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(54) **Fluid collection apparatus having an integrated lancet and reaction area**

Vorrichtung zum Sammeln von Flüssigkeiten mit integrierter Lanzette und Reaktionsbereich

Dispositif de collection de fluides avec lancette et domaine de réaction intégrée

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WO-A1-01/72220 WO-A1-02/49507
US-A- 5 801 057 US-A- 5 855 801
US-A- 6 102 927

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Description

FIELD OF THE INVENTION

[0001] The present invention relates generally to blood monitoring devices and, more particularly, to a fluid collection apparatus having an integrated lance and reaction area for use in determining one or more analytes in a body fluid.

BACKGROUND OF THE INVENTION

[0002] It is often necessary to quickly obtain a sample of blood and perform an analysis of the blood sample. One example of a need for quickly obtaining a sample of blood is in connection with a blood glucose monitoring system where a user must frequently use the system to monitor the user's blood glucose level.

[0003] Those who have irregular blood glucose concentration levels are medically required to self-monitor their blood glucose concentration level. An irregular blood glucose level can be brought on by a variety of reasons including illness, such as diabetes. The purpose of monitoring the blood glucose concentration level is to determine the blood glucose concentration level and then to take corrective action, based on whether the level is too high or too low, to bring the level back within a normal range. The failure to take corrective action can have serious implications. When blood glucose levels drop too low, a condition known as hypoglycemia, a person can become nervous, shaky, and confused. That person's judgment may become impaired and that person may eventually pass out. A person can also become very ill if their blood glucose level becomes too high, a condition known as hyperglycemia. Both conditions, hypoglycemia and hyperglycemia, are potentially life-threatening emergencies.

[0004] One method of monitoring a person's blood glucose level is with a portable, hand-held blood glucose testing device. A prior art blood glucose testing device 100 is illustrated in FIG. 1. The portable nature of these devices 100 enables the users to conveniently test their blood glucose levels wherever the user may be. The glucose testing device 100 contains a test sensor 102 to harvest the blood for analysis. The device 100 contains a switch 104 to activate the device 100 and a display 106 to display the blood glucose analysis results. In order to check the blood glucose level, a drop of blood is obtained from the body, usually from the fingertip, using a lancing device. A prior art lancing device 120 is illustrated in FIG. 2. The lancing device 120 contains a needle lance 122 to puncture the skin. Some lancing devices implement a vacuum to facilitate drawing blood. Once the requisite amount of blood is produced on the fingertip, the blood is harvested using the test sensor 102. The test sensor 102, which is inserted into a testing device 100, is brought into contact with the blood drop. The test sensor 102 is filled with blood and creates a color change or an elec-

trical current that is measured by the test device 100, which then determines the concentration of glucose in the blood. Once the results of the test are displayed on the display 106 of the test device 100, the test sensor 102 is discarded. Each new test requires a new test sensor 102.

[0005] One problem associated with many conventional testing systems is that the lance and the sensor are two separate, disposable pieces. Two separate pieces require more assembly work. This is time consuming for the user who must assemble the two disposable pieces prior to use. Also, because there are multiple pieces, there are more pieces for the user to keep track of, re-order, etc. Missing pieces may result in the test not being taken at the appropriate time, or it may result in an additional trip to the store, resulting in further inconvenience to the user.

[0006] Another problem associated with current testing devices is the difficulty in harvesting small samples when the sensor is separate from the lance. There is a trend in glucose testing towards smaller and smaller sample volumes. This trend is based on the assumption that there is a corresponding reduction in pain when less sample volume is acquired. As the sample volume is reduced, it becomes more difficult to manually manipulate the sensor in order to harvest the blood. This is especially true for people who may have vision impairments or other disabilities which may make it difficult to manipulate the sensor within a small area.

[0007] Another problem associated with obtaining small sample sizes is related to the precision needed to obtain the samples. When small amounts of blood are drawn by the lance, it is important that the entire sample or most of the sample be drawn into the testing device. When larger volumes of blood are drawn, it is less necessary to obtain all of the blood for the sensor. In small volume testing devices, it is advantageous to have the sensor located proximate to the puncture wound to maximize the amount of blood that is drawn into the sensor for testing. In current testing devices, where the sensor has to be manually moved to the puncture wound, it may be difficult to get close enough to the wound to obtain enough of the sample.

[0008] Another testing device has been developed for the collection of interstitial fluid (ISF) that utilizes an integrated lance and sensor. ISF is collected by piercing just below the skin before any nerve endings or any capillaries. Collecting ISF is sometimes desirable because there is minimal pain involved since it is above any nerve endings. In this device, the lance and sensor chamber is connected via a capillary channel, all of which are made by etching silicon wafers. This requires numerous steps to form. Furthermore, the lance needle is brittle and requires protection from production to final use. The lance needle and sensor are a single part, but a molded part and a cover are needed to house the integrated sensor for final packaging and use.

[0009] Other testing devices have been produced for

testing blood that utilize a sensor with a lance perpendicular to the sensor. In this arrangement, the sensor can be positioned to harvest a sample with the lance puncturing the body either through a hole in the sensor or adjacent to the tip of the sensor. When the sample is produced adjacent to the sensor, harvesting of the sample can be automatic and without user judgement. This approach requires precise alignment of both the lancet and the sensor either at the time of manufacture or at the time of use, preferably by the test device, to make it more convenient for the end user.

[0010] US 5,801,057 discloses a microsampling device for the measurement of glucose and other analyte in blood and a fabrication method for such microsampling device. For a device with one or without a window a patterned depression is etched in the top surface of a silicon wafer to design a micro sampler chamber, a needle bore and a vent channel. The patterned depression in the top surface of the wafer is covered with a cover glass. Subsequently the needle is etched out by removing the silicon and cover glass and a point on the needle is formed.

SUMMARY OF THE INVENTION

[0011] The present invention is a method of manufacturing a fluid collection apparatus that has an integrated lance and reaction area. The method includes providing a sheet of material and then coating the sheet with a photoresist in a pattern on one side of the sheet. The pattern defines a lance, a reaction area and a transfer area between the nose of the lance and the reaction area. At least one side of the sheet is placed in a solvent and is then corroded in areas not covered by the photoresist. The sheet is removed from the acid after a predetermined time to reveal an integrated lance, reaction area and transfer area.

[0012] The above summary of the present invention is not intended to represent each embodiment or every aspect of the present invention. This is the purpose of the Figures and the detailed description which follow.

BRIEF DESCRIPTION OF THE DRAWINGS

[0013] The foregoing and other advantages of the invention will become apparent upon reading the following detailed description and upon reference to the drawings.

FIG. 1 is a top view of a prior art blood glucose testing device.

FIG. 2 is a perspective view of a prior art lance.

FIG. 3a is a perspective view of a fluid collection apparatus according to one embodiment of the present invention.

FIG. 3b is a side view of the fluid collection apparatus of FIG. 3a.

FIG. 4a is a perspective view of a fluid collection apparatus according to another embodiment of the present invention.

FIG. 4b is a side view of the fluid collection apparatus of FIG. 4a.

FIG. 5 is a view of a first side of a sheet having a mask according to one embodiment of the present invention.

FIG. 6a is a view of a second side of a sheet having a mask according to one embodiment of the present invention.

FIG. 6b is a view of a second side of a sheet having a mask according to another embodiment of the present invention.

FIG. 7 is a view of a sheet having a plurality of fluid collection apparatuses according to one embodiment of the present invention.

FIG. 8 is an enlarged view of the circular cut out 8-8 taken from FIG. 7.

DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

[0014] FIG. 3a is a perspective view and FIG. 3b is a side view of a fluid collection apparatus 10 according to one embodiment of the present invention. The fluid collection apparatus 10 is designed to collect a body fluid, for example, blood, so the fluid may be tested for the concentration of a particular analyte, such as glucose. In describing the details of the operation of the fluid collection apparatus 10, the fluid described will be blood pricked from a user's skin and the analyte will be glucose. It is understood that the embodiment may also be used for other fluids and analytes and that these only serve as examples.

[0015] The fluid collection apparatus 10 includes a lid 10b and a body 10a (FIG. 3b). The body 10a has a reaction area 12, a lance 14, and a transfer area, such as a capillary channel 16 (FIG. 3a). According to one embodiment, the reaction area 12, the lance 14, and the capillary channel 16 are all formed of an integrated piece of metal, such as stainless steel. The lance 14 has a nose that is designed to be able to pierce a user's skin (e.g., from a finger tip) to obtain a sample of bleed. The nose may be a sharpened point, or it may be two sharpened points, located on opposite sides of the capillary channel 16. The capillary channel 16 couples the lance 14 to the reaction area 12, such that once the lance 14 pierces the skin of a user, the blood is drawn directly from the point of piercing, up through the capillary channel 16 and into the reaction area 12. The reaction area 12 contains a reagent 13 that is adapted to react with the blood that is drawn into the reaction area 12. A transparent lid (not shown) acts as a cover or top cover and is located over the top of the reaction area 12. Alternately, the reagent could be deposited on the inside surface of the transparent lid.

[0016] According to one embodiment, the fluid collection apparatus 10 can be used in conjunction with a photometric testing device (not shown), which measures a colorimetric reaction. In photometric testing, the reagent 13 used causes a change in color in the reaction area

12. The photometric testing device then measures the amount of color change. Photometric testing is described in more detail in commonly-owned U.S. Patent No. 5,611,999 entitled "Diffuse Reflectance Readhead".

[0017] In another embodiment of the fluid collection apparatus 10, an electrochemical testing device (not shown) is employed. The reaction area 12 includes a pair of electrodes 17. In electrochemical analysis, the change in current across the electrodes 17 caused by the reaction of the glucose and the reagent 13 creates an oxidation current at the electrodes 17, which is directly proportional to the user's blood glucose concentration. The current can be measured by an electrochemical testing device coupled to a pair of terminals (not shown) corresponding to the electrodes 17. The electrochemical testing device can then communicate to the user the blood glucose concentration. An example of an electrochemical test system is described in detail by commonly-owned U.S. Patent No. 5,723,284 entitled "Control Solution And Method For Testing The Performance Of An Electrochemical Device For Determining The Concentration Of An Analyte In Blood". It is also contemplated that other methods of testing the concentration of glucose in blood may be utilized.

[0018] According to the embodiment shown in FIG. 3a, the reaction area 12 has a thickness that is about half the thickness of the fluid collection apparatus 10, which is the thickness of the sheet of material. In these embodiments, the reaction area 12 is bounded on one side by a floor 18 in the fluid collection apparatus 10. These fluid collection apparatuses are also known as being two piece apparatuses. The two piece apparatuses include just the body 10a and the lid 10b (FIG. 3b).

[0019] In other embodiments, such as the one shown in FIGS. 4a and 4b, the fluid collection apparatus 10 is a three piece construction, including the body 10a, the lid 10b, and a second cover 10c. In these embodiments, the reaction area 12 has a thickness equal to the thickness of the fluid collection apparatus 10 and/or the sheet of material. The three piece construction is advantageous for an optical transmission design because the light source is on one side and the photodetector is on the other side of the reaction area 12.

[0020] Turning now to FIGS. 5-6b, the process for manufacturing the integrated fluid collection apparatus 10 will be described. As shown in FIG. 5, a first side 20 of a sheet of material 22 is coated (or masked) in a particular pattern 24 with a photoresist. The pattern 24 is in the shape of the fluid collection apparatus 10. A coating shown by the diagonal lines is formed around the reaction area 12, thus defining the reaction area 12. Similarly, the coating also does not cover the capillary channel 16 but, instead, defines the channel 16.

[0021] Turning now to FIGS. 6a and 6b, a second side 26 of the sheet 22 is coated with a photoresist. FIG. 6a is the manufacturing of the three piece apparatus, or the apparatus shown in FIG. 4a. In FIG. 6a, the coating on the second side 26 is in the pattern 24 of the first side 20. The reaction area 12 and the capillary channel 16

remain unmasked. The capillary channel could also be masked on the second side, but is not shown. In FIG. 6b, the photoresist is spread in a pattern 28 that extends over the whole shape of the fluid collection apparatus 10. In this embodiment, the reaction area 12 and the capillary channel 16 are coated. This pattern creates the two piece apparatus shown in FIG. 3a.

[0022] Once both sides of the sheet 22 have been appropriately coated (for being either a two piece or a three piece apparatus), the sheet 22 is then exposed using lithography. During lithography, the photoresist is hardened by exposing it to ultraviolet light. The sheet 22 is then placed in a solvent, such as an acid. The solvent mills or etches the uncoated portions of the material. The hardened photoresist protects the coated portion of the material from the acid. After a predetermined amount of time (i.e., time sufficient for the solvent to eat through the sheet), the material is removed from the solvent and cleaned.

[0023] Thus, the fluid collection apparatus 10 can be manufactured in only a few steps. Since the lance 14 and the reaction area 12 are one piece, they may be manufactured using this common chemical milling process. By making the lance 14, the capillary channel 16, and the reaction area 12 all one piece, the manufacturing time is reduced, as is the need for extra parts or machines to manufacture the different pieces.

[0024] In the embodiment shown in FIG. 6a, the reaction area 12 and the capillary channel 16 are being milled from both sides. Thus, after a predetermined time, the reaction area 12 and the capillary channel 16 are formed by the acid milling through the entire thickness of the material. This results in the fluid collection apparatus shown in FIG. 4a.

[0025] In the embodiment shown in FIG. 6b, the reaction area 12 and the capillary channel 16 are only left exposed on one side. Thus, the reaction area 12 and the capillary channel 16 will only be milled on one side. In this embodiment, if the sheet of material 22 is kept in the acid for the same amount of time as above, the fluid collection apparatus 10 will have a reaction area 12 and a capillary channel 16 that has half the thickness of the sheet 22. This method results in the fluid collection apparatus shown in FIG. 3a.

[0026] In another alternative embodiment of the fluid collection apparatus 10, the first side 20 of the sheet 22 may be milled using a first acid, while the second side 26 is milled using a second, different acid, having a different strength. This way, the acids can be used to create different thicknesses for the reaction area 12 and the capillary channel 16. For example, if a stronger acid is used on the first side 20 than on the second side 26, when the fluid collection apparatus 10 is finished being milled, the stronger acid will have eroded more than half of the sheet 22, thus the thickness of the reaction area 12 and the capillary channel 16 will be greater than half the thickness of the sheet 22. Conversely, if the weaker acid is used on the first side 20, the thicknesses of the

reaction area 12 and the capillary channel 16 will be less than half the thickness of the sheet 22.

[0027] In the embodiments described above, the fluid collection apparatus 10 typically has a width ranging from about 1,52 mm to about 2,29 mm (0.060 to about 0.090 inches) and a length ranging from about 3,05 mm to about 4,57 mm (0.120 to about 0.180 inches). The reaction area 12 is shown as generally circular and has a radius ranging from about 0,25 mm to about 0,76 mm (0.010 to about 0.030 inches), however, the shape can be oval, diamond, or of a shape to optimize the fluid flow into the reaction chamber. The capillary channel 16 has a width ranging from about 25 μm to about 127 μm (0.001 to about 0.005 inches). The fluid collection apparatus 10 is preferably made of metal, such as stainless steel.

[0028] Once the fluid collection apparatus 10 is created, the lid 10b is attached to one side of the fluid collection apparatus. The lid 10b may include the electrochemical electrodes 17 if electrochemical testing is taking place. Alternatively, the lid 10b may be a clear plastic window if optical testing is taking place. In the embodiments where the reaction area 12 and the collection capillary 16 have the same thickness as the material, the second cover 10c is also attached to a side of the fluid collection apparatus 10.

[0029] Now, the operation of the fluid collection apparatus 10 will be described. A user will pierce their skin (e.g., a finger tip) using the lance 14 located on the end of the fluid collection apparatus 10. As blood exits the laceration, the blood is drawn up into the capillary channel 16 through capillary action, and into the reaction area 12, where it mixes with the reagent 13, creating a measurable reaction as described above. After collecting the sample, the fluid collection apparatus 10 is used with a test device (not shown) to measure the reaction. The testing device may be a colorimetric spectrophotometer or current measuring for the electrochemical sensor as described above.

[0030] Turning now to FIG. 7, a sheet of material 28 with a plurality of fluid collection apparatuses 10 is depicted. FIG. 8 is an enlarged view of a portion of the sheet 28. In some embodiments, a plurality of fluid collection apparatuses 10 may be formed on each sheet 28 as shown in FIG. 7. The number of fluid collection apparatuses 10 on each sheet 28 may be modified to suit individual needs. By manufacturing numerous apparatuses 10 on one sheet, many apparatuses 10 can be dipped in the acid at the same time, which enables quick manufacturing of the fluid collection apparatus 10. It is advantageous to be able to mass produce the apparatuses since that decreases the time and cost of production. Also, there is less sheet of material that is wasted or that needs to be milled by the etchant, which also decreases the manufacturing cost since there is less excess material.

[0031] In other embodiments, the fluid collection apparatuses 10 are formed on a continuous web of material. The webs may be manufactured in rolls and continuously fed through the manufacturing machine. Utilizing a con-

tinuous web of material also allows for continuous manufacturing of the fluid collection apparatuses 10, which is advantageous since it decreases production costs.

Claims

1. A method of manufacturing an apparatus (10) for the collection of a bodily fluid having an integrated lance (14) and reaction area (12), comprising:

providing a sheet of material (22); and coating the sheet with a photoresist in a first pattern (24) on a first side (20) of the sheet (22), the first pattern (24) defining a lance (14), a reaction area (12) and a transfer area (16) between the nose of the lance (14) and the reaction area (12); placing at least the first side (20) of the sheet (22) in a solvent; corroding the sheet (22) in areas not covered by the photoresist; and removing the sheet (22) from the solvent after a predetermined time to reveal the integrated lance (14), the reaction area (12) and the transfer area (16).

2. The method of claim 1, further comprising coating on the second side (26) of the sheet (22) with a photoresist in a second pattern.
3. The method of claim 2, wherein the second pattern of the photoresist is substantially the same as the first pattern (24).
4. The method of claim 2, wherein the second pattern (28) of the photoresist has the same contour as the first pattern (24) and covers the entire area within said contour.
5. The method of one of the claims 1 to 4, wherein the step of placing the sheet (22) in an acid comprises covering both sides (20, 26) of the sheet (22) in the acid.
6. The method of claim 5, wherein the step of placing the sheet (22) in an acid comprises covering one side (20, 26) in one type of acid and covering the other side (20, 26) in a different type of acid, such that the acids corrode the sheet (22) at different rates.
7. The method of claim 1, wherein the photoresist is arranged on the sheet (28) so as to create a plurality of integrated lances (14), reaction areas (12) and transfer areas (16) between the noses of the lances (14) and the reaction areas (12).

8. The method of one of the claims 1 to 7, wherein the thickness of the reaction area (12) is about half the thickness of the sheet (22).
9. The method of one of the claims 1 to 7, wherein the thickness of the reaction area (12) is about equal the thickness of the sheet (22).
10. The method of one of the claims 1 to 9, wherein the sheet (22) is of metal.
11. The method of one of the claims 1 to 10, wherein the transfer area (16) has a width ranging from about 25 μm to about 127 μm .
12. The method of one of the claims 1 to 11, wherein the fluid collection apparatus has a length ranging from about 3,05 mm to about 4,57 mm.

Patentansprüche

1. Verfahren zum Herstellen einer Vorrichtung (10) zum Sammeln eines Körperfluids mit einer integrierten Lanze (14) und einem Reaktionsbereich (12), das Folgendes umfasst:

Vorsehen einer Materiallage (22); und Beschichten der Lage mit einem Photoresist in einem ersten Muster (24) auf einer ersten Seite (20) der Lage (22), wobei das erste Muster (24) eine Lanze (14), einen Reaktionsbereich (12) und einen Übertragungsbereich (16) zwischen der Nase der Lanze (14) und dem Reaktionsbereich (12) definiert;
Anordnen wenigstens der ersten Seite (20) der Lage (22) in einem Lösungsmittel;
Korrodiere die Lage (22) in Bereichen, die nicht mit dem Photoresist bedeckt sind; und
Entnehmen der Lage (22) aus dem Lösungsmittel nach einer vorgegebenen Zeit, um die integrierte Lanze (14), den Reaktionsbereich (12) und den Übertragungsbereich (16) freizulegen.

2. Verfahren nach Anspruch 1, das ferner das Beschichten der zweiten Seite (26) der Lage (22) mit einem Photoresist in einem zweiten Muster umfasst.
3. Verfahren nach Anspruch 2, wobei das zweite Muster des Photoresists im Wesentlichen gleich dem ersten Muster (24) ist.
4. Verfahren nach Anspruch 2, wobei das zweite Muster (28) des Photoresists den gleichen Umriss wie das erste Muster (24) besitzt und die gesamte Fläche innerhalb dieses Umrisses abdeckt.
5. Verfahren nach einem der Ansprüche 1 bis 4, wobei

der Schritt des Anordnens der Lage (22) in einer Säure das Bedecken beider Seiten (20, 26) der Lage (22) in der Säure umfasst.

6. Verfahren nach Anspruch 5, wobei der Schritt des Anordnens der Lage (22) in einer Säure das Bedecken einer Seite (20, 26) mit einem Typ Säure und das Bedecken der anderen Seite (20, 26) mit einem anderen Typ Säure umfasst, derart, dass die Säuren die Lage (22) mit verschiedenen Raten korrodieren.

7. Verfahren nach Anspruch 1, wobei der Photoresist auf der Lage (28) so angeordnet ist, dass er mehrere integrierte Lanzen (14), Reaktionsbereiche (12) und Übertragungsbereiche (16) zwischen den Nasen der Lanzen (14) und den Reaktionsbereichen (12) erzeugt.

8. Verfahren nach einem der Ansprüche 1 bis 7, wobei die Dicke des Reaktionsbereichs (12) etwa gleich der halben Dicke der Lage (22) ist.

9. Verfahren nach einem der Ansprüche 1 bis 7, wobei die Dicke des Reaktionsbereichs (12) etwa gleich der Dicke der Lage (22) ist.

10. Verfahren nach einem der Ansprüche 1 bis 9, wobei die Lage (22) aus Metall besteht.

11. Verfahren nach einem der Ansprüche 1 bis 10, wobei der Übertragungsbereich (16) eine Breite besitzt, die im Bereich von etwa 25 μm bis etwa 127 μm liegt.

12. Verfahren nach einem der Ansprüche 1 bis 11, wobei die Fluidsammelvorrichtung eine Länge besitzt, die im Bereich von etwa 3,05 mm bis etwa 4,57 mm liegt.

Revendications

1. Procédé de fabrication d'un dispositif (10) de collecte d'un fluide corporel comprenant une lance intégrée (14) et une zone de réaction (12), comprenant les étapes suivantes:

prévoir une feuille d'un matériau (22); et revêtir la feuille avec un photorésistant en formant un premier motif (24) sur un premier côté (20) de la feuille (22), le premier motif (24) définissant une lance (14), une zone de réaction (12) et une zone de transfert (16) entre le nez de la lance (14) et la zone de réaction (12);
placer au moins le premier côté (20) de la feuille (22) dans un solvant;
corroder la feuille (22) dans des régions qui ne sont pas couvertes par le photorésistant; et
sortir la feuille (22) du solvant après un temps prédéterminé afin de révéler la lance intégrée

(14), la zone de réaction (12) et la zone de transfert (16).

lecte de fluide est comprise dans la gamme d'environ 3,05 mm à environ 4,57 mm.

2. Procédé selon la revendication 1, comprenant en outre le revêtement du deuxième côté (26) de la feuille (22) avec un photorésistant en formant un deuxième motif. 5
3. Procédé selon la revendication 2, dans lequel le deuxième motif du photorésistant est sensiblement identique au premier motif (24). 10
4. Procédé selon la revendication 2, dans lequel le deuxième motif (28) du photorésistant présente le même contour que le premier motif (24) et recouvre la totalité de la surface à l'intérieur dudit contour. 15
5. Procédé selon l'une quelconque des revendications 1 à 4, dans lequel l'étape consistant à placer la feuille (22) dans un acide comprend le recouvrement des deux côtés (20, 26) de la feuille (22) avec l'acide. 20
6. Procédé selon la revendication 5, dans lequel l'étape consistant à placer la feuille (22) dans un acide comprend le recouvrement d'un premier côté (20, 26) avec un type d'acide, et le recouvrement de l'autre côté (20, 26) avec un type d'acide différent, de telle sorte que les acides corrodent la feuille (22) à des degrés différents. 25
30
7. Procédé selon la revendication 1, dans lequel le photorésistant est appliqué sur la feuille (28) de manière à créer une pluralité de lances intégrées (14), de zones de réaction (12) et de zones de transfert (16) entre les nez des lances (14) et les zones de réaction (12). 35
8. Procédé selon l'une quelconque des revendications 1 à 7, dans lequel l'épaisseur de la zone de réaction (12) correspond à environ la moitié de l'épaisseur de la feuille (22). 40
9. Procédé selon l'une quelconque des revendications 1 à 7, dans lequel l'épaisseur de la zone de réaction (12) est approximativement égale à l'épaisseur de la feuille (22). 45
10. Procédé selon l'une quelconque des revendications 1 à 9, dans lequel la feuille (22) est constituée de métal. 50
11. Procédé selon l'une quelconque des revendications 1 à 10, dans lequel la largeur de la zone de transfert (16) est comprise dans la gamme d'environ 25 μm à environ 127 μm . 55
12. Procédé selon l'une quelconque des revendications 1 à 11, dans lequel la longueur du dispositif de col-

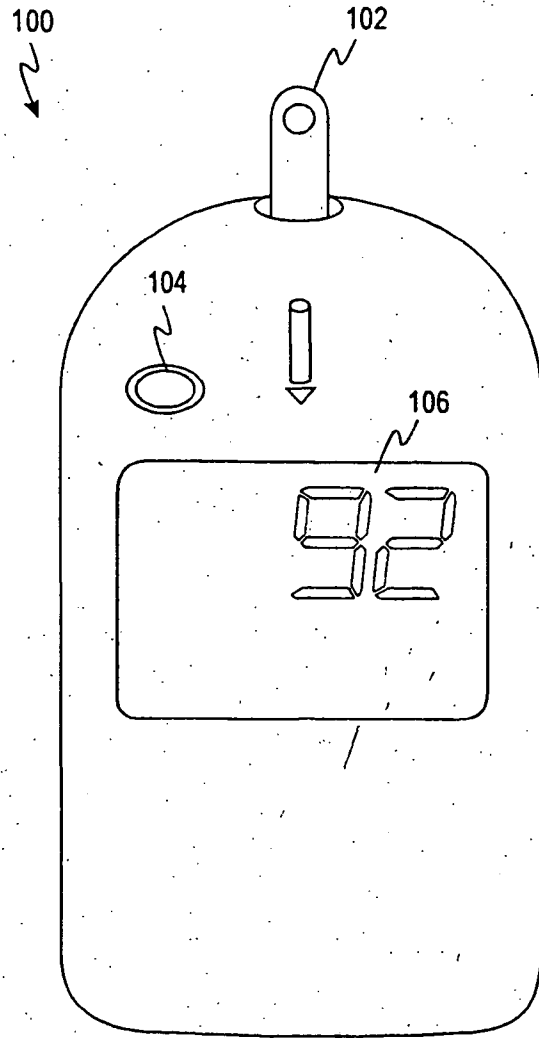


FIG. 1
(Prior Art)

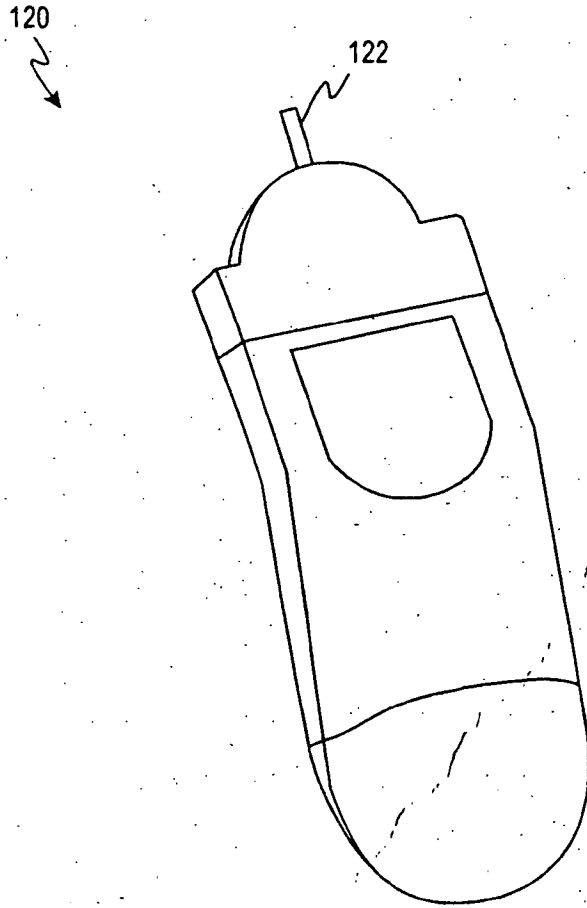


FIG. 2
(Prior Art)

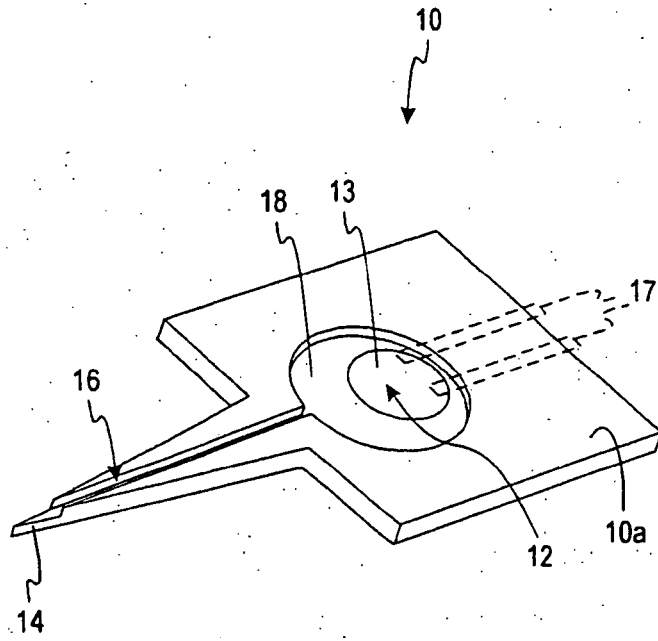


FIG. 3a

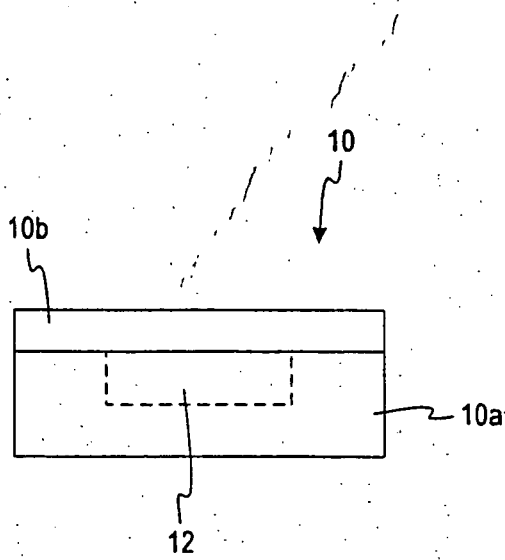


FIG. 3b

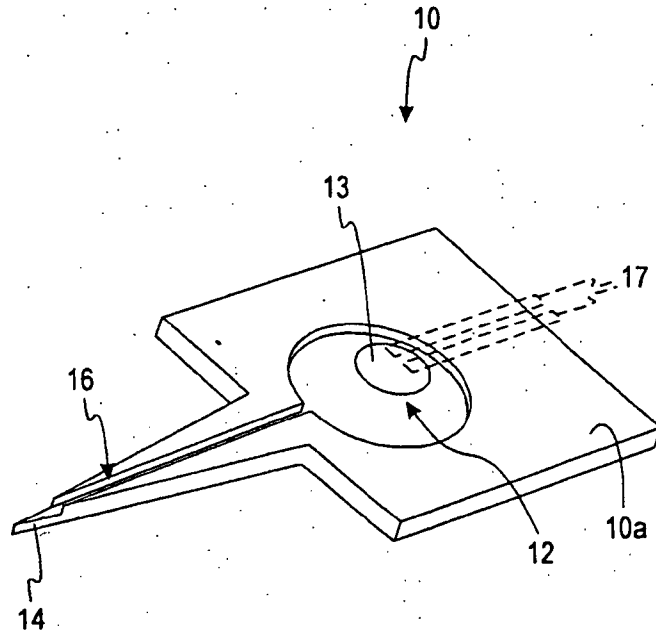


FIG. 4a

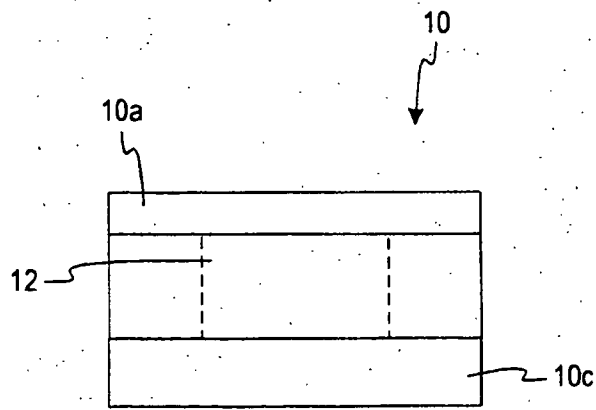


FIG. 4b

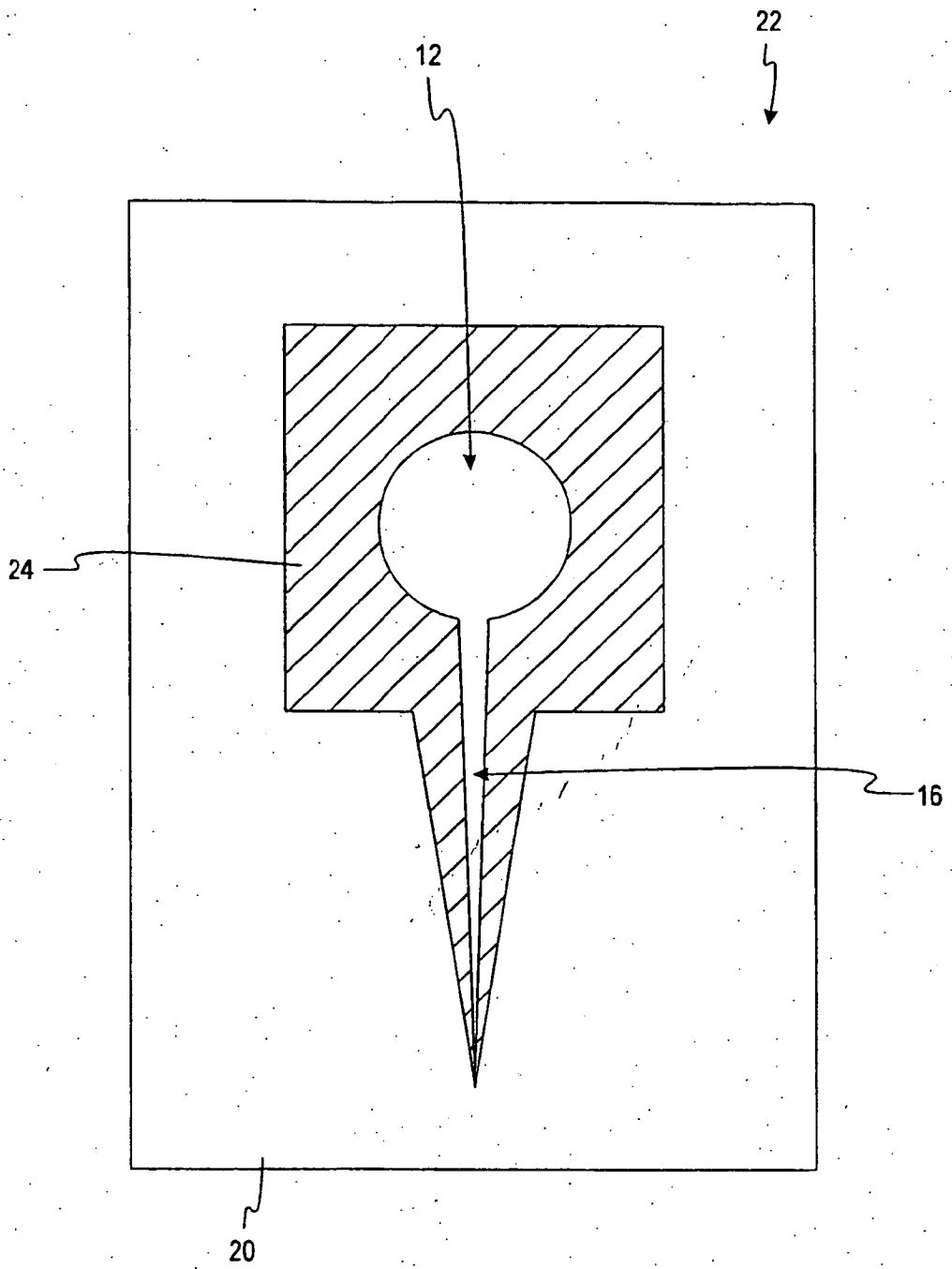


FIG. 5

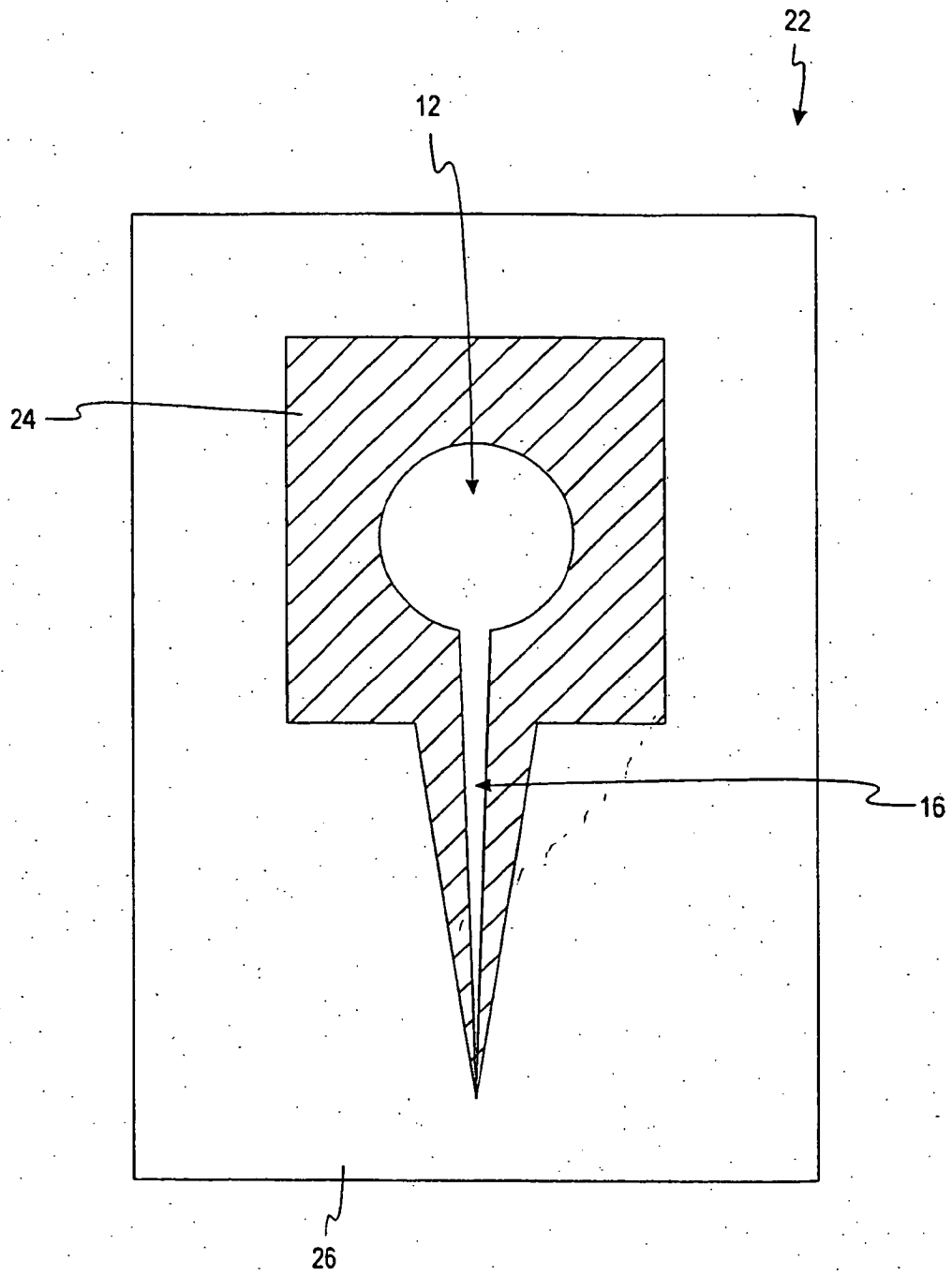


FIG. 6a

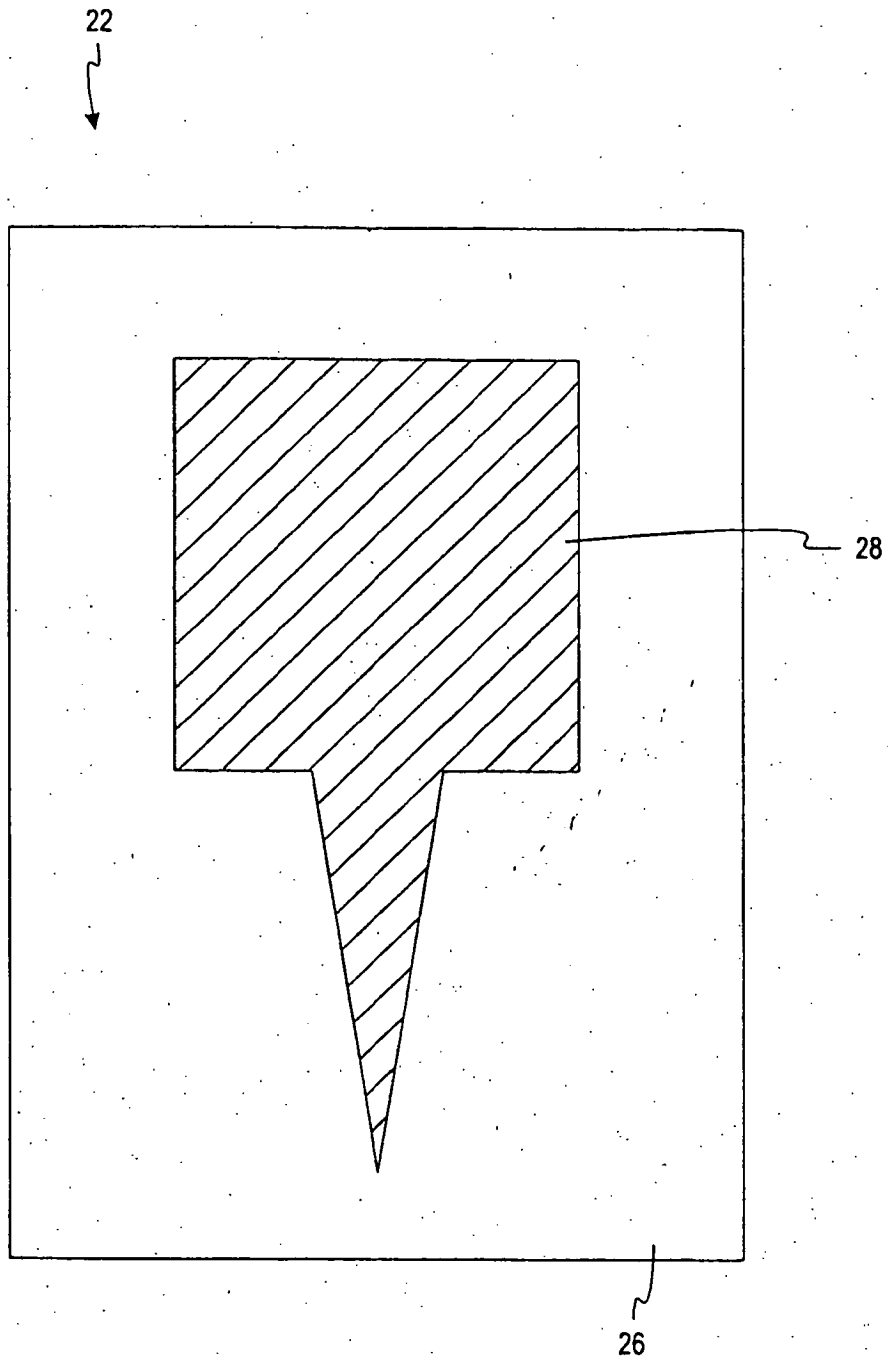


FIG. 6b

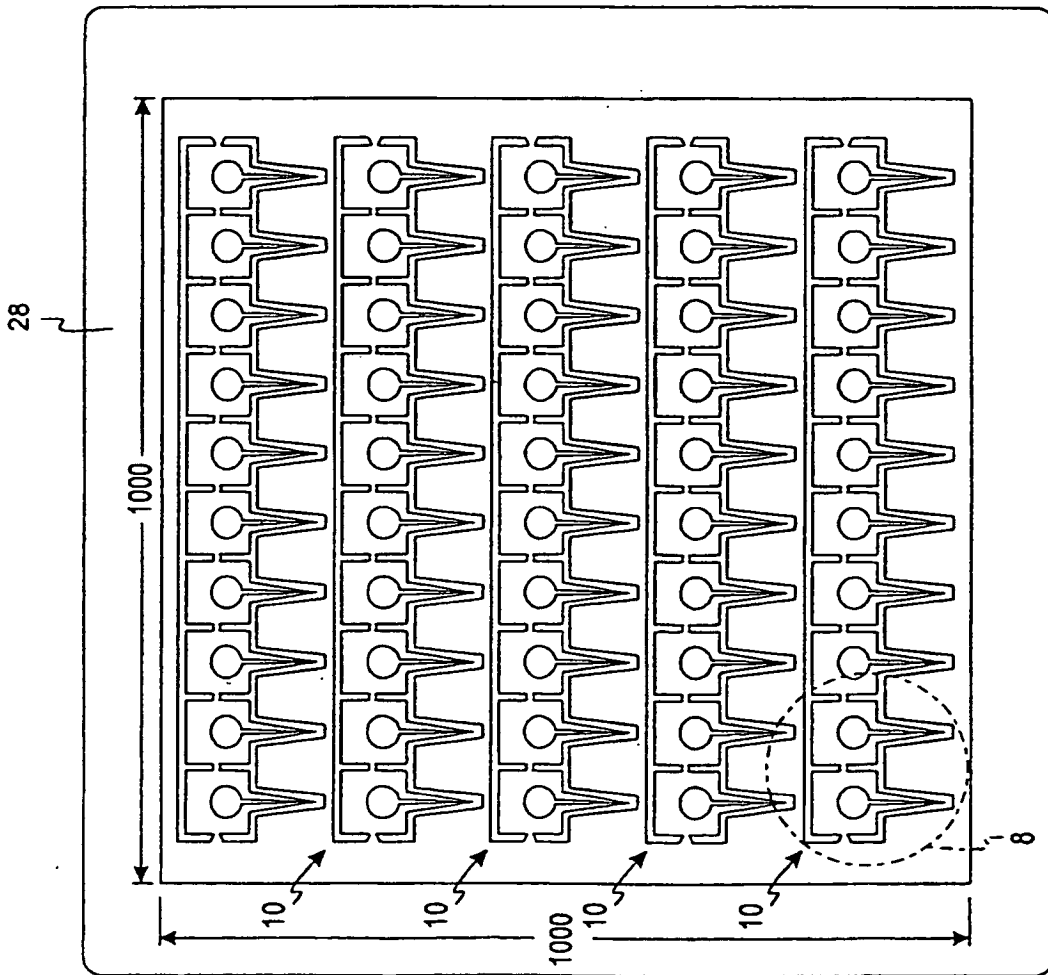


FIG. 7

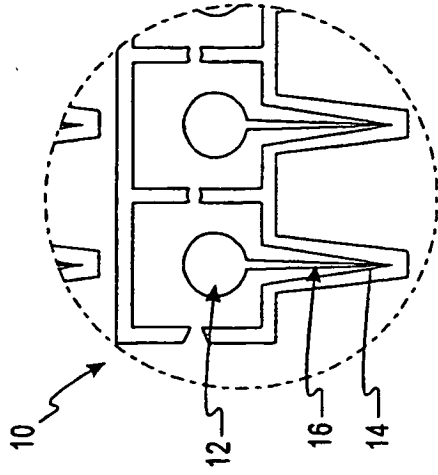


FIG. 8

REFERENCES CITED IN THE DESCRIPTION

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|----------------|---|---------|------------|
| 专利名称(译) | 流体收集设备具有集成的刺血针和反应区域 | | |
| 公开(公告)号 | EP1346686B1 | 公开(公告)日 | 2012-05-02 |
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| 优先权 | 60/361405 2002-03-05 US | | |
| 其他公开文献 | EP1346686A3 EP1346686A2 | | |
| 外部链接 | Espacenet | | |

摘要(译)

一种制造具有集成的喷枪和反应区域的流体收集设备的方法。该方法包括提供一片材料，然后在片材的一侧上用图案的光致抗蚀剂涂覆该片材。该图案定义了喷枪和反应区域。将片材的至少一面置于溶剂中。在未被光致抗蚀剂覆盖的区域中腐蚀片材后，将片材从溶剂中除去并显示出整合的喷枪和反应区域。

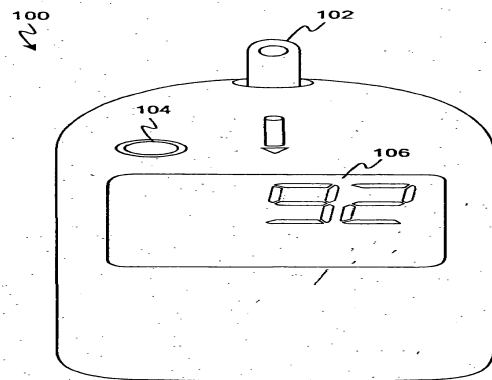


FIG. 1
(Prior Art)