



US 20020168290A1

(19) **United States**

(12) **Patent Application Publication**
Yuzhakov et al.

(10) **Pub. No.: US 2002/0168290 A1**

(43) **Pub. Date: Nov. 14, 2002**

(54) **PHYSIOLOGICAL SAMPLE COLLECTION
DEVICES AND METHODS OF USING THE
SAME**

filed on May 10, 2001. Provisional application No.
60/290,156, filed on May 10, 2001.

Publication Classification

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(51) **Int. Cl.⁷** **A61B 5/00**; G01N 31/22

(52) **U.S. Cl.** **422/56**; 600/583; 204/403.01;
422/82.05; 435/14

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(21) Appl. No.: **10/143,399**

(22) Filed: **May 9, 2002**

Related U.S. Application Data

(60) Provisional application No. 60/290,139, filed on May
10, 2001. Provisional application No. 60/290,140,

(57) **ABSTRACT**

Devices, systems and methods are provided for piercing the skin, accessing and collecting physiological sample therein, and measuring a characteristic, e.g., an analyte concentration, of the sampled physiological sample. The subject devices are in the form of a test strip which include a biosensor and at least one skin-piercing element which is a planar extension of a portion of the biosensor. At least one fluid pathway resides within a portion of the biosensor and within the skin-piercing element. The skin-piercing element has a space-defining configuration therein which acts as a sample fluid pooling area upon penetration into the skin. Systems are provided which include one or more test strip devices and a meter for making analyte concentration measurements. Methods for using the devices and systems are also provided.

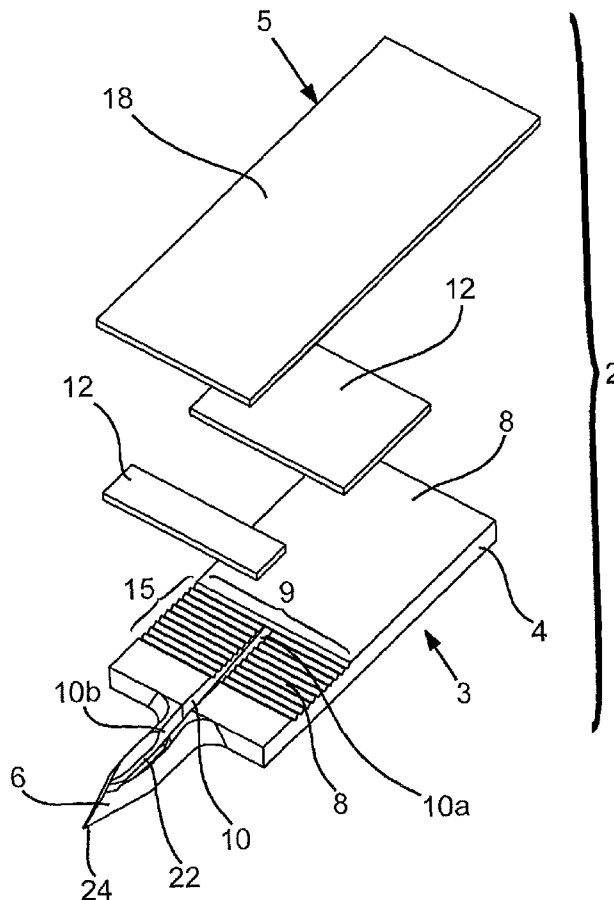


FIG. 1B

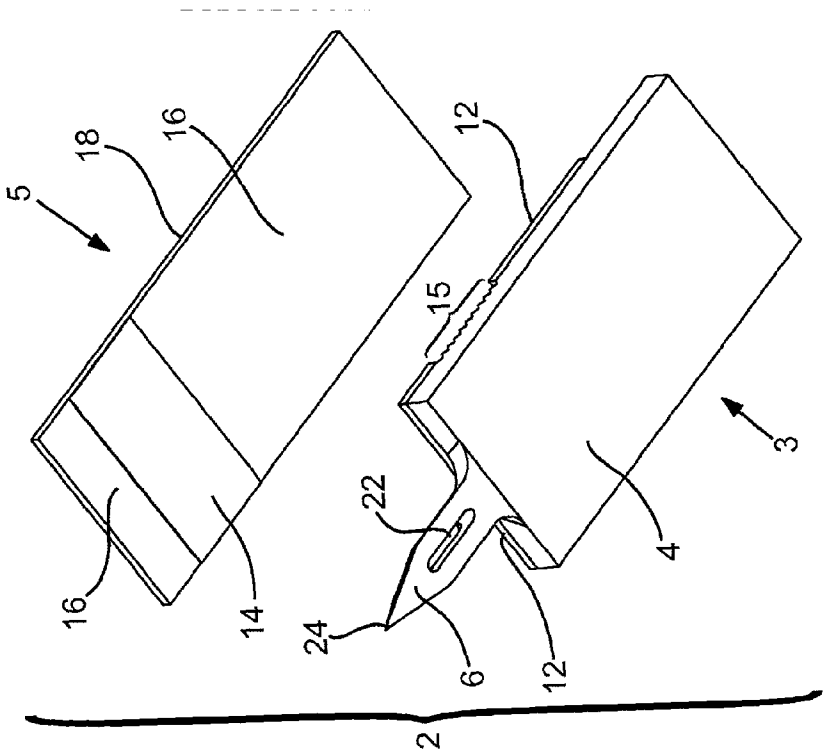
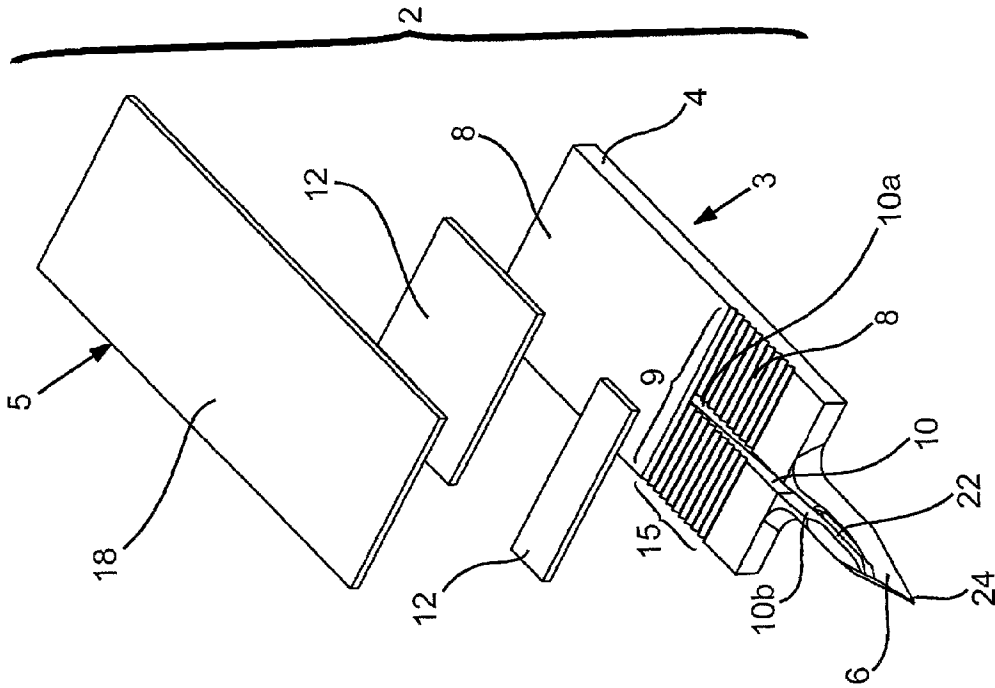


FIG. 1A



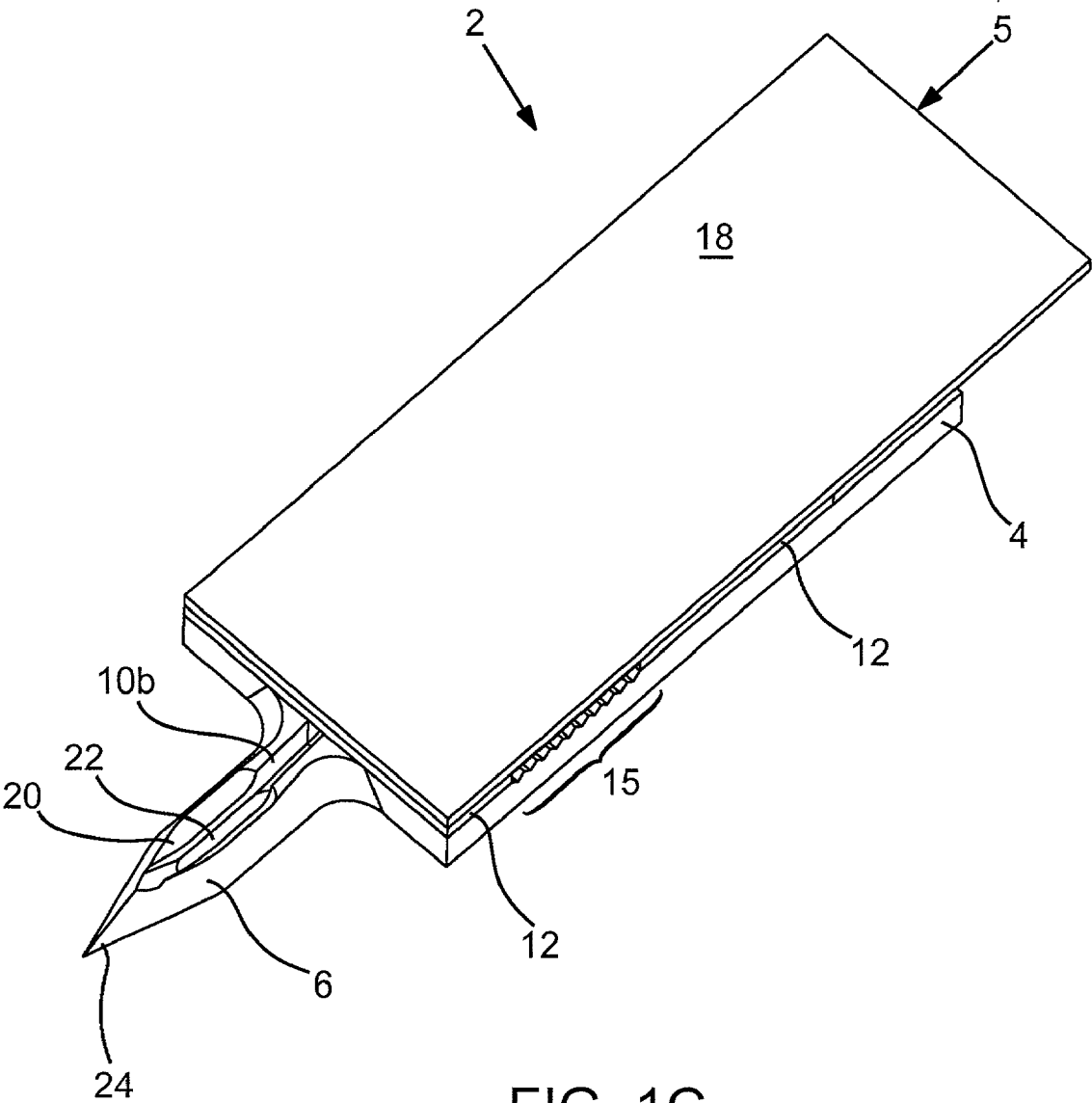


FIG. 1C

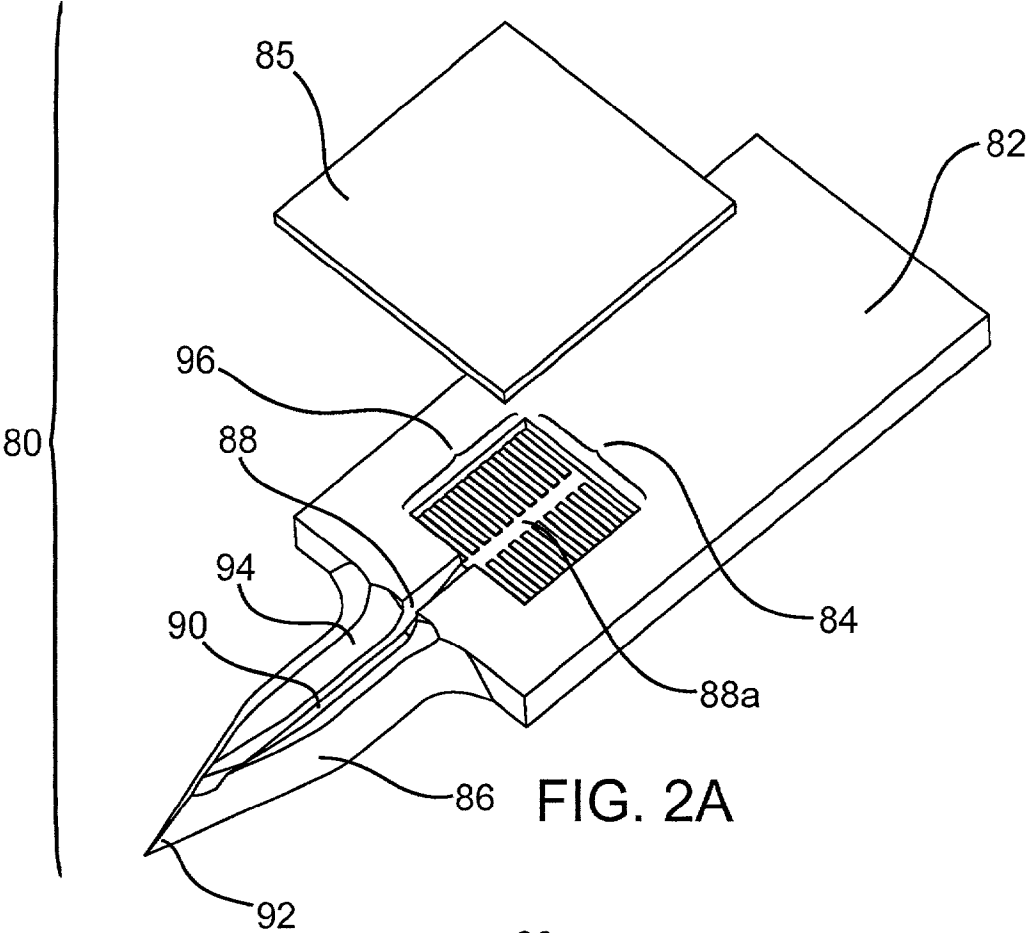


FIG. 2A

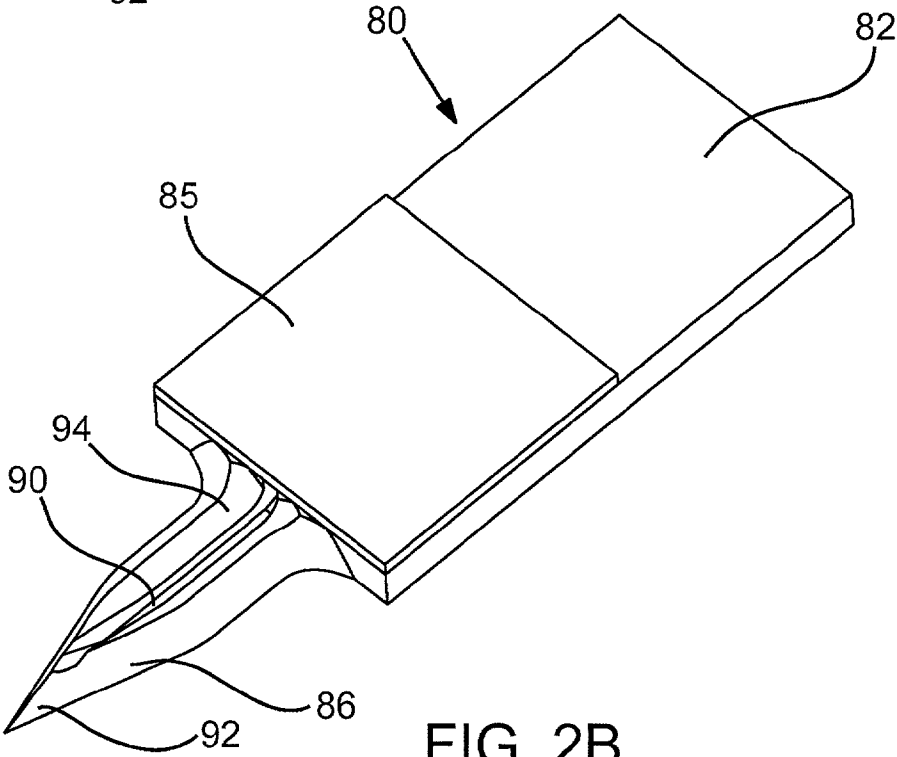
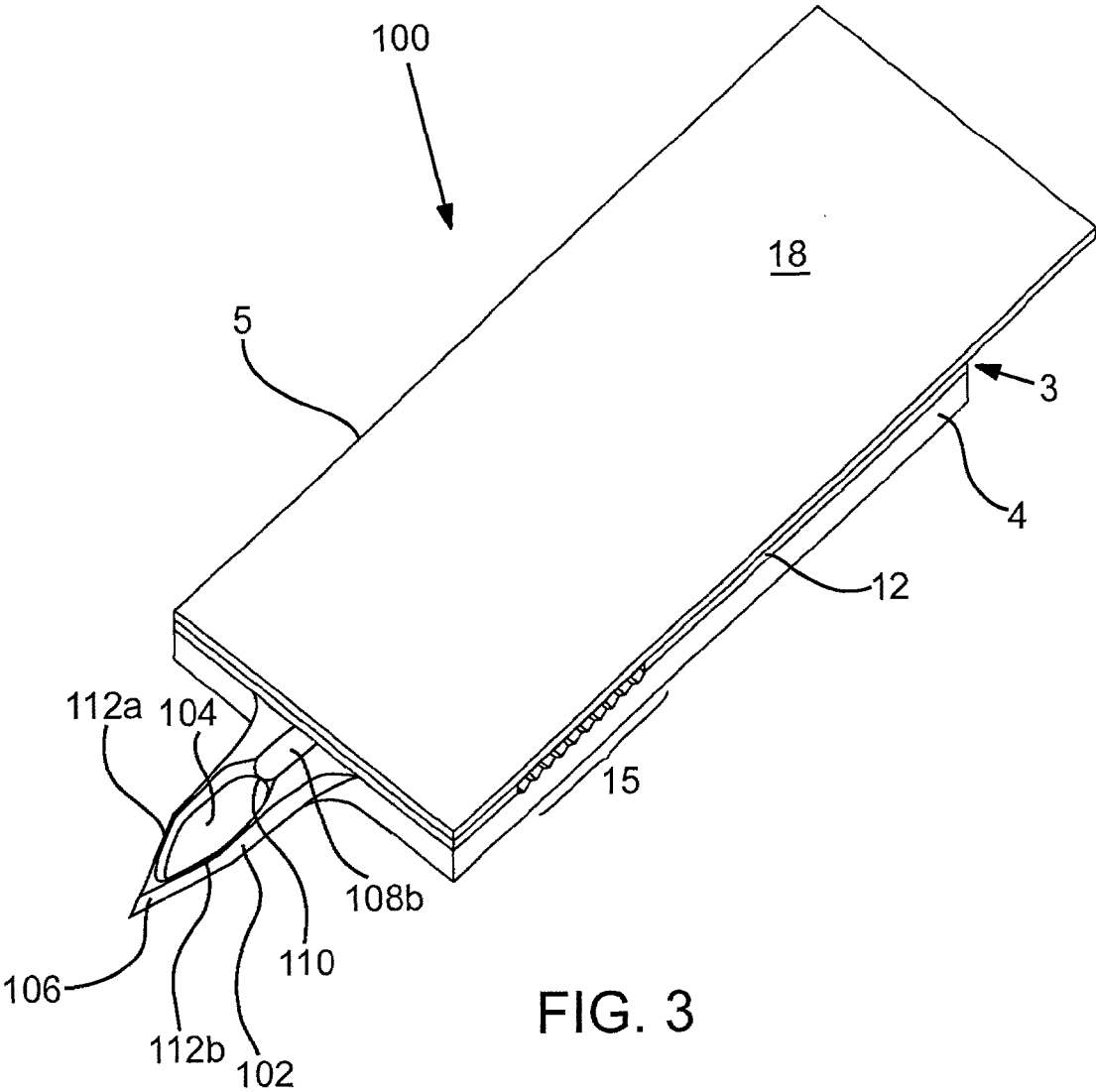


FIG. 2B



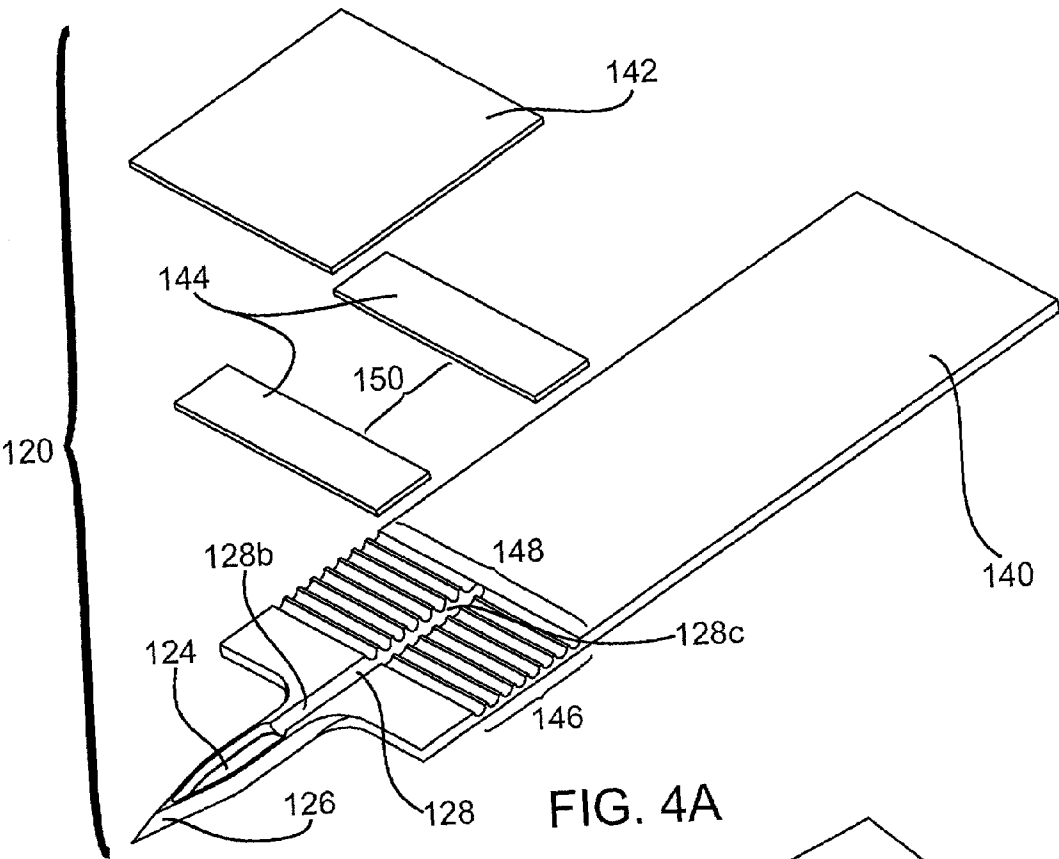


FIG. 4A

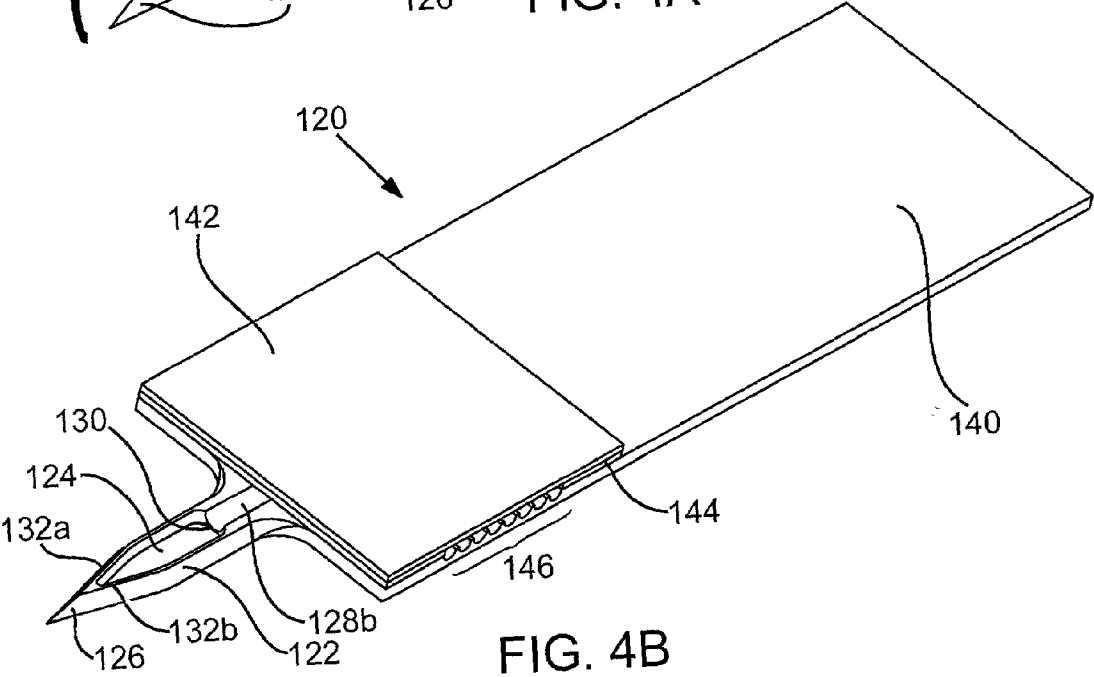
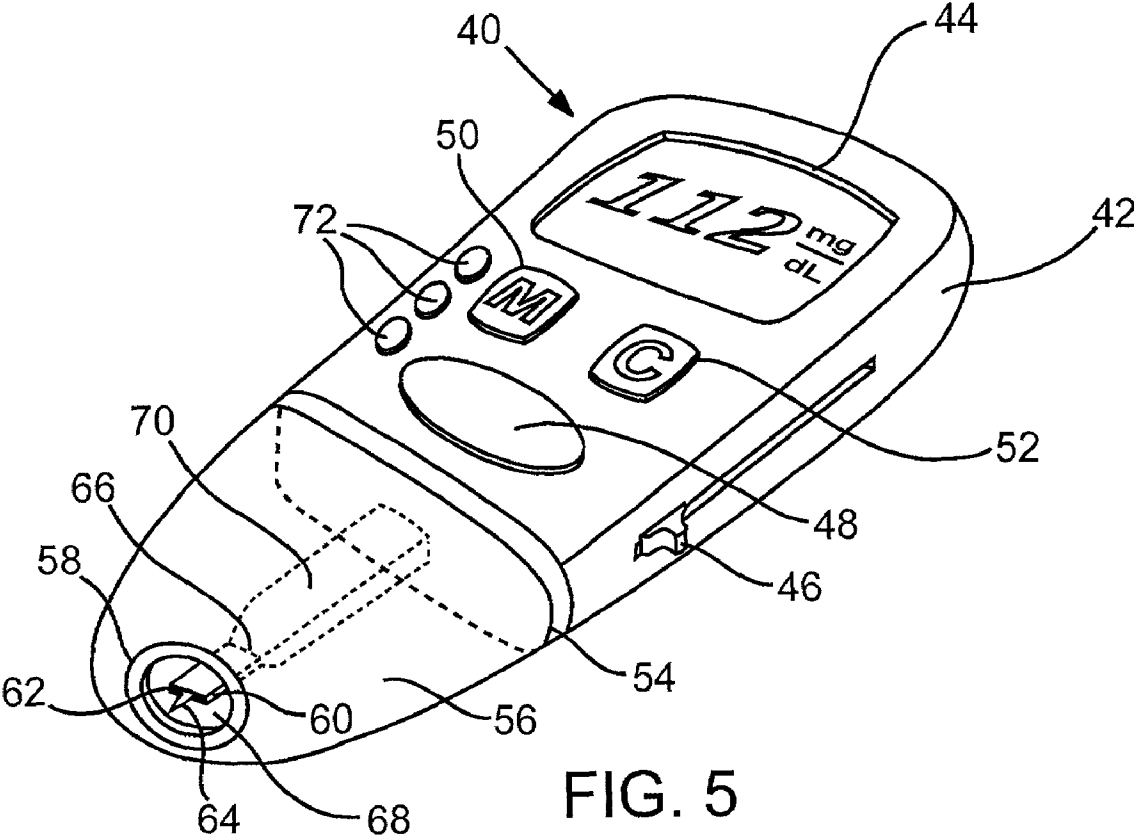


FIG. 4B



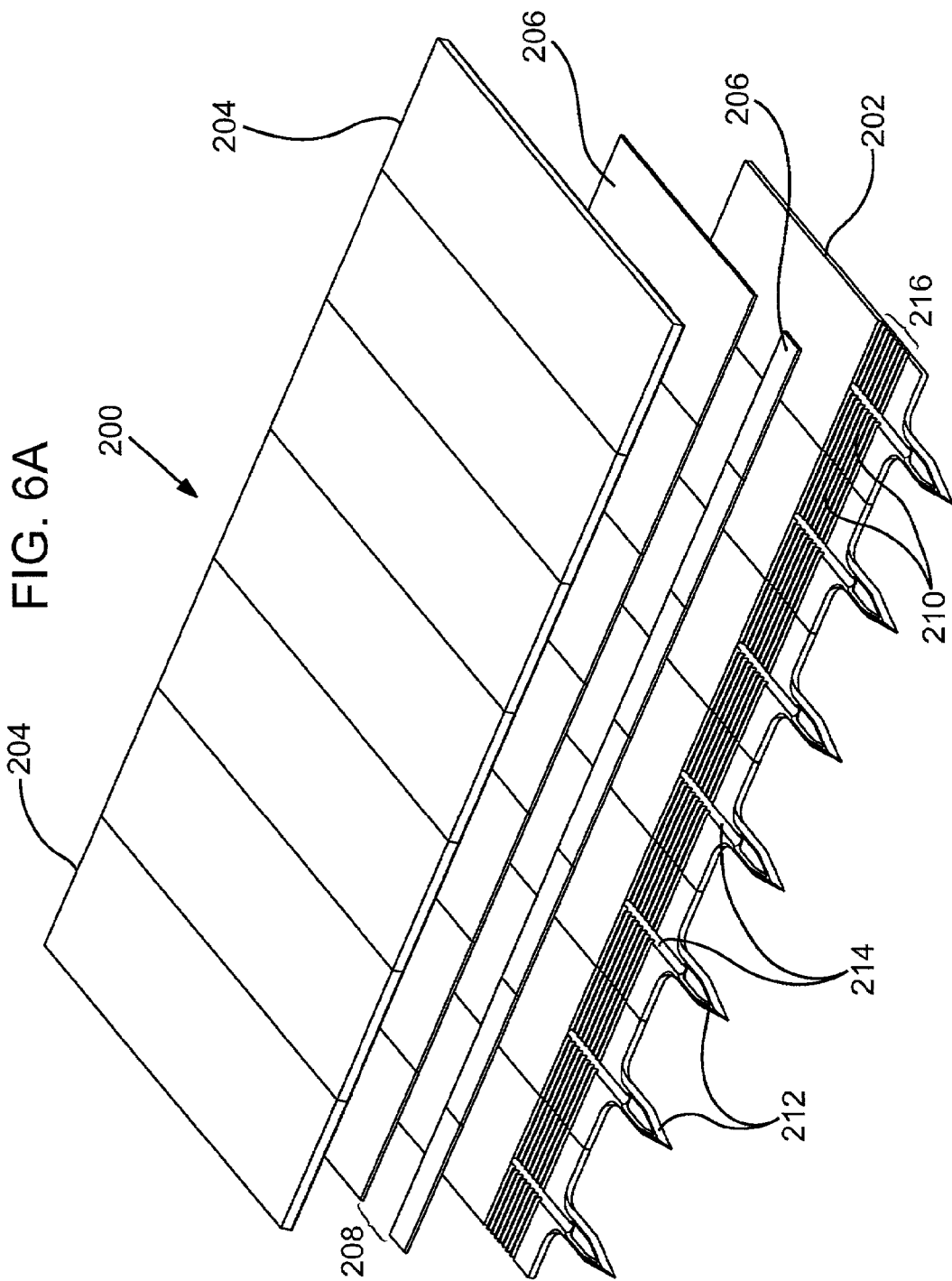


FIG. 6B

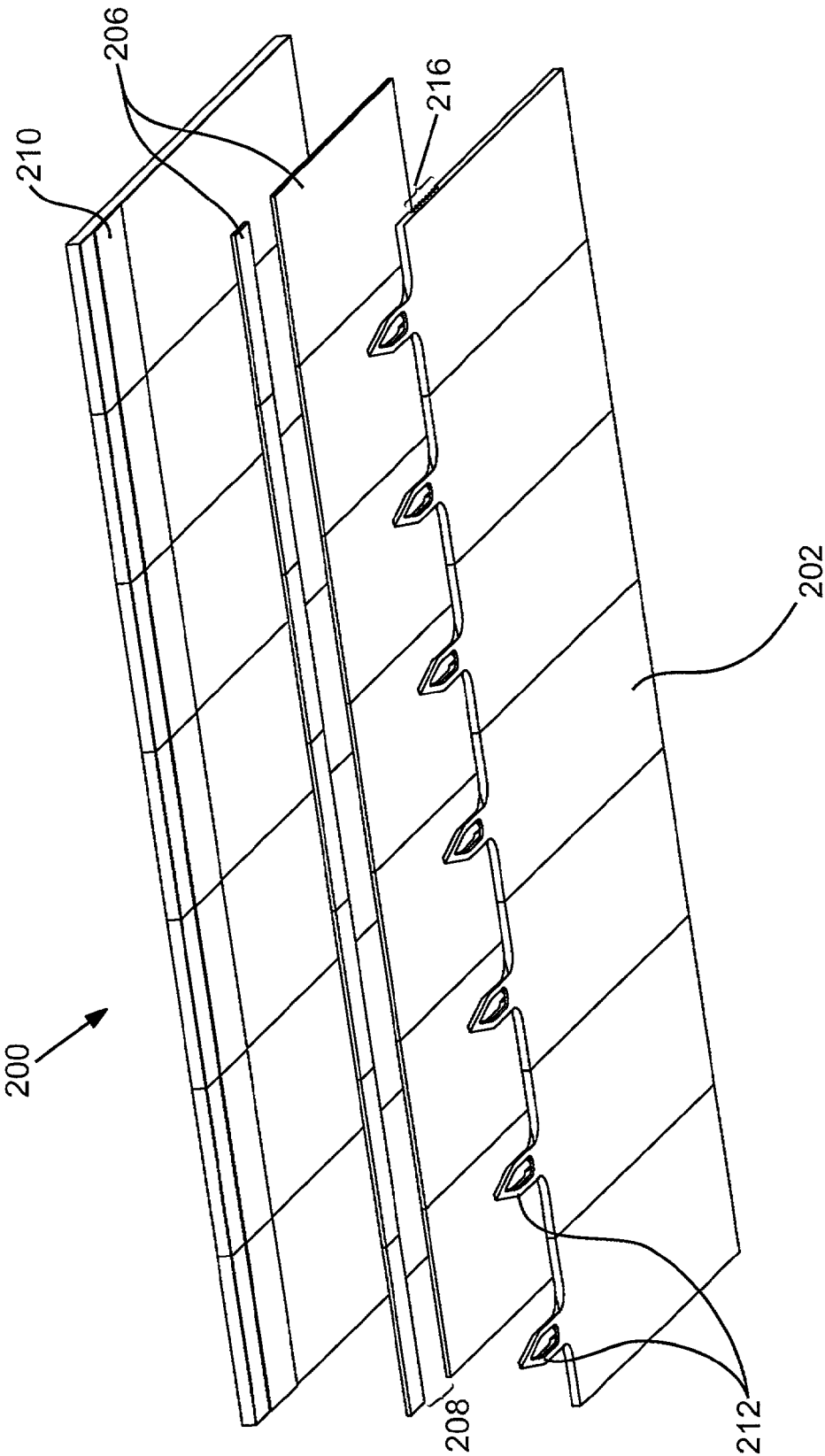


FIG. 6C

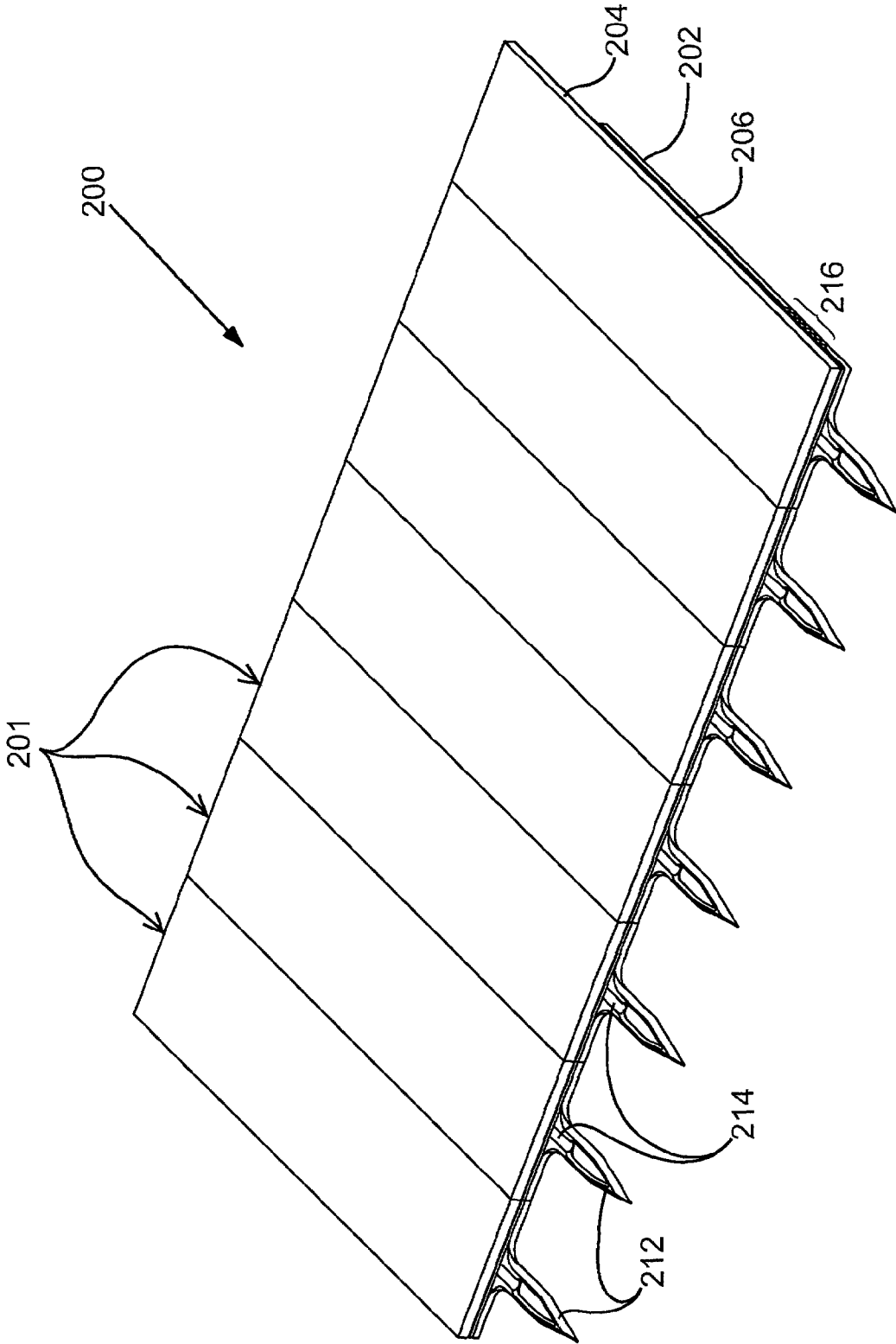
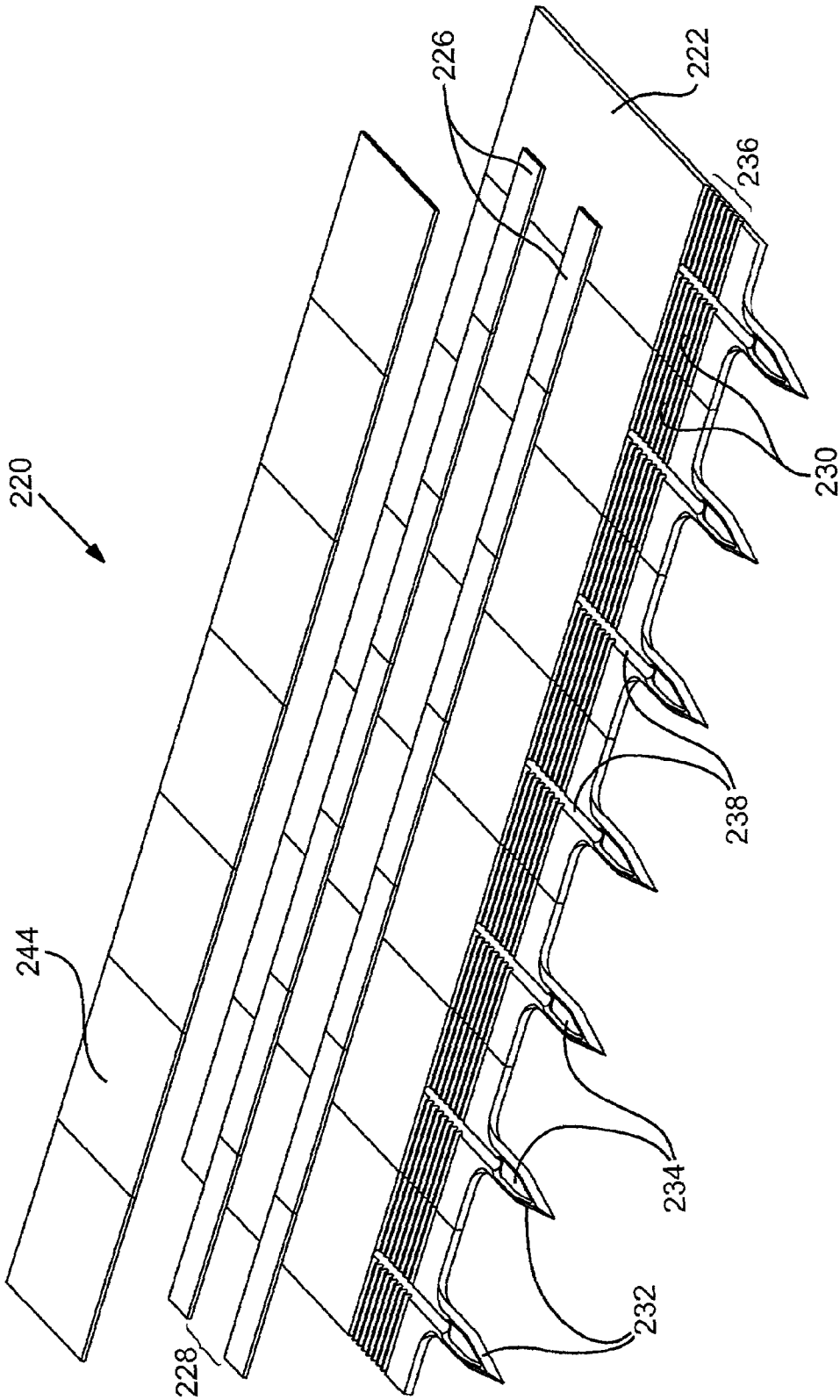


FIG. 7A



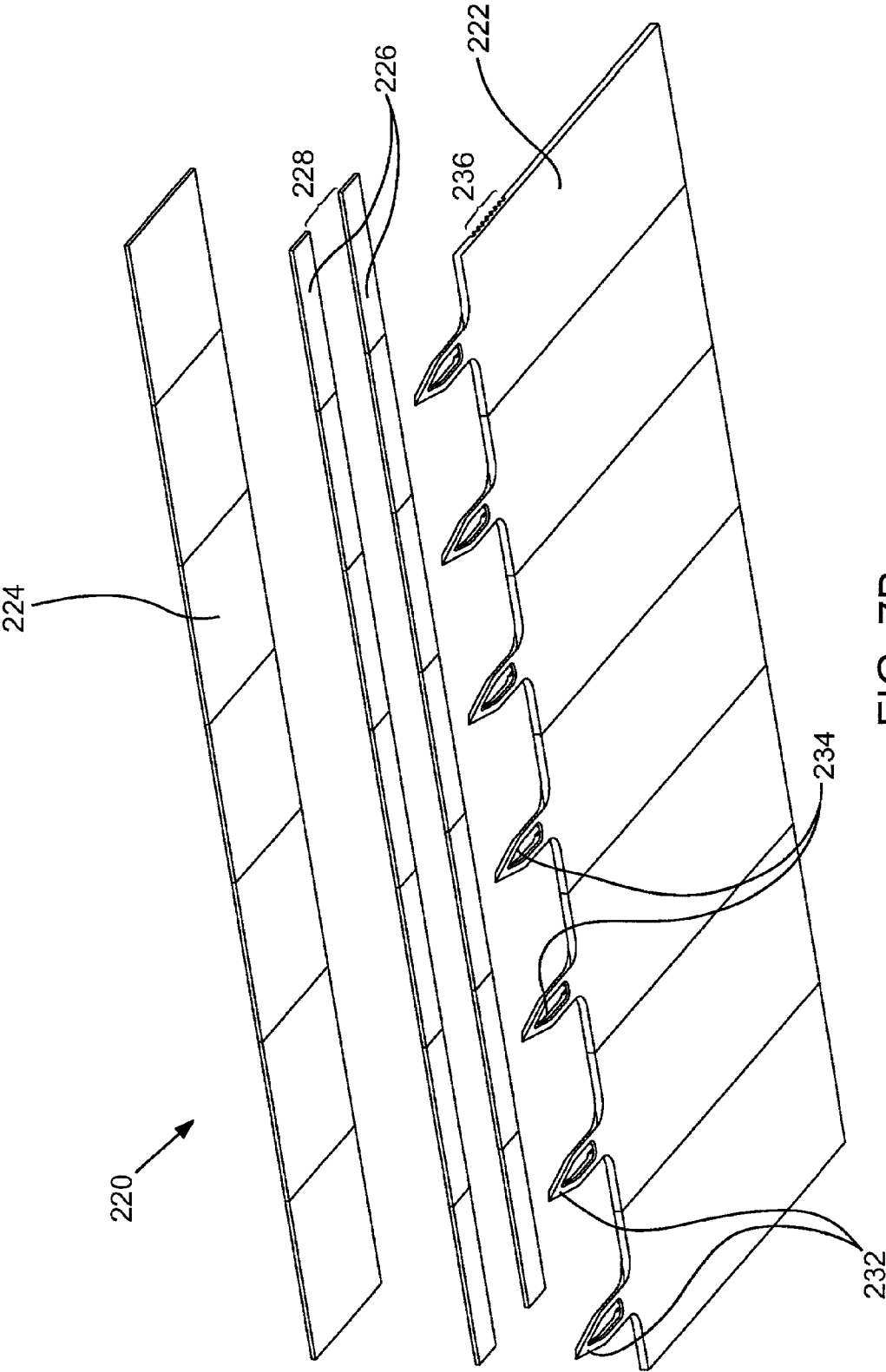
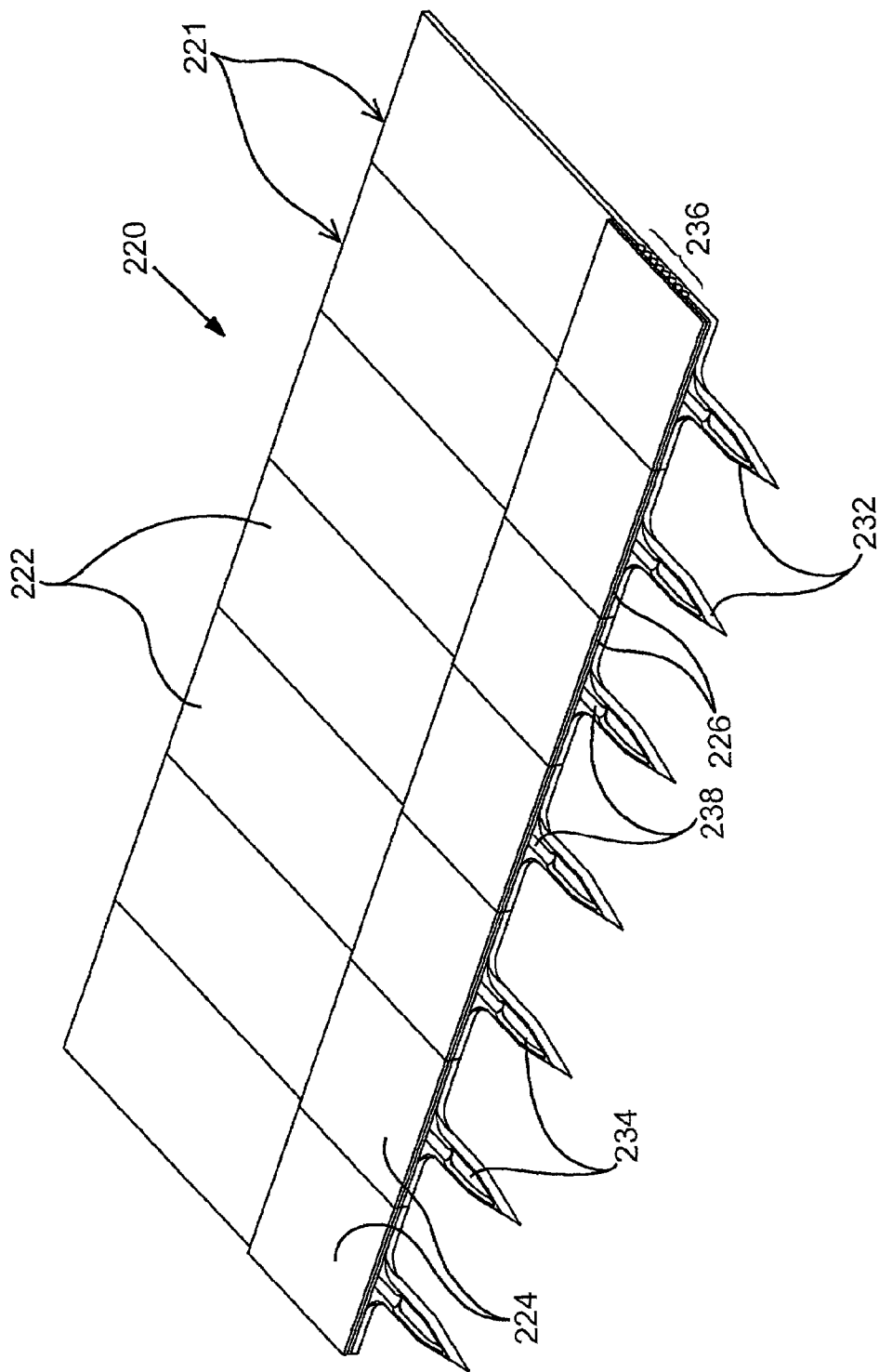


FIG. 7B

FIG. 7C



PHYSIOLOGICAL SAMPLE COLLECTION DEVICES AND METHODS OF USING THE SAME

FIELD OF THE INVENTION

[0001] The field of this invention is the collection of physiological samples and the determination of analyte concentrations therein.

BACKGROUND OF THE INVENTION

[0002] Analyte concentration determination in physiological samples is of ever increasing importance to today's society. Such assays find use in a variety of application settings, including clinical laboratory testing, home testing, etc., where the results of such testing play a prominent role in the diagnosis and management of a variety of disease conditions. Analytes of interest include glucose for diabetes management, cholesterol for monitoring cardiovascular conditions, and the like. In response to this growing importance of analyte concentration determination, a variety of analyte concentration determination protocols and devices for both clinical and home testing have been developed.

[0003] In determining the concentration of an analyte in a physiological sample, a physiological sample must first be obtained. Obtaining the sample often involves cumbersome and complicated devices which may not be easy to use or may be costly to manufacture. Furthermore, the procedure for obtaining the sample may be painful. For example, pain is often associated with the size of the needle used to obtain the physiological sample and the depth to which the needle is inserted. Depending on the analyte and the type of test employed, a relatively large, single needle or the like is often used to extract the requisite amount of sample.

[0004] The analyte concentration determination process may also involve a multitude of steps. First, a sample is accessed by use of a skin-piercing mechanism, e.g., a needle or lancet, which accessing may also involve the use of a sample collection mechanism, e.g., a capillary tube. Next, the sample must then be transferred to a testing device, e.g., a test strip or the like, and then oftentimes the test strip is then transferred to a measuring device such as a meter. Thus, the steps of accessing the sample, collecting the sample, transferring the sample to a biosensor, and measuring the analyte concentration in the sample are often performed as separate, consecutive steps with various device and instrumentation.

[0005] Because of these disadvantages, it is not uncommon for patients who require frequent monitoring of an analyte to simply become non-compliant in monitoring themselves. With diabetics, for example, the failure to measure their glucose level on a prescribed basis results in a lack of information necessary to properly control the level of glucose. Uncontrolled glucose levels can be very dangerous and even life threatening.

[0006] Attempts have been made to combine a lancing-type device with various other components involved in the analyte concentration determination procedure in order to simplify the assay process. For example, U.S. Pat. No. 6,099,484 discloses a sampling device which includes a single needle associated with a spring mechanism, a capillary tube associated with a pusher, and a test strip. An analyzer may also be mounted in the device for analyzing

the sample. Accordingly, the single needle is displaced toward the skin surface by un-cocking a spring and then retracting it by another spring. A pusher is then displaced to push the capillary tube in communication with a sample and the pusher is then released and the fluid is transferred to a test strip.

[0007] U.S. Pat. No. 5,820,570 discloses an apparatus which includes a base having a hollow needle and a cover having a membrane, whereby the base and cover are connected together at a hinge point. When in a closed position, the needle is in communication with the membrane and fluid can be drawn up through the needle and placed on the membrane of the cover.

[0008] There are certain drawbacks associated with each of the above devices and techniques. For example, the devices disclosed in the aforementioned patents are complex, thus decreasing ease-of-use and increasing manufacturing costs. Furthermore, as described, a single needle design may be associated with increased pain because the single needle must be large enough to extract the requisite sample size. Still further, in regards to the '484 patent, the steps of activating and retracting a needle and then activating and retracting a capillary tube adds still more user interaction and decreases ease-of-use.

[0009] As such, there is continued interest in the development of new devices and methods for use in the determination of analyte concentrations in a physiological sample. Of particular interest would be the development of integrated devices, and methods of use thereof, that are efficient, involve minimal pain, are simple to use and which may be used with various analyte concentration determination systems.

SUMMARY OF THE INVENTION

[0010] Devices, systems and methods are provided for piercing the skin, accessing and collecting physiological sample therein, and measuring a characteristic of the physiological sample. The subject devices include at least one microneedle or skin-piercing element integral with a test strip. More specifically, the subject test strips include a biosensor wherein the at least one skin-piercing element is structurally integral with the biosensor.

[0011] Each skin-piercing element has a space-defining configuration therein which, upon insertion into the skin, creates a space or volume within the pierced tissue. This space serves as a reservoir or pooling area within which bodily fluid is caused to pool while the skin-piercing element is in situ. A capillary channel or fluid pathway extending from the pooling space to within the test strip transfers fluid present pooled within the pooling space to the biosensor. In certain embodiments, the space-defining configuration is a recess within a surface of the skin-piercing element. Such a recess may have a concave configuration. In other embodiments, the space-defining configuration is an opening which extends transverse to a dimension of the skin-piercing element and occupies a substantial portion of a width or diameter dimension as well as a substantial portion of a length dimension of the microneedle.

[0012] In one embodiment of the subject test strip devices, the biosensor is an electrochemical biosensor having an electrochemical cell having two spaced-apart electrodes.

Each skin-piercing element or structure is provided as a parallel or planar extension of one of the electrodes, wherein the skin-piercing element and such electrode are preferably fabricated as a single, unitary piece or structure and are made of the same material.

[0013] In another embodiment of the test strip device, the biosensor is a photometric or colorimetric biosensor having a planar substrate defining a photometric matrix area covered by a photometric membrane, collectively configured for receiving a sample to be tested. With a photometric biosensor embodiment, each skin-piercing element or structure is provided as a planar extension of the substrate, wherein the skin-piercing element and such substrate are preferably fabricated as a single, unitary piece or structure and are made of the same material.

[0014] The extending skin-piercing element and the associated electrode (in electrochemical biosensors) or substrate (in photometric biosensors) define at least one pathway, wherein the proximal end of the at least one pathway resides within the electrode or substrate portion of the unitary piece and the distal end of the at least one pathway resides within the skin-piercing element or structure. At least a portion of the distal end of the at least one fluid pathway is open to the outside environment. Further, the distal end of the pathway is in fluid communication with the space-defining area of the skin-piercing element. The distal end of such pathway either extends into at least a portion of the space-defining area or terminates at the space-defining area. As such, the fluid pathway provides a capillary channel through which the fluid within the pooling volume defined by the skin-piercing element may be extracted and transferred to the biosensor portion of the test strip device for testing.

[0015] The subject systems include one or more subject test strip devices and a meter for receiving a subject test strip and for determining a characteristic of the sampled fluid, e.g., the concentration of at least one analyte in the sample, collected by within the test strip's biosensor. Moreover, such a meter may also provide means for activating and manipulating the test strip wherein the skin-piercing structure is caused to pierce the skin. Additionally, the meter may provide means for storing one or more subject test strips, or a cartridge containing a plurality of such test strips.

[0016] Also provided are methods for using the subject devices, as well as kits that include the subject devices and/or systems for use in practicing the subject methods. The subject devices, systems and methods are particularly suited for collecting physiological sample and determining analyte concentrations therein and, more particularly, glucose concentrations in blood, blood fractions or interstitial fluid.

[0017] The present invention further includes methods for fabricating the subject test strip devices, in which a microneedle or skin-piercing element is fabricated as an integral part of a biosensor having a test strip configuration. Such devices have wholly integrated functions including accessing the physiological fluid within the skin, extracting such fluid, transferring the fluid to a measurement area and providing the components necessary for the measurement of analyte concentration in the sample. In addition to fabricating wholly integrated test strip devices, the subject fabrication methods are ideal for the fabrication of such devices which have functionally and structurally complex compo-

nents, such as the microneedles mentioned above. For example, microneedles having intricate shapes or designs, multiple dimensions, small sizes and/or very sharp tips are producible with great repeatability with the subject fabrication methods. The subject methods are also versatile in that they can be used to fabricate biosensors having electrochemical or photometric configurations with certain variations in the fabrication processes. The subject fabrication methods may be used to fabricate individual test strip devices or a plurality of such test strip devices on a web, film or sheet of suitable material.

[0018] These and other objects, advantages, and features of the invention will become apparent to those persons skilled in the art upon reading the details of the methods and systems of the present invention which are more fully described below.

BRIEF DESCRIPTIONS OF THE DRAWINGS

[0019] FIG. 1A is an exploded top view of an embodiment of an electrochemical test strip device of the present invention.

[0020] FIG. 1B is a partially exploded bottom view of the electrochemical test strip device of FIG. 1A.

[0021] FIG. 1C is a perspective view of the assembled electrochemical test strip device of FIGS. 1A and 1B.

[0022] FIG. 2A is an exploded view of an embodiment of a colorimetric or photometric test strip device of the present invention.

[0023] FIG. 2B is a perspective view of the assembled colorimetric/photometric test strip device of FIG. 2A.

[0024] FIG. 3 is a perspective view of an electrochemical test strip device of the present invention having another embodiment of a skin-piercing element of the present invention.

[0025] FIG. 4A is an exploded view of another embodiment of a colorimetric or photometric test strip device of the present invention having the skin-piercing element of FIG. 3.

[0026] FIG. 4B is a perspective view of the assembled colorimetric/photometric test strip device of FIG. 4A.

[0027] FIG. 5 illustrates a system of the present invention which includes a meter and a subject test strip device configured to be received by the meter.

[0028] FIG. 6A is an exploded top view of a web of electrochemical test strip devices fabricated according to the methods of the present invention.

[0029] FIG. 6B is an exploded bottom view of the web of FIG. 6A.

[0030] FIG. 6C is a perspective view of the assembled web of FIGS. 6A and 6B.

[0031] FIG. 7A is an exploded top view of a web of the photometric/colorimetric test strip devices fabricated according to the methods of the present invention.

[0032] FIG. 7B is an exploded bottom view of the web of FIG. 7A.

[0033] FIG. 7C is a perspective view of the assembled web of FIGS. 7A and 7B.

[0034] FIG. 8 is planar view of a web layer for use with the webs of FIGS. 6 and 7.

DETAILED DESCRIPTION OF THE INVENTION

[0035] Before the present invention is described, it is to be understood that this invention is not limited to the particular embodiments described, as such may, of course, vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only, and is not intended to be limiting, since the scope of the present invention will be limited only by the appended claims.

[0036] Where a range of values is provided, it is understood that each intervening value, to the tenth of the unit of the lower limit unless the context clearly dictates otherwise, between the upper and lower limit of that range and any other stated or intervening value in that stated range is encompassed within the invention. The upper and lower limits of these smaller ranges may independently be included in the smaller ranges is also encompassed within the invention, subject to any specifically excluded limit in the stated range. Where the stated range includes one or both of the limits, ranges excluding either both of those included limits are also included in the invention.

[0037] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although any methods and materials similar or equivalent to those described herein can also be used in the practice or testing of the present invention, the preferred methods and materials are now described.

[0038] It must be noted that as used herein and in the appended claims, the singular forms "a", "an", and "the" include plural referents unless the context clearly dictates otherwise. Thus, for example, reference to "a test strip" includes a plurality of such test strips and reference to "the device" includes reference to one or more devices and equivalents thereof known to those skilled in the art, and so forth.

[0039] The publications discussed herein are provided solely for their disclosure prior to the filing date of the present application. Nothing herein is to be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention. Further, the dates of publication provided may be different from the actual publication dates which may need to be independently confirmed.

[0040] The present invention will now be described in detail. In further describing the present invention, various embodiments of the subject devices, including test strip devices having biosensors having either an electrochemical or a colorimetric/photometric configuration, will be described first followed by a detailed description of various microneedle configurations which are usable with either type of biosensor configuration. The subject systems which include a meter for use with the subject devices methods of using the subject test strip devices and systems will then be described followed by a description of the methods of fabricating the subject test strip devices. Finally, a brief

description is provided of the subject kits, which kits include the subject devices and systems for use in practicing the subject methods.

[0041] In the following description, the present invention will be described in the context of analyte concentration measurement applications; however, such is not intended to be limiting and those skilled in the art will appreciate that the subject devices, systems and methods are useful in the measurement of other physical and chemical characteristics of biological substances, e.g., blood coagulation time, blood cholesterol level, etc.

[0042] Test Strip Devices

[0043] As summarized above, the subject test strip devices include a biosensor and at least one skin-piercing element or microneedle which is structurally integral with the biosensor. The subject biosensor may have an electrochemical configuration, as illustrated in FIGS. 1A, 1B, 1C and FIG. 3 or a colorimetric or photometric (used interchangeably herein), as illustrated in FIGS. 2A and 2B and FIGS. 4A and 4B. Likewise, the subject skin-piercing elements make take on various configurations, wherein a first exemplary embodiment is illustrated in FIGS. 1A, 1B, 1C and 3 and a second exemplary embodiment is illustrated in FIGS. 2A, 2B, 4A and 4B.

[0044] In any embodiment, the subject test strip devices and biosensors are useful in the determination of a wide variety of different analyte concentrations, where representative analytes include, but are not limited to, glucose, cholesterol, lactate, alcohol, and the like. In many embodiments, the subject test strips are used to determine the glucose concentration in a physiological sample, e.g., interstitial fluid, blood, blood fractions, constituents thereof, and the like.

[0045] Electrochemical Test Strips

[0046] Referring now to FIGS. 1A, 1B, 1C and 3, wherein like reference numbers refer to like elements, two electrochemical test strip devices 2 and 100, respectively, of the present invention are illustrated. Test strips 2 and 100 have identical electrochemical biosensor configurations which are herein collectively described, however, their respective skin-piercing elements or microneedles 6 and 102, respectively, have different configurations. In each test strip device 2 and 100, the biosensor is defined by an electrochemical cell generally having two spaced-apart and opposing electrodes 3 and 5, respectively referred to herein as bottom electrode 3 and top electrode 5. At least the surfaces of electrodes 3 and 5 facing each other are comprised of a conductive layer 8 and 16, respectively, such as a metal.

[0047] In certain embodiments of the subject electrochemical biosensors, the electrodes are generally configured in the form of elongated rectangular strips but may be of any appropriate shape or configuration. Typically, the length of the electrodes ranges from about 0.5 to 4.5 cm and usually from about 1.0 to 2.8 cm. The width of the electrodes ranges from about 0.07 to 0.8 cm, usually from about 0.20 to 0.60 cm, and more usually from about 0.1 to 0.3 cm. The conductive layers and their associated substrate typically have a combined thickness ranging from about 100 to 500 μm and usually from about 125 to 250 μm .

[0048] The entire electrode may be made of the metal or made up of a substrate or backing 4 and 18, respectively on

the facing surfaces of which the metal layer **8** and **16**, respectively, is provided. In a particular embodiment, substrates **4** and **18** are made of a Mylar plastic film. The thickness of the inert backing material typically ranges from about 25 to 500 μm and usually from about 50 to 400 μm , while the thickness of the metal layer typically ranges from about 10 to 100 nm and usually from about 10 to 50 nm.

[0049] As mentioned above, electrodes **3** and **5** generally face each other and are separated by only a short distance, such that the spacing between the electrodes is extremely narrow. This minimal spacing is a result of the presence of a spacer layer **12** positioned or sandwiched between electrodes **3** and **5**. The thickness of spacer layer **12** may range from 10 to 750 μm and is often less than or equal to 500 μm , and usually ranges from about 25 to 175 μm . Spacer layer **12** preferably has double-sided adhesive to hold electrodes **3** and **5** together.

[0050] In certain embodiments, spacer layer **12** is configured or cut so as to provide a reaction zone or area **9**, where in many embodiments the volume of the reaction area or zone **9** typically has a volume in the range from about 0.01 to 10 μL , usually from about 0.1 to 1.0 μL and more usually from about 0.05 to 1.0 μL . However, the reaction area may include other areas of test strip **2** and **100** or be elsewhere all together, such as in a fluid pathway, described below in more detail, or the like. Spacer layer **12** may define any appropriately shaped reaction area **9**, e.g., circular, square, triangular, rectangular or irregular shaped reaction areas, and may further include side inlet and outlet vents or ports.

[0051] Regardless of where reaction zone **9** is located, in many embodiments, a redox reagent system or composition **14** is present within reaction zone **9**, where reagent system **14** is selected to interact with targeted components in the fluid sample during an assay of the sample. Redox reagent system **14** is deposited on the conductive layer **16** of top electrode **5** wherein, when in a completely assembled form (shown in FIG. 1C), redox reagent system **14** resides within reaction zone **9**. With such a configuration, bottom electrode **3** serves as a counter/reference electrode and top electrode **5** serves as the working electrode of the electrochemical cell. However, in other embodiments, depending on the voltage sequence applied to the cell, the role of the electrodes can be reversed such that bottom electrode **3** serves as a working electrode and top electrode **5** serves as a counter/reference electrode. In case of a double pulse voltage waveform, each electrode acts as a counter/reference and working electrode once during the analyte concentration measurement.

[0052] Reagent systems of interest typically include an enzyme and a redox active component (mediator). The redox component of the reagent composition, when present, is made up of one or more redox agents. A variety of different redox agents, i.e., mediators, is known in the art and includes: ferricyanide, phenazine ethosulphate, phenazine methosulfate, pheylenediamine, 1-methoxy-phenazine methosulfate, 2,6-dimethyl-1,4-benzoquinone, 2,5-dichloro-1,4-benzoquinone, ferrocene derivatives, osmium bipyridyl complexes, ruthenium complexes, and the like. In many embodiments, the redox active component of particular interest is ferricyanide, and the like. The enzyme of choice may vary depending on the analyte concentration which is to be measured. For example, suitable enzymes for the assay of glucose in whole blood include glucose oxidase or dehy-

drogenase (NAD or PQQ based). Suitable enzymes for the assay of cholesterol in whole blood include cholesterol oxidase and esterase.

[0053] Other reagents that may be present in the reaction area include buffering agents (e.g., citraconate, citrate, malic, maleic, phosphate, "Good" buffers and the like); divalent cations (e.g., calcium chloride, and magnesium chloride); surfactants (e.g., Triton, Macol, Tetric, Silwet, Zonyl, and Pluronic); and stabilizing agents (e.g., albumin, sucrose, trehalose, mannitol and lactose).

[0054] Examples of electrochemical biosensors suitable for use with the subject invention include those described in copending U.S. application Ser. Nos. 09/333,793; 09/497,304; 09/497,269; 09/736,788 and 09/746,116, the disclosures of which are herein incorporated by reference.

[0055] Colorimetric/Photometric Test Strips

[0056] Referring now to FIGS. 2A, 2B, 4A and 4B, wherein like reference numbers refer to like elements, two photometric/colorimetric test strip devices **80** and **120**, respectively, of the present invention are illustrated. Test strips devices **80** and **120** have different photometric/colorimetric biosensor configurations and their respective skin-piercing elements or microneedles **86** and **122**, respectively, have different configurations. More specifically, a portion of test strip device **80** is made of an inert material while the corresponding portion of test strip device **120** is made of a metal material.

[0057] In test strip device **80** of FIGS. 2A and 2B, the colorimetric or photometric (herein used interchangeably) biosensor is generally made up of at least the following components: a support element or substrate **82** made of an inert material, a matrix area **84** for receiving a sample, a reagent composition (not shown as a structural component) within matrix area **84** that typically includes one or more members of an analyte oxidation signal producing system, an air venting port (not shown) and a top transparent layer **85** which covers at least matrix area **84**. In other embodiments, top layer **85** may be a membrane containing a reagent composition impregnated therein while the matrix area **84** may or may not contain reagent composition.

[0058] The inert material of support substrate **82** provides a physical structure to enable test strip **80** to be inserted into a meter without undue bending or kinking. Substrate **82**, and thus test strip **80**, is typically in the form of a substantially rectangular or square-like strip. Typically, the length of the substrate **82** is from about 1 to 1000 mm, usually from about 10 to 100 mm and more usually about 20 to 60 mm. Typically, the width of substrate **82** is from about 1 to 100 mm, usually from about 1 to 10 mm and more usually from about 5 to 7 mm. Typically, the height or thickness of substrate **82** is from about 0.01 to 1 mm, usually from about 0.1 to 1 mm and more usually from about 0.1 to 0.2 mm.

[0059] Matrix area **84** defines an inert area, preferably a recessed area, formed within a surface of substrate **82** wherein all four sides of matrix area **84** are bordered by substrate **82**. Matrix area **84** provides an area for deposition of the sampled physiological fluid and for the various members of the signal producing system, described infra, as well as for the light absorbing or chromogenic product produced by the signal producing system, i.e., the indicator, as well as provides a location for the detection of the

light-absorbing product produced by the indicator of the signal producing system. In such an embodiment, top layer **85** is transparent so that the color intensity of the chromogenic product resulting from the reaction between the target analyte and the signal producing system can be measured. Transparent layer **85** may, for example, be made of clear thin polyester.

[0060] This approach, in which the reagent is loaded into matrix area **84** and the biosensor is covered with a transparent film **85**, is useful in color generation systems that use an enzyme independent of oxygen, such as NAD-, or PQQ-based glucose dehydrogenase.

[0061] In yet another embodiment, top layer **85** is one that is permissive of aqueous fluid flow and is sufficiently porous, i.e., provides sufficient void space, for the chemical reactions of the signal producing system to take place. In principle, the nature of porous membrane **85** is critical to the subject test strips in that it should support an aqueous fluid flow both lateral and across the membrane thickness. Ideally, the membrane pore structure would not support red blood cell flow to the surface of the membrane being interrogated, i.e., the color intensity of which is a subject of the measurement correlated to analyte concentration. As such, the dimensions and porosity of test strip **80** may vary greatly, where matrix area **84** may or may not have pores and/or a porosity gradient, e.g. with larger pores near or at the sample application region and smaller pores at the detection region. Materials from which matrix membrane **85** may be fabricated vary, include polymers, e.g. polysulfone, polyamides, cellulose or absorbent paper, and the like, where the material may or may not be functionalized to provide for covalent or non-covalent attachment of the various members of the signal producing system.

[0062] While test strip device **120** of **FIGS. 4A and 4B** has a substrate **140** having a size and shape similar to substrate **82**, has a membrane **142** which has a configuration similar to transparent layer **85** and employs the same signal producing system as the test strip device of **FIGS. 2A and 2B**, there are certain notable differences between the two test strip devices. First, substrate **140** is made of a metal material rather than an inert material. Additionally, matrix **148** is not recessed within substrate **140** and extends across the complete width of substrate **140**. Further, test strip **120** has a double-sided adhesive layer **144** situated between substrate **140** and membrane **142**. Double-sided adhesive layer **144** has a cut-out portion **150** which corresponds to the area covered by matrix **148** and defines a deposition area as described above with respect to matrix area **84**. The double-sided adhesive layer **144** holds membrane **142** attached to substrate **140**.

[0063] A number of different matrices have been developed for use in various analyte detection assays, which matrices may differ in terms of materials, dimensions and the like, where representative matrices usable with the photometric/colorimetric test strip devices of the present invention include, but are not limited to, those described in U.S. Pat. Nos. 4,734,360; 4,900,666; 4,935,346; 5,059,394; 5,304,468; 5,306,623; 5,418,142; 5,426,032; 5,515,170; 5,526,120; 5,563,042; 5,620,863; 5,753,429; 5,573,452; 5,780,304; 5,789,255; 5,843,691; 5,846,486; 5,968,836 and 5,972,294; the disclosures of which are herein incorporated by reference.

[0064] The one or more members of the signal producing system produce a detectable product in response to the presence of analyte, which detectable product can be used to derive the amount of analyte present in the assayed sample. In the subject test strips, the one or more members of the signal producing system are associated, e.g., covalently or non-covalently attached to, at least a portion of (i.e., the detection region) the matrix, and in many embodiments to substantially all of the matrix. The signal producing system is an analyte oxidation signal producing system. By analyte oxidation signal producing system is meant that in generating the detectable signal from which the analyte concentration in the sample is derived, the analyte is oxidized by a suitable enzyme to produce an oxidized form of the analyte and a corresponding or proportional amount of hydrogen peroxide. The hydrogen peroxide is then employed, in turn, to generate the detectable product from one or more indicator compounds, where the amount of detectable product generated by the signal measuring system, i.e. the signal, is then related to the amount of analyte in the initial sample. As such, the analyte oxidation signal producing systems present in the subject test strips are also correctly characterized as hydrogen peroxide based signal producing systems.

[0065] As indicated above, the hydrogen peroxide based signal producing systems include an enzyme that oxidizes the analyte and produces a corresponding amount of hydrogen peroxide, where by corresponding amount is meant that the amount of hydrogen peroxide that is produced is proportional to the amount of analyte present in the sample. The specific nature of this first enzyme necessarily depends on the nature of the analyte being assayed but is generally an oxidase or dehydrogenase. As such, the first enzyme may be: glucose oxidase (where the analyte is glucose), or glucose dehydrogenase either using NAD or PQQ as cofactor; cholesterol oxidase (where the analyte is cholesterol); alcohol oxidase (where the analyte is alcohol); lactate oxidase (where the analyte is lactate) and the like. Other oxidizing enzymes for use with these and other analytes of interest are known to those skilled in the art and may also be employed. In those preferred embodiments where the reagent test strip is designed for the detection of glucose concentration, the first enzyme is glucose oxidase. The glucose oxidase may be obtained from any convenient source, e.g. a naturally occurring source such as *Aspergillus niger* or *Penicillium*, or recombinantly produced.

[0066] The second enzyme of the signal producing system is an enzyme that catalyzes the conversion of one or more indicator compounds into a detectable product in the presence of hydrogen peroxide, where the amount of detectable product that is produced by this reaction is proportional to the amount of hydrogen peroxide that is present. This second enzyme is generally a peroxidase, where suitable peroxidases include: horseradish peroxidase (HRP), soy peroxidase, recombinantly produced peroxidase and synthetic analogs having peroxidative activity and the like. See, e.g., Y. Ci, F. Wang; *Analytica Chimica Acta*, 233 (1990), 299-302.

[0067] The indicator compound or compounds, e.g., substrates, are ones that are either formed or decomposed by the hydrogen peroxide in the presence of the peroxidase to produce an indicator dye that absorbs light in a predetermined wavelength range. Preferably the indicator dye absorbs strongly at a wavelength different from that at which the sample or the testing reagent absorbs strongly. The

oxidized form of the indicator may be a colored, faintly-colored, or colorless final product that evidences a change in color of the testing side of the membrane. That is to say, the testing reagent can indicate the presence of glucose in a sample by a colored area being bleached or, alternatively, by a colorless area developing color.

[0068] Indicator compounds that are useful in the present invention include both one- and two-component chromogenic substrates. One-component systems include aromatic amines, aromatic alcohols, azines, and benzidines, such as tetramethyl benzidine-HCl. Suitable two-component systems include those in which one component is MBTH, an MBTH derivative (see for example those disclosed in U.S. patent application Ser. No. 08/302,575, incorporated herein by reference), or 4-aminoantipyrine and the other component is an aromatic amine, aromatic alcohol, conjugated amine, conjugated alcohol or aromatic or aliphatic aldehyde. Exemplary two-component systems are 3-methyl-2-benzothiazolinone hydrazone hydrochloride (MBTH) combined with 3-dimethylaminobenzoic acid (DMAB); MBTH combined with 3,5-dichloro-2-hydroxybenzene-sulfonic acid (DCHBS); and 3-methyl-2-benzothiazolinonehydrazone N-sulfonyl benzenesulfonate monosodium (MBTHSB) combined with 8-anilino-1 naphthalene sulfonic acid ammonium (ANS). In certain embodiments, the dye couple MBTHSB-ANS is preferred.

[0069] In yet other embodiments of colorimetric test strips, signal producing systems that produce a fluorescent detectable product (or detectable non-fluorescent substance, e.g. in a fluorescent background) may be employed, such as those described in Kiyoshi Zaitzu, Yosuke Ohkura, New fluorogenic substrates for Horseradish Peroxidase: rapid and sensitive assay for hydrogen peroxide and the Peroxidase, *Analytical Biochemistry* (1980) 109, 109-113. Examples of such colorimetric reagent test strips suitable for use with the subject invention include those described in U.S. Pat. Nos. 5,563,042; 5,753,452; 5,789,255, herein incorporated by reference.

[0070] Skin-Piercing Elements/Microneedles

[0071] Referring to test strips **2** and **80** of **FIGS. 1 and 2**, respectively, as well as to test strips **100** and **120** of **FIGS. 3 and 4**, respectively, the various configurations of skin-piercing element/microneedles of the present invention will now be discussed in greater detail. As discussed above, test strip device **100** includes an electrochemical biosensor configuration similar to that of test strip **2** of **FIG. 1** while test strip device **120** includes a colorimetric/photometric biosensor configuration similar to that of test strip **80** of **FIG. 2**; however, the test strip devices **100** and **120** of **FIGS. 3 and 4**, respectively, differ from those of **FIGS. 1 and 2** in that they have a different microneedle configuration. In both embodiments, the microneedles extend from a substrate of the respective test strip. Specifically, in the electrochemical test strip device embodiments of **FIGS. 1 and 3**, the microneedle may extend from either of one of the two substrates, i.e., biosensor electrodes, wherein the microneedle and such associated electrode are integrated with each other.

[0072] Any suitable shape of skin-piercing element may be employed with the subject test strip devices, as long as the shape enables the skin to be pierced with minimal pain to the patient. For example, the skin-piercing element may have a

substantially flat or planar configuration, or may be substantially cylindrical-like, wedge-like or triangular in shape such as a substantially flattened triangle-like configuration, blade-shaped, or have any other suitable shape. The cross-sectional shape of the skin-piercing element, or at least the portion of skin-piercing element that is penetrable into the skin, may be any suitable shape, including, but not limited to, substantially rectangular, oblong, square, oval, circular, diamond, triangular, star, etc. Additionally, the skin-piercing element may be tapered or may otherwise define a point or apex at its distal end. Such a configuration may take the form of an oblique angle at the tip or a pyramid or triangular shape or the like.

[0073] The dimensions of the skin-piercing element may vary depending on a variety of factors such as the type of physiological sample to be obtained, the desired penetration depth and the thickness of the skin layers of the particular patient being tested. Generally, the skin-piercing element is constructed to provide skin-piercing and fluid extraction functions and, thus, is designed to be sufficiently robust to withstand insertion into and withdrawal from the skin. Typically, to accomplish these goals, the ratio of the penetration length (defined by the distance between the base of the skin-piercing element and its distal tip) to diameter (where such diameter is measured at the base of the skin-piercing element) is from about 1 to 1, usually about 2 to 1, more usually about 5 to 1 or 10 to 1 and oftentimes 50 to 1.

[0074] The total length of the skin-piercing elements generally ranges from about 1 to 30,000 microns, usually from about 100 to 10,000 microns and more usually from about 1,000 to 3,000 microns. The penetration length of the skin-piercing elements generally ranges from about 1 to 5000 microns, usually about 100 to 3000 microns and more usually about 1000 to 2000 microns. The height or thickness of skin-piercing elements **6** and **86**, at least the thickness of the distal portion of the skin-piercing element, typically ranges from about 1 to 1000 microns, usually from about 10 to 500 microns and more usually from about 50 to 250 microns. The outer diameter at the base generally ranges from about 1 to 2000 microns, usually about 300 to 1000 microns and more usually from about 500 to 1000 microns. In many embodiments, the outer diameter of the distal tip generally does not exceed about 100 microns and is generally less than about 20 microns and more typically less than about 1 micron. However, it will be appreciated by one of skill in the art that the outer diameter of the skin-piercing element may vary along its length or may be substantially constant.

[0075] Each of the skin-piercing elements of the test strip devices of **FIGS. 1-4** has a space-defining configuration or structure therein which, upon insertion into the skin, creates a space or volume within the pierced tissue. This space serves as a reservoir within which bodily fluid is caused to pool in situ prior to being transferred to the biosensor portion of the subject test strip devices. Generally, the space-defining configurations of the present invention create or define a space within the pierced tissue having a volume at least as great as the available fluid volume in the reaction zone of the biosensor. Such space or volume ranges from about 10 to 1,000 nL, and more usually from about 50 to 250 nL. Such volume occupies a substantial portion of the entire volume occupied by the structure of the skin-piercing element, and

ranges from about 50% to 99% and more usually from about 50% to 75% of the entire volume occupied by the skin piercing element.

[0076] Two exemplary configurations of the microneedle of the present invention are illustrated; however, such examples are not intended to be limiting. As illustrated in **FIGS. 1 and 2**, the microneedle's space-defining configuration is a recess **20** or **94** within a surface, e.g., the top surface, of skin-piercing structure **6** and **86**. In many embodiments, recesses **20** and **94** have concave configurations wherein the depth of the recess is in the range from about 1 to 1000 microns and more usually from about 50 to 250 microns. Microneedles **6** and **86** may further be characterized by an opening **22** and **90**, respectively, in the microneedle structure to further expose the pooling area defined by recess **22** and **86** to the outside environment, thereby increasing the volume and flow rate of body fluid into the pooling area.

[0077] In other embodiments, as illustrated in **FIGS. 3 and 4**, the space-defining configuration is an opening **104** and **124**, respectively, which extends transverse to a dimension, e.g., width or thickness, of skin-piercing elements **102** and **122**, respectively. In skin-piercing embodiments having more annular cross-sections, such opening transverses a diameter of the skin-piercing element. In the illustrated embodiments, openings **104** and **124** each occupy a substantial portion of the width of their respective skin-piercing elements **102** and **122**, as well as a substantial portion of a length dimension of their respective test strips **100** and **120**. Openings **104** and **124** define sidewalls **112a** and **112b** and sidewalls **132a** and **132b**, respectively, of microneedles **100** and **120**, which have a thickness sufficient to maintain the structure of the microneedle when subject to normal forces.

[0078] The recesses **20** and **94** and openings **104** and **124** each define a space or volume within the overall space or volume occupied by the respective structure of the skin-piercing element. Such space or volume creates a corresponding space or volume within skin tissue upon penetration into the skin, which acts as a sample fluid collection reservoir wherein fluid released upon penetration is pooled within the space. Such configuration is advantageous over conventional skin-piercing needles (i.e., a hollow needle or one having a closed outer surface defining an internal fluid transport lumen) which normally plug or close most of the pierced blood capillaries within skin upon penetration in such a way that body fluid cannot be extracted while the needle is still inserted within the skin. On the other hand, the open-spaced microneedle configurations and structures of the present invention create a free or open volume inside the skin which exposes a significant portion of blood capillaries pierced by the microneedle tip, referenced as **24**, **92**, **106** and **126** in **FIGS. 1-4**, respectively. As such, the availability of a greater volume of body fluid can be provided with a tip that is smaller and/or sharper than conventional microneedles, thereby reducing pain. The greater availability of body fluid also results in a faster collection rate of sampling.

[0079] Sample Fluid Extraction Channels and Sub-Channels

[0080] The subject test strip devices further include a sample fluid transfer or extraction pathway or channel, referenced as **10**, **88**, **108** and **128** in **FIGS. 1, 2, 3 and 4**, respectively, which extends from the open space of the

respective microneedle to within the biosensor. At least a portion of the proximal end of the pathway resides within the biosensor portion of the test strip device. The distal end of the pathway may terminate just proximal to the microneedle structure (see **FIGS. 2A and 2B**) or may have a portion which resides within the skin-piercing structure (see **FIGS. 1A, 1C, 3 and 4**). In the latter configuration, such distal portion may be exposed to the outside environment.

[0081] In the test strip device of **FIG. 1**, bottom electrode **3** and microneedle **6** host a sample fluid transfer pathway or channel **10**, wherein the proximal end **10a** of pathway **10** resides within bottom electrode **3**, specifically within reaction zone **9**, and a portion of distal end **10b** of pathway **10** resides within skin-piercing element or structure **6**. Similarly, colorimetric test strip device **80** of **FIG. 2**, substrate **82** and skin-piercing element **86** host a fluid transfer pathway or channel **88**, wherein the proximal end **88a** of pathway **88** resides within substrate **82**, specifically within matrix area **84**. However, unlike pathway **10**, the distal end of pathway **88** terminates proximal to skin-piercing element **86**. Test strip devices **100** and **120** of **FIGS. 3 and 4**, respectively, host fluid pathways **108** and **128**, respectively, of which only the distal ends **108b** and **128b** are visible in the Figures. The distal ends **108b** and **128b** extend within a portion of microneedles **102** and **122**, respectively, and their distal openings **110** and **130**, respectively, terminate at associated openings **104** and **124**.

[0082] The pathways or channels of the present invention are preferably dimensioned so as to exert a capillary force on fluid within the pooling area defined by the open space portion of the microneedle, and draws or wicks physiological sample to within the reaction zone or matrix area of the biosensor. As such, the diameter or width of a single fluid channel or pathway does not exceed 1000 microns and will usually be about 100 to 200 microns in diameter. This diameter may be constant along its length or may vary. In certain embodiments, the fluid pathway may further include one or more agents to facilitate sample collection. For example, one or more hydrophilic agents may be present in the fluid pathway, where such agents include, but are not limited to types of surface modifiers or surfactants such as MESA, Triton, Macol, Tetriconic, Silwet, Zonyl and Pluronic.

[0083] As illustrated in the devices of **FIGS. 1 and 2**, channel **10** and **88**, respectively, may further include one or a plurality of sub or side branches or sub-channels **15** and **96**, respectively, which laterally extend from the proximal portion of the respective channel to within a portion or the entirety of the reaction zone **9** or matrix area **94**. Such sub-channels **15** and **96** are created by forming ridges or ribs in the respective substrates **4** and **82**, and/or the metal layer **3** which forms bottom electrode **3** of electrochemical test strip **2**. These ridges could be formed during the microneedle microfabrication process. In test strip **2**, electrode **5** acts as a cover over the ridges to form sub-channels **15**. Similarly, in test strip **80**, the matrix membrane or a clear film (not shown) acts as a cover over the ridges to form sub-channels **96**. Sub-channels **15** and **96** each have diameters sufficient to provide a capillary force on fluid residing within channels **10** and **88**, respectively. As such, the sub-channels facilitate the filling of reaction zone **9** and matrix area **84** with the sampled fluid. Sub-channels **15** and **96** have cross-sectional diameters in the range from about 1 to 200 microns and more usually from about 20 to 50 microns. In the illustrated

embodiment, capillary branches **15** and **96** extend perpendicularly from channel **10** and **88**, respectively; however, they may extend angularly from their respective channels.

[0084] Systems

[0085] As mentioned above, the subject devices may be used in the context of a subject system, which generally includes a system capable of obtaining a physiological sample and determining a property of the sample, where determining the property of interest may be accomplished automatically by an automated device, e.g., a meter. The subject system is more particularly described herein in the context of analyte concentration determination. Accordingly, as illustrated in **FIG. 5**, the analyte concentration determination system of the subject invention includes at least one test strip device **60** (having either an electrochemical or colorimetric configuration as described above) having at least one subject skin-piercing element **64**, as described above, associated therewith, and a meter **40**. The subject test strip devices, whether electrochemical or colorimetric, are configured and adapted to be inserted into meter **40**. More specifically, as illustrated in **FIG. 6**, test strip device **60** has a first end **62** and a second end **66**, wherein the skin-piercing element **64** is associated with first end **62** and at least the second end **66** is configured for insertion into a meter **40**.

[0086] Meter **40** preferably has an ergonomically-designed housing **42** having dimensions which allow it to be comfortably held and manipulated with one hand. Housing **42** may be made of a metal, plastic or other suitable material, preferably one that is light weight but sufficiently durable. The distal portion **56** of housing **42** provides an aperture **68** through which test strip device **60** is translatable from a retracted position within meter **40** to an extended position wherein at least a portion of the test strip microneedle extends a distance distally from aperture **68**. Distal portion **56** further defines a chamber in which test strip device **60** is received within a test strip receiving mechanism **70** of meter **40**. Test strip device **60** may be inserted into meter **40** by removing distal housing portion **56** from housing **42** and inserting test strip device **60** into test strip receiving mechanism **70**. Alternatively, test strip device **60** may be inserted into meter **40** and received into mechanism **70** via aperture **58**. Preferably, distal housing portion **56** is transparent or semi-transparent to allow the user to visually confirm proper engagement between test strip device **60** and receiving area **70** prior to conducting the analyte concentration assay, as well as to visualize the test site and to visually confirm the filling of strip **60** with body fluid during the assay. When test strip device **60** is properly seated within receiving mechanism **70**, the biosensor with test strip device **60** operatively engages with the meter's testing components. In other words, with electrochemical test strip embodiments, the electrodes of the biosensor operatively engage with the meter's electronics; and with colorimetric test strip embodiments, the matrix area having a signal producing system is operatively aligned with the meter's optical components. The meter's electronics or optical componentry, upon sensing when the reaction zone or matrix area, respectively, within test strip device **60** is filled with the sampled fluid, supplies an input signal to the test strip biosensor and receives an output signal therefrom which is representative of the sample fluid characteristic being measured.

[0087] Circumferentially positioned about aperture **68** is a pressure ring **58**, the distal surface of which is applied to the

skin and encircles the piercing site within the skin during a testing procedure. The compressive pressure exerted on the skin by pressure ring **58** facilitates the extraction of body fluids from the surrounding tissue and the transfer of such fluid into test strip device **60**.

[0088] Distal housing portion **56** is itself in movable engagement with meter **40** wherein distal housing portion **56** is slightly translatable or depressible along the longitudinal axis of meter **40**. Between distal housing portion **56** and the proximal portion of housing **42**, is a pressure sensor **54** which senses and gauges the amount of pressure exerted on distal housing portion **56** when compressing pressure ring **58** against the skin. Pressure sensor **54** is an electrical type sensor which may be of the kind commonly known in the field of electronics. Pressure sensor indicators **72**, in electrical communication with pressure sensor **54**, are provided to indicate the level of pressure being applied to distal housing portion **56** so that the user may adjust the amount of pressure being applied, if necessary, in order to apply an optimal pressure.

[0089] In many embodiments, meter **40** has a display **44**, such as an LCD display, for displaying data, such as input parameters and test results. Additionally, meter **40** has various controls and buttons for inputting data to the meter's processing components and for controlling the piercing action of test strip device **60**. For example, lever **46** is used to retract test strip device **60** to a loaded position within meter **40** and thereby pre-load a spring mechanism (not shown) for later, on-demand extension or ejection of test strip device **60** from aperture **68** by means of depressing button **48**. When distal housing portion **56** is properly positioned on the skin, such ejection of test strip device **60** causes microneedle **64** to instantaneously pierce the skin for accessing the body fluid therein. Buttons **50** and **52**, when depressed, input signals to the meter's processing components indicating whether the measurement to be made is for testing/information purposes (and for recovering the test results from a memory means within the meter's electronics) or for calibration purposes, respectively.

[0090] Optionally, meter **40** may further be configured to receive and retain a replaceable cartridge containing a plurality of the subject test strip devices. After using a test strip device, meter **40** may either eject the used test strip from the meter or store them for disposal at a later time. Such a configuration eliminates the necessary handling of test strips, thereby minimizing the likelihood of damage to the strip and inadvertent injury to the patient. Furthermore, because manual handling of the test strips is eliminated, the test strips may be made much smaller thereby reducing the amount of materials required, providing a cost savings.

[0091] The meter disclosed in U.S. patent application Ser. No. _____, entitled "Minimal Procedure Analyte Test System," having attorney docket no. LIFE-054 and filed on the same day herewith, is of particular relevance and is suitable for use with the subject invention. Additionally, certain aspects of the functionality of meters suitable for use with the subject systems are disclosed in U.S. Pat. No. 6,193,873, as well as in copending, commonly owned U.S. application Ser. Nos. 09/497,304, 09/497,269, 09/736,788, 09/746,116 and 09/923,093, the disclosures of which herein incorporated by reference. Of course, in those embodiments using a colorimetric assay system, a spectrophotometer or optical

meter will be employed, where certain aspects of the functionality of such meters suitable for use are described in, for example, U.S. Pat. Nos. 4,734,360, 4,900,666, 4,935,346, 5,059,394, 5,304,468, 5,306,623, 5,418,142, 5,426,032, 5,515,170, 5,526,120, 5,563,042, 5,620,863, 5,753,429, 5,773,452, 5,780,304, 5,789,255, 5,843,691, 5,846,486, 5,968,836 and 5,972,294, the disclosures of which are herein incorporated by reference.

[0092] Methods

[0093] As summarized above, the subject invention provides methods for determining a characteristic of the sample, e.g., the concentration of an analyte in a sample. The subject methods find use in the determination of a variety of different analyte concentrations, where representative analytes include glucose, cholesterol, lactate, alcohol, and the like. In many embodiments, the subject methods are employed to determine the glucose concentration in a physiological sample.

[0094] While in principle the subject methods may be used to determine the concentration of an analyte in a variety of different physiological samples, such as urine, tears, saliva, and the like, they are particularly suited for use in determining the concentration of an analyte in blood or blood fractions, and more particularly in whole blood or interstitial fluid.

[0095] The subject methods will now be described in detail with reference to Figures. In practicing the subject methods, at least one subject test strip device as described above, is provided, and a subject microneedle 6 thereof is inserted into a target area of skin. Typically, the skin-piercing element is inserted into the skin of a finger or forearm for about 1 to 60 seconds, usually for about 1 to 15 seconds and more usually for about 1 to 5 seconds. Depending on the type of physiological sample to be obtained, the subject skin-piercing element 6 may be penetrated to various skin layers, including the dermis, epidermis and the stratum corneum, but in many embodiments will penetrate no farther than the subcutaneous layer of the skin.

[0096] While the subject test strips may be handled and inserted into the skin manually, the subject test strips are preferably used with the hand-held meter 40 of FIG. 5. As such, a test strip device 60 is either initially inserted into test strip receiving mechanism 70 either through aperture 68 or by temporarily removing distal portion 56 of housing 42 and placing the test strip into receiving mechanism 70 of meter 40. Alternatively, test strip device 60 may be provided pre-loaded within receiving mechanism 70. Still yet, as mentioned above, test strip device 60 may be collectively pre-loaded with a plurality of like test strips in a test strip cartridge (not shown). In such an embodiment, the cartridge is removably engageable with meter 40. Used strips may be automatically disposed of, e.g., either ejected from the meter or deposited into a separate compartment within the cartridge, while an unused test strip is automatically removed from the cartridge and inserted into receiving area 70 of meter 40.

[0097] Once test strip device 60 is properly received within mechanism 70, mechanism 70 may then be spring loaded or cocked by means of lever 46 of meter 40. As such, mechanism 70 and, thus test strip device 60, is in a retracted position. Meter 40 is then positioned substantially perpen-

dicular to the targeted skin surface wherein distal housing portion 56, and more specifically pressure ring 58, is caused to contact the target skin area. Some compressive pressure may be manually applied to the target skin area, i.e., by pressing the distal end of meter 40 against the target skin area, to ensure that skin-piercing element 64 is properly inserted into the skin. By applying such pressure, a counter force causes distal housing portion 56 to press back upon pressure sensor 54 of meter 40. The relative amount (i.e., high, normal and low) of counter pressure is then measured and displayed by pressure sensor indicators 72. Preferably, the amount of pressure applied should generally be in the "normal" range. Indicators 72 inform the user as to when too much or too little pressure is being applied. When indicators 72 indicate that the applied pressure is "normal", the user may then depress the spring-release button 48. Due to the spring force released, receiving/carrying mechanism 70 and test strip device 60 are caused to thrust forward thereby causing skin-piercing element 65 to extend from aperture 68 and puncture the targeted skin area.

[0098] Whether by manual means or by use of meter 40, the penetration of skin-piercing element 64 into the skin creates a fluid sample pooling area (defined by the recess or opening within skin-piercing element) adjacent the fluid pathway, as described above, within element 64. Sample fluid enters the pooling area via the open-space configuration, e.g., recess or opening, within skin piercing element 64, and from the opposite side of skin-piercing element 46. The pooled sample fluid is then transferred via the fluid pathway by at least a capillary force exerted on the pooled fluid to the reaction zone or matrix within the biosensor of the test strip device 60. As mentioned above, the transfer of fluid may be further facilitated by exerting physical positive pressure circumferentially around the penetration site by means of a pressure ring 58 or by applying a source of negative pressure through the fluid channel thereby vacuuming the body fluid exposed to the distal end of the channel. The fluid entering the fluid pathway enters into the distal portion of the pathway first and then proceeds by capillary force (or by applied vacuum pressure) to within the proximal portion of the pathway which resides within the reaction zone or the matrix area. The fluid is then caused to translate laterally through the reaction zone or matrix area via sub-channels 15 or 96, respectively, wherein the entire available volume within the reaction zone or matrix area may be filled with the sample fluid.

[0099] Once meter 40 senses that the reaction zone or matrix area is completely filled with the sample of body fluid, the meter electronics or optics are activated to perform analysis of the extracted sample. At this point, the meter may be removed by the patient from the penetration site or kept on the skin surface until the test results are shown on the display. Meter 40 may alternatively or additionally include means for automatically retracting the microneedle strip from the skin once the reaction cell is filled with the body fluid sample.

[0100] When the biosensor reaction zone or matrix area is completely filled with the sample fluid, the concentration of the analyte of interest in the sampled fluid is determined. With an electrochemical based analyte concentration determination assay, an electrochemical measurement is made using counter/reference and working electrodes. The electrochemical measurement that is made may vary depending

on the particular nature of the assay and the meter with which the electrochemical test strip is employed, e.g., depending on whether the assay is coulometric, amperometric or potentiometric. Generally, the electrochemical measurement will measure charge (coulometric), current (amperometric) or potential (potentiometric), usually over a given period of time following sample introduction into the reaction area. Methods for making the above described electrochemical measurement are further described in U.S. Pat. Nos. 4,224,125; 4,545,382; and 5,266,179; as well as in International Patent Publications WO 97/18465 and WO 99/49307; the disclosures of which are herein incorporated by reference. Following detection of the electrochemical measurement or signal generated in the reaction zone as described above, the presence and/or concentration of the analyte present in the sample introduced into the reaction zone is then determined by relating the electrochemical signal to the amount of analyte in the sample.

[0101] For a colorimetric or photometric analyte concentration determination assay, sample applied to a subject test strip, more specifically to a reaction area of a test strip, is allowed to react with members of a signal producing system present in the reaction zone to produce a detectable product that is representative of the analyte of interest in an amount proportional to the initial amount of analyte present in the sample. The amount of detectable product, i.e., signal produced by the signal producing system, is then determined and related to the amount of analyte in the initial sample. With such colorimetric assays, optical-type meters are used to perform the above mentioned detection and relation steps. The above described reaction, detection and relating steps, as well as instruments for performing the same, are further described in U.S. Pat. Nos. 4,734,360; 4,900,666; 4,935,346; 5,059,394; 5,304,468; 5,306,623; 5,418,142; 5,426,032; 5,515,170; 5,526,120; 5,563,042; 5,620,863; 5,753,429; 5,773,452; 5,780,304; 5,789,255; 5,843,691; 5,846,486; 5,968,836 and 5,972,294; the disclosures of which are herein incorporated by reference. Examples of such colorimetric or photometric reagent test strips suitable for use with the subject invention include those described in U.S. Pat. Nos. 5,563,042; 5,753,452; 5,789,255, herein incorporated by reference.

[0102] Test Strip Device Fabrication Methods

[0103] As mentioned above, the skin-piercing elements of the present invention are preferably fabricated with a corresponding substrate (for colorimetric embodiments) or a substrate/electrode combination (for electrochemical embodiments), as a single, unitary piece or structure and made of the same material. Alternatively, the skin-piercing elements may be manufactured as separate components or pieces which are then affixed or attached to a corresponding substrate or substrate/conductive layer combination by any suitable means, for example, an adhesive commonly used in the art.

[0104] The test strip devices may be fabricated according to the present invention using any convenient techniques including, but not limited to, microreplication techniques including injection molding, photo-chemical etching (PCE), microstamping, embossing, and casting processes.

[0105] Because the test strip devices of the present invention are planar, the devices may be fabricated from and processed on one or more webs, films or sheets of suitable

material. Such web-based manufacturing of the subject test strip devices provide significant cost advantage over more conventional methods in which test strips and the like are produced one at a time. FIGS. 6A-C and 7A-C illustrate such webs of fabricated test strip devices having electrochemical and photometric/colorimetric configurations, respectively.

[0106] While the following discussion of the subject fabrication methods is in the context of web-based manufacturing, the techniques discussed may also be used to make singular test strip devices. Additionally, while only certain fabrication techniques are emphasized, those skilled in the art will recognize that other known fabrication techniques may also be used which enable low cost manufacturing when desiring to form small structures having intricate features, such as the microneedles described above and the sample fluid channels and sub-channels within the reaction area of the subject test strip devices.

[0107] Fabrication of Electrochemical Test Strip Devices

[0108] The electrochemical test strip device webbing **200** of FIGS. 6A-C includes a plurality of individual test strip devices **201** (shown fully assembled in FIG. 6C) fabricated in a side-to-side arrangement along the length of the webbing. Each test strip device **201** includes two spaced-apart electrodes, bottom electrode **202** and top electrode **204**, and an insulating space layer **206** there between. Spacer layer **206** has a cut-out portion **208** which defines the reaction zone of the electrochemical biosensor containing a redox reagent system. A microneedle **212**, shown having a configuration similar to that of microneedle **122** of FIGS. 4A and 4B, extends from and is planar with bottom electrode **202**. Formed within a portion of bottom electrode **202** and a proximal portion of microneedle **212** is a channel **214** for transporting fluid pooled within the opening of microneedle **212**. Extending laterally from both sides of channel **214** are a plurality of sub-channels **216** for facilitating the transfer and distribution of sampled fluid to within the reaction zone of the electrochemical biosensor.

[0109] Electrodes **202** and **204**, as well as the associated microneedles, may be made entirely of metal or may be made up of an inert substrate or a support structure covered by a metal layer. Where the electrodes are primarily made of metal, photochemical etching (PCE) (also known as photochemical milling, chemical milling and photoetching) or microstamping techniques are suitable fabrication techniques.

[0110] With photochemical etching, suitable metals include, but are not limited to, aluminum, copper, gold, platinum, palladium, iridium, silver, titanium, tungsten, carbon and stainless steels. Fabrication may be done on sheets or continuous coils of metals. Such sheet provides a thin metal base for the etching process and generally has a thickness in the range from about 10 to 1,000 μm and more typically from about 50 to 150 μm . A photoresistant layer is then applied to one or both sides of the metal base as desired. Next, lithography techniques are used to precisely define the geometries that will be etched partially into, e.g., the fluid channels **214** and sub-channels **216**, or etched completely through, e.g., the openings in the microneedles, the metal base. Specifically, the base metal is selectively masked to protect areas of the metal which are not to be etched and to expose areas of the metal which are to be etched.

[0111] Etching is accomplished by an electrochemical dissolution process wherein an acid substance is applied to

the surface of the base metal and a current is conducted through the metal. The areas of the metal surface which are not masked are then dissolved by the acid. After the etching step, the photoresist layer is stripped from the surface of the metal part, and, as illustrated in **FIG. 8**, the sheet **300** remains having a series of completely fabricated microneedles **302** and associated space-defining configurations **312**, fluid transfer channels **304** and sub-channels **306**. The portion **308** from which the bottom substrates are to be cut, remains a continuous area of metal while the area **310** of sheet **300** has been cut etched away completely.

[0112] Microstamping, another technique suitable for fabricating all-metal electrodes or those made out of a very strong plastic material, involves the use of dies which have been precisely machined such as by electro-discharge machining (EDM). Long sheets or webbings of a substrate metal, such as those metals commonly used in PCE processing, are continuously or semi-continuously fed into a stamping press between die sets to selectively blank (i.e., punch holes in), coin (i.e., deform one side of the metal) and/or deform the metal substrate from both sides. This stamping process can be performed at a rate of 1,200 strokes per minute and can produce multiple electrodes per stroke.

[0113] Where the electrodes **202** and **204** include an inert substrate material, hot embossing and injection molding techniques are suitable for fabrication of the subject devices particularly when the substrate material is a plastic. The substrate material is sufficiently rigid to provide structural support to the electrode and to the electrochemical test strip as a whole. Such suitable materials include polymers (plastics) and inorganic materials such as silicon, ceramic, glass, and the like. Suitable polymers include, for example, polyester, e.g., polyethylene terephthalate (PET), glycol modified polyethylene terephthalate (PETG); polyimide, e.g., polyetherimide; polycarbonate; cellophane (regenerated cellulose); fluorinated polymer, e.g., polyvinyl fluoride, perfluoroalkoxy and fluorinated ethylene propylene copolymers; ionomer; polyamide, e.g., nylon 6, nylon 6,6, nylon 11, nylon 12; polyethylene and its copolymers; polystyrene and its copolymers; polypropylene and its copolymers; polymethylpentene; polyvinyl chloride and its copolymers; polysulfone; polyvinylidene chloride and its copolymers; and polymer composites reinforced with minerals or nanoparticles. A preferred material for the substrate is a Mylar plastic film.

[0114] With hot embossing, a precursor material such as a suitable thermoplastic precursor material having a thickness in the range of about 25 to 650 microns, usually from about 50 to 625 microns and more usually from about 75 to 600 microns is placed into an embossing apparatus, where such an apparatus includes a mold having features, often times a negative image of the features, of the skin-piercing element. The precursor material is then compressed by the mold under heat and a suitable compression force. Usually, a temperature in the range from about 20° C. to 1500° C. is used, usually from about 100° C. to 1000° C. and more usually from about 200° C. to 500° C. Heat is applied for about 0.1 to 1000 seconds, usually for about 0.1 to 100 seconds and more usually for about 0.1 to 10 seconds. The compression force is usually applied in the range from about 1 to 50 GPa is used, usually from about 10 to 40 GPa and more usually from about 20 to 30 GPa. The compression force is applied for about 0.1 to 100 seconds, usually for

about 0.1 to 10 seconds and more usually for about 0.1 to 1 second. The heat and compression force may be applied at the same or different times. After the material is cooled, it is removed from the apparatus, and post processing may then occur.

[0115] Next, the upper side of the bottom substrate and the underside of the top substrate are metallized by vacuum sputtering or screen printing a conductive layer of metal over such substrates. The conductive layer may extend to cover the microneedle(s) **212** and, as such, the microneedle(s) functions as part of the associated electrode. More specifically, in certain electrochemical biosensor embodiments, the conductive material which is deposited over an inert substrate to form an electrode is also deposited over the sample fluid pathway or channel including the portion of the associated skin piercing element into which the fluid pathway extends. Suitable metals for the conductive layer include palladium, gold, platinum, silver, iridium, stainless steel and the like, or a metal oxide, such as indium doped tin oxide, or carbon, e.g., conductive carbon ink. In a preferred embodiment, the metal layer of electrode(s) **202** is gold and the metal layer electrode(s) **204** is palladium. An additional insulating layer may be printed on top of this conductive layer which exposes a precisely defined pattern of the electrode.

[0116] By means of any of the above fabrication techniques, bottom electrode(s) **202** functions as the counter/reference electrode and top electrode(s) **204** functions as the working electrode within the electrochemical cell. After fabrication of the electrodes, a redox reagent system is selected and deposited within the reaction zone **210** of bottom electrode(s) **202**. Such deposition may be accomplished with slot coating, needle coating or ink jet printing techniques, which are well known in the art. The redox reagent system may also be deposited within the sample extraction channel. Optionally, the conductive surface of electrode **202** may be subsequently treated with a hydrophilic agent to facilitate transport of a fluid sample through the sample extraction channel and into the reaction zone **210**. Suitable hydrophilic agent components include, for example, apoly(oxyethylene-co-oxypropylene) block polymer having the trade name Pluorinic™ F68, sodium dioctylsulfosuccinate having the trade name Aerosol™ OT 100%, octylphenoxypolyethoxy(9-10)ethanol having the trade name TRITON™ X-100, polyoxyethylene(20)sorbitan monolaurate having the trade name TWEEN™ 20, and polyoxyethylene(20)sorbitan monooleate having the trade name TWEEN™ 80, and 2-mercaptoethanesulfonic acid, sodium salt (MESA). In another embodiment redox reagent system may be deposited at the top electrode, i.e. layer **204** at the area corresponding to the zone **210** of the bottom layer **202** by the same deposition techniques. In yet another embodiment a redox system can be deposited on both electrodes, i.e., on layers **202** and **204** aligned so that the reagent coated chemistries face one another.

[0117] As mentioned above, electrodes **202** and **204** (and their respective webs) are separated by a spacer layer **206**, or a web of such spacer layer, positioned or sandwiched between electrodes **102** and **104**, or between their web structures. Spacer layer **106** may be fabricated from any convenient material, where representative suitable materials include polyethylene terephthalate, glycol modified (PETG), polyimide, polycarbonate, and the like. Both surfaces of

spacer layer **106** have an adhesive to allow it to adhere to the respective electrodes. By process known in web-based manufacturing, all three layers are aligned in a stacked relationship and laminated together into assembled web **200** which is then cut into singulated test strip devices **201**.

[0118] Fabrication of Photometric/Colorimetric Test Strip Devices

[0119] Many of the same techniques and processes, discussed above, for fabricating the electrochemical test strip devices of the present invention may also be used to fabricate the photometric/colorimetric test strip devices of the present invention.

[0120] Referring now to FIGS. 7A-C, the fabrication of the photometric/colorimetric devices of the present invention is described. A webbing **220** (shown assembled in FIG. 7C) includes a plurality of individual test strip devices **221** fabricated in a side-to-side arrangement along the length of webbing **220**. Such test strip devices **221** have a metal substrate configuration as described above with respect to FIGS. 4A and 4B. However, the subject fabrication techniques also apply to photometric test strip devices having inert material substrates as described above with respect to FIGS. 2A and 2B.

[0121] Webbing **220** is formed of at least three layers of sheets, a metal substrate sheet **222**, a membrane sheet **224** and a double-sided adhesive layer **226** there between. Double-sided adhesive layer **226** has a cut-out portion **228** which aligns with the matrix area **230** of the photometric biosensor which contains a signal producing system. A plurality of microneedles **232**, shown having a configuration similar to that of microneedle **122** of FIGS. 4A and 4B, extend from and are planar with substrate sheet **222**. Formed within a portion of each substrate **222** and a proximal portion of microneedle **232** is a channel **238** for transporting fluid pooled within the opening **234** of each microneedle **232**. Extending laterally from both sides of each channel **238** are a plurality of sub-channels **230** for facilitating the transfer and distribution of sampled fluid to within matrix **236** of the photometric biosensor.

[0122] As mentioned above, substrate sheet **222** as well as the associated microneedles **232** are made of metal, but may be made up of an inert material. Where the substrate made of metal, photochemical etching (PCE) and microstamping are suitable fabrication techniques. As with the electrochemical test strip devices, suitable metals for the substrate include, but are not limited to, aluminum, copper, gold, platinum, palladium, iridium, silver, titanium, tungsten, carbon and stainless steels. The metal sheet provides a thin metal base for the etching process and generally has a thickness in the range from about 10 to 1,000 μm and more typically from about 50 to 150 μm . A photoresistant layer is then applied to one or both sides of the metal base as desired. Next, lithography techniques are used to precisely define the geometries that will be etched partially into, e.g., the fluid channels **238** and sub-channels **230**, or etched completely through, e.g., the openings **234** in the microneedles **232**, the metal base. Specifically, the base metal is selectively masked to protect areas of the metal which are not to be etched and to expose areas of the metal which are to be etched. The electrochemical dissolution process of sheet **222** is as described above with respect to the electrochemical test strip devices of FIGS. 6A-6C, producing a sheet having the configuration of sheet **300** of FIG. 8.

[0123] Where the substrate sheet **222** is to be made of an inert substrate material, hot embossing and injection molding techniques, as described above with respect to fabrication of the electrochemical test strip devices, may be used for fabrication of the subject photometric test strip devices. The substrate material is sufficiently rigid to provide structural support to the electrode and to the electrochemical test strip as a whole. Such suitable inert materials for making support substrate sheet **222** include but are not limited to polyolefins, e.g., polyethylene or polypropylene, polystyrene or polyesters.

[0124] After fabrication of substrate **222**, a signal producing system, as described above, is selected and deposited within matrix **230**. Such deposition may be accomplished with slot coating, needle coating or ink jet printing techniques, which are well known in the art. The signal producing system may also be deposited within the sample extraction channels **238**. Optionally, the surface of matrices **230** as well as channels **238** may be subsequently treated with a hydrophilic agent having a surfactant to facilitate transport of a fluid sample through the sample extraction channel **238** and into the matrix **230**.

[0125] As mentioned above, substrate sheet **222** and membrane sheet **224** are separated by a double-sided adhesive layer **226**. Double-sided adhesive layer **226** may be fabricated from any convenient material, where representative suitable materials include polyethylene terephthalate, glycol modified polyethylene terephthalate (PETG), polyimide, polycarbonate, and the like. Both surfaces of spacer layer **226** have an adhesive to allow it to adhere to substrate **222** and membrane sheet **224**. In embodiments where substrate sheet **222** is made of an inert material, a spacer layer is not used. Instead, the side of membrane sheet **224** which is to contact substrate sheet **222** is provided with an adhesive coating, thereby allowing it to adhere to substrate sheet **222**. By processes known in web-based manufacturing, all layers, i.e., two, three or more as the case may be, are aligned in a stacked relationship and laminated together into assembled web **230** which is then cut into singulated photometric test strip devices **221**.

[0126] Kits

[0127] Also provided by the subject invention are kits for use in practicing the subject methods. The kits of the subject invention include at least one subject test strip device, oftentimes a plurality of test strip devices, where the at least one test strip device comprises at least one skin-piercing element. The kits may also include a reusable or disposable meter that may be used with disposable test strip devices. When a plurality of test strip devices is provided, they may be collectively packaged within a cartridge, which may be reusable or disposable. Certain kits may include various types of test strip devices, e.g., electrochemical and/or colorimetric test strip devices. Such various test strip devices may contain the same or different reagents. Finally, the kits may further include instructions for using the subject test strip devices and meters in the determination of an analyte concentration in a physiological sample. These instructions may be present on one or more of the packaging, label inserts, containers in the kits, and the like.

[0128] It is evident from the above description and discussion that the above described invention provides a simple, quick, safe and convenient way to obtain a physi-

ological sample and determine an analyte concentration thereof. The above described invention provides a number of advantages, including ease of use, decreased testing times, efficiency and minimal pain. As such, the subject invention represents a significant contribution to the art.

[0129] All publications and patents cited in this specification are herein incorporated by reference as if each individual publication or patent were specifically and individually indicated to be incorporated by reference. The citation of any publication is for its disclosure prior to the filing date and should not be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention.

[0130] Although the foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, it is readily apparent to those of ordinary skill in the art in light of the teachings of this invention that certain changes and modifications may be made thereto without departing from the spirit or scope of the appended claims.

What is claimed is:

1. A skin-piercing element for piercing the skin and accessing body fluid therein, said skin-piercing element comprising:

an opening within said skin-piercing element wherein said opening occupies a substantial portion of a width, diameter or length dimension of said skin-piercing element; and

a fluid pathway in fluid communication with said opening, wherein a pooling area is created within the skin by said opening upon insertion of said skin-piercing element into the skin.

2. The skin-piercing element of claim 1 wherein said opening has a volume in the range from about 50 to 500 nL.

3. The skin-piercing element of claim 1 wherein said opening occupies from about 50% to 95% of the volume occupied by said skin-piercing element.

4. The skin-piercing element of claim 1 wherein said skin-piercing element comprises a plastic material.

5. The skin-piercing element of claim 1 wherein said fluid pathway is dimensioned to apply a capillary force on fluid present within said pooling area.

6. The skin-piercing element of claim 1 further comprising a recess within a surface of said skin-piercing element, wherein said recess is in fluid communication with said opening.

7. The skin-piercing element of claim 6 wherein said recess has a concave configuration.

8. A test strip device comprising:

a biosensor for determining a characteristic of a physiological fluid;

at least one microneedle integral with and extending from said biosensor; said microneedle comprising an opening which occupies a substantial portion of a width, diameter or length dimension of said microneedle; and

a fluid pathway extending from said biosensor to said microneedle wherein said fluid pathway is in fluid communication with said opening and said biosensor.

9. The test strip device according to claim 8, wherein said biosensor has an electrochemical configuration.

10. The test strip device according to claim 9, wherein said biosensor comprises at least two electrodes and wherein said at least one microneedle is a planar extension of one of said at least two electrodes.

11. The test strip device according to claim 10 wherein said electrode from which said microneedle extends comprises a conductive material formed on a substrate material, and said microneedle is formed from said substrate material.

12. The test strip device according to claim 11 wherein said microneedle is further formed from said conductive material.

13. The test strip device according to claim 10 wherein said microneedle and said associated electrode are formed from a unitary structure.

14. The test strip device according to claim 8 wherein said microneedle comprises a metal material.

15. The test strip device according to claim 8 wherein said microneedle comprises an inert material.

16. The test strip device according to claim 10, wherein said biosensor further comprises a spacer layer between said at least two electrodes.

17. The test strip device according to claim 10 wherein said test strip further comprises a reaction zone between said electrodes and a redox reagent system contained at least within said reaction zone.

18. The test strip device according to claim 17 wherein said redox reagent system is further contained within at least a portion of said fluid pathway.

19. The test strip device according to claim 17 wherein a proximal portion of said fluid pathway resides within said reaction zone.

20. The test strip device according to claim 8, wherein said biosensor has a colorimetric configuration.

21. The test strip device according to claim 20 wherein said biosensor comprises a substrate having a matrix area defined therein and a membrane covering said matrix area.

22. The test strip device according to claim 21, wherein said matrix area contains an optical signal producing system.

23. The test strip device according to claim 21 wherein said membrane is porous.

24. The test strip device according to claim 21 wherein said membrane comprises a nonporous transparent film

25. The test strip device according to claim 8 comprising a plurality of microneedles.

26. The test strip device according to claim 8 further comprising a plurality of sub-channels extending from and in fluid communication with said fluid pathway.

27. A system for determining the concentration of at least one analyte in a physiological sample, said system comprising:

at least one test strip device according to claim 8, and

a meter for automatically determining the concentration of analyte in the physiological sample, wherein said meter is configured for receiving said test strip device.

28. The system according to claim 27, further comprising a test strip cartridge for containing a plurality of said test strip devices, said cartridge configured for releasable engagement with said meter.

29. The system according to claim 28, wherein said cartridge comprises a compartment for holding test strips devices which have been used.

30. The system according to claim 27, wherein said meter is handheld.

31. The system of according to claim 27, wherein said meter comprises a housing, an aperture at a distal end of said housing and a test strip-receiving mechanism within said housing for operatively receiving said at least one test strip device.

32. The system according to claim 31, wherein said meter further comprises means for spring-loading said test strip device in a retracted position within said distal end of said housing and means for releasing said at least one test strip from said spring-load wherein said at least one test strip is rapidly extended from said aperture.

33. The system according to claim 31, wherein said distal end of said housing is made of transparent or semi-transparent material.

34. The system according to claim 31, wherein said meter further comprises a pressure sensor for detecting and measuring pressure against said aperture.

35. The system according to claim 34, wherein said meter further comprises a pressure sensor indicator for indicating the pressure measured by said pressure sensor.

36. The system according to claim 27 wherein said meter further comprises a data display.

37. The system according to claim 27 wherein said meter further comprises a source of negative pressure for applying a vacuum through said fluid pathway for facilitating the transfer of physiological sample exposed to said pathway to within said test strip.

38. A method for collecting physiological fluid sample from skin, said method comprising:

providing at least one skin-piercing element comprising:

- (i) an opening which occupies a substantial portion of a width, diameter or length dimension of said skin-piercing element and
- (ii) a fluid pathway in fluid communication with said opening;

inserting said at least one skin-piercing element into the skin, wherein a pooling area is created within the skin by said opening and said physiological fluid pools within the pooling area; and

collecting by means of said fluid pathway said pooled physiological fluid from within the skin.

39. The method according to claim 38, wherein said step of inserting comprises inserting said at least one skin-piercing element no deeper than the subcutaneous layer of the skin.

40. The method according to claim 38, wherein said step of inserting comprises inserting said at least one skin-piercing element into the skin for about 1 to 60 seconds.

41. The method according to claim 38, wherein said step of collecting comprises exerting a capillary force on said pooled physiological fluid.

42. The method according to claim 38, wherein said at least one skin-piercing element is integral with a biosensor for determining the concentration of at least one analyte in said physiological fluid.

43. The method according to claim 42, further comprising the steps of:

transferring said collected physiological fluid through said at least one fluid pathway to said biosensor; and
determining the concentration of said at least one analyte.

44. The method according to claim 43, wherein said step of determining the analyte concentration further comprises employing a meter.

45. The method according to claim 43, wherein said step of determining the analyte concentration is performed by electrochemical means.

46. The method according to claim 43, wherein said step of determining the analyte concentration is performed by colorimetric means.

47. The method according to claim 46, wherein said step of determining is performed by fluorescent measuring means.

48. The method according to claim 38, wherein said physiological fluid is blood and said analyte is glucose.

49. The method according to claim 38 wherein said pooling area has a volume which is about 50% to 99% of the volume occupied by said skin piercing element.

50. The method according to claim 49 wherein said pooling area has a volume which is about 50% to 75% of the volume occupied by said skin piercing element.

51. A method for collecting a sample of physiological fluid, said method comprising the steps of:

penetrating the skin to access said physiological fluid;

creating a pooling area within said skin, wherein said pooling has a volume within the range from about 10 to 1,000 nL;

allowing said access physiological fluid to pool within said pooling area; and

exerting a capillary force on said pooled physiological fluid.

52. The method of claim 51 further comprising the step of extracting said pooled physiological sample to biosensor outside the skin.

53. The method of claim 51 wherein said pooling area has a volume within the range from about 50 to 250 nL.

54. A method for determining the concentration of at least one analyte within a physiological fluid sample, said method comprising the steps of:

(a) providing the system of claim 27 wherein said test strip device is operatively received within a distal end of said meter;

(b) spring-loading said test strip device within said meter;

(c) operatively contacting said distal end of said meter with a targeted skin surface;

(d) releasing said spring-loaded test strip device, wherein said targeted skin surface is pierced by said microneedle;

(f) creating a pooling area within the skin adjacent said microneedle whereby said physiological fluid pools within said pooling area; and

(g) collecting said pooled physiological fluid from within the skin by means of said fluid pathway.

55. The method according to claim 54 further comprising the step of applying optimal pressure against said target skin surface with said distal end of said meter.

56. The method according to claim 55 wherein said step of applying optimal pressure comprises the steps of:

sensing the pressure applied;

indicating the amount of said sensed pressure; and

adjusting said applied pressure if necessary according to said indicated amount of pressure.

57. The method according to claim 54 wherein said step of collecting said pooled physiological fluid comprises exerting a capillary force on said pooled physiological fluid by means of said fluid pathway.

58. The method according to claim 54 wherein said step of collecting further comprises applying pressure about the microneedle piercing site.

59. The method according to claim 54 wherein said step of collecting further comprises applying a negative pressure to said pooled physiological fluid.

60. The method according to claim 54 wherein said test strip device is visualized during one or more steps of said method.

61. The method according to claim 60 wherein said one or more steps include steps (a), (b), (c) or (d).

62. The method according to claim 54 wherein said step of providing comprises the step of inserting said test strip into said distal end of said meter.

63. The method according to claim 54 wherein said step of inserting comprises inserting said test strip through said aperture.

64. The method according to claim 54 wherein said step of providing comprises removing a distal portion of said meter and inserting said test strip into said receiving means within said distal end of said meter.

65. A kit for determining at least one target analyte concentration of a physiological sample, said kit comprising a system according to claim 27.

66. The kit according to claim 65, wherein said meter is disposable.

67. A kit according to claim 65 further comprising instructions for using said system.

68. A kit for determining at least one target analyte concentration of a physiological sample, said kit comprising a plurality of test strips according to claim 8.

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