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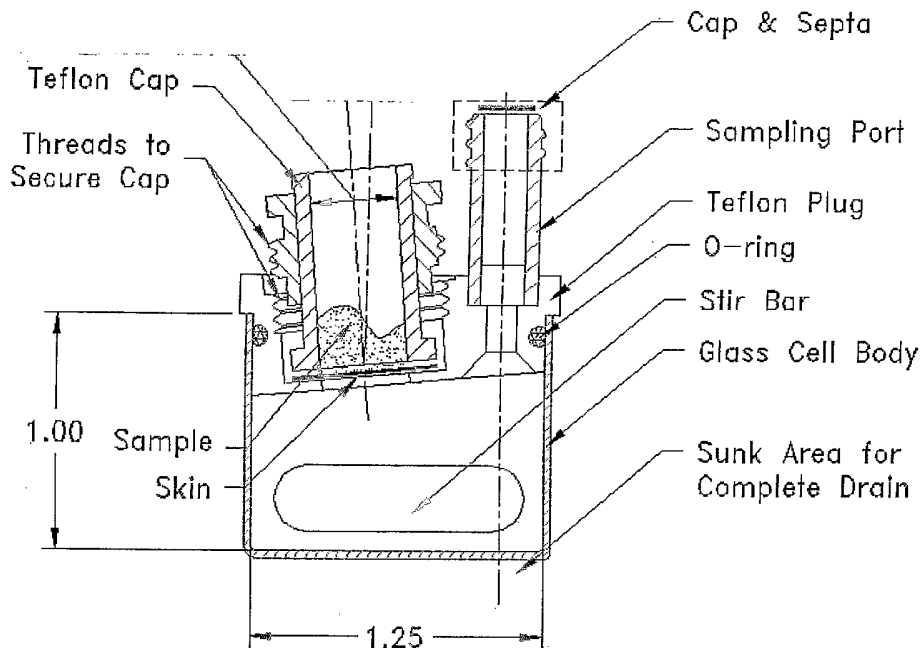
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[Continued on next page]

(54) Title: STATIC DIFFUSION CELL FOR DIFFUSION SAMPLING SYSTEMS



(57) Abstract: A static diffusion cell for use in a diffusion sampling apparatus. The diffusion cell comprises a single-chamber receptor compartment, a donor compartment a diffusion membrane, and a sampling arm. The diffusion membrane is positionned over a first outlet of the receptor compartment between the receptor and donor compartments. A second outlet of the receptor compartment forms a sampling arm and a bubble trap. To facilitate the removal of the bubbles to the second outlet the top surface of the receptor compartment may be slanted or the outlets may be connected with a bubble channel.



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STATIC DIFFUSION CELL FOR DIFFUSION SAMPLING SYSTEMS**BACKGROUND OF THE INVENTION**

5 Field of the Invention: The present invention relates to static diffusion cells useful in automated and manual diffusion sampling systems as well as assay methods that utilize diffusion sampling systems that include one or more diffusion cells according to the present invention. In particular, the present invention provides a static diffusion cell that includes a single chambered
10 receptor compartment, which design reduces or eliminates the disadvantages associated with diffusion cells having multi-chambered receptor compartments and allows for improved sampling systems and assay methods.

15 Description of the Related Art: *In vitro* membrane diffusion systems with automated sampling are widely available for flow-through diffusion cells. However, applicants are presently aware of only two commercial systems that provide both static diffusion cells and automated sampling. Hanson Research Corp. of Chatsworth, CA, sells the Hanson MicroettePlus™ Transdermal
20 Diffusion System, and Logan Instruments Corp. of Somerset, VT, sells the Logan System-902 and Logan System-912 Automated Transdermal Sampling Systems. Logan also sells an upgraded system that includes a cell design similar to the 902-system and an XYZ robot for automatic sampling. Although, the diffusion systems available from Hanson Research and Logan
25 Instruments exhibit differences in design, the systems available from both companies includes a large amount of small diameter tubing and pumps (peristaltic or syringe) that move fluids through multiple compartments within the systems.

30 As illustrated in FIG. 1, in the Logan system, tubing is part of the cell design. The receptor compartment consists of three chambers linked together by small diameter tubing. The diffusion membrane is positioned on the diffusion chamber, which includes a stir bar, a sampling port that allows introduction of a sampling probe, and a water jacket with an inlet and an outlet that facilitate

circulation of water around the diffusion chamber to maintain the diffusion chamber at a desired temperature. A material or formulation to be evaluated is placed over the diffusion membrane, and samples are collected from the collection chamber or flow cell using a suitable collection apparatus. The
5 third chamber is used as a bubble trap. A peristaltic pump continuously circulates the receptor medium between the three chambers to maintain adequate mixing. After receptor medium flows out of the diffusion chamber and flows through both the collection chamber and the bubble trap, the receptor medium returns to the diffusion chamber through a media return.

10 As illustrated in FIG. 2, in the Hanson MicroettePlus™ Transdermal Diffusion System, the diffusion cell consists of a single chamber, but the input arm of the receptor chamber is connected by tubing to a syringe chamber (Microette unit) and the output arm to a central sample collection chamber. Samples
15 from the receptor chambers are collected in the central sample collection chamber by positive displacement initiated by the syringe unit. Therefore, the MicroettePlus™ Transdermal Diffusion System also utilizes multiple chambers interconnected by tubing.

20 The designs of the systems described above can make cell set-up, handling, and cleaning difficult and can lead to inaccurate experimental data. In particular, the relatively extensive use of tubing in the systems available from Logan Instruments Corp. and Hanson Research Corp. introduces several potential problems. For example, the tubing can clog or leak, and the use of
25 tubing can result in variable or inaccurate calculations of cell volume.

Perhaps even more problematic is binding of media constituents, such as the material to be assayed, to the tubing or leaching of chemicals from the tubing into the receptor medium. Both the binding of materials from the receptor medium to the tubing and the leaching of materials from the tubing into the
30 receptor medium can lead to an inaccurate assay of the amount or type of materials that diffuse through the diffusion membrane and into the receptor medium.

Further, systems designed according to those available from Logan Instruments Corp. and Hanson Research Corp. permit the accumulation of air bubbles under the diffusion membrane even after receptor medium has been degassed. Accumulation of air bubbles under the diffusion membrane causes a reduction in the area of the diffusion membrane in contact with the receptor medium, thereby reducing the effective diffusion area. Such a reduction in diffusion area can, in turn, result in inaccurate experimental data. However, removing air bubbles from the systems available from Logan Instruments Corp. and Hanson Research Corp. system is tedious at set-up and very difficult or even unattainable once a diffusion experiment has started. Accordingly, when these systems are used, they typically require supervision to ensure that air bubbles do not accumulate under the diffusion membrane, which supervision defeats, at least in part, the purpose of having an automated sampling process.

SUMMARY OF THE INVENTION

In one aspect, the present invention is directed to a new design for a diffusion cell that can be used in conjunction with automated or manual sampling systems. In particular, the present invention provides a diffusion cell that integrates a diffusion chamber, sampling chamber, and bubble trap into a single receptor compartment. The cell design of the present invention allows for a diffusion cell that is completely free of tubing.

In another aspect, the present invention includes a diffusion sampling system that incorporates one or more diffusion cells according to the present invention.

In yet another aspect, the present invention an assay method that utilizes a diffusion sampling system according to the present invention.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 provides a schematic illustration of a Logan System-902 Transdermal Sampling System showing the three-chamber diffusion cell.

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FIG. 2 provides a schematic illustration of the Hanson MicroettePlus™ Transdermal Diffusion System.

FIG. 3 provides a schematic illustration of a diffusion cell according to the present invention, wherein the sampling arm and bubble trap are part of the receptor compartment.

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FIG. 4 provides a schematic illustration of a diffusion cell according to the present invention, wherein the first opening of the receptor compartment is positioned on the side of the cell, preventing accumulation of air bubbles under the diffusion membrane.

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FIG. 5 provides a schematic illustration of a diffusion cell according to the present invention, wherein the first opening creates a diffusion area that is tilted upward toward the second opening, which works to prevent accumulation of air bubbles under the diffusion membrane

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FIG. 6 and FIG. 7 provide schematic illustrations of diffusion cells according to the present invention, wherein the diffusion cells include a top section and a bottom section and the bottom section can be removed and made of different sizes to provide receptor compartments of different volumes.

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FIG. 8 provides a schematic illustration of an alternative embodiment of a diffusion cell according to the present invention that includes a top section and a bottom section.

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FIG. 9 provides a schematic illustration of a bubble channel that can be formed between the first opening and the second opening in the receptor compartment of a diffusion cell according to the present invention.

5 FIG. 10 depicts one example of arranging the static diffusion cells in a mounting apparatus, wherein the sampling ports are lined up in multiple parallel rows.

FIG. 11 depicts another example of arranging the static diffusion cells in a
10 mounting apparatus, wherein the sampling ports are lined up in a single row.

FIG. 12 depicts sampling heads to match a mounting apparatus as illustrated in Figures 10 and 11, respectively.

15 FIG. 13 illustrates a reinforcing ring for use with a diffusion membrane in a diffusion cell of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

20 In one aspect, the present invention is directed to a new design for a diffusion cell that can be used in conjunction with automated or manual sampling systems. In particular, the present invention provides a diffusion cell that integrates a diffusion chamber, sampling chamber, and bubble trap into a single receptor compartment. The cell design of the present invention allows
25 for a diffusion cell that is completely free of tubing. Moreover, the design of a diffusion cell according to the present invention eliminates the need for a pump for circulating the fluid within various chambers.

It is believed that the design of the diffusion cell of the present invention has
30 several advantages over diffusion cell designs that include multiple compartments, wherein the receptor medium flows into the different compartments through tubing that places the compartments in fluid communication. For example, relative to systems requiring the interconnection

of multiple compartments via lengths of tubing, the diffusion cells of the present invention can be easily removed and replaced within a diffusion apparatus, which eases experimental set-up and cleaning. Moreover, in diffusion systems including multiple chambers interconnected by tubing, calculation of the cell
5 volume includes assessment of the receptor fluid within the tubing, which makes an accurate determination of cell volume difficult. In contrast, because diffusion cells of the present invention include a single-chambered receptor compartment, diffusion cells of the present invention allow easier and more accurate determination of cell volume. The absence of tubing in a diffusion cell
10 according to the present invention also eliminates potential tube leaks and clogs as well as the problems that result from binding of receptor medium materials to tube walls or leaching of chemicals from the tubing into the receptor medium. Diffusion cells designed according to the present invention are also ease the tasks of replenishing receptor medium and maintaining the
15 receptor medium at a constant volume within the diffusion cell.

Further, unlike existing sampling systems (e.g., the Hanson and the Logan systems described earlier), the diffusion cell design of the present invention allows for relatively easy removal of bubbles appearing under the surface of the
20 diffusion membrane during the course of the experiment. In preferred embodiments, the static diffusion cell of the present invention is designed to automatically reduce the possibility of accumulation of air bubbles at the surface of the diffusion membrane exposed to the receptor compartment.

Each embodiment of the diffusion cell of the present invention illustrated herein is shown without a water jacket. Existing diffusion cells are often designed to include a water jacket, which serves to control the temperature of the diffusion cell by circulating water of a desired temperature around at least a portion of the receptor compartment. Though the diffusion cell of the present invention
25 can be designed to include a water jacket, if desired, presently preferred embodiments do not include such a feature. Eliminating the water jacket further simplifies the design of the diffusion cell, and, in particular, eliminates the need for the tubing and fixtures to support such a temperature regulating
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system. In addition, control of the temperature within the diffusion cells of the present invention can be achieved through alternative means. For example, temperature control of a diffusion cell according to the present invention can be achieved by positioning the diffusion cell within a mounting block that is
5 regulated to a desired temperature and is designed to accommodate one or more diffusion cells.

A first embodiment of the diffusion cell of the present invention is illustrated in FIG. 3. The diffusion cell of the present invention includes a single-chamber
10 receptor compartment, a donor compartment, a diffusion membrane, and a sampling arm. As can be seen in FIG. 3, the diffusion membrane is positioned over the first outlet and once the diffusion cell is assembled, a top surface of the diffusion membrane forms at least a portion of the bottom surface of the donor compartment, and a bottom surface of the diffusion membrane forms at
15 least a portion of the top surface of the receptor compartment. The diffusion membrane used in a diffusion cell according to the present invention may be any natural or synthetic material suitable for application in a diffusion cell. Natural membranes useful in a diffusion cell according to the present invention include, but are not limited to, skin, mucosal membranes, cornea, and epithelial
20 membranes (e.g., intestinal, colonic, or nasal epithelial membranes). In a preferred embodiment, the diffusion membrane is formed of animal or human skin.

In order to ensure the diffusion membrane is properly held in place between the
25 donor and receptor compartments, the diffusion membrane may be positioned over or disposed between a device or component that reinforces the diffusion membrane and allows the diffusion membrane to be securely held in place without undesired damage to the membrane. As is illustrated in FIG. 13, in preferred embodiments, a reinforcing ring, such as a washer or gasket, is
30 positioned under the diffusion membrane or over the diffusion membrane. Alternatively, two reinforcing rings may be positioned both over and under the diffusion membrane. A reinforcing ring used in a diffusion cell of the present invention can be formed of any suitable material, such as a polymer material.

In addition, a cover material, such as a cover fabric (shown in FIG. 13), can be positioned over the diffusion membrane and test material.

5 A test material is deposited in the donor compartment in contact with the top surface of the diffusion membrane. The test material includes one or more constituents, such as one or more drugs, to be tested for permeability or flux across the diffusion membrane. Though in preferred embodiments the diffusion cell of the present invention is used to evaluate the flux of one or more drugs contained in a test material across the diffusion membrane, diffusion
10 cells according to the present invention are not so limited in use. A diffusion cell according to the present invention can be used to evaluate the permeability or flux of virtually any desired substance across a chosen diffusion membrane. The test material, therefore, can include a wide range of substances or formulations. For example, the test material may simply be a desired amount
15 of a particular compound at a chosen purity. Alternatively, the test material may include a formulation of two or more materials, such as a liquid formulation (e.g., a solution, suspension, emulsion, etc.) or a lotion, cream, gel, or other semi-solid formulation. Even further, the test material may include a drug delivery device, such as a transdermal therapeutic device, designed to delivery
20 a chosen substance, such as one or more therapeutic agents.

The receptor compartment of a diffusion cell of the present invention may be designed according to any desired size or shape. However, the geometry of the receptor compartment is preferably designed for efficient stirring of the
25 receptor medium. The receptor compartment typically includes a magnetic stir bar or other suitable means to ensure proper stirring of the receptor medium throughout the diffusion cell, which reduces the possibility of forming stagnant diffusion layers near the diffusion membrane. A receptor compartment of a diffusion cell according to the present invention also includes a first outlet and a
30 second outlet.

The first outlet may be formed to any size and shape that allows positioning of the diffusion membrane over the first opening. For example, where the

receptor compartment is designed to accommodate a 5-30 ml volume of receptor medium, the first opening is preferably sized from about 0.7 to about 5 cm². However, the size and shape of the first opening can be varied, as desired, to allow the use of differently sized diffusion membranes and to suit any particular test conditions. As can be seen in FIG. 3, the first opening can also be adapted to facilitate positioning of the diffusion membrane and association of the donor compartment with the receptor compartment. The first outlet and donor compartment of the diffusion cell illustrated in FIG. 3, are designed with a flange that facilitate clamping of the donor compartment over the diffusion membrane and the first opening of the receptor compartment. Of course, it is to be understood that the donor compartment can be associated with the receptor compartment using any suitable mechanism, not just a clamp. For example, the donor compartment can be associated with the receptor compartment using a threaded connection, a male-female connection, a snap-fit connection, or through a friction or interference fit. Both the donor compartment and the receptor compartment, including the first opening, can be adapted as necessary to facilitate the use of a desired mechanism for the association of the donor compartment with the receptor compartment.

The second outlet of the receptor compartment serves as a sampling arm and a bubble trap. The second outlet may be sized and shaped according to any desired configuration providing a suitable sampling arm. Typically, the opening of the second outlet will be smaller than that of first outlet. The second outlet includes a cap or a seal having a septum that can be penetrated by a sampling needle. Typically the opening of the second outlet will be circular in shape and will be about 0.5 cm in diameter. In preferred embodiments, the second outlet is sized and shaped so that HPLC vial caps with a septum can be used to cap the second outlet.

The sampling arm formed at the second outlet also serves as a bubble trap, allowing the removal of bubbles that form under the diffusion membrane. Depending on the design of the diffusion cell of the present invention, bubbles are removed either by tilting the cell and forcing the bubbles from under the

diffusion membrane and into the sampling arm, or the diffusion cell may be designed such that bubbles forming within the cell automatically migrate into the sampling arm formed by the second outlet. If large bubbles accumulate in the sampling arm and the total receptor medium level decreases, more
5 receptor medium can be easily added through the sampling arm to keep the total receptor medium level constant.

The donor compartment of a diffusion cell according to the present invention is positioned over the first opening of the receptor compartment and can be sized
10 and shaped as desired to meet any experimental need. The donor compartment is designed to contain the test material. A cap or seal, such as a screw cap, septum, or the like, may be provided over the donor compartment and may be necessary where the test material is a liquid or low viscosity or where evaporation or contamination of the test material are to be reduced or
15 eliminated. As is indicated above, the donor compartment can be associated with the receptor compartment using any suitable mechanism. For example, the donor compartment may be clamped to the receptor compartment or, alternatively, the donor compartment may be associated with the receptor compartment using a threaded connection, a male-female connection, a snap-fit connection or by a friction or interference fit. Therefore, the design of one or
20 more components forming the donor compartment can be adapted to facilitate association of the donor compartment with the receptor compartment according to any desired mechanism. In each embodiment, the mechanism for associating the donor compartment with the receptor compartment maintains
25 the donor compartment in close contacting relationship with the diffusion membrane or the first opening of the receptor compartment. Though the means for associating the donor compartment with the receptor compartment may form an adequate seal between the two compartments and the diffusion membrane, one or more additional sealing elements, such as one or more O-rings or gaskets, may be used where the diffusion membrane, donor
30 compartment, or receptor compartment interface with one or more other components of the diffusion cell.

FIG. 4, FIG. 5 and FIG. 8 illustrate embodiments of the diffusion cell of the present invention that facilitate the automatic migration of bubbles from the underside of the diffusion membrane positioned at the first outlet of the receptor compartment to the bubble trap formed at the second outlet of the receptor compartment. These designs permit bubbles that may be formed near the diffusion membrane to move towards the bubble trap formed by the second outlet without requiring removal of the diffusion cell from a diffusion apparatus or manipulation of the diffusion cell by a human operator (*e.g.*, tilting the diffusion cell to move bubbles into the bubble trap).

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As shown in FIG. 4, the first outlet of the receptor compartment can be positioned on a side of the receptor compartment instead of the top. Designing the diffusion cell in this manner effectively rotates the arrangement of the diffusion membrane from a roughly horizontal position to a roughly vertical position. Because the diffusion membrane is positioned horizontally on one side of the receptor compartment, any bubbles formed within the receptor compartment will tend to rise away from the surface of the diffusion membrane to the top of the receptor compartment, where the second outlet and bubble trap are located. In addition, if desired, the top surface of the receptor chamber can form an incline that raises toward the second outlet, thereby further increasing the likelihood that any bubbles formed within the receptor compartment will automatically migrate into the bubble trap formed by the second outlet. Where the first outlet is positioned at one side of the diffusion cell, it is necessary that the association of the diffusion membrane, donor compartment and receptor compartment form a seal that prevents leaking of the receptor medium from within the receptor compartment. Any suitable mechanism for associating the donor and receptor compartments can be used. The necessary seal at the interface between the diffusion membrane, donor compartment and receptor compartment can be formed solely by the mechanism associating the components, or, alternatively, one or more additional sealing members may be included to provide the desired seal. The mechanisms already described for associating the donor compartment and receptor compartment, as well as the additional sealing members described

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herein are also suitable for use in a diffusion cell designed according to the embodiment illustrated in FIG. 4.

FIG. 5 and FIG. 8 illustrate diffusion cells according to the present invention wherein both the first and second outlets of the receptor compartment are located at the top of the receptor compartment, but the first outlet is designed such that the portion of the top surface of the receptor compartment formed by the bottom surface of the diffusion membrane inclines upward toward the second outlet. This tilt or inclination of the top surface of the receptor compartment toward the second outlet of the receptor compartment works to automatically direct any bubbles that form or come to rest on the bottom surface of the diffusion membrane toward the bubble trap formed by the second outlet.

In a particularly preferred embodiment, the diffusion cell of the present invention is not only designed with receptor compartment having a top surface that inclines upward toward the second opening, but the diffusion cell also includes a channel connecting the first and second outlets. The channel can simply be a depression formed in the top surface of the receptor compartment that extends between the first and second outlets. An illustration of such a channel is provided in FIG. 9. Typically, the first outlet of the receptor compartment will not be designed such that the bottom surface of the diffusion membrane is flush with the other portions of the top surface of the receptor compartment. Because of this, bubbles may be trapped at the step formed where the bottom surface of the diffusion membrane interfaces with the remainder of the top surface of the receptor compartment. Forming a channel between the first and second outlet reduces any step formed at this interface and thereby further facilitates the automatic migration of bubbles from the bottom surface of the diffusion membrane to the bubble trap formed by the second outlet.

The volume of receptor medium contained within a diffusion cell according to the present invention must be considered when designing a diffusion study.

Commonly, receptor compartment volumes for diffusion cells range from about 5-30 ml. However, diffusion cells having virtually any desired volume of receptor media can be designed. When it is anticipated that the material to be assayed from the receptor media exhibits a low permeability or flux across the diffusion membrane, relatively small volumes of receptor medium are desirable and may be necessary so that the concentration of the material to be assayed within the receptor medium can be above the limit of detection for the assay method used within a reasonable amount of time. However, when it is anticipated that the material to be assayed from the receptor media exhibits a high permeability or flux across the diffusion membrane, a relatively large volumes of receptor media are desirable and may be necessary to maintain sink conditions in the receptor compartment throughout the diffusion study. Therefore, the volume of the receptor compartment can be varied to provide any particular volume of receptor medium useful for a chosen diffusion study.

To better facilitate tailoring the volume of the receptor compartment to a chosen diffusion study, the diffusion cell of the present invention can be designed having separable top and bottom sections (shown in FIG. 6 – FIG.8). The top section of such an embodiment is formed as a cap or a plug, includes the first outlet of the receptor compartment and integrates an area for association of the donor compartment with the receptor compartment. A bottom surface of the top section also forms at least a portion of the top surface of the receptor compartment. Preferably, the top section also incorporates the second outlet of the receptor compartment and, as a result, the sample arm and bubble trap of the diffusion cell. The bottom section is typically a cup-, well-, or tube-shaped reservoir that is removable from the top section of the diffusion cell. The shape or geometry of the bottom section is preferably designed for efficient stirring of the receptor medium. A sealed receptor compartment of a given volume is formed when the top and bottom sections of the diffusion cell are properly associated. By varying the volume of the bottom section of a diffusion cell having separable top and bottom sections, the volume of the receptor compartment can be readily tailored to meet the needs of a desired diffusion study.

Where the diffusion cell of the present invention is designed with separable top and bottom sections, the two sections of the diffusion cell can be associated using any suitable means. In a preferred embodiment (illustrated in FIG. 8) the top and bottom section are designed to fit together with a friction or interference fit. In such an embodiment, the top section may include a lip, flange, or other feature that ensures the top section is inserted into the bottom section to a desired depth, providing a receptor compartment having a desired volume. Moreover, to ensure that an adequate seal is formed between the top and bottom sections, the top or bottom section can incorporate one or more sealing members, such as one or more O-rings or gaskets. Where the top and bottom sections of a diffusion cell of the present invention are designed to be associated with a friction or interference fit, the force required to associate and dissociate the top and bottom sections should be at least sufficient to maintain the two sections together during a diffusion test and to form an adequate seal where the top and bottom sections interface.

It should be understood that top and bottom sections of a diffusion cell of the present invention might also be associated using any other suitable mechanism. For example, the top and bottom sections of a diffusion cell according to the present invention can be associated using a clamp. Alternatively, the top and bottom sections of a diffusion cell of the present invention can be associated using, for example, a threaded connection, a male-female connection or a snap-fit connection. The design of the top and bottom sections of a diffusion cell of the present invention is flexible and can be altered to accommodate the use of virtually any suitable connection mechanism.

The various different components of diffusion cells of the present invention can be fabricated using materials well known in the art. However, in preferred embodiments, at least a portion of the receptor compartment is formed of a glass material. Where the diffusion cell according to the present invention includes a top and bottom section, it is preferable to form the top section of TEFLON® and the bottom section of glass. However, any other materials that

are suitable for application in a diffusion cell of the present invention may also be used. For example, instead of TEFLON®, the top section of a diffusion cell of the present invention can be formed of a metal or metal alloy, a polymer material or glass. Moreover, instead of glass, the bottom section of a diffusion cell of the present invention can be formed of a metal or metal alloy or a polymer material. A material is suitable for use in the fabrication of a diffusion cell of the present invention provided it is capable of withstanding exposure to the anticipated test conditions without physical failure and without contaminating the receptor media or retaining undesirable amounts of the material to be assayed.

The diffusion cells of the present invention can be used to conduct a variety of assays used in the art. For example, diffusion cells of the present invention can be used to test various drug dosage forms, including transdermal dosage forms, oral dosage forms, ocular dosage forms such as transdermal or transmucosal patches, tablets, semi-solid dosage forms, gel formulations, pastes, ointments, emulsions, suspension, drops and the like.

Diffusion cells according to the present invention can also be used for mechanistic studies, e.g. transport studies through epithelia that focus on various parameters, such as vehicle effect, ionic strength, markers and the like. The diffusion cells of the present invention can also be used for non-passive diffusion assessments that involve active transport mechanisms, which include iontophoresis, sonophoresis, and the like.

The diffusion cells of the present invention can be used to form a diffusion sampling system. A diffusion sampling system of the present invention includes one or more diffusion cells according to the present invention positioned within a mounting apparatus, such as a mounting block, that allows for stirring of the receptor media in the diffusion cells and serves to maintain the diffusion cells at a desired temperature. Where a mounting block is used, the mounting block is preferably formed of a conductive metal, such as an aluminum alloy or stainless steel, that can be maintained at a substantially

uniform temperature throughout the mounting block and allows the use of magnetic stir bars within the receptor compartment. The diffusion cells included in a diffusion sampling system of the present invention are positioned within the mounting apparatus such that sampling ports of each diffusion cell are readily accessible. In preferred embodiments, the diffusion cells include a keying feature that require the diffusion cells to be positioned within the mounting apparatus such that the sample ports are aligned in a configuration that facilitates easy sampling. Once positioned within the mounting apparatus, the diffusion cells can be sampled manually or automatically.

Where automatic sampling of multiple diffusion cells is desired, the diffusion sampling system of the present invention can include, for example, a robot with multidirectional flexibility, such as an XYZ. XYZ robots are often used in high throughput screening applications and are capable of controlled, programmable movements in all directions along the XYZ axes. XYZ robots can also be provided with sampling heads that allow the simultaneous sampling of several cells (generally up to 6 to 12 cells simultaneously). Depending on the positioning of the diffusion cells in the mounting apparatus, different robot sampling heads can be used. Examples of two designs for the blocks and the matching sampling heads are shown in Figs. 10-12. The static diffusion cells of the present invention are also adapted for manual sampling.

The above-described exemplary embodiments are intended to be illustrative in all respects, rather than restrictive, of the present invention. Thus the present invention is capable of many variations in detailed implementation that can be derived from the description contained herein by a person skilled in the art. All such variations and modifications are considered to be within the scope and spirit of the present invention.

CLAIMS

5 We claim:

1. A diffusion cell comprising a receptor and donor compartment, wherein the receptor compartment is a single-chambered compartment that incorporates a sampling arm and a bubble trap.

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2. A diffusion cell comprising a receptor and donor compartment, wherein the receptor compartment is a single-chambered compartment that includes a first outlet and a second outlet, a diffusion membrane and the donor compartment are positioned over the first outlet, and the second outlet forms a

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3. The diffusion cell of claim 2, wherein the first outlet is formed on a side of the receptor compartment and the second outlet is formed at a top surface of the receptor compartment.

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4. The diffusion cell of claim 2, wherein a top surface of the receptor compartment inclines upward toward the second opening.

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5. The diffusion cell of claim 2, wherein a bottom surface of the diffusion membrane forms at least a portion of the top surface of the receptor compartment and the first outlet of the receptor chamber is formed such that the portion of the top surface of the receptor compartment formed by the bottom surface of the diffusion membrane inclines upward toward the second outlet.

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6. The diffusion cell of claim 5, wherein the first outlet and the second outlet are formed at the top surface of the receptor compartment and a bubble channel extends between the first and second outlet.

7. The diffusion cell of claim 2, wherein the receptor compartment includes a top surface, the first outlet and the second outlet are formed in the top surface, and a bubble channel extends between the first and second outlet.

5 8. The diffusion cell of claim 1 or 2, wherein the diffusion cell is formed of a top section and a bottom section and the top and bottom sections are separable.

9. The diffusion cell of claim 8, wherein the top section of the diffusion cell
10 comprises the first outlet of the receptor compartment.

10. The diffusion cell of claim 9, wherein the top section of the diffusion cell comprises the first outlet and the second outlet of the receptor compartment.

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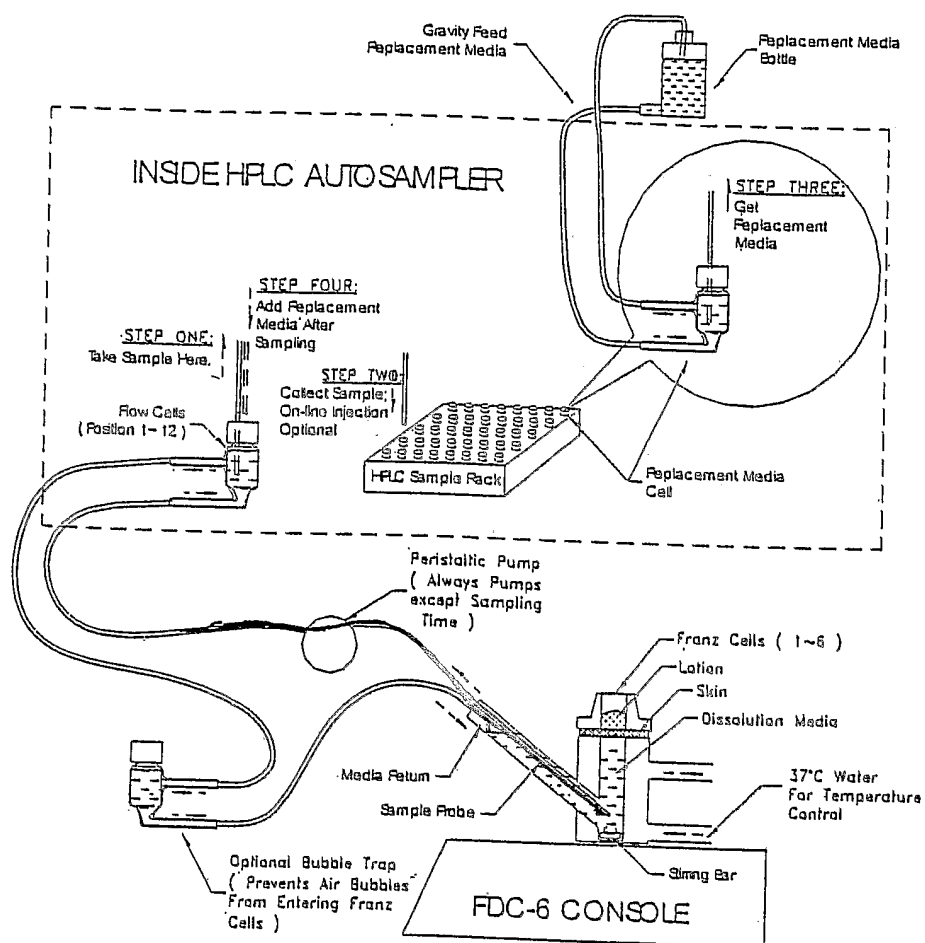


FIG 1

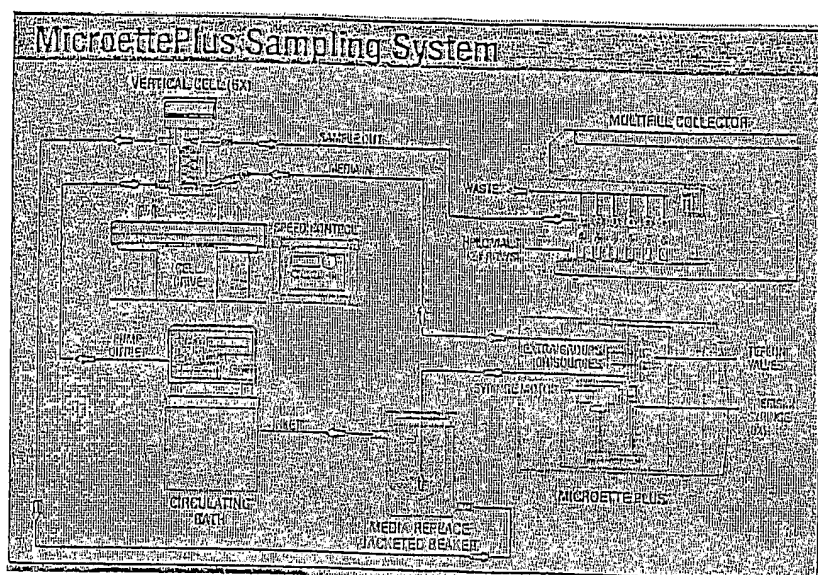


FIG. 2

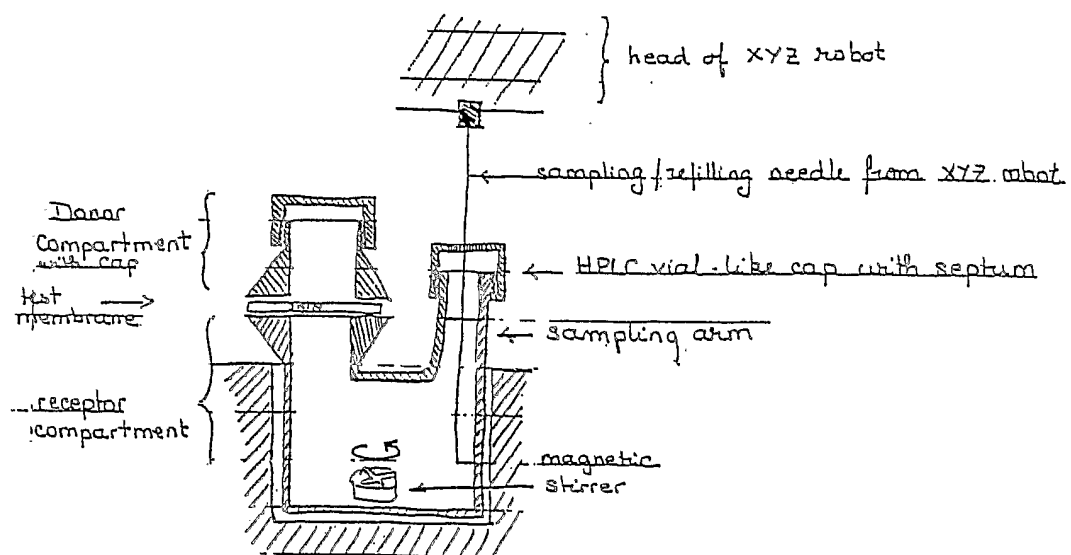


FIG 3

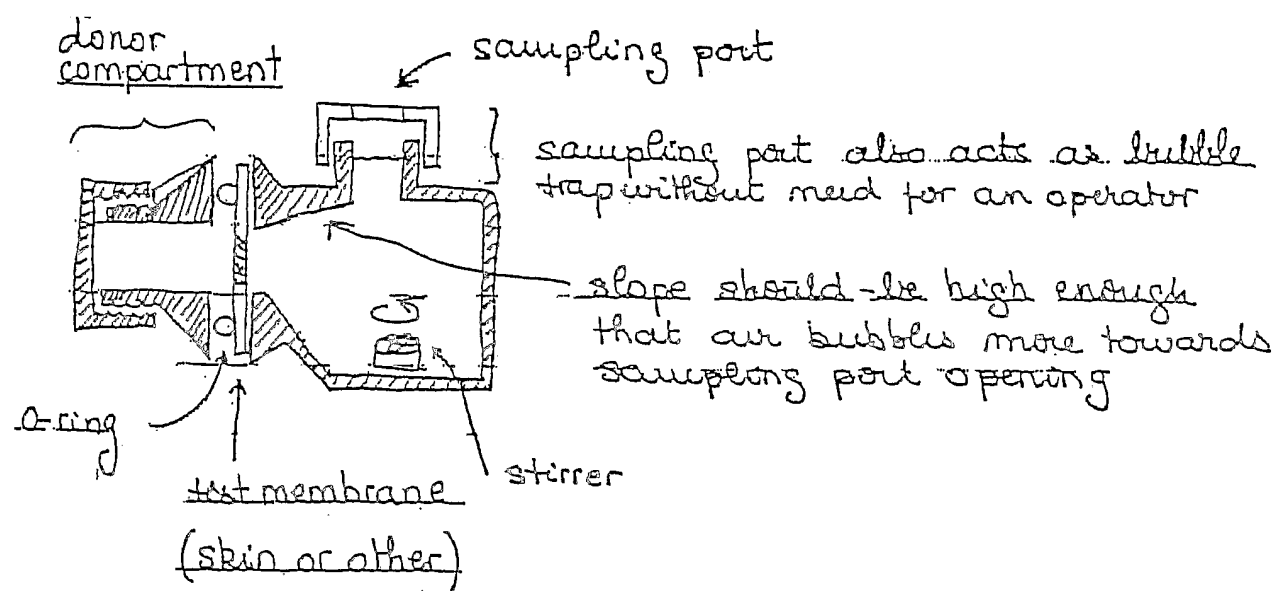


FIG. A

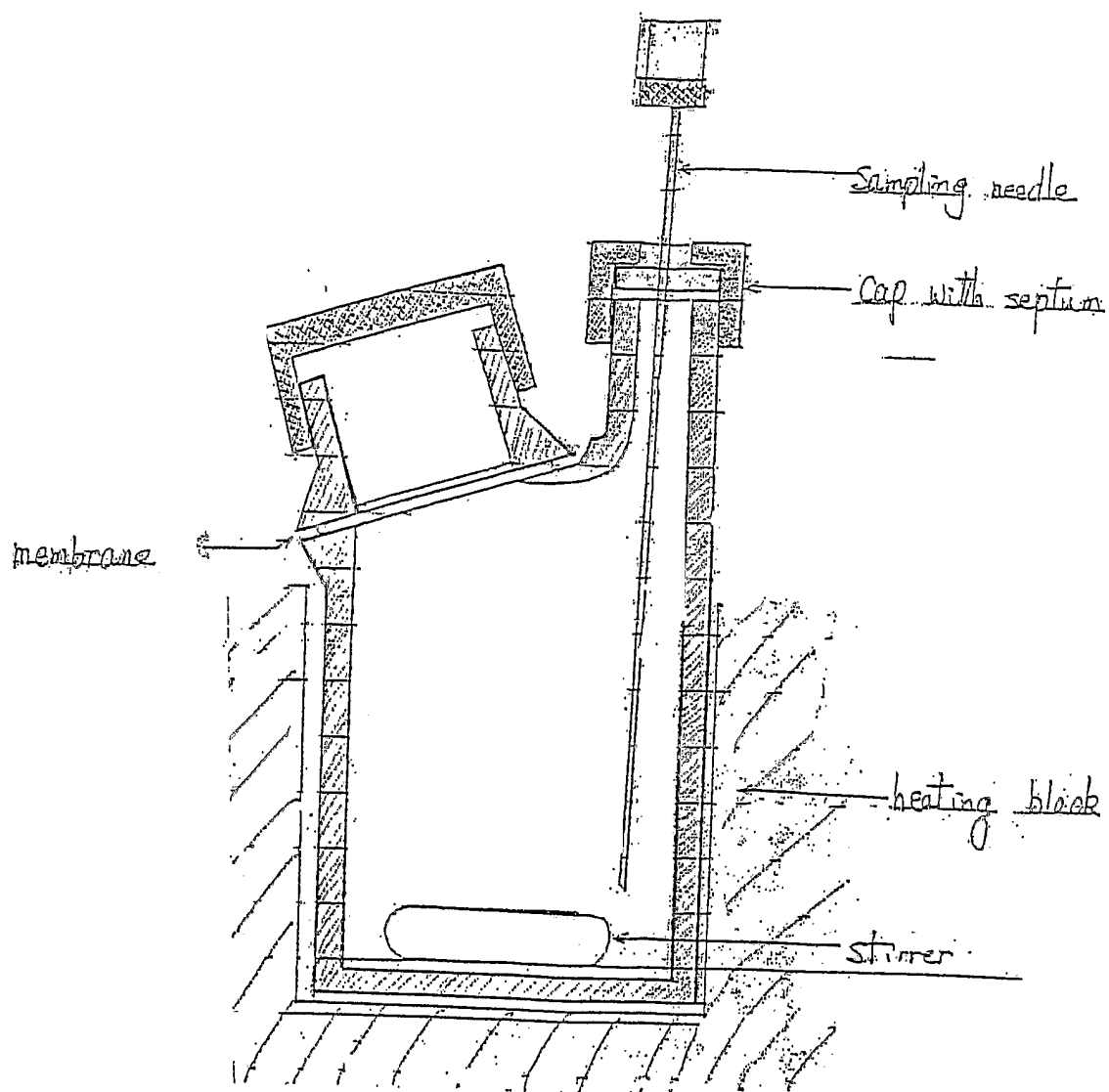


FIG. 5

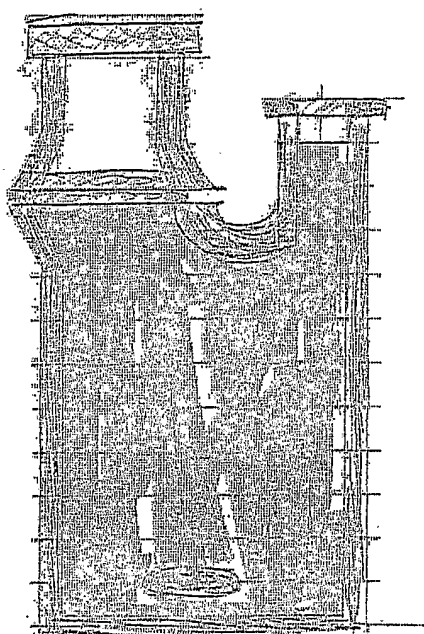


FIG. 6

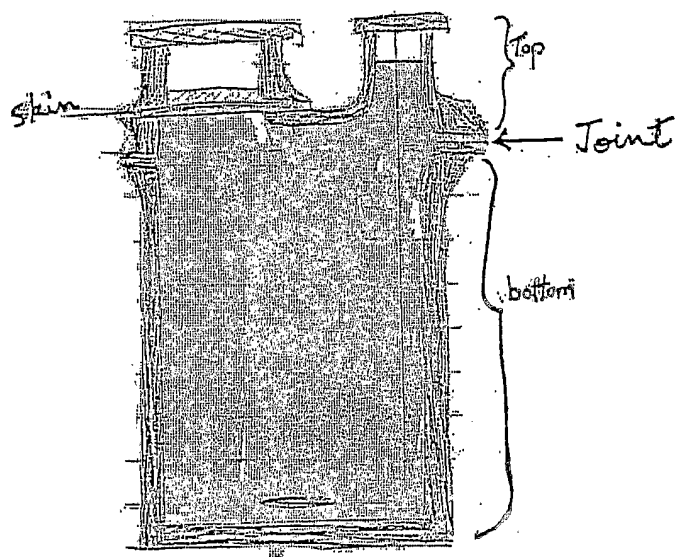


FIG. 7

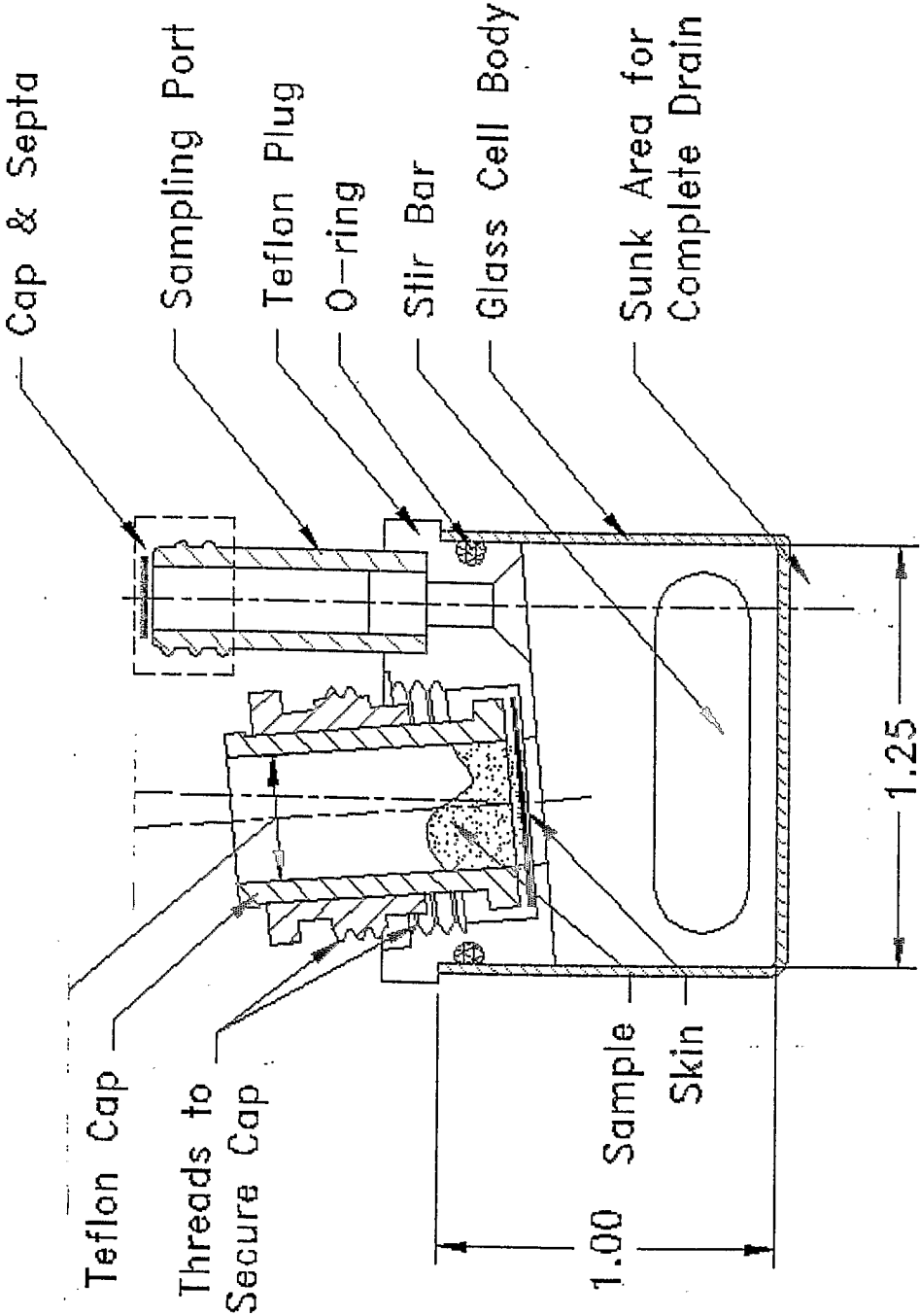


FIG. 8

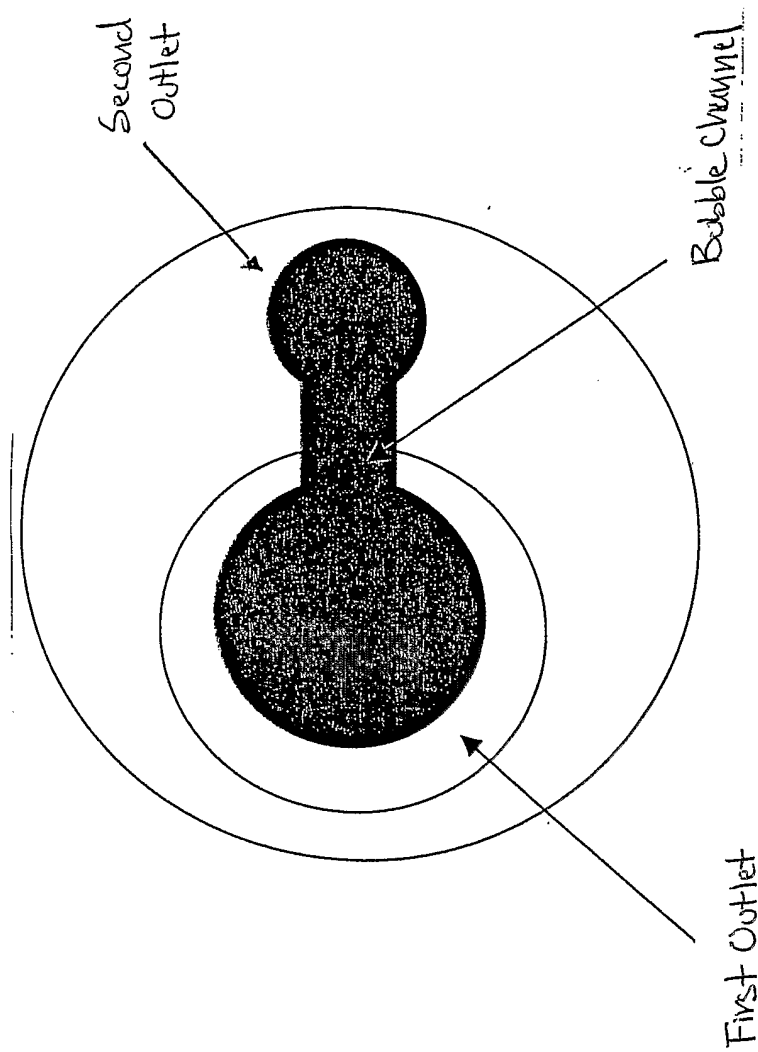


FIG. 9

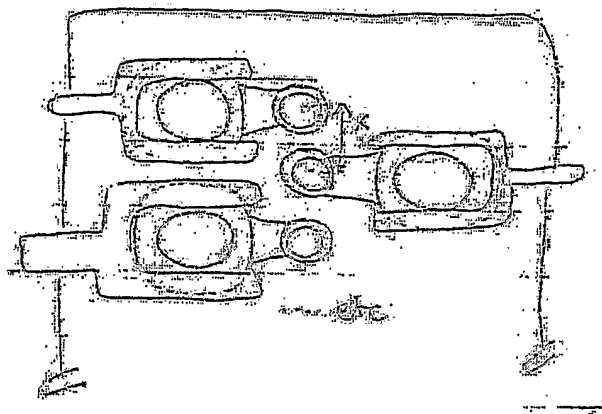


FIG. 10

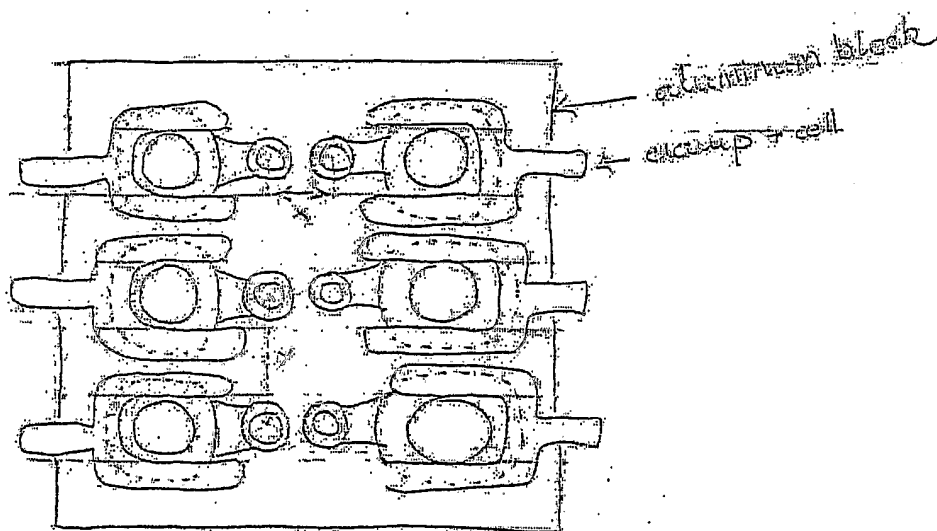


FIG. 11

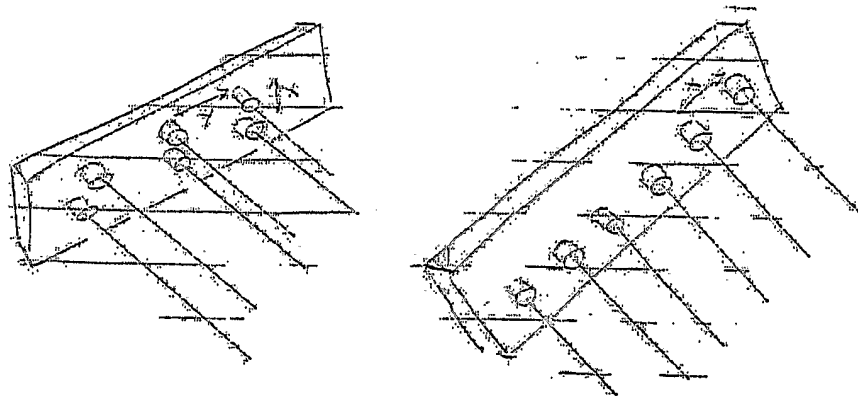
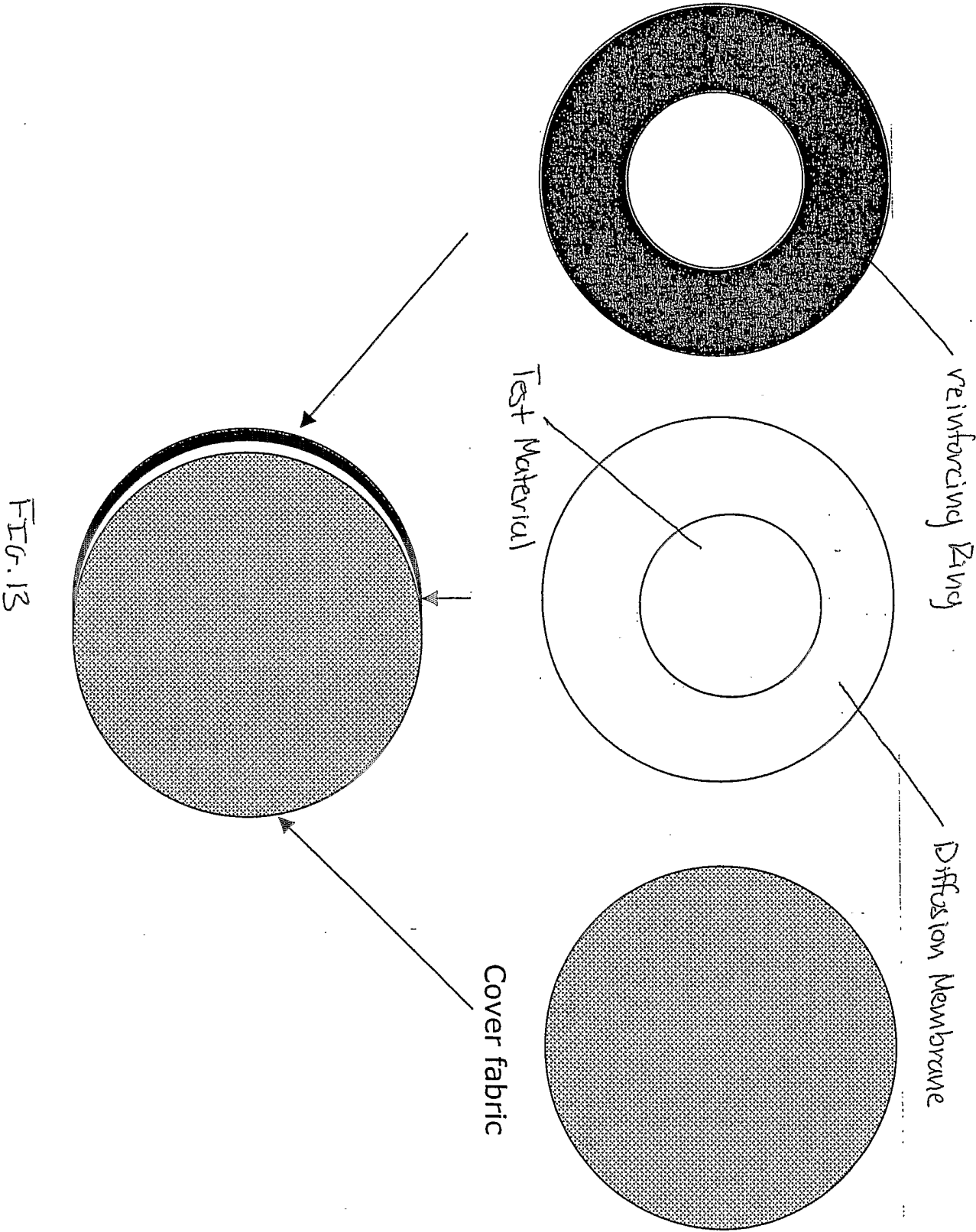


FIG. 12



INTERNATIONAL SEARCH REPORT

International Application No
T/US2004/009193

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 G01N13/00

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 G01N B01L

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

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

EPO-Internal, WPI Data, BIOSIS, COMPENDEX, EMBASE, INSPEC, IBM-TDB

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 6 521 191 B1 (SCHENK DIRK ET AL) 18 February 2003 (2003-02-18)	1-3,8-10
Y	column 1, line 46 - line 52 column 2, line 35 - line 51 column 2, line 58 - column 3, line 4 column 3, line 10 - line 26 column 4, line 3 - line 11 figures 2,6	4
Y	US 4 667 504 A (HOBSON DAVID W) 26 May 1987 (1987-05-26) column 3, line 7 - line 35 column 3, line 45 - line 50 figure 3	4

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

20 July 2004

Date of mailing of the international search report

03/08/2004

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Timonen, T

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US2004/009193

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4 740 309 A (HIGUCHI TAKERU) 26 April 1988 (1988-04-26) column 3, line 29 - line 60 figure 2 ---	1-10
A	US 5 198 109 A (HANSON WILLIAM A ET AL) 30 March 1993 (1993-03-30) column 3, line 57 -column 4, line 45 figure 2 ---	1-10
A	US 4 594 884 A (GRABOWSKI LUCILLE A ET AL) 17 June 1986 (1986-06-17) the whole document ---	1-10
A	I.J. BOSMAN ET AL.: "Novel diffusion cell for in vitro transdermal permeation, compatible with automated dynamic sampling" JOURNAL OF PHARMACEUTICAL AND BIOMEDICAL ANALYSIS, vol. 14, no. 8-10, June 1996 (1996-06), pages 1015-1023, XP002288878 the whole document ---	1-10
A	S.C. CHATTARAJ ET AL.: "A simple diffusion cell to monitor drug release from semi-solid dosage forms" INTERNATIONAL JOURNAL OF PHARMACEUTICS, vol. 120, no. 1, June 1995 (1995-06), pages 119-124, XP002288879 the whole document -----	1-10

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US2004/009193

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 6521191	B1	18-02-2003	DE 29911790 U1 EP 1067382 A2 US 6294134 B1	30-09-1999 10-01-2001 25-09-2001
US 4667504	A	26-05-1987	NONE	
US 4740309	A	26-04-1988	NONE	
US 5198109	A	30-03-1993	NONE	
US 4594884	A	17-06-1986	NONE	