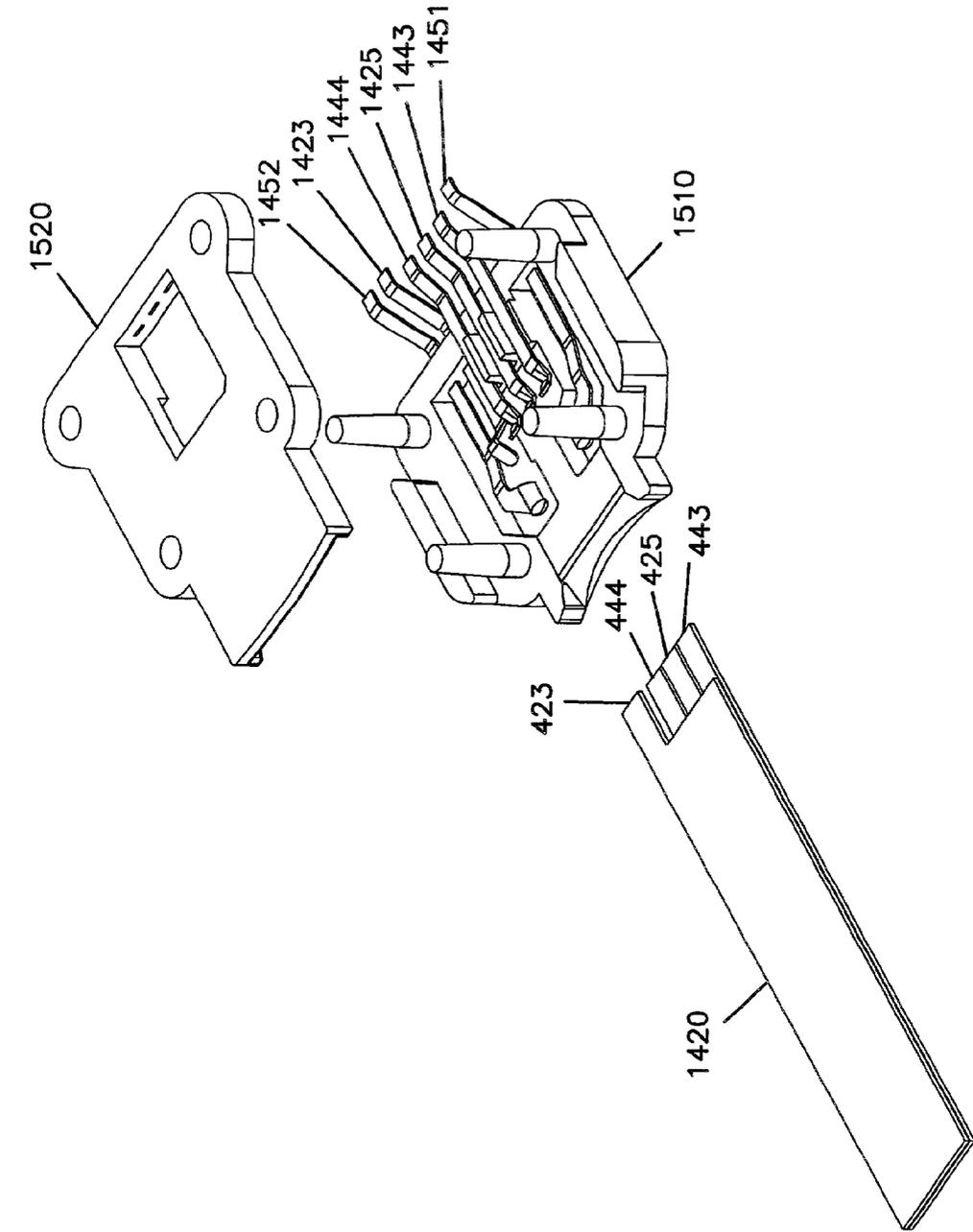


FIG. 18A

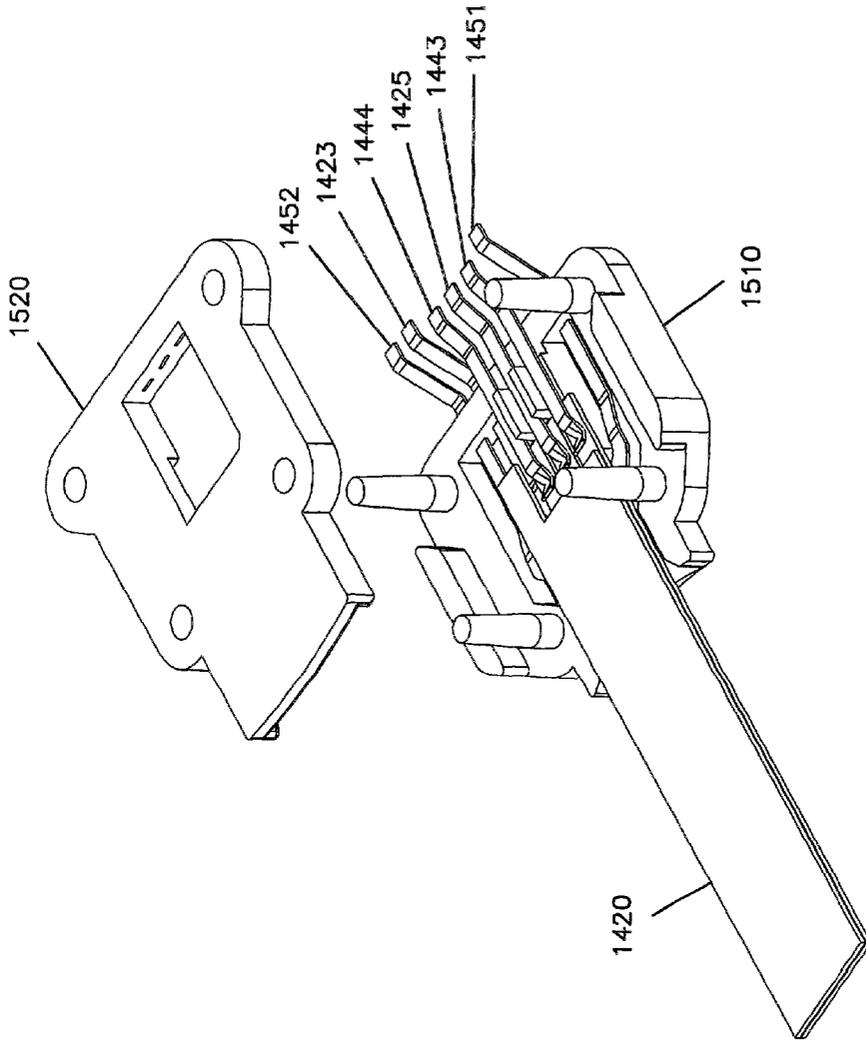


FIG. 18B

FIG. 19A

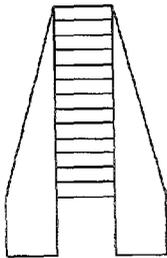


FIG. 19B

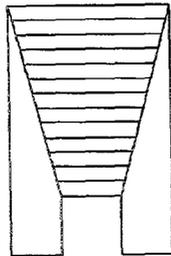


FIG. 19C

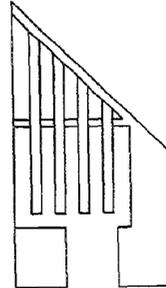


FIG. 19D

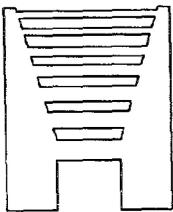


FIG. 19E

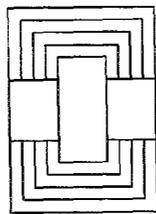


FIG. 19F

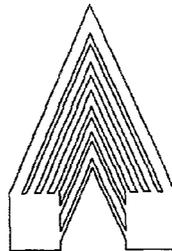


FIG. 19G



FIG. 19H



FIG. 19I

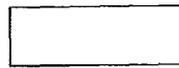


FIG. 19J



FIG. 19K

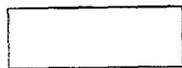


FIG. 19L



FIG. 20

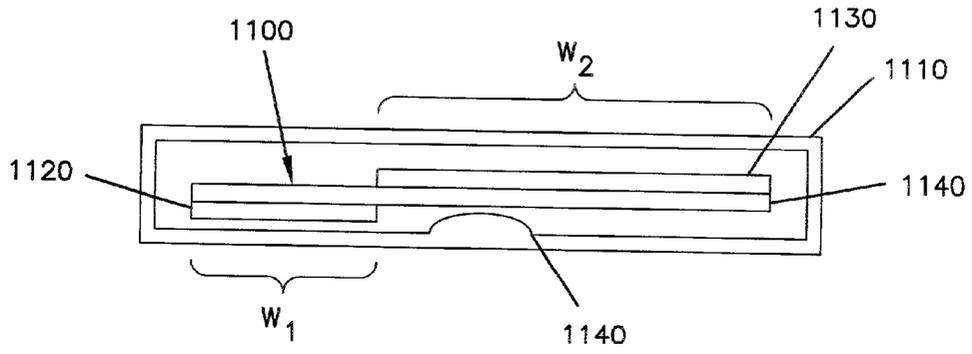
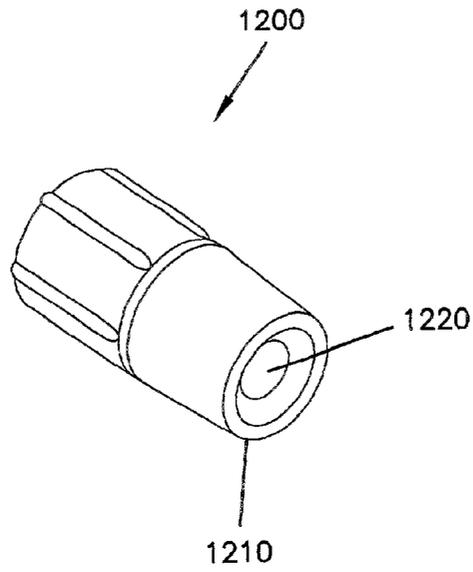


FIG. 21



SMALL VOLUME IN VITRO ANALYTE SENSOR AND METHODS

FIELD OF THE INVENTION

[0001] This invention relates to analytical sensors for the detection of bioanalytes in a small volume sample, and methods of making and using the sensors.

BACKGROUND

[0002] Analytical sensors are useful in chemistry and medicine to determine the presence and concentration of a biological analyte. Such sensors are needed, for example, to monitor glucose in diabetic patients and lactate during critical care events.

[0003] Currently available technology measures bioanalytes in relatively large sample volumes, e.g., generally requiring 3 microliters or more of blood or other biological fluid. These fluid samples are obtained from a patient, for example, using a needle and syringe, or by lancing a portion of the skin such as the fingertip and "milking" the area to obtain a useful sample volume. These procedures are inconvenient for the patient, and often painful, particularly when frequent samples are required. Less painful methods for obtaining a sample are known such as lancing the arm or thigh, which have a lower nerve ending density. However, lancing the body in the preferred regions typically produces submicroliter samples of blood, because these regions are not heavily supplied with near-surface capillary vessels.

[0004] It would therefore be desirable and very useful to develop a relatively painless, easy to use blood analyte sensor, capable of performing an accurate and sensitive analysis of the concentration of analytes in a small volume of sample.

[0005] It would also be desirable to develop methods for manufacturing small volume electrochemical sensors capable of decreasing the errors that arise from the size of the sensor and the sample.

SUMMARY OF THE INVENTION

[0006] The sensors of the present invention provide a method for the detection and quantification of an analyte in submicroliter samples. In general, the invention includes a method and sensor for analysis of an analyte in a small volume of sample by, for example, coulometry, amperometry and/or potentiometry. A sensor of the invention preferably utilizes a non-leachable or diffusible redox mediator. The sensor also includes a sample chamber to hold the sample in electrolytic contact with the working electrode. In many instances, the sensor also contains a non-leachable or diffusible second electron transfer agent.

[0007] In a preferred embodiment, the working electrode faces a counter electrode, forming a measurement zone within the sample chamber, between the two electrodes, that is sized to contain no more than about 1 μL of sample, preferably no more than about 0.5 μL , more preferably no more than about 0.32 μL , still more preferably no more than about 0.25 μL , and most preferably no more than about 0.1 μL of sample.

[0008] In one embodiment of the invention, a sensor, configured for insertion into an electronic meter, is provided

with a working electrode and a counter electrode, and a conductive insertion monitor which provides electrical contact with the electronic meter if the sensor is properly inserted into the meter. The conductive insertion monitor is configured and arranged to close an electrical circuit when the sensor is properly inserted into the electronic connector.

[0009] In another embodiment of the invention, a sensor is provided with a plurality of contacts, each contact having a contact pad, which is a region for connection with an electronic meter. The plurality of contacts and contact pads are disposed on a substrate having a length and a width, and each contact pad has a contact pad width taken parallel to the width of the substrate. The sum of the contact pad widths is greater than the width of the substrate. In a preferred embodiment, six electrical connections are made with six contact pads on the sensor but in a width that is approximately the width of four contact pads. For example, a working electrode, three counter electrodes (e.g., one counter electrode and two indicator electrodes), and two insertion trace connections each have a contact pad; connection can be made to each of these six contact pads in the same width of the contact pads of the working electrode and three counter electrodes.

[0010] The present invention also includes an electrical connector, for providing electrical contact between a sensor and an electrical meter or other device. The electrical connector has a plurality of contact structures, each which has a proximal contact end for electrical connection to a sensor contact, and a distal end for electrical connection to the electrical device. In one embodiment, a plurality of first contact structures extend longitudinally parallel from the distal to the proximal end. Additionally, one or more second contact structures extend longitudinally next to the first contact structures, from the distal end past the proximal end of the first contact structures, and angle toward a longitudinal center line of the connector. Contact to the sensor is then made via the proximal contact ends.

[0011] Preferably, the electrical connector has at least two second contact structures extending longitudinally past the proximal end of the first contact structures and angling toward the longitudinal center line of the connector. After the angled or bent portion, the proximal contact ends of the second contact structures of one embodiment make electrical contact with a single conductive surface of a sensor, such as a conductive insertion monitor. In another aspect, the first contact structures can be configured and arranged to contact one or more working and/or counter electrodes of a sensor, and the second contact structures are configured and arranged to contact one or more conductive insertion monitors.

[0012] The sensors of the present invention can be configured for side-filling or tip-filling. In addition, in some embodiments, the sensor may be part of an integrated sample acquisition and analyte measurement device. The integrated sample acquisition and analyte measurement device can include the sensor and a skin piercing member, so that the device can be used to pierce the skin of a user to cause flow of a fluid sample, such as blood, that can then be collected by the sensor. In at least some embodiments, the fluid sample can be collected without moving the integrated sample acquisition and analyte measurement device.

[0013] In one embodiment, the sensor is connected with an electrical device, to provide a processor coupled to the

sensor. The processor is configured and arranged to determine, during electrolysis of a sample in the sample chamber, a series of current values. The processor determines a peak current value from the series of current values. After the current values decrease below a threshold fraction of the peak current values, slope values are determined from the current values and represent a linear function of the logarithm of current values over time. The processor determines, from the slope values, an extrapolation slope. From the extrapolated slope and the measured current values, the processor determines an amount of charge needed to electrolyze the sample and, from that amount of charge, the concentration of the analyte in the sample.

[0014] One method of forming a sensor, as described above, includes forming at least one working electrode on a first substrate and forming at least one counter or counter/reference electrode on a second substrate. A spacer layer is disposed on either the first or second substrates. The spacer layer defines a chamber into which a sample can be drawn and held when the sensor is completed. A redox mediator and/or second electron transfer agent can be disposed on the first or second substrate in a region that will be exposed within the sample chamber when the sensor is completed. The first and second substrates are then brought together and spaced apart by the spacer layer with the sample chamber providing access to the at least one working electrode and the at least one counter or counter/reference electrode. In some embodiments, the first and second substrates are portions of a single sheet or continuous web of material. The invention includes particularly efficient and reliable methods for the manufacture of these sensors.

[0015] One such efficient and reliable method includes providing an adhesive having first and second surfaces covered with first and second release liners and then making detailed cuts through the first release liner and the adhesive but not through the second release liner. These cuts define one or more sample chamber regions. A portion of the first release liner is removed to expose a portion of the first adhesive surface, which leaves a remaining portion of the first release liner over the sample chamber regions. This exposed first adhesive surface is applied to a first substrate having one or more conductive traces disposed thereon. The second release liner is removed together with the adhesive and the first release liner of the sample chamber regions in order to expose the second adhesive surface. The second adhesive surface is then applied to a second substrate having one or more conductive traces disposed thereon. This method forms a sensor having a sample chamber corresponding to one of the sample chamber regions.

[0016] These and various other features which characterize the invention are pointed out with particularity in the attached claims. For a better understanding of the invention, its advantages, and objectives obtained by its use, reference should be made to the drawings and to the accompanying description, in which there is illustrated and described preferred embodiments of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

[0017] Referring now to the drawings, wherein like reference numerals and letters indicate corresponding structure throughout the several views:

[0018] FIG. 1 is a schematic view of a first embodiment of an electrochemical sensor in accordance with the present

invention having a working electrode and a counter electrode facing each other and having a sample chamber;

[0019] FIG. 2 is an exploded view of the sensor shown in FIG. 1, the layers illustrated individually;

[0020] FIG. 3 is a schematic view of a second embodiment of an electrochemical sensor in accordance with the principles of the present invention having a working electrode and a counter electrode facing each other and having an extended sample chamber;

[0021] FIG. 4 is a top view of a third embodiment of an electrochemical sensor in accordance with the principles of the present invention, this sensor includes multiple working electrodes;

[0022] FIG. 5A illustrates a top view of a first substrate with a working electrode for use in a fourth embodiment of a sensor according to the invention;

[0023] FIG. 5B illustrates a bottom view of a second substrate (inverted with respect to FIG. 5A) with counter electrodes placement over and opposite the substrate of FIG. 5A;

[0024] FIG. 5C illustrates a top view of a spacer for placement between the first substrate of FIG. 5A and the second substrate of FIG. 5B;

[0025] FIG. 6A illustrates a top view of a first film with a working electrode for use in a fifth embodiment of a sensor according to the invention;

[0026] FIG. 6B illustrates a top view of a spacer for placement on the first film of FIG. 6A;

[0027] FIG. 6C illustrates a bottom view of a second film (inverted with respect to FIGS. 6A and 6B) with counter electrodes placement over the spacer of FIG. 6B and first film of FIG. 6A;

[0028] FIG. 7A illustrates a top view of a first film with a working electrode for use in a sixth embodiment of a sensor according to the invention;

[0029] FIG. 7B illustrates a top view of a spacer for placement on the first film of FIG. 7A;

[0030] FIG. 7C illustrates a bottom view of a second film (inverted with respect to FIGS. 7A and 7B) with counter electrodes placement over the spacer of FIG. 7B and first film of FIG. 7A;

[0031] FIG. 8A illustrates a top view of a first film with a working electrode for use in a seventh embodiment of a sensor according to the invention;

[0032] FIG. 8B illustrates a top view of a spacer for placement on the first film of FIG. 8A;

[0033] FIG. 8C illustrates a bottom view of a second film (inverted with respect to FIGS. 8A and 8B) with counter electrodes placement over the spacer of FIG. 8B and first film of FIG. 8A;

[0034] FIG. 9A illustrates a top view of a first film with a working electrode for use in an eighth embodiment of a sensor according to the invention;

[0035] FIG. 9B illustrates a top view of a spacer for placement on the first film of FIG. 9A;

[0036] FIG. 9C illustrates a bottom view of a second film (inverted with respect to FIGS. 9A and 9B) with counter electrodes placement over the spacer of FIG. 9B and first film of FIG. 9A;

[0037] FIG. 10A illustrates a top view of a first film with a working electrode for use in a ninth embodiment of a sensor according to the invention;

[0038] FIG. 10B illustrates a top view of a spacer for placement on the first film of FIG. 10A;

[0039] FIG. 10C illustrates a bottom view of a second film (inverted with respect to FIGS. 10A and 10B) with counter electrodes placement over the spacer of FIG. 10B and first film of FIG. 10A;

[0040] FIG. 11A illustrates a top view of a first film with a working electrode for use in a tenth embodiment of a sensor according to the invention;

[0041] FIG. 11B illustrates a top view of a spacer for placement on the first film of FIG. 11A;

[0042] FIG. 11C illustrates a bottom view of a second film (inverted with respect to FIGS. 11A and 11B) with counter electrodes placement over the spacer of FIG. 11B and first film of FIG. 11A;

[0043] FIG. 12A illustrates a top view of a first film with a working electrode for use in an eleventh embodiment of a sensor according to the invention;

[0044] FIG. 12B illustrates a top view of a spacer for placement on the first film of FIG. 12A;

[0045] FIG. 12C illustrates a bottom view of a second film (inverted with respect to FIGS. 12A and 12B) with counter electrodes placement over the spacer of FIG. 12B and first film of FIG. 12A;

[0046] FIG. 13A illustrates a top view of one embodiment of a sheet of sensor components, according to the invention;

[0047] FIG. 13B illustrates a top view of another embodiment of a sheet of sensor components, according to the invention;

[0048] FIGS. 14A through 14F illustrate cross sectional views of a sequential process of providing a sample chamber in a spacer layer;

[0049] FIG. 14G is a bottom view of FIG. 14B;

[0050] FIG. 15A is a graphical representation of the current over time used to calculate the concentration of an analyte;

[0051] FIG. 15B is a graphical representation of the log of current over time used to calculate the concentration of an analyte;

[0052] FIG. 16A is a top perspective view of a sensor positioned for insertion within an electrical connector device in accordance with the present invention;

[0053] FIG. 16B is an exploded view of the electrical connector device of FIG. 16A;

[0054] FIG. 17A is a top perspective view of a sensor fully positioned within the electrical connector device of FIG. 16A;

[0055] FIG. 17B is an exploded view of the electrical connector device of FIG. 17A;

[0056] FIG. 18A is a bottom perspective view of the electrical connector device of FIGS. 16A and 16B;

[0057] FIG. 18B is a bottom perspective view of the electrical connector device of FIGS. 17A and 17B;

[0058] FIGS. 19A through 19L illustrate examples of suitable conductive pathways between contact pads;

[0059] FIG. 20 illustrates a cross-sectional view looking from inside a connector to a sensor of the invention disposed within the connector; and

[0060] FIG. 21 illustrates a perspective view of a ring for use with a lancet device.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

[0061] As used herein, the following definitions define the stated term:

[0062] An "air-oxidizable mediator" is a redox mediator that is oxidized by air, preferably so that at least 90% of the mediator is in an oxidized state upon storage in air either as a solid or as a liquid during a period of time, for example, one month or less, and, preferably, one week or less, and, more preferably, one day or less.

[0063] "Amperometry" includes steady-state amperometry, chronoamperometry, and Cottrell-type measurements.

[0064] A "biological fluid" is any body fluid in which the analyte can be measured, for example, blood (which includes whole blood and its cell-free components, such as, plasma and serum), interstitial fluid, dermal fluid, sweat, tears, urine and saliva.

[0065] "Coulometry" is the determination of charge passed or projected to pass during complete or nearly complete electrolysis of the analyte, either directly on the electrode or through one or more electron transfer agents. The charge is determined by measurement of charge passed during partial or nearly complete electrolysis of the analyte or, more often, by multiple measurements during the electrolysis of a decaying current and elapsed time. The decaying current results from the decline in the concentration of the electrolyzed species caused by the electrolysis.

[0066] A "counter electrode" refers to one or more electrodes paired with the working electrode, through which passes an electrochemical current equal in magnitude and opposite in sign to the current passed through the working electrode. The term "counter electrode" is meant to include counter electrodes which also function as reference electrodes (i.e. a counter/reference electrode) unless the description provides that a "counter electrode" excludes a reference or counter/reference electrode.

[0067] An "effective diffusion coefficient" is the diffusion coefficient characterizing transport of a substance, for example, an analyte, an enzyme, or a redox mediator, in the volume between the electrodes of the electrochemical cell. In at least some instances, the cell volume may be occupied by more than one medium (e.g., the sample fluid and a polymer film). Diffusion of a substance through each medium may occur at a different rate. The effective diffusion

coefficient corresponds to a diffusion rate through this multiple-media volume and is typically different than the diffusion coefficient for the substance in a cell filled solely with sample fluid.

[0068] An “electrochemical sensor” is a device configured to detect the presence of and/or measure the concentration of an analyte via electrochemical oxidation and reduction reactions. These reactions are transduced to an electrical signal that can be correlated to an amount or concentration of analyte.

[0069] “Electrolysis” is the electrooxidation or electroreduction of a compound either directly at an electrode or via one or more electron transfer agents (e.g., redox mediators and/or enzymes).

[0070] The term “facing electrodes” refers to a configuration of the working and counter electrodes in which the working surface of the working electrode is disposed in approximate opposition to a surface of the counter electrode. In at least some instances, the distance between the working and counter electrodes is less than the width of the working surface of the working electrode.

[0071] A compound is “immobilized” on a surface when it is entrapped on or chemically bound to the surface.

[0072] An “indicator electrode” is an electrode that detects partial or complete filling of a sample chamber and/or measurement zone.

[0073] A “layer” is one or more layers.

[0074] The “measurement zone” is defined herein as a region of the sample chamber sized to contain only that portion of the sample that is to be interrogated during an analyte assay.

[0075] A “non-diffusible,” “non-leachable,” or “non-releasable” compound is a compound which does not substantially diffuse away from the working surface of the working electrode for the duration of the analyte assay.

[0076] The “potential of the counter/reference electrode” is the half cell potential of the reference electrode or counter/reference electrode of the cell when the solution in the cell is 0.1 M NaCl solution at pH7.

[0077] A “redox mediator” is an electron transfer agent for carrying electrons between the analyte and the working electrode, either directly, or via a second electron transfer agent.

[0078] A “reference electrode” includes a reference electrode that also functions as a counter electrode (i.e., a counter/reference electrode) unless the description provides that a “reference electrode” excludes a counter/reference electrode.

[0079] A “second electron transfer agent” is a molecule that carries electrons between a redox mediator and the analyte.

[0080] A “surface in the sample chamber” is a surface of a working electrode, counter electrode, counter/reference electrode, reference electrode, indicator electrode, a spacer, or any other surface bounding the sample chamber.

[0081] A “working electrode” is an electrode at which analyte is electrooxidized or electroreduced with or without the agency of a redox mediator.

[0082] A “working surface” is the portion of a working electrode that is covered with non-leachable redox mediator and exposed to the sample, or, if the redox mediator is diffusible, a “working surface” is the portion of the working electrode that is exposed to the sample.

[0083] The small volume, in vitro analyte sensors of the present invention are designed to measure the concentration of an analyte in a portion of a sample having a volume no more than about 1 μL , preferably no more than about 0.5 μL , more preferably no more than about 0.32 μL , still more preferably no more than about 0.25 μL , and most preferably no more than about 0.1 μL of sample.

[0084] The analyte of interest is typically provided in a solution or biological fluid, such as blood or serum.

[0085] Referring to the Drawings in general, and in particular FIGS. 1 and 2, a small volume, in vitro electrochemical sensor 20 of the invention generally includes a working electrode 22 on a first substrate 32, a counter (or counter/reference) electrode 24 on a second substrate 34, and a sample chamber 26. The sample chamber 26 is configured so that when a sample is provided in the chamber, the sample is in electrolytic contact with both the working electrode 22, the counter electrode 24 and any reference electrodes or indicator electrodes that may be present. This allows electrical current to flow between the electrodes to effect the electrolysis (electrooxidation or electroreduction) of the analyte. A spacer 33 is positioned between first substrate 32 and second substrate 34 to provide a spacing between electrodes 22, 24 and to provide sample chamber 26 in which the sample to be evaluated is housed.

[0086] Working Electrode

[0087] The working electrode 22 may be formed from a molded carbon fiber composite or it may include an inert non-conducting base material, such as polyester, upon which a suitable conducting layer is deposited. The conducting layer typically has relatively low electrical resistance and is typically electrochemically inert over the potential range of the sensor during operation. Suitable conducting layers include gold, carbon, platinum, ruthenium dioxide, palladium, and conductive epoxies, such as, for example, ECCO-COAT CT5079-3 Carbon-Filled Conductive Epoxy Coating (available from W. R. Grace Company, Woburn, Mass.), as well as other non-corroding materials known to those skilled in the art. The electrode (e.g., the conducting layer) is deposited on the surface of the inert material by methods such as vapor deposition or printing. Preferably, the electrode is printed onto the base material.

[0088] The inert non-conducting base material, is also referred to as a substrate, base, or the like. This base material is typically an electrically non-conducting material, e.g., any insulating material, that is not capable of carrying electric charge or current. Examples of materials usable as the base material for sensors of the present invention include polyesters, polyethylene (both high density and low density), polyethylene terephthalate, polycarbonate, vinyls, and the like. The base material can be treated with a primer or other such coating to improve the adhesion of the electrodes thereon.

[0089] A tab 23' can be provided on the end of the working electrode 22 for easy connection of the electrode to external electronics (not shown) such as a voltage source or current

measuring equipment. Contact pad **23**, which is connected to working electrode **22**, such as an extension from the working electrode, can be positioned on tab **23'**.

[**0090**] To prevent electrochemical reactions from occurring on portions of the working electrode not coated by the mediator, when a non-leachable mediator is used, a dielectric or other insulating material can be deposited on the electrode over, under, or surrounding the region with the redox mediator. Suitable dielectric materials include waxes and non-conducting organic polymers, such as polyethylene. The dielectric material can also cover a portion of the redox mediator on the electrode. The covered portion of the redox mediator will not contact the sample, and, therefore, will not be a part of the electrode's working surface.

[**0091**] Sensing Chemistry

[**0092**] In addition to the working electrode **22**, sensing chemistry materials are provided in the sample chamber **26** for the analysis of the analyte. This sensing chemistry preferably includes a redox mediator and a second electron transfer mediator, although in some instances, one or the other may be used alone. The redox mediator and second electron transfer agent can be independently diffusible or non-leachable (i.e., non-diffusible) such that either or both may be diffusible or non-leachable. Placement of sensor chemistry components can depend on whether they are diffusible or non-leachable. For example, non-leachable and/or diffusible component(s) typically form a sensing layer on the working electrode. Alternatively, one or more diffusible components can be disposed on any surface in the sample chamber prior to the introduction of the sample. As another example, one or more diffusible component(s) are placed in the sample prior to introduction of the sample into the sensor.

[**0093**] If the redox mediator is non-leachable, then the non-leachable redox mediator is typically disposed on the working electrode **22** as a sensing layer. In an embodiment having a redox mediator and a second electron transfer agent, if the redox mediator and second electron transfer agent are both non-leachable, then both of the non-leachable components are disposed on the working electrode **22** as a sensing layer.

[**0094**] If, for example, the second electron transfer agent is diffusible and the redox mediator is non-leachable, then at least the redox mediator is disposed on the working electrode **22** as a sensing layer. The diffusible second electron transfer agent need not be disposed on a sensing layer of the working electrode, but can be disposed on any surface of the sample chamber, including within the redox mediator sensing layer, or can be placed in the sample. If the redox mediator is diffusible, then the redox mediator can be disposed on any surface of the sample chamber or can be placed in the sample. If both the redox mediator and second electron transfer agent are diffusible, then the diffusible components can be independently or jointly disposed on any surface of the sample chamber and/or placed in the sample (i.e., each diffusible component need not be disposed on the same surface of the sample chamber or placed in the sample).

[**0095**] The redox mediator, whether it is diffusible or non-leachable, mediates a current between the working electrode **22** and the analyte and enables the electrochemical

analysis of molecules which may not be suited for direct electrochemical reaction on an electrode. The mediator functions as an electron transfer agent between the electrode and the analyte.

[**0096**] Analytes that can be interrogated include, for example, glucose, acetyl choline, amylase, bilirubin, cholesterol, chorionic gonadotropin, creatine kinase (e.g., CK-MB), creatine, DNA, fructosamine, glucose, glutamine, growth hormones, hormones, ketones, lactate, peroxide, prostate-specific antigen, prothrombin, RNA, thyroid stimulating hormone, and troponin. The concentration of analytes such as drugs or medication, such as, for example, antibiotics (e.g., gentamicin, vancomycin, and the like), digitoxin, digoxin, drugs of abuse, theophylline, and warfarin, can also be determined. Assays suitable for determining the concentration of DNA and/or RNA are disclosed in U.S. patent applications Ser. Nos. 09/138,888 and 09/145,776 and described in PCT Application PCT/US99/14460, all which are incorporated herein by reference.

Redox Mediators

[**0097**] Although any organic or organometallic redox species can be used as a redox mediator, one type of suitable redox mediator is a transition metal compound or complex. Examples of suitable transition metal compounds or complexes include osmium, ruthenium, iron, and cobalt compounds or complexes. In these complexes, the transition metal is coordinatively bound to one or more ligands. The ligands are typically mono-, di-, tri-, or tetradentate. The most preferred ligands are heterocyclic nitrogen compounds, such as, for example, pyridine and/or imidazole derivatives. Multidentate ligands may include multiple pyridine and/or imidazole rings. Alternatively, metallocene derivatives, such as, for example, ferrocene, can be used. An example of a suitable mediator is $[\text{Os}(4\text{-}(\text{N}\text{-}(6\text{-aminohexyl})\text{aminobipyridine})(1,1'\text{-dimethyl-2,2'-biimidazole})_2)]\text{Cl}_3$.

[**0098**] The redox mediators can be diffusible redox mediators or non-leachable redox mediators, such as non-leachable redox polymers. For additional information on redox mediators, see, for example, U.S. patent application Ser. No. 09/295,962, filed Apr. 21, 1999, and PCT published application WO 98/35225, both which are incorporated herein by reference.

Second Electron Transfer Agent

[**0099**] In a preferred embodiment of the invention, the sensor includes a redox mediator and a second electron transfer agent which is capable of transferring electrons to or from the redox mediator and the analyte. The second electron transfer agent can be diffusible or can be non-leachable (e.g., entrapped in or coordinatively, covalently, or ionically bound to a redox polymer). One example of a suitable second electron transfer agent is an enzyme which catalyzes a reaction of the analyte. For example, a glucose oxidase or glucose dehydrogenase, such as pyrroloquinoline quinone glucose dehydrogenase (PQQ), is used when the analyte is glucose. A lactate oxidase fills this role when the analyte is lactate. Other enzymes can be used for other analytes.

[**0100**] Counter Electrode

[**0101**] Counter electrode **24**, as illustrated in **FIGS. 1 and 2**, can be constructed in a manner similar to working

electrode **22**. Counter electrode **24** may also be a counter/reference electrode. Alternatively, a separate reference electrode may be provided in contact with the sample chamber. Suitable materials for the counter/reference or reference electrode include, for example, Ag/AgCl or Ag/AgBr printed on a non-conducting base material or silver chloride on a silver metal base. The same materials and methods may be used to make the counter electrode as are available for constructing the working electrode **22**, although different materials and methods may also be used. Preferably, the counter or counter/reference electrode is printed on an insulating base material. A tab **25'**, on which contact pad **25** is disposed, can be provided for making convenient connection to the external electronics (not shown), such as a coulometer, potentiostat, or other measuring device.

[0102] Optionally, a non-conductive filler material, such as a non-conductive ink, can be formed adjacent a counter electrode, or between multiple counter electrodes to provide a planar surface along the path of travel of the sample fluid in the sample chamber. The non-conductive filler material preferably creates a smooth surface to facilitate filling of the sample chamber by capillary action and/or for reducing the likelihood that air bubbles will become entrapped near the counter electrodes. This non-conductive material can be colored or colorless and may be formed on the substrate by printing or other techniques. The non-conductive material may be deposited prior to or subsequent to the formation of the counter electrode. In one embodiment, a non-conductive ink is used to fill in the area between multiple 12.3 micrometer (0.5 mil) thick counter electrodes. In another embodiment, a non-conductive ink is used to fill the area between multiple 6.4 micrometer (0.25 mil) thick counter electrodes. Generally, no filler ink is needed for thickness less about 6.4 micrometers, and in some sensor designs, no filler ink is needed for 6.4 micrometer thick counter electrodes.

[0103] Electrode Configuration

[0104] In one embodiment of the invention, working electrode **22** and counter electrode **24** are disposed opposite to and facing each other to form a facing electrode pair as depicted in **FIGS. 1 and 2**. In this preferred configuration, the sample chamber **26** is typically disposed between the two electrodes. For this facing electrode configuration, it is preferred that the electrodes are separated by a distance of no more than about 0.2 mm (e.g., at least one portion of the working electrode is separated from one portion of the counter electrode by no more than 200 μm), preferably no more than 100 μm , and most preferably no more than 50 μm .

[0105] The electrodes need not be directly opposing each other; they may be slightly offset. Furthermore, the two electrodes need not be the same size. Preferably, the working electrode **22** extends the width of sensor **20** and counter electrode **24** is a portion or all of that width. Either of working electrode **22** or counter electrode **24** may include more than one electrode, such as shown in **FIG. 4** (as counter electrodes **324, 344**) and in **FIG. 5B** (as counter electrodes **424, 441, 442**). Counter electrode **24** can also be formed with tines in a comb shape. Other configurations of both the counter electrode and working electrode are within the scope of the invention. However, for this particular embodiment, the separation distance between at least one portion of the working electrode and some portion of the counter electrode preferably does not exceed the limits specified hereinabove.

[0106] In another embodiment of the invention, the working and counter electrodes are coplanar. In this case, the sample chamber is in contact with both electrodes and is bounded on the side opposite the electrodes by a non-conducting inert base or substrate. Suitable materials for the inert base include non-conducting materials such as polyester.

[0107] Other configurations of the inventive sensors are also possible. For example, the two electrodes can be formed on surfaces that make an angle to each other. One such configuration would have the electrodes on surfaces that form a right angle. Another possible configuration has the electrodes on a curved surface such as the interior of a tube. For example, the working and counter electrodes can be arranged so that they face each other from opposite sides of the tube. This is another example of a facing electrode pair. Alternatively, the electrodes can be placed near each other on the tube wall (e.g., one on top of the other or side-by-side). In any configuration, the two electrodes must be configured so that they do not make direct electrical contact with each other, to prevent shorting of the electrochemical sensor.

[0108] Referring again to **FIGS. 1 and 2**, the electrodes **22, 24** extend from the sample chamber **26** to the other end of the sensor **20** as electrode extensions called "traces". Each trace provides a contact pad **23, 25** for providing electrical connection to a meter or other device to allow for data and measurement collection, as will be described later. Preferably, each contact pad **23, 25** is positioned on a tab **23', 25'** that extends from each non-conducting base substrate **32, 34**. In one embodiment, a tab has more than one contact pad positioned thereon. In a second embodiment, a single contact pad is used to provide a connection to one or more electrodes; that is, multiple electrodes are coupled together and are connected via one contact pad.

[0109] A spacer **33** can be used to keep the electrodes apart when the electrodes face each other as depicted in **FIGS. 1 and 2**; spacer **33** is clearly seen in **FIG. 2**. The spacer is typically constructed from an inert non-conducting material such as pressure-sensitive adhesive, polyester, Mylar™, Kevlar™ or any other strong, thin polymer film, or, alternatively, a thin polymer film such as a Teflon™ film, chosen for its chemical inertness. In addition to preventing contact between the electrodes, the spacer **33** can function as a portion of the boundary for the sample chamber **26**. Other spacers include layers of adhesive and double-sided adhesive tape (e.g., a carrier film with adhesive on opposing sides of the film). Adhesive may be applied, for example by coating, onto a polymeric material to provide spacer **33**.

[0110] Sample Chamber

[0111] The sample chamber **26** is typically defined by a combination of the electrodes **22, 24**, the substrates **32, 34**, and a spacer **33** as shown in **FIGS. 1 and 2**. A measurement zone is contained within this sample chamber and is the region of the sample chamber that contains only that portion of the sample that is interrogated during the analyte assay. In the embodiment of the invention illustrated in **FIGS. 1 and 2**, sample chamber **26** is the space between the two electrodes **22, 24** and their non-conductive base substrates **32, 34**, bounded by spacer **33**. In this embodiment, the sample chamber has a volume that is preferably no more than about 1 μL , more preferably no more than about 0.32 μL , and still more preferably no more than about 0.25 μL .

[0112] In the embodiment of the invention depicted in FIGS. 1 and 2, the measurement zone has a volume that is approximately equal to the volume of the sample chamber. In a preferred embodiment the measurement zone includes 80% of the sample chamber, 90% in a more preferred embodiment, and about 100% in a most preferred embodiment.

[0113] In another embodiment of the invention, shown in FIG. 3, sensor 220 has a working electrode 222 on non-conducting substrate base 232, a counter electrode 224 on non-conducting substrate base 234, and a spacer 233 therebetween. A contact pad 223 extends from working electrode 222, and likewise, a contact pad 225 extends from counter electrode 224. Sample chamber 226 (defined by base 232, 234 and spacer 233) extends the length of sensor 220 and includes much more space than the region proximate electrodes 222, 224. In this embodiment, the measurement zone, corresponding to the region containing the portion of the sample which will be interrogated, is the portion of sample chamber 226 bounded by the working surface of the working electrode 222 and counter electrode 224. In this embodiment, the measurement zone has a volume that is preferably no more than about 1 μL , more preferably no more than about 0.32 μL , still more preferably no more than about 0.25 μL , and most preferably no more than about 0.1 μL of sample.

[0114] In both of the embodiments discussed above, the thickness of the sample chamber and of the measurement zone correspond typically to the thickness of spacer 33, 233 (e.g., the distance between the electrodes in FIGS. 2 and 3, or the distance between the electrodes and the inert base in an embodiment where the electrodes are co-planar). The spacer can be, for example, an adhesive or double-sided adhesive tape or film. Examples of useable adhesives include urethanes, acrylates, acrylics, latexes, rubbers, and other known adhesive materials. Preferably, this thickness is small to promote rapid electrolysis of the analyte, as more of the sample will be in contact with the electrode surface for a given sample volume. In addition, a thin sample chamber helps to reduce errors from diffusion of analyte into the measurement zone from other portions of the sample chamber during the analyte assay, because diffusion time is long relative to the measurement time. Typically, the thickness of the sample chamber is between about 50 and about 200 micrometers.

[0115] Sorbent Material

[0116] The sample chamber can be empty before the sample is placed in the chamber, or, in some embodiments, the sample chamber can include a sorbent material (shown in FIG. 3 as sorbent 50) to sorb and hold a fluid sample during the measurement process. Suitable sorbent materials include polyester, nylon, cellulose, and cellulose derivatives such as nitrocellulose. The sorbent material facilitates the uptake of small volume samples by a wicking action which can complement or, preferably, replace any capillary action of the sample chamber. In addition to or alternatively, a portion or the entirety of the wall of the sample chamber may be coated by a surfactant, which is intended to lower the surface tension of the fluid sample and improve fluid flow within the sample chamber. An example of a useable surfactant is available under the tradename "Zonyl FSO" from Dupont of Wilmington, Del.

[0117] Methods other than the wicking action of a sorbent can be used to transport the sample into the sample chamber or measurement zone. Examples of such methods for transport include the application of pressure on a sample to push it into the sample chamber, the creation of a vacuum by a pump or other vacuum-producing method in the sample chamber to pull the sample into the chamber, capillary action due to interfacial tension of the sample with the walls of a thin sample chamber, as well as the wicking action of a sorbent material.

[0118] The entire sensor assembly is held firmly together to ensure that the sample remains in contact with the electrodes and that the sample chamber and measurement zone maintain the same volume. This is an important consideration in the coulometric analysis of a sample, where measurement of a defined sample volume is needed.

[0119] Alternative Sensor Designs

[0120] FIGS. 4 through 12 illustrate alternative sensor designs, both tip-filling and side-filling. Referring to FIG. 4, a sensor 320 has a working electrode 322, a counter electrode 324, a second counter electrode 344 (which may provide a fill indicator function, as described below), and a sample chamber 326 extending along at least a portion of the length of sensor 320 and optionally including a sorbent 350.

[0121] FIGS. 5A through 5C illustrate three layers that when assembled, provide a preferred embodiment of a sensor. FIG. 5A has a working electrode 422 on substrate 432. Working electrode 422 has a trace 453 extending from sample chamber 426 to tab 423' on which is contact pad 423; contact pad 423 connects the sensor to a meter or other measurement equipment. FIG. 5B, (inverted with respect to FIG. 5A to show the electrode side up), has a counter electrode 424 on substrate 434, and also includes a first indicator electrode 441 and a second indicator electrode 442. Counter electrode 424 has a trace 454, first indicator electrode 441 has trace 451, and second indicator electrode 442 has trace 452, that end at contact pads 425, 443, 444, respectively, on tab 425'. Spacer 433 in FIG. 5C defines sample chamber 426 and provides spacing between the electrodes when the two substrates 432, 434 are positioned opposite and facing one another.

[0122] FIGS. 6A through 6C also illustrate three layers that, when assembled, provide a sensor. In FIG. 6A, a working electrode 502 is formed on first substrate 500. The working electrode 502 includes a contact pad 503 for connection with external electronics; this contact pad 503 is connected to working electrode 502 by trace 552. A spacer 504, shown in FIG. 6B, such as a layer of adhesive or a double-sided tape defines a channel 506 to produce a sample chamber for the sensor. Two counter (or counter/reference) electrodes 510, 512 are formed on a second substrate 508, as shown in FIG. 6C (inverted with respect to FIG. 6A to show the electrode side up). This multiple counter electrode arrangement can provide a fill indicator function, using counter electrode 512, as described below. Each counter electrode 510, 512 has a contact region or pad 511, 513 for connection with external electronics; these contact pads 511, 513 are connected to counter electrodes 510, 512 by traces 551, 553. The second substrate 508 is inverted and placed over the first substrate 500, with the spacer 504 between, so that the working electrode 502 and the two counter electrodes 510, 512 are facing in the region of the channel 506.

[0123] In some instances, the counter electrode **510** nearest an entrance **514** (FIG. **6B**) of the channel **506** has a surface area within the sample chamber that is at least two times larger than the other counter electrode **512**, and may be at least five or ten times larger. The non-leachable or diffusible redox mediator and/or second electron transfer agent can be provided on either the first or second substrates **500**, **508** in a region corresponding to the channel **506**, as described above.

[0124] The working electrode and counter electrodes can be formed to cover the entire channel region (except for a small space between the two counter electrodes). In this embodiment, the sample chamber and measurement zone are effectively the same and have the same volume. In other embodiments, the measurement zone has, for example, 80% or 90% of the volume of the sample chamber. It will be understood that similar sensors could be made using one counter electrode or three or more counter electrodes. It will also be understood that multiple working electrodes may also be provided on the sensor.

[0125] FIGS. **7A**, **7B** and **7C** illustrate a side-filling sensor arrangement. FIG. **7A** shows a first substrate **520** with a working electrode **522**. FIG. **7B** illustrates a spacer **524** defining a channel **526**. FIG. **7C** (inverted with respect to FIG. **7A** and **7B**) illustrate a second substrate **528** with three counter (or counter/reference) electrodes **530**, **532**, **534**. This multiple counter electrode arrangement can provide a fill indicator function, as described below. An indentation **536**, **538** or recessed or indented portion can be formed at either edge of the opening to channel **526** to facilitate the drawing of fluid into the sensor. This configuration can aid in wicking or capillary filling of the channel (i.e., sample chamber). This configuration can also reduce the likelihood that the user will inadvertently block the channel during collection of the sample, which could occur by pressing the tip of the sensor edgewise against the skin.

[0126] FIGS. **8A**, **8B**, and **8C** illustrate another example of a side-filling sensor arrangement. FIG. **8A** illustrates a first substrate **540** with a working electrode **542**. FIG. **8B** illustrates a spacer **544** defining a channel **546**. FIG. **8C** (inverted with respect to FIGS. **8A** and **8B**) illustrates a second substrate **548** with three counter (or counter/reference) electrodes **550**, **552**, **554**.

[0127] FIGS. **9A**, **9B**, and **9C** illustrate another example of a tip-filling sensor arrangement. FIG. **9A** illustrates a first substrate **560** with a working electrode **562**. FIG. **9B** illustrates a spacer **564** defining a channel **566**. FIG. **9C** (inverted with respect to FIGS. **9A** and **9B**) illustrates a second thin film substrate **568** with two counter (or counter/reference) electrodes **570**, **572**. This multiple counter electrode arrangement can provide a fill indicator function, as described below. A vent hole **574** (indicated as a shaded region in FIG. **9C**) is provided through the second substrate. In the illustrated embodiment, this vent hole **574** is made through only the substrate **568** that carries the counter electrode(s) and, optionally, the spacer **564**. In this embodiment, the vent hole can be formed by, for example, die cutting a portion of the substrate. This die cut can remove a portion of at least one counter electrode, but a sufficient amount of the counter electrode should remain for contact with the sample in the channel and for electrical connection to a contact at the other end of the sensor. In another

embodiment, the vent hole **574** can be made through all of the layers or through the first substrate and not the second substrate.

[0128] Another embodiment is illustrated in FIGS. **10A**, **10B**, and **10C**, with a different shape. This sensor includes a first substrate **579** with at least one working electrode **580**, as illustrated in FIG. **10A**. The sensor also includes a spacer **581** with a channel **582** formed in the spacer **581**, as shown in FIG. **10B**. The sensor further includes a second substrate **583** with two counter electrodes **584**, **585**, as shown in FIG. **10C** (inverted with respect to FIGS. **10A** and **10B**). A venting aperture **586** is cut typically through all of the layers and extends from a side of the sensor. In some embodiments, the venting aperture and the front portion **587** of the sensor are simultaneously cut with a reproducible distance between the venting aperture and the front portion **587** of the sensor to provide a reproducible length for the channel **582** and the working electrode **580**. This tip-filling sensor arrangement optionally includes an indentation **588** or recessed or indented portion that can be formed at the filling opening of the channel **582** to facilitate the drawing of fluid into the sensor.

[0129] FIGS. **11A**, **11B**, and **11C** illustrate another example of a side-filling sensor arrangement. FIG. **11A** illustrates a first substrate **640** with a working electrode **642**. FIG. **11B** illustrates a spacer **644** defining a channel **646**. FIG. **11C** (inverted with respect to FIGS. **11A** and **11B**) illustrates a second substrate **648** with three counter (or counter/reference) electrodes **650**, **652**, **654**. This multiple counter electrode arrangement can provide a fill indicator function, as described below. The length of the channel **646** is typically defined by the two parallel cuts along the sides **656**, **658** of the sensors.

[0130] Because of the straight sides, these sensors (and also those shown in FIGS. **1**, **2** and **5**) can be manufactured adjacent to one another, as illustrated in FIG. **13A**. Such positioning during manufacture produces less waste material. Another optional processing advantage of the adjacent sensor manufacturing process is that the redox mediator and/or second electron transfer agent can be readily disposed in the channel by striping a continuous stream of these components along a row or column of adjacent sensors. This can result in better efficiency and less waste of expensive reagents such as the redox mediator and/or second electron transfer agent, as compared to other techniques, such as individually placing these components within the individual channels.

[0131] FIGS. **12A**, **12B**, and **12C** illustrate another sensor configuration. This sensor includes a first substrate **600** with at least one working electrode **602**, as illustrated in FIG. **12A**. The sensor also includes a spacer **604** with a channel **606** formed in the spacer **604**, as shown in FIG. **12B**. The sensor further includes a second substrate **608** with two counter electrodes **610**, **612**, as shown in FIG. **12C** (inverted with respect to FIGS. **12A** and **12B**). This multiple counter electrode arrangement may provide a fill indicator function, as described below. The sensor can also include, for example, an indicator, such as a slot **614** or an extension **616** from the body of the sensor that indicates to the user which side of the sensor should be placed adjacent to the sample. Surface printing can also or alternatively be applied to the sensor to act as an indicator. Some indication mechanism

can be particularly important where the sensor reading is only correct when the sample enters from a particular side.

[0132] Multiple Electrode Sensors and Calibration Thereof

[0133] Multiple electrode sensors can be used for a variety of reasons. For example, multiple electrodes can be used to test a variety of analytes using a single sample. One embodiment with multiple electrodes has one or more sample chambers, each of which contains one or more working electrodes, with each working electrode defining a different measurement zone. If the redox mediator is non-leachable, one or more of the working electrodes can have the appropriate chemical reagents, for example, an appropriate enzyme, to test a first analyte and one or more of the remaining working electrodes can have the appropriate chemical reagents to test a second analyte, and so on. For example, a multiple electrode sensor might include a working electrode having glucose oxidase disposed thereon to determine glucose concentration and another working electrode can have lactate oxidase disposed thereon to determine lactate concentration.

[0134] Multiple electrodes could also be used to improve the precision of the resulting readings. The measurements from each of the working electrodes (all of which are detecting the same analyte) can be averaged or otherwise combined together to obtain a more precise or reliable reading. In some cases, measurements could be rejected if the difference between the value and the average exceeds a threshold limit. This threshold limit could be, for example, determined based on a statistical parameter, such as the standard deviation of the averaged measurements. The average could then be recalculated while omitting the rejected values. In addition to using multiple electrode sensors to increase precision, multiple measurements could be made at each electrode and averaged together to increase precision. This technique could also be used with a single electrode sensor to increase precision.

[0135] One example of a multiple electrode sensor that can be used to accurately determine the volume of the measurement zones of the electrode pairs and that is also useful in reducing noise is presented herein. In this example, one of the working electrodes is prepared with a non-leachable redox mediator and a non-leachable second electron transfer agent (e.g., an enzyme). Another working electrode includes non-leachable redox mediator, but no second electron transfer agent on the electrode. An optional third working electrode has no redox mediator and no second electron transfer agent bound to the electrode. A similar configuration can be constructed using diffusible redox mediator and/or diffusible second electron transfer agent although diffusible components are not limited to being disposed on the working electrode. Preferably, the distance between the working electrodes is sufficient that redox mediator and/or enzyme do not substantially diffuse between electrodes within the measurement period (e.g., in the time period from introduction of the same sample into the sample chamber to the end of the measurement).

[0136] The sensor error caused by the redox mediator being in a non-uniform oxidation state prior to the introduction of the sample can be measured by concurrently electrolyzing the sample in the measurement zones that are proximate working and counter electrodes. At the first

working electrode, the analyte is electrolyzed to provide the sample signal. At the second working electrode, the analyte is not electrolyzed because of the absence of the second electron transfer agent (assuming that a second electron transfer agent is necessary). However, a charge will pass (and a current will flow) due to the electrolysis of the redox mediator that was in a mixed oxidation state (i.e., some redox centers in the reduced state and some in the oxidized state) prior to the introduction of the sample and/or the shuttling of a diffusible redox mediator between the working electrode and the counter electrode. The small charge passed using this second working electrode can be subtracted from the charge passed using the first electrode pair to substantially remove the error due to the oxidation state of the redox mediator and/or to remove the background current caused by a diffusible redox mediator. This procedure also reduces the error associated with other electrolyzed interferents, such as ascorbate, urate, and acetaminophen, as well as errors associated with capacitive charging and faradaic currents.

[0137] The thickness of the sample chamber can be determined by measuring the capacitance, preferably in the absence of any fluid, between an electrode and its corresponding counter electrode. The capacitance of an electrode pair depends on the surface area of the electrodes, the interelectrode spacing, and the dielectric constant of the material between the plates. The dielectric constant of air is unity which typically means that the capacitance of this electrode configuration is a few picofarads (or about 100-1000 picofarads if there is fluid between the electrode and counter electrode given that the dielectric constant for most biological fluids is approximately 75). Thus, since the surface area of the electrodes are known, measurement of the capacitance of the electrode pair allows for the determination of the thickness of the measurement zone to within about 1-5%.

[0138] Other electrode configurations can also use these techniques (i.e., capacitance measurements and coulometric measurements in the absence of a critical component) to reduce background noise and error due to interferents and imprecise knowledge of the volume of the interrogated sample. Protocols involving one or more working electrode(s) and counter electrode(s) and one or more of the measurements described above can be developed and are within the scope of the invention. For example, only one electrode pair is needed for the capacitance measurements, however, additional electrodes can be used for convenience.

[0139] Fill Indicator

[0140] When using a sample chamber that is filled with 1 μ L or less of fluid, it is often desirable to be able to determine when the sample chamber is filled. FIGS. 6A through 6C illustrate a sensor having a fill indicator structure. In particular, FIG. 6A illustrates a first substrate 500 upon which a working electrode 502 is printed. A spacer 504 (FIG. 6B), such as, for example, a layer of adhesive or a double-sided tape, is formed over the first substrate 500 and working electrode 502 with a channel 506 formed in the layer to provide a sample chamber. A second substrate 508 is printed with two counter electrodes 510, 512, as shown in FIG. 6C (inverted with respect to FIGS. 6A and 6B to show the electrode side up). Preferably, the counter electrode 510 nearest an entrance 514 of the channel 506 has a surface area

within the sample chamber that is at least two times larger than the other counter electrode **512**, and preferably at least five or ten times larger.

[0141] The sensor can be indicated as filled, or substantially filled, by observing a signal between the second counter electrode **512** and the working electrode **502** as the sensor fills with fluid. When fluid reaches the second counter electrode **512**, the signal from that counter electrode will change. Suitable signals for observing include, for example, voltage, current, resistance, impedance, or capacitance between the second counter electrode **512** and the working electrode **502**. Alternatively, the sensor can be observed after filling to determine if a value of the signal (e.g., voltage, current, resistance, impedance, or capacitance) has been reached indicating that the sample chamber is filled.

[0142] In alternative embodiments, the counter electrode or working electrode can be divided into two or more parts and the signals from the respective parts observed to determine whether the sensor has been filled. In one example, the working electrode is in a facing relationship with the counter electrode and the indicator electrode. In another example, the counter electrode, working electrode, and indicator electrode are not in a facing relationship, but are, for example, side-by-side. Typically, the indicator electrode is further downstream from a sample inlet port than the working electrode and counter electrode.

[0143] For side-fill sensors, such as those illustrated in FIGS. 5, 7, 8 and 11, an indicator electrode can be disposed on each side of the primary counter electrode. This permits the user to fill the sample chamber from either the left or right side with an indicator electrode disposed further upstream. This three-electrode configuration is not necessary. Side-fill sensors can also have a single indicator electrode and, preferably, some indication as to which side should be placed in contact with the sample fluid.

[0144] Alternately or additionally, two indicator electrodes, used in combination with one counter/reference electrode, detect when the sample chamber begins to fill and when the sample chamber has been filled to prevent partial filling of the sample chamber. The two indicator electrodes are optionally held at a different potential than the counter/reference electrode. The start and completion of filling of the sample chamber is indicated by the flow of current between the indicator and counter/reference electrodes.

[0145] In other instances, the potential of each of the counter/reference electrodes may be the same. When the potential at all three counter/reference electrodes is the same, for example, 0 volts, then as the measurement zone begins to fill, the fluid allows for electrical current flow between a working electrode and the first counter/reference electrode, causing a current at the first counter/reference electrode due to the reaction of the analyte with the enzyme and the mediator. When the fluid reaches the third counter/reference electrode, another current may be measured similar to the first counter/reference electrode indicating that the measurement zone is full. When the measurement zone is full, the three counter/reference electrodes are optionally shorted together or their signals can be added or otherwise combined.

[0146] The indicator electrode can also be used to improve the precision of the analyte measurements. The indicator

electrode may operate as a working electrode or as a counter electrode or counter/reference electrode. In the embodiment of FIGS. 6A through 6C, the indicator electrode **512** can act as a second counter or counter/reference electrode with respect to the working electrode **502**. Measurements from the indicator electrode/working electrode can be combined (for example, added or averaged) with those from the first counter/reference electrode/working electrode to obtain more accurate measurements. In one embodiment, the indicator electrode operates as a second working electrode with the counter electrode or counter/reference electrode. In another embodiment, the indicator electrode operates as a second working electrode with a second counter electrode or counter/reference electrode. In still another embodiment, the indicator electrode operates as a second counter electrode or counter/reference electrode with a second working electrode.

[0147] The sensor or a sensor reader can include a sign (e.g., a visual sign or auditory signal) that is activated in response to the indicator electrode to alert the user that the measurement zone has been filled. The sensor or a sensor reader can be configured to initiate a reading when the indicator electrode indicates that the measurement zone has been filled with or without alerting the user. The reading can be initiated, for example, by applying a potential between the working electrode and the counter electrode and beginning to monitor the signals generated at the working electrode.

[0148] General Method for Manufacturing Sensors

[0149] Referring now to FIGS. 13A and 13B, one example of a method for making thin film sensors is described with respect to the sensor arrangement displayed in FIGS. 5A through 5C, although this method can be used to make a variety of other sensor arrangements, including those described before. When the three layers of FIGS. 5A through 5C are assembled, a sensor **420** is formed.

[0150] In FIGS. 13A and 13B, a substrate **1000**, such as a plastic substrate, is moving in the direction indicated by the arrow. The substrate **1000** can be an individual sheet or a continuous roll on a web. Multiple sensors **420** can be formed on a substrate **1000** as sections **1022** that have working electrodes **422** (FIG. 5A) thereon and sections **1024** that have counter electrode **424** and indicator electrodes **441**, **442** (FIG. 5B). These working, counter and indicator electrodes are electrically connected to their corresponding traces and contact pads. Typically, working electrode sections **1022** are produced on one half of substrate **1000** and counter electrode sections **1024** are produced on the other half of substrate **1000**. In some embodiments, the substrate **1000** can be scored and folded to bring the sections **1022**, **1024** together to form the sensor. In some embodiments, as illustrated in FIG. 13A, the individual working electrode sections **1022** can be formed next to or adjacent each other on the substrate **1000**, to reduce waste material. Similarly, individual counter electrode sections **1024** can be formed next to or adjacent each other. In other embodiments, the individual working electrode sections **1022** (and, similarly, the counter electrode sections **1024**) can be spaced apart, as illustrated in FIG. 13B. The remainder of the process is described for the manufacture of multiple sensors, but can be readily modified to form individual sensors.

[0151] Carbon or other electrode material (e.g., metal, such as gold or platinum) is formed on the substrate **1000** to provide a working electrode **422** for each sensor. The carbon or other electrode material can be deposited by a variety of methods including printing a carbon or metal ink, vapor deposition, and other methods. The printing may be done by screen printing, gravure roll printing, transfer printing, and other known printing methods. Trace **453** and contact pad **423** are preferably applied together with working electrode **422**, but may be applied in a subsequent step.

[0152] Similar to the working electrode **422**, the counter electrode(s) **424**, **441**, **442** (shown in FIG. 5B) are formed on the substrate **1000**. The counter electrodes are formed by providing carbon or other conductive electrode material onto the substrate. In one embodiment, the material used for the counter electrode(s) is a Ag/AgCl ink. The material of the counter electrode(s) may be deposited by a variety of methods including printing or vapor deposition. The printing may be done by screen printing, gravure roll printing, transfer printing, and other known printing methods. Traces **454**, **451**, **452** and contact pads **425**, **443**, **444** are preferably applied together with counter electrodes **424**, **441**, **442**, but may be applied in a subsequent step.

[0153] Preferably, multiple sensors **420** are manufactured simultaneously; that is, the working electrodes, including their traces and contact pads, for a plurality of sensors are produced (e.g., printed) on a polymer sheet or web, and simultaneously or subsequently, the counter electrodes, and their traces and contact pads, for a plurality of sensors are produced (e.g., printed). The working electrode(s) and counter electrode(s) can be formed on separate substrates that are later positioned opposite one another so that the electrodes face each other. Alternately, to simplify registration of the substrates, the working electrodes can be formed on a first half of a substrate sheet of web and the counter electrodes are formed on a second half of the substrate sheet or web so that the sheet or web can be folded to superimpose the working and counter electrodes in a facing arrangement.

[0154] To provide a sample chamber **426**, a spacer **433** is formed over at least one of the substrate/working electrode and substrate/counter electrode(s). The spacer **433** is preferably an adhesive spacer, such as a single layer of adhesive or a double-sided adhesive tape (e.g., a polymer carrier film with adhesive disposed on opposing surfaces). Suitable spacer materials include adhesives such as urethanes, acrylates, acrylics, latexes, rubbers and the like.

[0155] A channel, which will result in the sample chamber, is provided in the spacer **433**, either by cutting out a portion of the adhesive spacer or placing two adhesive pieces in close proximity but having a gap therebetween. The adhesive can be printed or otherwise disposed on the substrate according to a pattern which defines the channel region. The adhesive spacer can be optionally provided with one or more release liners prior to its incorporation into the sensor. The adhesive can be cut (e.g., die-cut or slit) to remove the portion of the adhesive corresponding to the channel prior to disposing the spacer on the substrate.

[0156] In one method of the invention, illustrated in FIGS. 14A through 14G, the adhesive includes a first and a second release liner. Prior to applying the adhesive to a first substrate/working electrode or substrate/counter electrode(s), the adhesive is "kiss-cut" through the first release liner and

at least a portion of the underlying adhesive, preferably the entire adhesive but not through the second release liner, to form one or more sections, one of which will eventually result in a sample chamber. The first release liner is removed from the adhesive over its length but is retained over the sample chamber section. The exposed adhesive is then applied to the first substrate/working electrode or substrate/counter electrode(s). On removal of the second release liner, the adhesive positioned within the sample chamber sections is also removed, leaving a channel or opening in the adhesive spacer. The second substrate/working electrode or substrate/counter electrode(s) is then applied to the adhesive to form the layered sensor structure.

[0157] FIGS. 14A through 14G describe in detail the processing of an adhesive layer by a "kiss-cut". An adhesive layer that will eventually become spacer **433** is preferably obtained from an adhesive structure including an adhesive layer **1030** between a first release liner **1020** and a second release liner **1040**. A blade (e.g., a knife or die blade) cuts through the first release liner **1020** and at least a portion of the adhesive layer **1030**, but not through the second release liner **1040**, as shown in FIG. 14B and 14G, to form release liner sections **1020a**, **1020b**, and adhesive sections **1030a**, **1030b**. The space occupied by **1030b** is the area that will result in the sample chamber of the sensor and can be referred to as "sample chamber region". A cut configuration such as shown in FIG. 14G allows removal of one release liner section **1030a**, leaving behind release liner section **1030b**. The liner could be cut to provide individual strips extending the entire length of the liner; this would then need removal of each individual liner strip.

[0158] The release liner section **1030a** is removed to provide the adhesive configuration shown in FIGS. 14C. The exposed adhesive is positioned over and securely adhered to the substrate **1050**, as illustrated in FIG. 14D. The adhesive sections **1030a** without the release liner will adhere to the substrate **1050**. As shown in FIG. 14E, the second release liner **1040** is removed, pulling the cut and non-adhered section of adhesive **1030b** and the first liner **1020b** with it; this results in adhesive areas **1030a** providing spacer layer sections **433a**, **433b**, with sample chamber **426** therebetween, as illustrated in FIG. 14F. The second substrate is then positioned over the adhesive layer and laminated to the first substrate via the spacer **433**. The thickness of the spacer **433** typically determines the spacing between the working and counter electrodes, and thus impacts the volume of the sample chamber **426**. Preferably, the thickness does not vary more than $\pm 5\%$ over the individual sensor and/or among individual sensors in a batch. Use of the "kiss-cut" method for manufacturing the layered sensor structure preserves the integrity of the sample chamber walls during manufacturing, thereby providing for more reliable and reproducible sample chamber volumes.

[0159] The non-leachable or diffusible redox mediator and/or second electron transfer agent are disposed onto the substrate in at least the sample chamber regions. If either or both of these components is non-leachable, that component or components is preferably disposed on the working electrode. If either or both of these components is diffusible, that component or components can be disposed on any surface of the substrate in the channel region. The redox mediator and/or second electrode transfer agent can be disposed independently or together on the substrate prior to or after

disposition of the spacer. The redox mediator and/or second electrode transfer agent may be disposed by a variety of methods including, for example, screen printing, ink jet printing, spraying, painting, striping along a row or column of aligned and/or adjacent electrodes, and the like. Other components can be deposited separately or together with the redox mediator and/or second electrode transfer agent; these components can include, for example, surfactants, polymers, polymer films, preservatives, binders, buffers, and cross-linkers.

[0160] After disposing the spacer, redox mediator, second electron transfer agent, sensing layers, and the like, the first and second substrates (having the working and counter electrodes thereon) are positioned opposite each other to form the sensor. The faces of the substrate are joined by the adhesive of the spacer. Preferably, after bringing the faces together, individual sensors are cut out from the web of sensors using a variety of methods including, for example, die cutting, slitting, or otherwise cutting away the excess substrate material and separating the individual sensors. In some embodiments, a combination of cutting or slitting methods is used. As another alternative, the individual sensor components can first be cut out of the substrates and then brought together to form the sensor by adhesively joining the two components, such as by using the spacer adhesive.

[0161] The sides of the sensor can be straight to allow the sensor to be cut out from the remainder of the substrate and/or from other sensors by slitting the substrate in parallel directions using, for example, a gang arbor blade system. The edges of the sensor can define edges of the sample chamber and/or measurement zone. By accurately controlling the distance between cuts, variability in sample chamber volume can often be reduced. In some instances, these cuts are preferably parallel to each other, as parallel cuts are typically the easiest to reproduce.

[0162] Sensor Connection to Electrical Device

[0163] Referring generally now to **FIGS. 16A and 16B, 17A and 17B, and 18A and 18B**, the assembled sensor of **FIGS. 5A through 5C** (as will be referred to as sensor **1420**) is typically coupled to a meter or other electrical device by electrical connector **1500** which is configured to couple with and contact the end of the sensor **1420** at contact pads **423, 425, 443, 444**. The sensor meter typically includes a potentiostat or other component to provide a potential and/or current for the electrodes of the sensor. The sensor reader also typically includes a processor (e.g., a microprocessor or hardware) for determining analyte concentration from the sensor signals. The sensor meter also includes a display or a port for coupling a display to the sensor. The display displays the sensor signals and/or results determined from the sensor signals including, for example, analyte concentration, rate of change of analyte concentration, and/or the exceeding of a threshold analyte concentration (indicating, for example, hypo- or hyperglycemia).

[0164] One example of a suitable connector is shown in **FIGS. 16A and 16B, 17A and 17B, and 18A and 18B**. Connector **1500** (which is used to connect a sensor to a meter or other electrical device) is generally a two part structure, having top portion **1510** and bottom portion **1520** (see **FIG. 16B**). Positioned between and secured by top portion **1510** and bottom portion **1520** are contact leads

1423, 1425, 1443, 1444 that provide electrical connection between the sensor **1420** and a meter. These leads **1423, 1425, 1443, 1444** have proximal ends to physically contact pads **423, 425, 443, 444** (in **FIGS. 5A and 5B**, respectively) and distal ends to connect to any attached meter. The end of the sensor **1420** which has the contact pads can be slid into or mated with the connector **1500** by placing the sensor **1420** into slide area **1530**, which provides a support for and retains sensor **1420**. It is typically important that the contact structures of the connector **1500** make electrical contact with the correct pads of the sensor so that the working electrode and counter electrode(s) are correctly coupled to the meter.

[0165] One optional feature is an insertion monitor **1450**, that is illustrated in **FIGS. 16A and 16B**, which are top views of the sensor prior to insertion into connector **1500**. This conductive insertion monitor is positioned on the non-conductive base substrate and has a contact pad for electrical contact with a connector. The insertion monitor is configured and arranged to close an electrical circuit between two contact structures **1451** and **1452** when the sensor is properly inserted into the connector. Proper insertion into the connector **1500** means that the sensor strip **1420** is inserted right side up, that the correct end of the strip is inserted into the connector, and that the strip is inserted far enough into the connector that reliable electrical connections are made between the electrodes' contact pads on the strip and the corresponding contact structures of the connector. Preferably, no closed circuit is made unless all electrode pads have properly contacted the contact structures of connector **1500**. The insertion monitor may have shapes other than a stripe across the width of the sensor; for example, other designs include an individual dot, a grid pattern, or may include stylistic features, such as words or letters.

[0166] Because this insertion monitor **1450** is not at the end with the contact regions for the electrodes, the insertion monitor **1450** does not require additional width space on the sensor. The width of the contact pads **443, 425, 444, 423** is defined as the width on which a lead could be placed that would result in an electrical connection; typically, the contact width is the width of the exposed contact area. In one embodiment, six contact lead structures on the connector (i.e., **1451, 1443, 1425, 1444, 1423, 1452**) can contact sensor **1420** in the same width as the four contact pads (i.e., **443, 425, 444, 423**). This concept of having contact points on the sensor that occupy more width than the width of the sensor may be used for any number of contact points; this may be used with or without an insertion monitor.

[0167] As a particular example, four leads **1443, 1425, 1444, 1423** make contact with contact pads **443, 425, 444, 423**. If each lead and/or contact pad is one millimeter wide, a sensor of at least 4 mm wide is needed to make contact. Additional leads, such as those for insertion monitor **1450** (i.e., contact leads **1451, 1452**), can make contact by having leads **1451, 1452** extend along the side of leads **1443** and **1423** and then angle in toward the center of the strip **1420** after the point where leads **1443, 1425, 1444, 1423** contact strip **1420**. The insertion monitor stripe leads **1451, 1452** cross side edges of sensor **1420** to make contact with the sensor, thus not requiring additional sensor width.

[0168] The contact structures are parallel and non-overlapping and they run longitudinally from the distal end to the proximal end. The lead structures **1443, 1425, 1444, 1423**

terminate at their proximal end, but lead structures **1451**, **1452** continue longitudinally past the proximal end of lead structures **1443**, **1425**, **1444**, **1423**. Once past the proximal end, lead structures **1452**, **1452** angle in toward the center of the sensor strip.

[**0169**] The insertion monitor can also be used to encode information on a test strip. The encoded information can be, for example, calibration information for that manufacturing lot or for that specific strip. Such calibration information may relate to the sensitivity of the strip or to the y-intercept of its calibration curve. In one embodiment, the insertion monitor comprises two or more contact pads for connection to a meter. The two or more contact pads are connected to each other by a conductive material, such as a conductive ink. The resistance of conductive path between the two or more contact pads is related to the encoded information. As an example of discrete calibration values, resistance values in a given range can correspond to one calibration setting, and resistance values in a different range can correspond to a different calibration setting. As an example of continuous calibration values, the calibration setting may be a continuous function of the resistance. Examples of suitable conductive pathways between contact pads are shown in **FIGS. 19A through 19L**.

[**0170**] In **FIGS. 19A-19F**, multiple conductive pathways with different resistances are printed on the strip. The resistance of the pathway is varied by cutting or scoring some or all of the conductive pathways so that they do not carry charge. In **FIGS. 19G-19L**, the resistance is controlled by the width or length of the conductive path. While it is preferred to provide this encoded information on the insertion monitor, it should be recognized that the insertion monitor function and the encoding of information can also be implemented separately using separate conductive traces on the strip.

[**0171**] In another embodiment to ensure proper insertion of a sensor into a meter, the meter may include a raised area or bump that prevents or hinders the insertion of the sensor in an improper direction, as shown in **FIG. 20**. To ensure proper insertion of the sensor **1100** into a connector **1110**, the connector **1110** may include a raised area **1140** that prevents or hinders the insertion of the sensor in an improper direction. For example, the width, w_2 , of the contact region of the second substrate **1130** may be wider than the width, w_1 , of the contact region of the first substrate **1120**. In this instance, the raised area **1140** is positioned to allow sensor **1100** to be slid into the connector so that the first substrate **1120** is next to the surface **1150** from which the raised area **1140** protrudes, but would prevent or hinder having the second substrate **1130** next to the surface **1150** from which the raised area **1140** protrudes. Objects other than a raised area can also be used to guide the user in correct introduction of the sensor into the meter.

[**0172**] Integrated Sample Acquisition and Analyte Measurement Device

[**0173**] An analyte measurement device constructed according to the principles of the present invention typically includes a sensor, as described hereinabove, combined with a sample acquisition apparatus to provide an integrated sampling and measurement device. The sample acquisition apparatus typically includes, for example, a skin piercing member, such as a lancet, that can be injected into a patient's

skin to cause blood flow. In a preferred embodiment, the integrated sample acquisition and analyte measurement device comprises a lancing instrument that holds a lancet and measurement strip. The lancing instrument preferably requires active cocking. By requiring the user to cock the device prior to use, the risk of inadvertently triggering the lancet is minimized. Preferably, the lancing instrument will also permit the user to adjust the depth of penetration of the lancet into the skin. Such devices are already commercially available from companies such as Boehringer Mannheim and Palco. This feature allows users to adjust the lancing device for differences in skin thickness, skin durability, and pain sensitivity across different sites on the body and across different users.

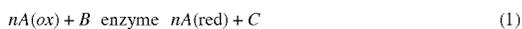
[**0174**] Typically, a larger sample of body fluid such as blood or interstitial fluid is expressed when pressure is applied around a site where a hole has been created in the skin. A sample acquisition device having a protruding ring, such as illustrated in **FIG. 21**, may be used. Ring **1200** circumferentially surrounds the lancing site to create increased pressure and extract a larger sample out of the wound. A lancet would protrude from the center **1220** of ring **1200**. Ring **1200** has a surface area **1210** that presses against the skin, thus increasing the tension on the skin and often providing a bump of skin. Skin that is more taut, and even extended, provides a large volume of fluid from the wound. The increased surface area **1210** and tighter skin provided by ring **1200**, decreases the amount of pain experienced by the patient during the lancing. Further, the tendency to bruise is typically decreased.

[**0175**] In one embodiment, the lancing instrument and the meter are integrated into a single device. To operate the device the user need only insert a disposable cartridge containing a measurement sensor and lancing device into the integrated device, cock the lancing instrument, press it against the skin to activate it, and read the result of the measurement. Such an integrated lancing instrument and test reader simplifies the testing procedure for the user and minimizes the handling of body fluids.

[**0176**] Operation of the Sensor and Meter

[**0177**] An electrochemical sensor of the invention can be operated with or without applying a potential across the electrodes. In one embodiment, the electrochemical reaction occurs spontaneously and a potential need not be applied between the working and counter electrodes. In another embodiment, a potential, which may or may not remain constant, is applied between the working and counter electrodes. The magnitude of the required potential depends on the redox mediator used. The potential at which the electrode poises itself, or where it is poised by applying an external bias, and where the analyte is electrolyzed, is typically such that the electrochemical reaction is driven to or near completion, but it is, preferably, not oxidizing enough to result in significant electrochemical reaction of interferents, such as urate, ascorbate, and acetaminophen, that can affect the signal measured. For non-leachable redox mediators, the potential is typically between about -350 mV and about $+400$ mV versus the standard calomel electrode (SCE). Preferably, the potential of the redox mediator is more negative than $+100$ mV, more preferably the potential is more negative than 0 mV, and most preferably the potential is about -150 mV versus SCE.

[0178] When an external potential is applied, it can be applied either before or after the sample has been placed in the sample chamber. If the measurement zone includes only a portion of the sample chamber then the potential is preferably applied after the sample has come to rest in the sample chamber to prevent electrolysis of sample passing through the measurement zone as the sample chamber is filling. Alternatively, in the case where the measurement zone includes most or all of the sample chamber, the potential can be applied before or during the filling of the sample chamber without affecting the accuracy of the assay. When the potential is applied and the sample is in the measurement zone, an electrical current will flow between the working electrode and the counter electrode. The current is a result, at least in part, of the electrolysis of the analyte in the sample. This electrochemical reaction occurs via the redox mediator and the optional second electron transfer agent. For many biomolecules, B, the process is described by the following reaction equations:



[0179] Biochemical B is oxidized to C by redox mediator A in the presence of an appropriate enzyme. Then the redox mediator A is oxidized at the electrode. Electrons are collected by the electrode and the resulting current is measured. The measured current may also include a background current resulting in a measured background charge, due, at least in part, to the shuttling of a diffusible redox mediator between the working electrode and the counter electrode. This background current can be minimized or accounted for, as described in PCT Application PCT/US99/23425, incorporated herein by reference.

[0180] A variety of techniques can be used to determine the concentration of an analyte in a sample. These techniques include, for example, coulometry, amperometry, and potentiometry. To determine the concentration of analyte (e.g., glucose) in the sample by coulometry, the charge passing or projected to pass between the working electrode(s) and counter electrode(s) during electrolysis of the analyte is determined. Knowledge of the charge and the volume of the sample chamber permit the calculation of the concentration of the electrolyzed analyte in the sample. This charge can be determined by several methods.

[0181] For example, the charge can be measured directly. This can be accomplished using a coulometer and known coulometric techniques. Typically, the charge is measured during the complete or nearly complete electrolysis of the analyte.

[0182] As another example, the charge can be determined from measurements of the electrolysis current, i , over time, t . A series of currents ($i_x, i_{x+1}, i_{x+2}, \dots$) is measured for a series of times ($t_x, t_{x+1}, t_{x+2}, \dots$). The current can then be integrated (e.g., numerically integrated using known numerical methods) to give the charge. In some instances, current can be measured during the entire electrolysis. In other instances, current can be extrapolated after a period of time and the charge determined using a combination of the measured currents and extrapolated currents.

[0183] Extrapolation of current utilizes a projected relationship between current and time during at least a portion of the electrolysis. Typically, when the system is diffusion limited, the natural logarithm of the current is linearly related to the time. This typically occurs after a period of time during which the sensor "equilibrates". FIG. 15A is an example of a graph of current versus time for electrolysis of an analyte, such as glucose, in one of the previously described sensors; FIG. 15B is a graph of the natural logarithm of current versus time for the same data. After the sample completely fills the sample chamber, the current increases to a peak current value, designated as "P" in FIGS. 15A and 15B, during which time the sensor is equilibrating. The peak current typically occurs during a period of time in which the system is kinetically, rather than diffusively, limited. Typically, the current then begins to decrease, although, in some instances, further increases may occur before the current becomes diffusion limited. Eventually, the current values enter a region, designated as "L" in FIG. 15B, where there is a linear relationship between the natural logarithm of current and time. The remaining current needed to electrolyze a remaining portion of the analyte can then be extrapolated using estimation methods, such as nonlinear or, preferably, linear estimation methods (e.g., linear least squares techniques). The region where extrapolation occurs is designated as "E" in FIG. 15B and the extrapolation is indicated as a solid line.

[0184] For this discussion, the current values being referred to are the absolute value of the actual current, thus, the sign of the current measurement is ignored. For example, the peak current is the largest deviation from zero current, whether positive or negative.

[0185] During the period when the system is diffusion limited the relationship between the natural logarithm of the current and time is linear. The remaining charge required to completely electrolyze the analyte remaining in the sample can be reliably estimated from the slope of this line. Care must be taken to ensure that the system is in the diffusion limited range before performing the estimation. One method of ensuring that the system is in the diffusion limited regime includes observing the current values until the peak current, i_{peak} ("P") is achieved. Current values continue to be observed until the current drops below a threshold value, i_{thresh} , that is typically a fraction of the peak current. For example, the threshold value can be one half, one third, or one quarter of the peak current (e.g., $i_{\text{thresh}} = j * i_{\text{peak}}$ where j is, for example, 0.5, 0.33, or 0.25). The threshold value is chosen based on the characteristics of the sensor so that one has a high degree of confidence that the system will be diffusion limited at this time.

[0186] Many other methods may be used to ensure that the system is in a diffusion limited regime. For example, one could observe the current values after i_{peak} until an inflection point is reached and the second derivative of the natural log of the current turns positive. At this point the system is typically in a diffusion limited regime.

[0187] After achieving the threshold value, slope values are determined using natural logarithms of two or more of the measured current values (e.g., $m_x = (\log(i_x) - \log(i_{x-y})) / (t_x - t_{x-y})$ where m_x is the slope at time t_x , i_x is the current at time t_x , and i_{x-y} is the current at time t_{x-y} before t_x). Although the slope can be determined using natural loga-

gorithms of any two current values (e.g., current values measured one directly after the other), preferably the slope is determined using natural logarithms of multiple consecutive current values or current values that are spaced apart by at least about ten (i.e., $y=10$), and, preferably, at least about thirty (i.e., $y=30$), current values. This procedure can reduce the effect of random noise in the system. Optionally, the consistency of multiple slope values (e.g., three or five or ten slope values) can be observed for verification. For example, the system can look for ten monotonically decreasing slope values before accepting the slope value for use in the estimation.

[0188] The determined slope value is used to extrapolate the charge required to completely electrolyze the analyte remaining in the sample. That is, the current is extrapolated to zero current. Known extrapolation techniques (e.g., linear extrapolation techniques) can be used. The total charge required to electrolyze the analyte is determined by adding the measured charge and extrapolated charge.

[0189] Current values (or other measured values) can also be used to monitor the operation of the sensor and meter. For example, ratios of current values can be checked to determine if there has been some failure of the measurement (e.g., the sensor has been pulled out of the measuring device, etc.). The acceptable ratio range will typically depend on the type and components of the sensor, the rate at which measurements are made, and the acceptable noise level. As an example, an acceptable ratio range could be, for example, 2:1 to 1:2 for immediately adjacent measurements.

[0190] The discussion above has focused on the extrapolation of charge required to complete electrolysis of an analyte in a sample using the slope of the natural logarithm of current versus time under diffusion limited conditions. Those skilled in the art will recognize that this slope is related to a Effective Diffusion Coefficient, and that other linear or non-linear mathematical functions related to the Effective Diffusion Coefficient can also be used to extrapolate the charge.

[0191] The extrapolation of current measurements and the determination of charge passed or projected to pass, as well as the concentration of the analyte, and other functions can be performed by a processor with or without a storage medium, in which the desired procedures are performed by software, hardware, or a combination thereof. According to another embodiment, these same procedures are accomplished using discrete or semi-programmable hardware configured, for example, using a hardware descriptive language, such as Verilog. In yet another embodiment, the procedures are performed using a processor having at least one look-up table arrangement with data stored therein to represent the complete result or partial results of the above equations based on a given set of input data.

[0192] Current measurements can be made by a variety of methods. For example, the current measurements can be made by measuring the current directly using any known techniques and devices.

[0193] Another method of measuring current includes discharging a known amount of charge into the sample (by electrolysis of the analyte) and measuring the time required for the discharge. Current is then calculated as the quotient of charge and discharge time. As an example, a capacitor can

be charged by circuitry in the meter and then coupled to the working or counter electrode(s) to discharge by electrolysis of the analyte. The time to discharge to a threshold level can be measured using, for example, a clock circuit that is part of the processing circuitry of the meter. The use of a clock circuit permits very accurate measurements of time. This can be advantageous over designs that directly measure current or charge and must convert these analog measurements to processible digital representations using the more expensive A/D (analog-to-digital) converters.

[0194] The accuracy of the charge and time measurement, which provides the current values, affects the accuracy of the final analyte concentration measurement. Although it is desired to use a high-quality and highly accurate measuring device, the capacitor used to store the charge can be a high-quality, expensive capacitor that has a precisely known capacitance. Alternatively, an accurate charge measurement can be obtained by using a calibrating resistor in parallel with the capacitor. During a calibration period, the capacitor discharges through the resistor and the current through the resistor or the time constant of the discharge can be determined. This can then be used to determine the charge stored in the capacitor. A high-quality, very accurate resistor is typically less expensive than a similarly accurate capacitor, and will provide the same amount of accuracy to the charge measurement.

[0195] The charge, Q , no matter how determined, is then used to calculate the concentration of the analyte (C_A) by the following equation (when the redox mediator is non-leachable):

$$C_A = Q/nFV \quad (3a)$$

[0196] where n is the number of electron equivalents required to electrolyze the analyte, F is Faraday's constant (approximately 96,500 coulombs per equivalent), and V is the volume of sample in the measurement zone. When using a diffusible mediator, the concentration of the analyte can be obtained from the following equation:

$$C_A = (Q_{\text{tot}} - Q_{\text{back}})/nFV \quad (3b)$$

[0197] where Q_{tot} is the total charge transferred during the measurement and Q_{back} is the amount of charge transferred that was not due to the analyte, e.g., charge transferred by the shuttling of the diffusible mediator between the working electrode and the counter electrode. In at least some instances, the sensor is constructed so that the background charge is at most 5 times the size of the charge generated by electrolysis of an amount of analyte. Preferably, the background signal is at most 200%, 100%, 50%, 25%, 10%, or 5% of the charge generated by electrolysis of the analyte.

[0198] One example of a method for determining the ratio of background signal to signal generated by electrolysis of the analyte is described as follows. If the shuttling of the redox mediator is not disabled by the applied potential, the charge that results from the shuttling may be represented by the following formula:

$$Q_{\text{back}} = (AFD_M C_M d)(t_M)$$

[0199] where A is the area of the working electrode; F is Faraday's constant (96,500 coulombs/equivalent); D_M is the effective diffusion coefficient of the redox mediator; C_M is the concentration of the redox mediator in the measurement zone; d is the distance separating facing electrodes; t is the

amount of time for the measurement; and n_M is the number of electrons gained or lost by the redox mediator.

[0200] Additionally, the charge of the analyte, when the analyte is electrooxidized to about 90% completion in the measurement period, may be represented by the following formula:

$$Q_G = Ad(0.90)C_G n_G F$$

[0201] where A is the area of the working electrode; d is the distance separating facing electrodes; C_G is the concentration of glucose; n is the number of electrons needed to electrolyze the analyte (e.g., 2 electrons per glucose molecule); and F is Faraday's constant. For glucose, when C_G is 5 mM (or 5×10^{-6} moles/cm³), t is 60 seconds, n_G is 2, and n_M is 1, the ratio of charge from the redox mediator to the charge from electrooxidation of the analyte may be represented by the following formula:

$$\frac{Q_{\text{Back}}/Q_G}{(6.7 \times 10^9)} = \frac{(D_M C_M / d^2)(t n_M (0.9 n_G C_G))}{(D_M C_M / d^2) \times 5}$$

[0202] For example, if the ratio of Q_{Back}/Q_G is 5, then $(D_M C_M)/d^2$ is 7.5×10^{-7} moles/(cm³ sec). Also for example, if the ratio of Q_{Back}/Q_G is 1, then $(D_M C_M)/d^2$ is 1.5×10^{-7} moles/(cm³ sec). Still another example, if the ratio is 0.1, then $(D_M C_M)/d^2$ is 1.5×10^{-8} moles/(cm³ sec). Thus, depending on the ratio desired, a sensor may be configured to have the desired ratio by choosing D_M , C_M , and d accordingly.

[0203] The ratio can be affected, for example, by reducing the concentration of the redox mediator (i.e., C_M may be reduced). Alternatively, or additionally, the diffusion of the redox mediator may be reduced. Other sensor configurations are also suitable for controlling the ratio of background signal to signal generated by the analyte and will be described below.

[0204] The background charge, Q_{back} , can be accounted for in a variety of ways. Q_{back} can be made small, for example, by using only limited amounts of diffusible redox mediator; by providing a membrane over the counter electrode that limits diffusion of the redox mediator to the counter electrode; or by having a relatively small potential difference between the working electrode and the counter electrode. Other examples of sensor configurations and methods suitable for reducing Q_{back} include having a redox mediator reaction rate at the working electrode that is significantly faster than that at the counter electrode; immobilizing the redox mediator on the working electrode; having the redox mediator become immobilized on the counter or counter/reference electrode upon its reaction at the counter or counter/reference electrode; or slowing the diffusion of the redox mediator.

[0205] For coulometric measurements, at least 20% of the analyte is electrolyzed, preferably at least 50%, more preferably at least 80%, and even more preferably at least 90% of the analyte is electrolyzed. In one embodiment of the invention, the analyte is completely or nearly completely electrolyzed. The charge can then be calculated from current measurements made during the electrochemical reaction, and the concentration of the analyte is determined using equation (3a) or (3b). The completion of the electrochemical reaction is typically signaled when the current reaches a steady-state value. This indicates that all or nearly all of the analyte has been electrolyzed. For this type of measurement, at least 90% of the analyte is typically electrolyzed, prefer-

ably, at least 95% of the analyte is electrolyzed and, more preferably, at least 99% of the analyte is electrolyzed.

[0206] For coulometry, it is typically desirable that the analyte be electrolyzed quickly. The speed of the electrochemical reaction depends on several factors, including the potential that is applied between the electrodes and the kinetics of reactions (1) and (2). Other significant factors include the size of the measurement zone. In general, the larger the potential, the larger the current through the cell and therefore, the faster the reaction will typically occur. However, if the potential is too large, other electrochemical reactions may introduce significant error in the measurement. Typically, the potential between the electrodes as well as the specific redox mediator and optional second electron transfer agent are chosen so that the analyte will be almost completely electrolyzed in less than 5 minutes, based on the expected concentration of the analyte in the sample. Preferably, the analyte will be almost completely electrolyzed within about 2 minutes and, more preferably, within about 1 minute.

[0207] Although coulometry has the disadvantage of requiring the volume of the measured sample be known, coulometry is a preferred technique for the analysis of the small sample because it has the advantages of, for example, no temperature dependence for the measurement, no enzyme activity dependence for the measurement, no redox-mediator activity dependence for the measurement, and no error in the measurement from depletion of analyte in the sample. As already described above, coulometry is a method for determining the amount of charge passed or projected to pass during complete or nearly complete electrolysis of the analyte. One coulometric technique involves electrolyzing the analyte on a working electrode and measuring the resulting current between the working electrode and a counter electrode at two or more times during the electrolysis. The electrolysis is complete when the current reaches a steady state. The charge used to electrolyze the sample is then calculated by integrating the measured currents over time and accounting for any background signal. Because the charge is directly related to the amount of analyte in the sample there is no temperature dependence of the measurement. In addition, the activity of the enzyme does not affect the value of the measurement, but only the time required to obtain the measurement (i.e., less active enzyme requires a longer time to achieve complete electrolysis of the sample) so that decay of the enzyme over time will not render the analyte concentration determination inaccurate. And finally, the depletion of the analyte in the sample by electrolysis is not a source of error, but rather the objective of the technique. (However, the analyte need not be completely electrolyzed if the electrolysis curve is extrapolated from the partial electrolysis curve based on well-known electrochemical principles.)

[0208] It may be desirable in some instances to utilize non-coulometric assays, such as amperometric or potentiometric measurement techniques. "Amperometry" and "chronopotentiometry" refer to taking a potentiometric measurement at one or more points in time. These measurement techniques are useful when the volume of the measured sample is unknown; the volume of the sample in the measurement zone of a small volume sensor (i.e., no more than one microliter) may be difficult to accurately reproduce if the manufacturing tolerances of one or more dimensions of the

measurement zone have significant variances. For information regarding using non-coulometric measuring, such as amperometric and potentiometric techniques, see, for example, U.S. application Ser. No. 09/295,962, filed Apr. 21, 1999, incorporated herein by reference in its entirety.

[0209] Heating of Sample

[0210] The sample can be heated to increase the rate of diffusion, oxidation, or reduction of the analyte. This heating can be accomplished by a variety of techniques including placing the sensor in a heated environment or applying a heating unit to the sensor.

[0211] Another technique includes providing a thermal heating element, such as, for example, a wire or an ink that is capable of converting electrical energy into heat energy, on the sensor. This wire or ink can be applied, for example, on the opposite side of a base material, such as a polymer film, from one or more of the working, counter, reference, or counter/reference electrodes, or applied around the periphery of the working, counter, reference, or counter/reference electrodes. In some instances, the sample is heated up to 5 to 20° C. above an initial temperature. In other instances, the temperature of the sample may not be known but a constant amount of power or current is applied to the wire or ink.

[0212] General Examples

[0213] The invention will be further characterized by the following general examples. These examples are not meant to limit the scope of the invention which has been fully set forth in the foregoing description. Variations within the concepts of the invention are apparent to those skilled in the art.

[0214] One method for manufacturing the sensor shown in FIGS. 5A through 5C, having a working electrode, multiple counter/reference electrodes, and a side filled sample chamber, is described below. According to the method described herein, a plurality of sensors are produced and then separated.

[0215] A first non-conducting substrate, preferably polyester, is provided. A conductive carbon ink, such as that available from Ercon of Wareham, Mass. under the trade designation "Graphite #G4491", is screen printed onto a first half of the first polymeric substrate in the pattern shown in FIG. 5A and dried, to provide the working electrode 422, trace 453 and contact pad 423. Once the substrate web is cut to the final size, each working electrode 422 will extend across the width of the sensor and have a trace 453 that extends to the far end of the sensor. As an example, a sensor can be made with a width of about 6.1 mm (0.240 inch). Typical sensor widths are about 2 mm to 20 mm, preferably about 4 mm to 10 mm, but wider or narrower sensors can be used as desired.

[0216] A sensing layer, as described in U.S. patent application Ser. No. 09/295,962, comprising a diffusible mediator and diffusible glucose dehydrogenase enzyme, is applied to the working electrode in a sample chamber region. The sensing layer can be provided on the electrode prior to or subsequent to providing the sample chamber. On the other half of the polyester substrate, a conductive silver/silver chloride ink, such as that commercially available from Ercon under the trade designation "Silver/Silver Chloride #R414", is screen printed onto a second half of the first

polymeric substrate in the pattern shown in FIG. 5B and dried. In the embodiment shown in FIG. 5B, the Ag/AgCl ink provides a first indicator or reference electrode 441, a counter electrode 424, and a second indicator electrode 442. Counter electrode may extend across the entire sensor surface, or may extend only a portion thereof. As an example, the width of a counter electrode is about 1 mm to 20 mm, preferably about 2 to 10 mm. In one sensor configuration, the counter electrode 424 has a width of about 3.71 mm (0.146 inch). The indicator electrodes can have a width, for example, of about 0.1 mm to 5 mm, preferably about 0.25 mm to about 1 mm.

[0217] In one sensor configuration, working electrode, such as working electrode 422 shown in FIG. 5A, extends the width of the sensor, which is about 6.1 mm (0.240 inch). A counter electrode, such as counter electrode 424, has a width of about 3.71 mm (0.146 inch). Indicator electrodes, such as indicator electrodes 441, 442 in FIG. 5B, each can have a width of about 0.51 mm (0.020 inch). A spacing of 0.30 mm (0.012 inch) is present between these indicator electrodes and the counter electrode. The indicator electrodes may be each set back from the eventual side edge of the sensor strip, for example, by approximately 0.38 mm (0.015 inch), so that the electrodes 441, 442 are not damaged, for example, smeared, streaked, chipped or the like, by the process of slitting the substrate into individual sensors.

[0218] The invention has been described with reference to various specific and preferred embodiments and techniques. However, it will be apparent to one of ordinary skill in the art that many variations and modifications may be made while remaining within the spirit and scope of the invention.

[0219] All publications and patent applications in this specification are indicative of the level of ordinary skill in the art to which this invention pertains. All publications and patent applications are herein incorporated by reference to the same extent as if each individual publication or patent application was specifically and individually incorporated by reference.

We claim:

1. A sensor comprising:

- a first substrate having a first major surface and a second major surface opposing the first major surface;
- a second substrate having a first major surface and a second major surface opposing the first major surface, the first and second substrates being disposed so that the first major surface of the first substrate is in facing relationship with the first major surface of the second substrate;
- a working electrode disposed on the first major surface of the first substrate;
- a counter electrode disposed on the first major surface of one of the first substrate and the second substrate; and
- an insertion monitor disposed on the second major surface of one of the first substrate and the second substrate, the insertion monitor being configured and arranged to provide a path for electrical current between at least two contact leads of a meter for indication that the sensor has been properly inserted into the meter.

2. The sensor according to claim 1, wherein the insertion monitor is disposed on the second major surface of the first substrate.

3. The sensor according to claim 1, wherein the counter electrode is disposed on the first major surface of the second substrate.

4. The sensor according to claim 1, wherein the sensor has a width, and the insertion monitor is a conductive stripe extending across the width of the sensor.

5. The sensor according to claim 1, wherein the insertion monitor has two or more contact regions for electrical contact with the meter.

6. The sensor of claim 1, wherein the insertion monitor is configured and arranged to provide encoded information about the strip.

7. A sensor strip comprising:

a first substrate having a first major surface and a second major surface opposing the first major surface;

a second substrate having a first major surface and a second major surface opposing the first major surface, the first and second substrates being disposed so that the first major surface of the first substrate is in facing relationship with the first major surface of the second substrate;

a working electrode disposed on the first major surface of the first substrate;

a counter electrode disposed on the first major surface of one of the first substrate and the second substrate; and

a plurality of contact pads disposed on the first and second substrates, at least one of the contact pads being in electrical communication with the working electrode and at least one other contact pad being in electrical communication with the counter electrode, wherein each contact pad has a contact pad width measured parallel to a width of the first and second substrates, and wherein the sum of the contact pad widths is greater than the width of the first substrate and greater than the width of the second substrate.

8. The sensor strip of claim 7 wherein the plurality of contact pads are disposed on at least three of the following surfaces: the first major surface of the first substrate, the second major surface of the first substrate, the first major surface of the second substrate, and the second major surface of the second substrate.

9. A connector comprising:

a plurality of contact structures for releasable connection to an electrochemical analyte sensor, wherein each contact lead comprises

a proximal end for electrical connection to a sensor contact, and

a distal end for electrical connection to an electrical device; and

wherein the plurality of contact leads comprises

one or more first contact leads extending longitudinally from the distal end to the proximal end, and

one or more second contact leads extending longitudinally from the distal end past the proximal end of the one or more first contact leads and angling toward a longitudinal center line of the sensor.

10. The connector of claim 9, wherein the connector comprises at least two second contact leads to make electrical contact with a single conductive surface of the sensor.

11. The connector of claim 10, wherein the first contact leads are configured and arranged to contact at least one of a working electrode and a counter electrode of a sensor, and wherein the second contact leads are configured and arranged to contact an insertion monitor of a sensor.

12. A method for the manufacture of a sensor comprising the steps of:

providing an adhesive having first and second surfaces covered with first and second release liners;

cutting through the first release liner but not through the second release liner to define a plurality of sample chamber regions;

removing a portion of the first release liner to expose a portion of the first adhesive surface leaving a remaining portion of the first release liner over the sample chamber regions;

applying the exposed first adhesive surface to a first substrate;

removing the second release liner together with the adhesive and first release liner of the sample chamber regions to expose the second adhesive surface and to expose the first substrate in the sample chamber region; and

applying a second substrate over the second adhesive surface forming a sample chamber in the sample chamber region, wherein a plurality of conductive traces are disposed on the first substrate, the second substrate, or a combination thereof, the conductive traces defining at least one working electrode and at least one counter electrode, and the sample chambers are disposed to allow an analyte sample disposed in the sample chambers to be electrolyzed by a current flowing between the working electrode and the counter electrode; and

separating a plurality of electrochemical sensors, each electrochemical sensor comprising at least one of the working electrodes, at least one of the counter electrodes, and at least one of the sample chambers.

13. The method of claim 12 wherein the working electrodes are disposed on one of the substrates and the counter electrodes are disposed on the other substrate.

14. A method for determining a concentration of an analyte in a body fluid sample, the method comprising steps of:

disposing a body fluid sample in a sample chamber of an electrochemical sensor, the electrochemical sensor further comprising at least one working electrode and at least one counter electrode for electrolyzing an analyte in the body fluid sample in the sample chamber of the electrochemical sensor;

electrolyzing at least a portion of an analyte in the body fluid sample by causing a current to flow between the at least one working electrode and the at least one counter electrode;

determining, during the electrolyzing of the analyte, a plurality of current values, at least some of the current

values being obtained while the electrolysis of the analyte in the sensor is substantially diffusion limited;

determining, from the current values obtained while the electrolysis of the analyte in the sensor is substantially diffusion limited, parameters for extrapolation of a relationship between the current values and time;

determining from the plurality of current values the amount of an actual charge consumed in the partial electrolysis of the analyte in the sample;

determining, from the parameters for extrapolation of the relationship between the current values and time, an extrapolated charge required to electrolyze the analyte remaining in the sample;

determining the concentration of the analyte in the body fluid sample from the actual charge and the extrapolated charge.

15. The method of claim 14, wherein the step of determining parameters for extrapolation of the relationship between current values and time comprises determining a value related to a diffusion coefficient of the analyte in the sensor.

16. The method of claim 14, wherein the step of determining parameters for extrapolation of the relationship between current values and time comprises determining a slope of a natural logarithm of current values versus time obtained while the electrolysis of the analyte in the sensor is substantially diffusion limited.

17. An electrochemical sensor device comprising:

(a) an electrochemical sensor comprising

(i) at least one working electrode,

(ii) at least one counter electrode, and

(iii) a sample chamber for holding a body fluid sample in electrolytic contact with the at least one working electrode and at least one counter electrode; and

(b) a processor coupled to the electrochemical sensor, the processor being configured and arranged to

(i) electrolyze at least a portion of an analyte in the body fluid sample by causing a current to flow between the at least one working electrode and at least one counter electrode;

(ii) determine, during the electrolyzing of the analyte, a plurality of current values, at least some of the current values being obtained while the electrolysis of the analyte in the sensor is substantially diffusion limited;

(iii) determine, from the current values obtained while the electrolysis of the analyte in the sensor is substantially diffusion limited, parameters for extrapolation of a relationship between the current values and time;

(iv) determine the amount of an actual charge consumed in the partial electrolysis of the analyte in the sample;

(v) determine an extrapolated charge required to electrolyze the analyte remaining in the sample; and

(vi) determine, from the actual charge and the extrapolated charge, the concentration of the analyte in the body fluid sample.

18. A method of determining an amount of current used to electrolyze a portion of an analyte in a body fluid sample disposed in an electrochemical sensor, the method comprising:

discharging an amount of charge into the body fluid sample disposed in the electrochemical sensor and electrolyzing the analyte;

determining an amount of time needed to discharge the amount of charge; and

determining the current used to electrolyze the portion of the analyte using the amount of charge and the amount of time.

19. The method of claim 18 wherein the charge is provided by the discharge of a capacitor.

20. The method of claim 19, further comprising calibrating the capacitor by discharging the capacitor through a resistor of known resistance and measuring either a current or a time constant associated with the discharge.

21. A n electrochemical sensor device, comprising:

(a) an electrochemical sensor comprising

(i) at least one working electrode,

(ii) at least one counter electrode, and

(iii) a sample chamber for holding a body fluid sample in electrolytic contact with the at least one working electrode and at least one counter electrode; and

(b) a measuring device comprising

(i) a capacitor configured and arranged for coupling to the at least one working electrode to repeatedly deliver an amount of charge by repeatedly discharging, for electrolyzing at least a portion of an analyte in a body fluid disposed in the sample chamber,

(ii) a clock to measure discharge times of the capacitor, and

(iii) a processor to determine a concentration of the analyte in the body fluid based on the discharge times.

22. A sensor comprising:

a first substrate having a first major surface and a second major surface opposing the first major surface;

a second substrate having a first major surface and a second major surface opposing the first major surface, the first and second substrates being disposed so that the first major surface of the first substrate is in facing relationship with the first major surface of the second substrate;

a working electrode disposed on the first major surface of the first substrate;

a counter electrode disposed on the first major surface of one of the first substrate and the second substrate; and

a conductive trace positioned on the second major surface of one of the first substrate and the second substrate, the conductive trace electrically connected to two or more contact pads for electrical contact with a connector, the conductive trace configured and arranged to close an electrical circuit, and wherein conductive trace has a resistance that carries information related to the sensor.

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