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(54) **POLYMER MICROFILTRATION DEVICES, METHODS OF MANUFACTURING THE SAME AND THE USES OF THE MICROFILTRATION DEVICES**

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(73) Assignee: **Creatv MicroTech, Inc.**, Potomac, MD (US)

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Related U.S. Application Data

(63) Continuation-in-part of application No. PCT/US11/30966, filed on Apr. 1, 2011, Continuation-in-part of application No. PCT/US12/66390, filed on Nov. 21, 2012.

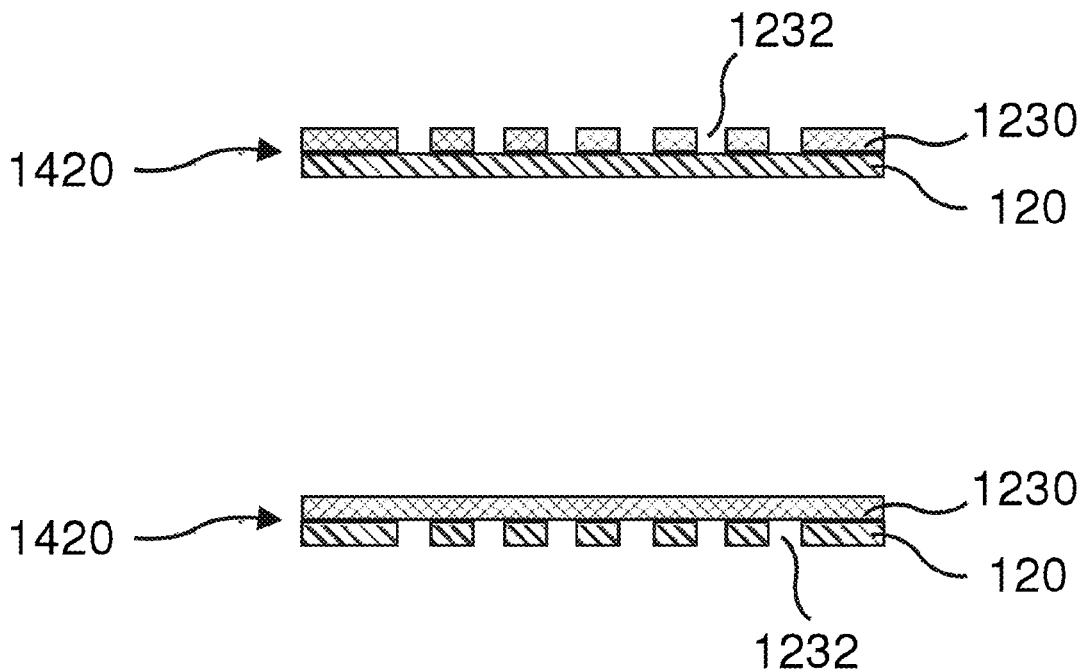
(60) Provisional application No. 61/562,404, filed on Nov. 21, 2011, provisional application No. 61/618,641, filed on Mar. 30, 2012, provisional application No. 61/654,636, filed on Jun. 1, 2012.

Publication Classification

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G01N 1/40 (2006.01)
B01D 29/00 (2006.01)
(52) **U.S. Cl.**
CPC *G01N 1/4005* (2013.01); *B01D 29/00* (2013.01)
USPC *435/6.11*; 210/490; 210/767; 29/428; 435/287.1; 435/7.1; 435/34; 435/39; 435/325

(57) **ABSTRACT**

A microfilter comprising a polymer layer formed from photo-definable dry film, and a plurality of apertures each extending through the polymer layer. A microfilter comprising two or more polymer layers formed from photo-definable dry film, and a plurality of apertures or open areas each extending through the polymer layer. Methods of forming apertures in one or more layers of photo-definable dry film are also disclosed. Filter holder designs and methods appropriate to hold microfilters to collect the rare cells and to perform of assays in the filter holder are provided. Microfiltration chip designs and methods appropriate to collect the rare cells and to perform assays in the microfluidic chips are provided. The invention also describes the use of the microfilter, filter holder and microfilter chips to collect rare cells from body fluids and perform assays, and these rare cells can be used for medical and biological research applications.



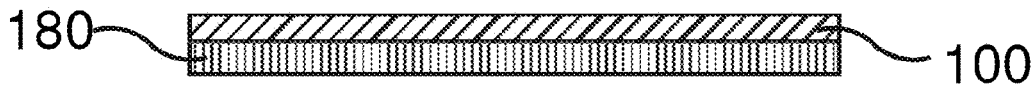


FIG. 1A

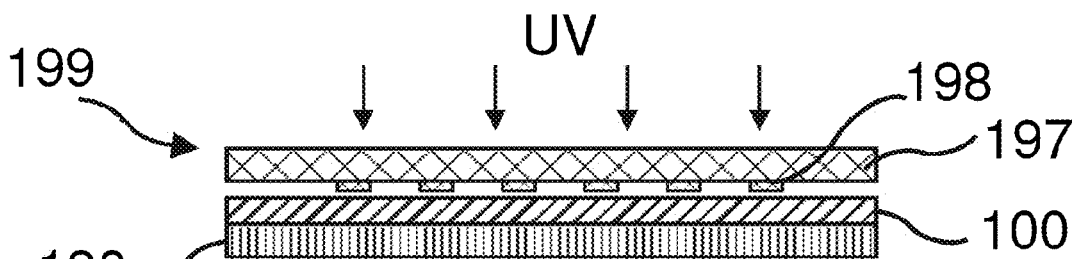


FIG. 1B

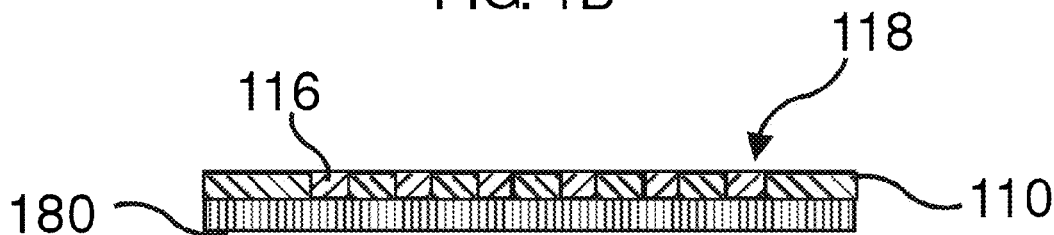


FIG. 1C

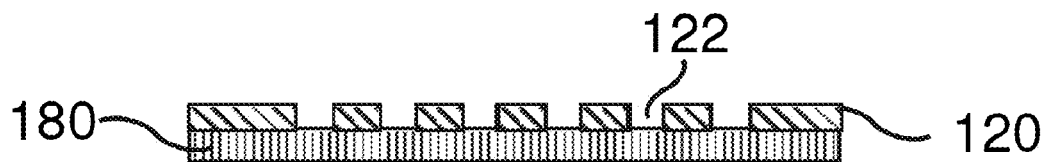


FIG. 1D

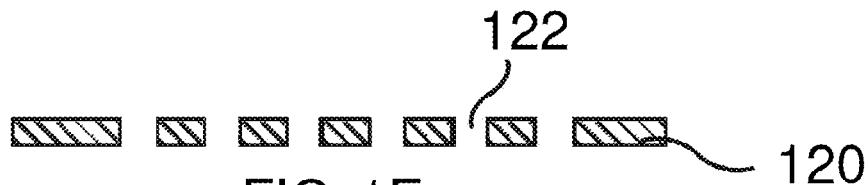


FIG. 1E

FIG. 2A

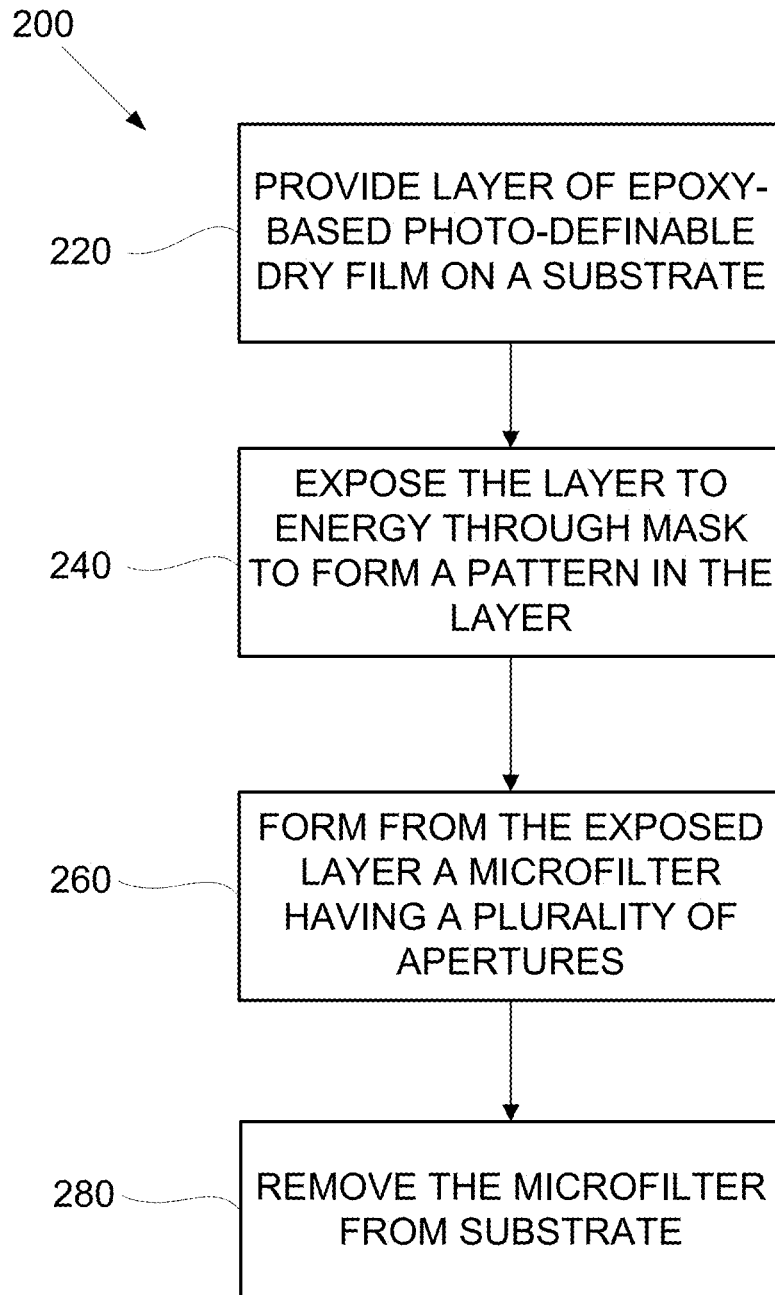


FIG. 2B

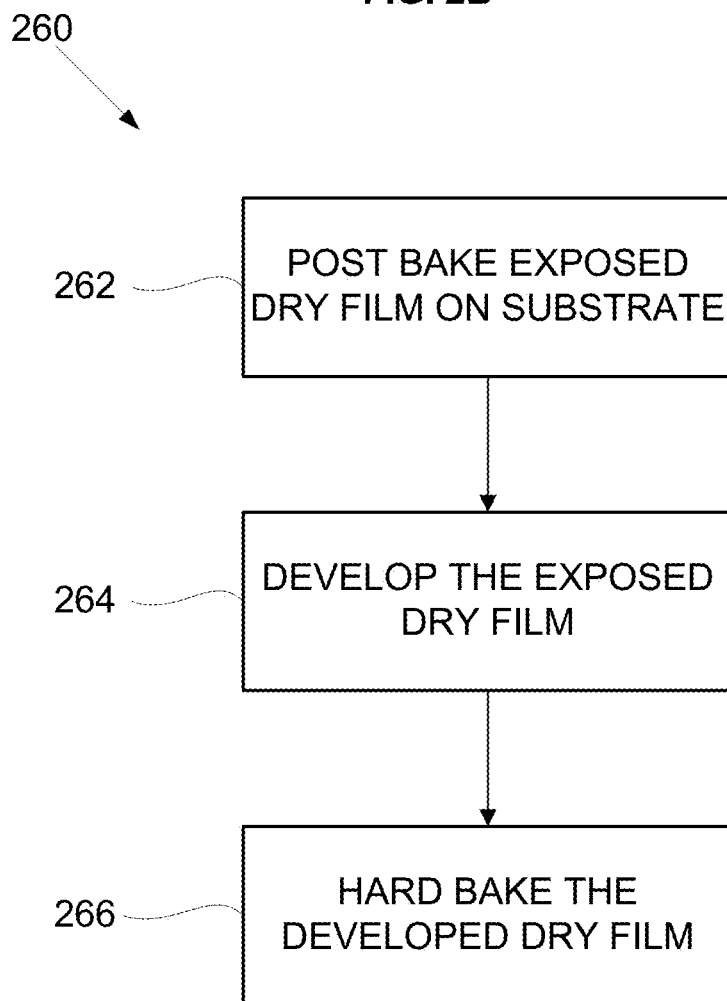




FIG. 3A

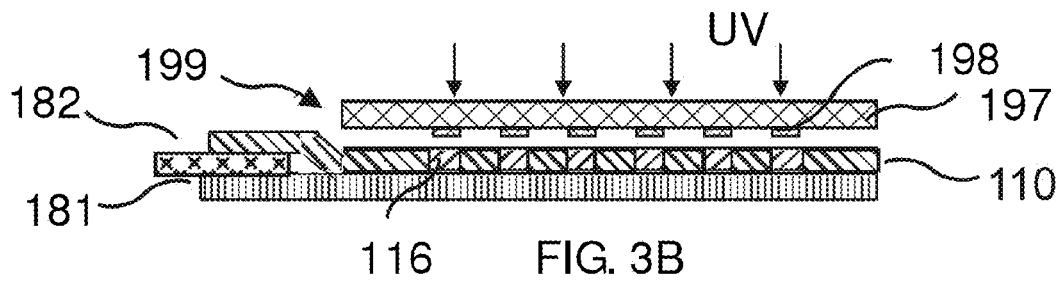


FIG. 3B



FIG. 3C



FIG. 3D

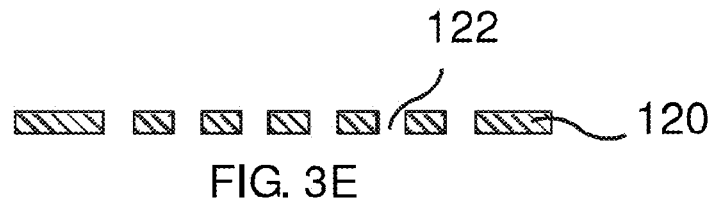
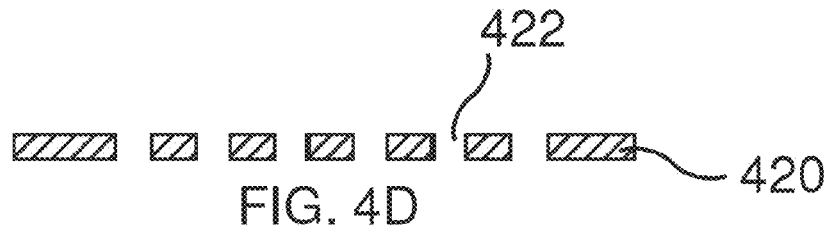
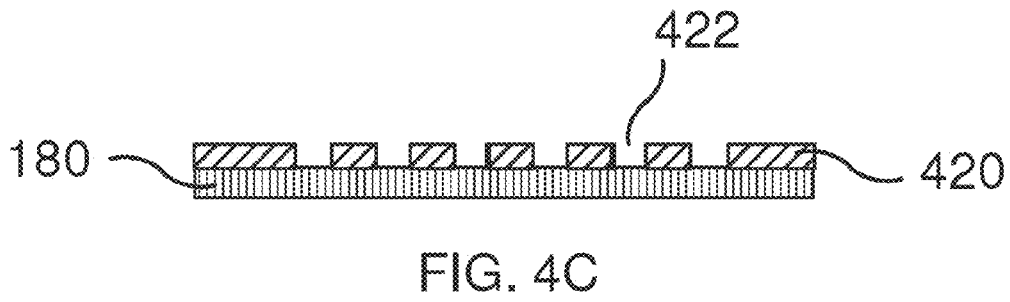
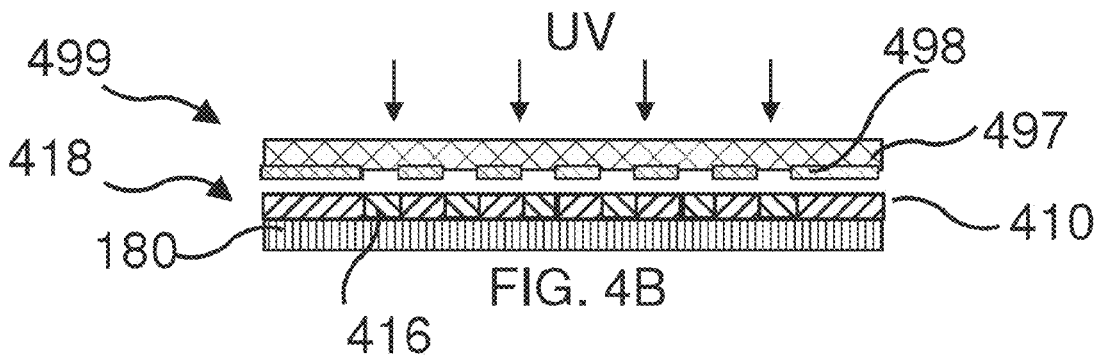


FIG. 3E



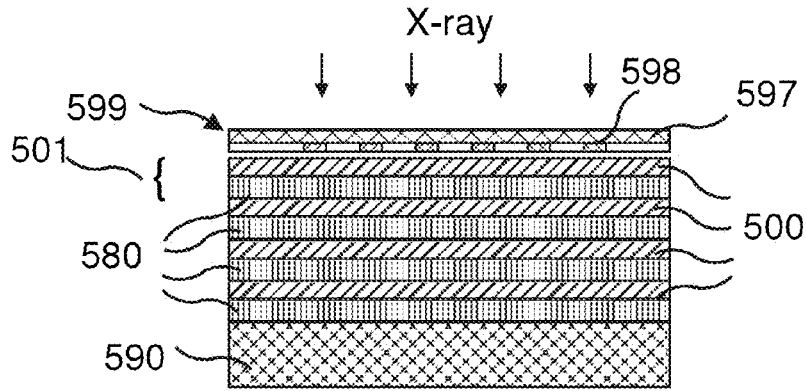


FIG. 5A

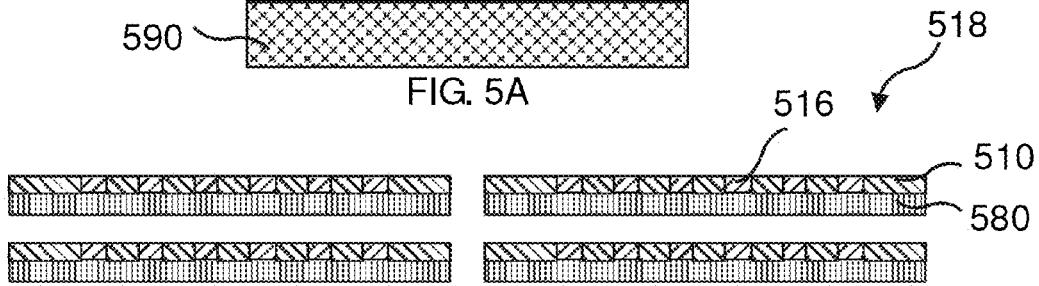


FIG. 5B

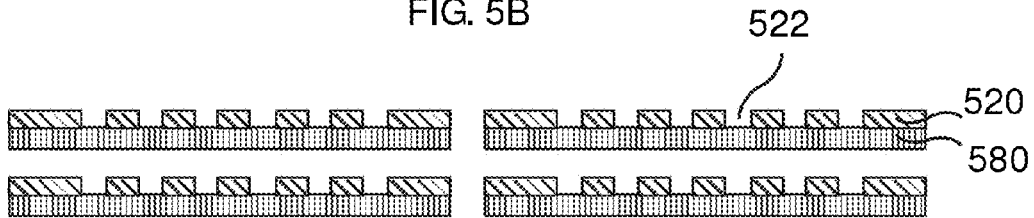


FIG. 5C

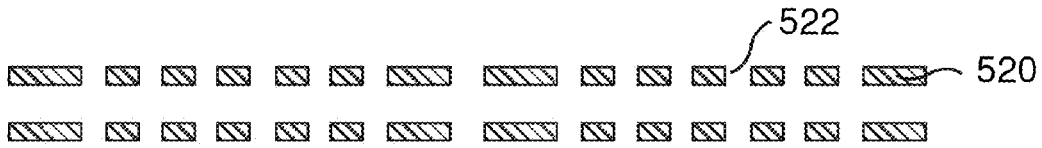


FIG. 5D

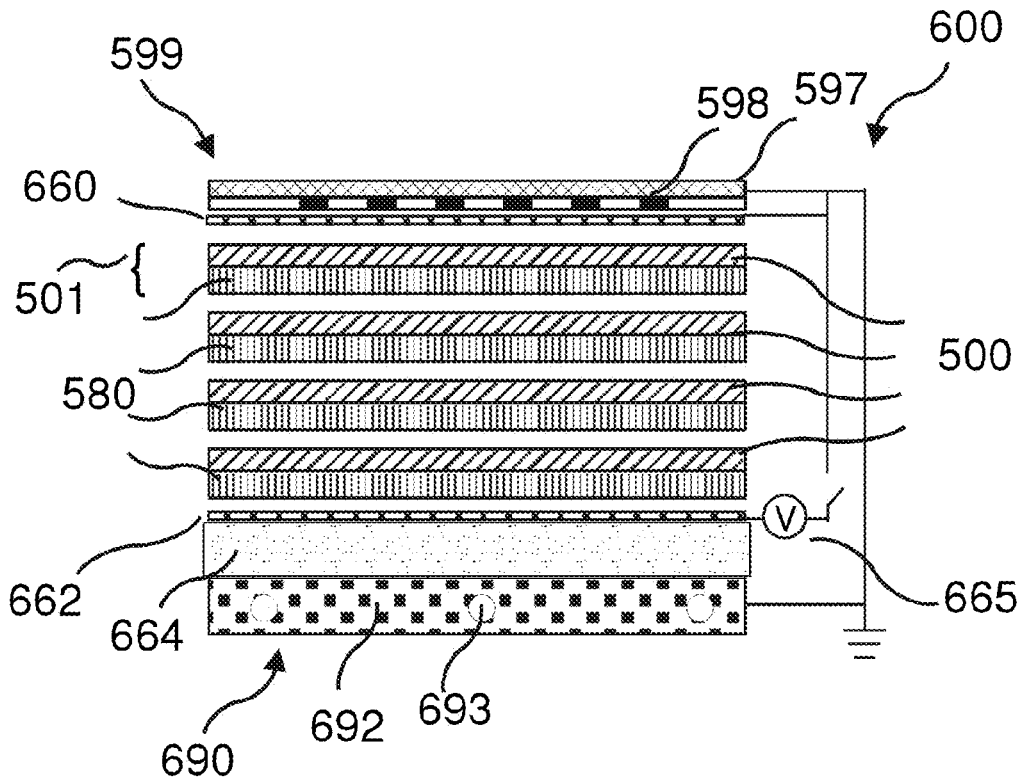


FIG. 6A

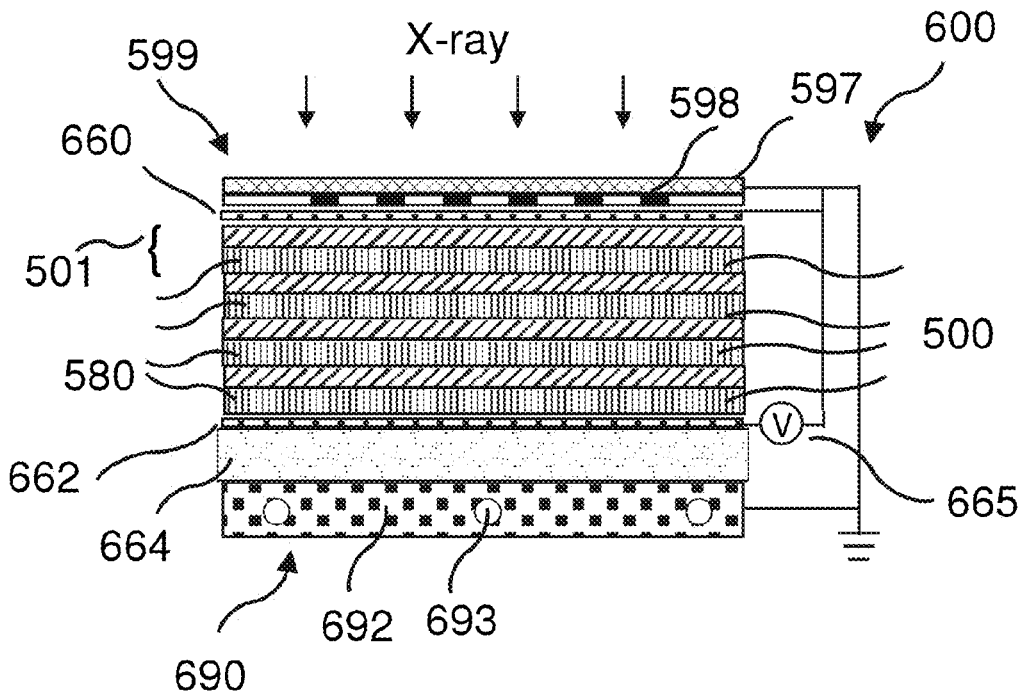


FIG. 6B

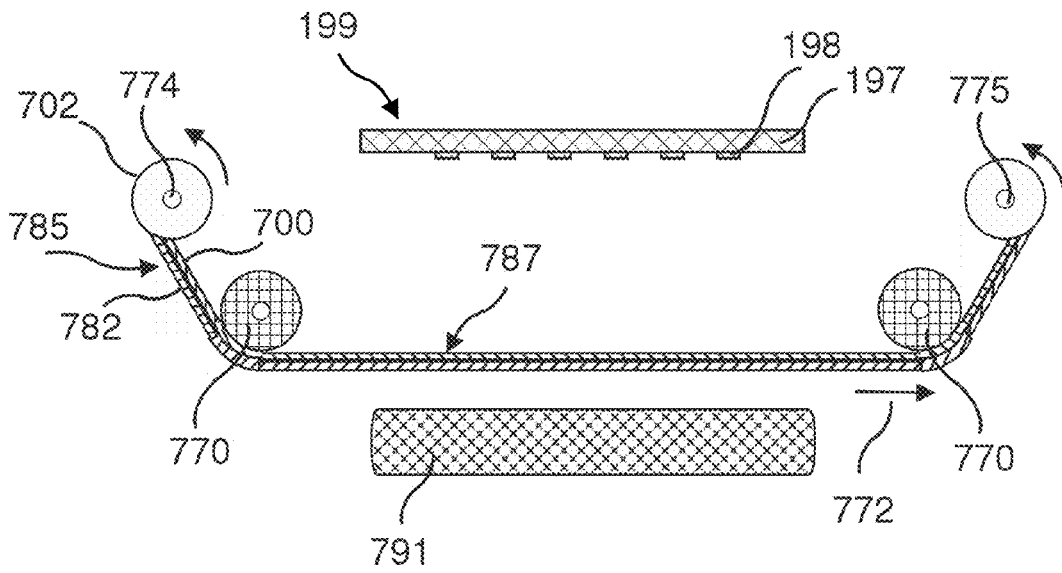


FIG. 7A

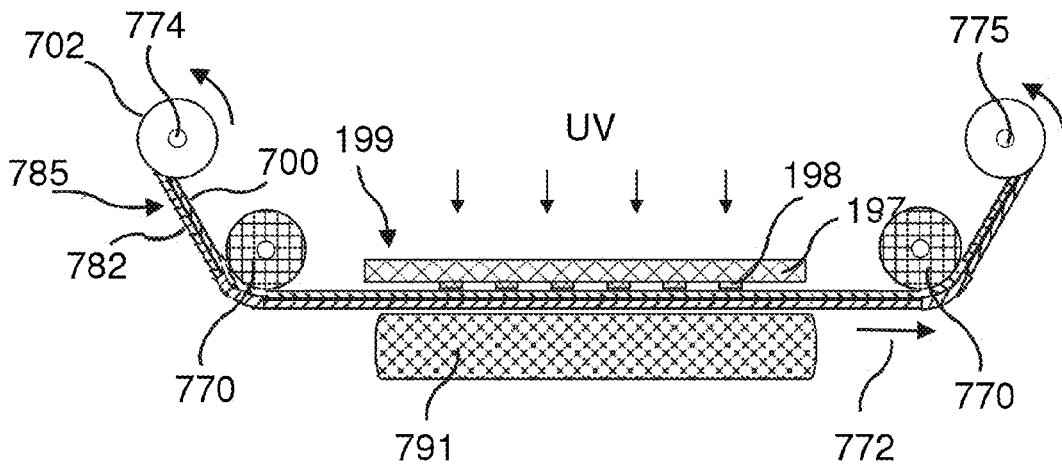


FIG. 7B

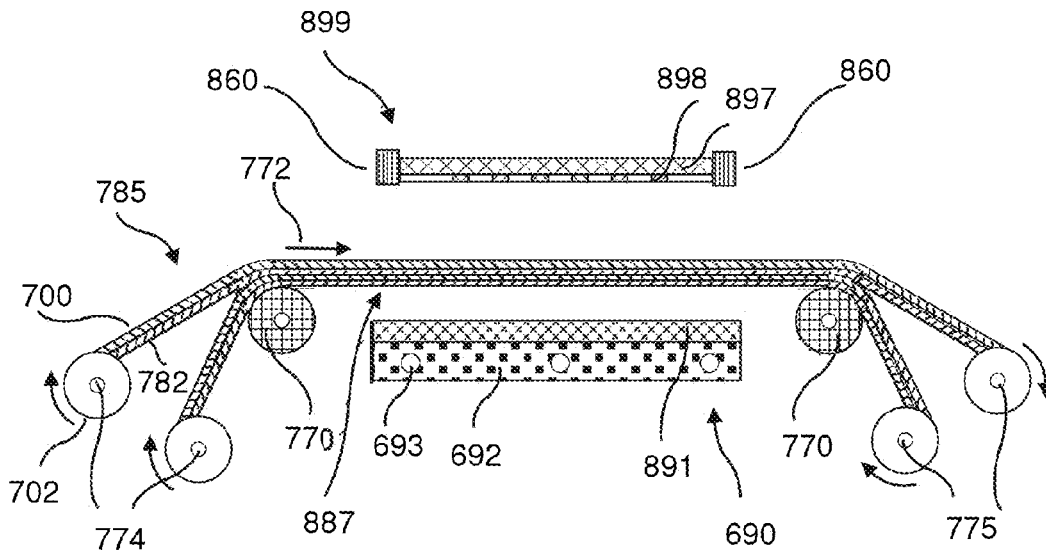


FIG. 8A

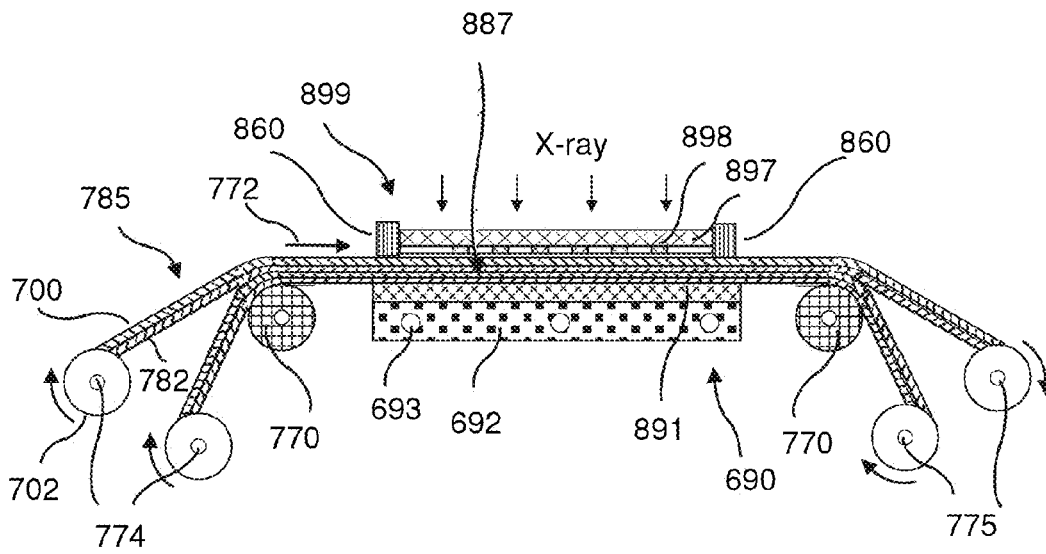


FIG. 8B

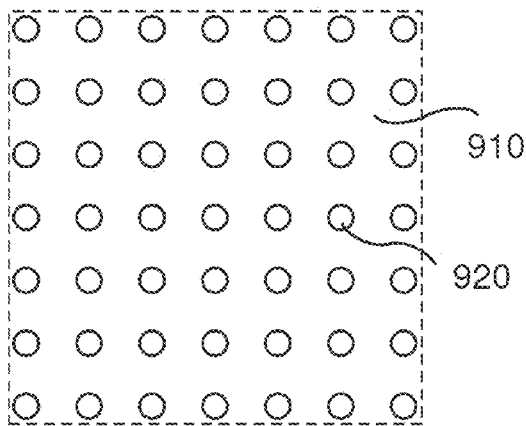


FIG. 9A

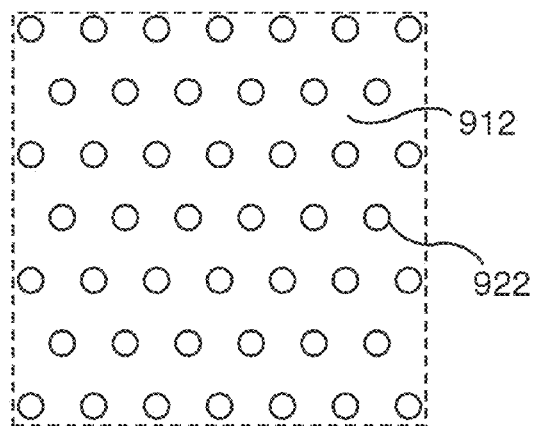


FIG. 9B

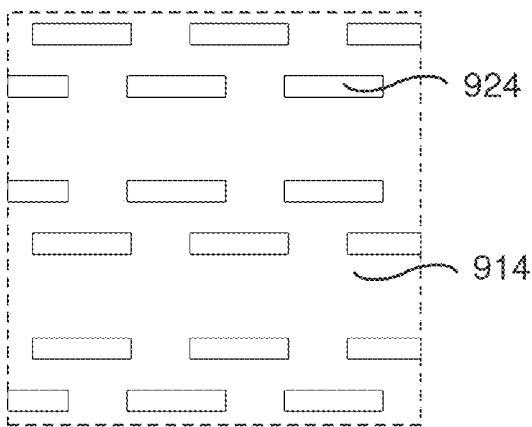


FIG. 9C

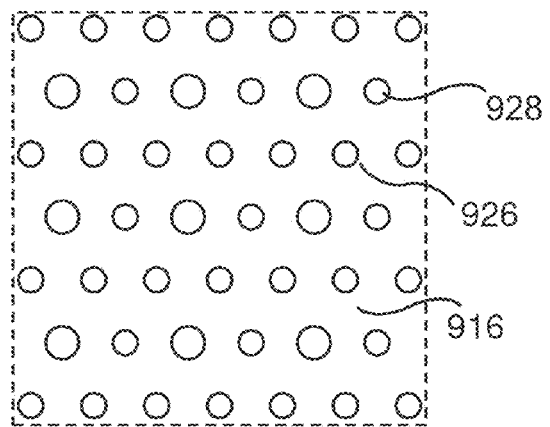


FIG. 9D

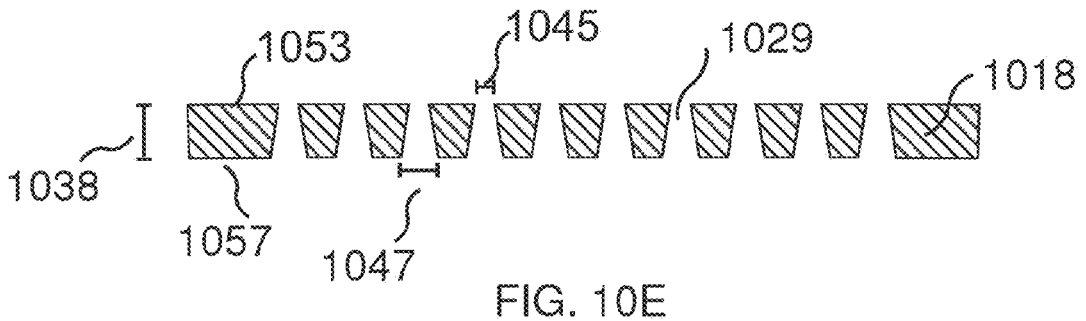
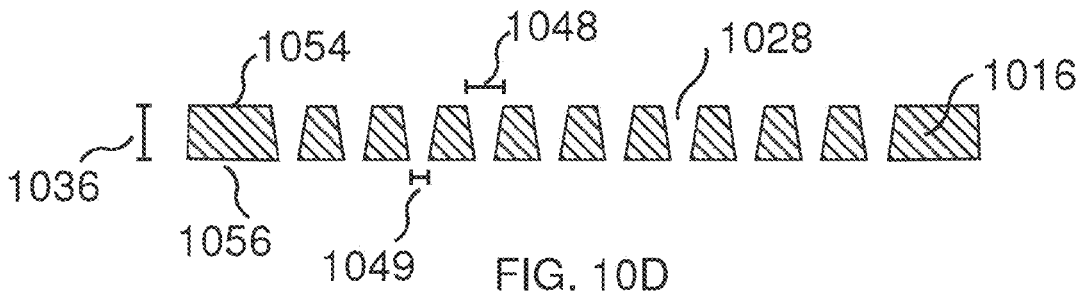
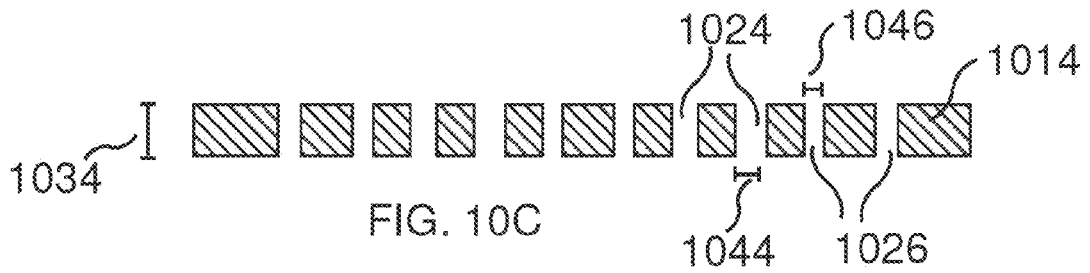
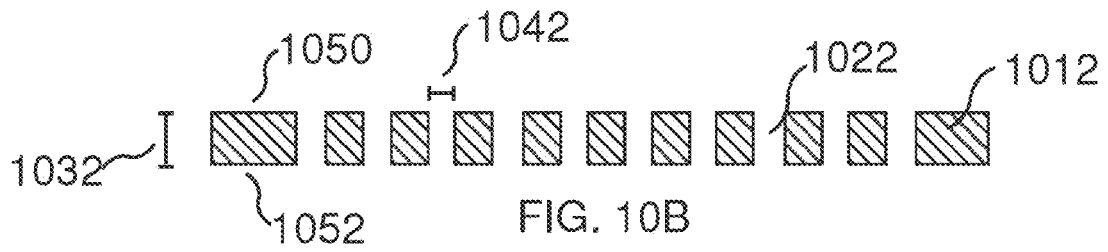
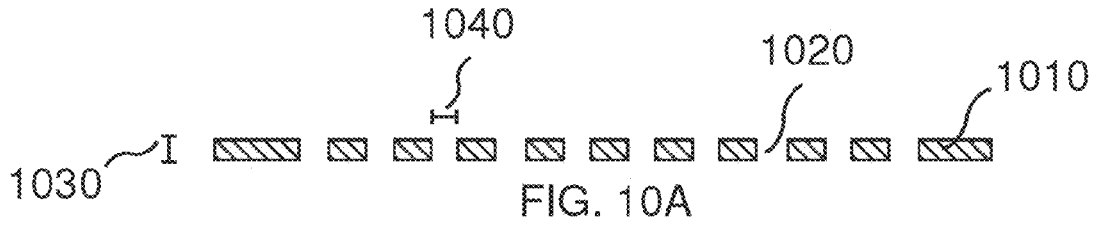
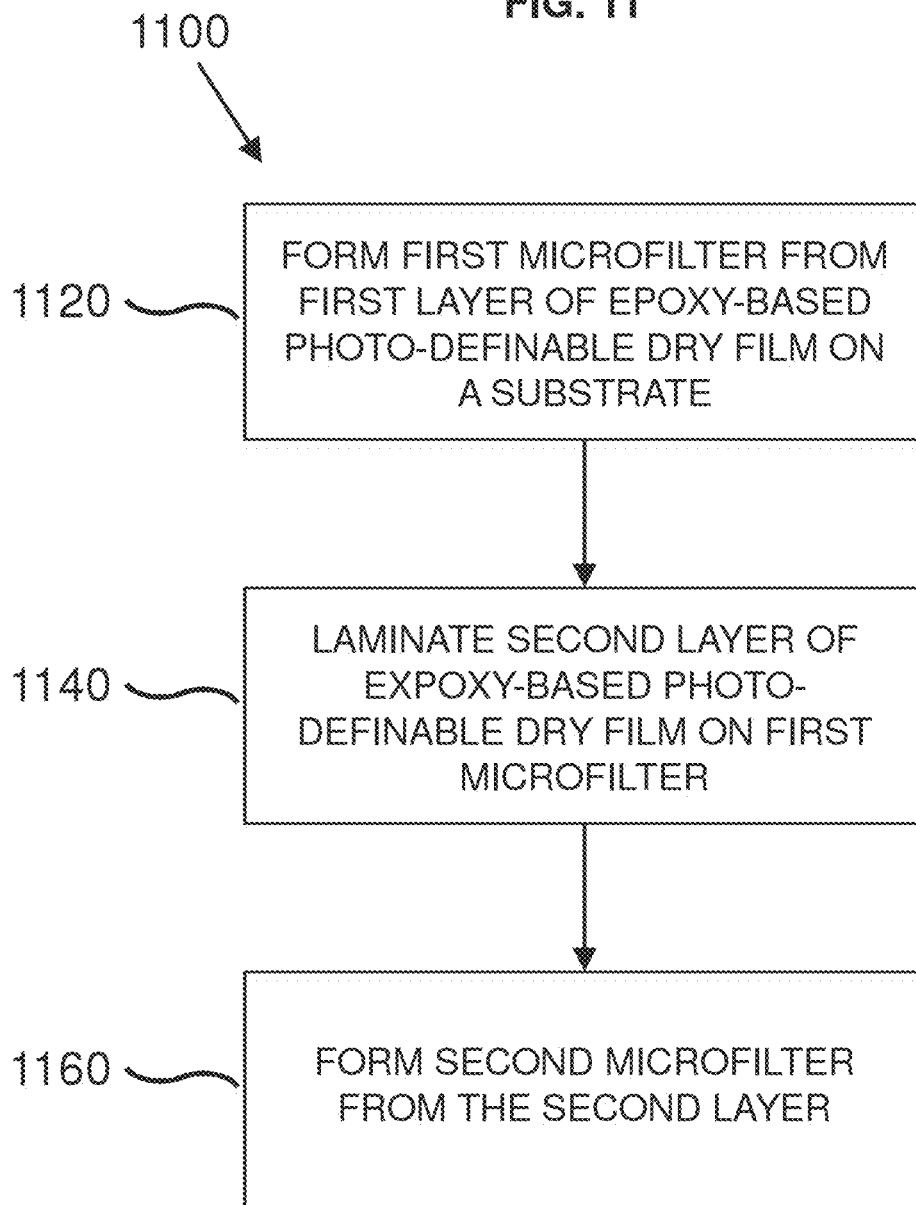


FIG. 11



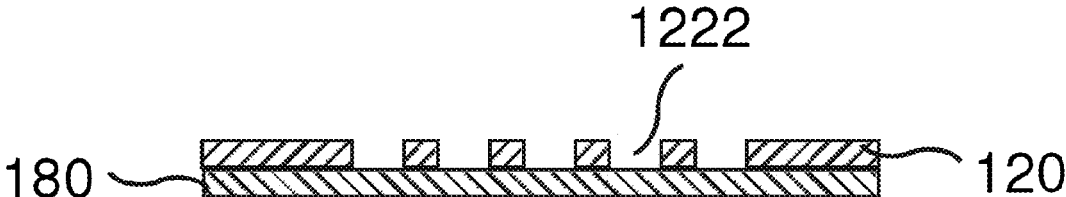


FIG. 12A

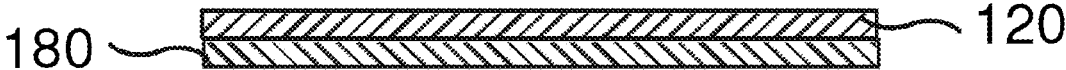
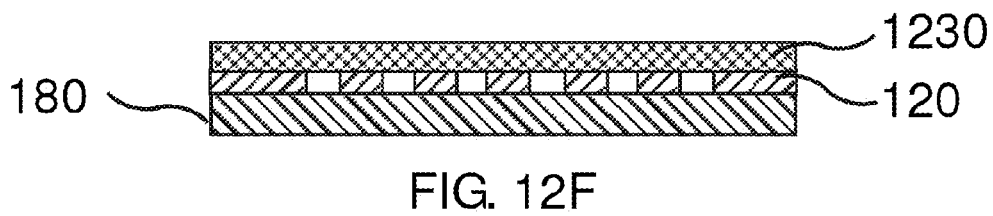
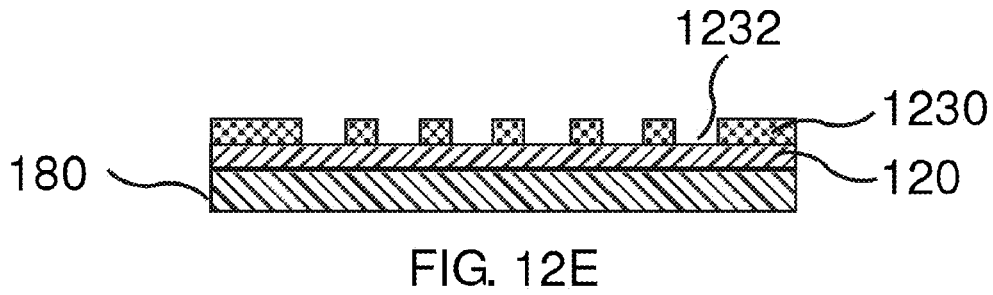
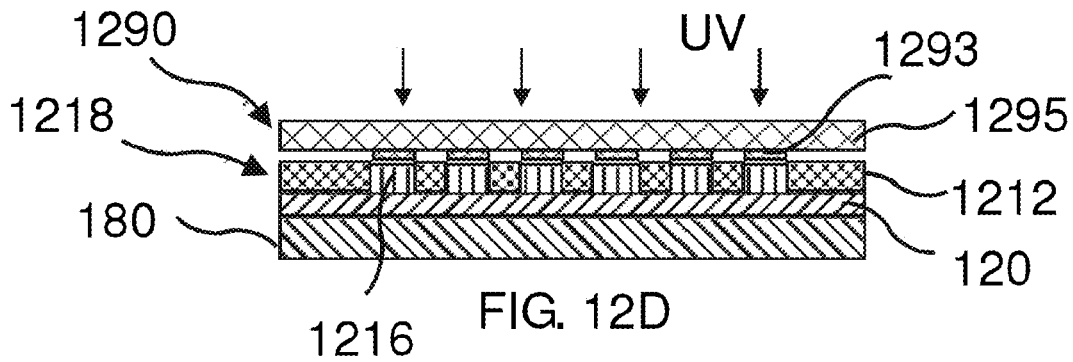
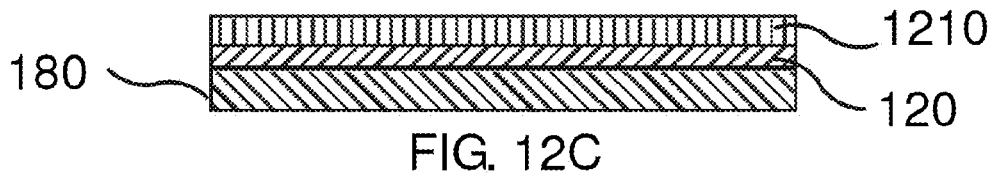


FIG. 12B



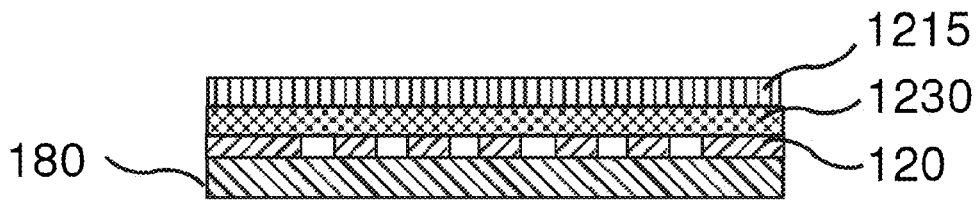


FIG. 12G

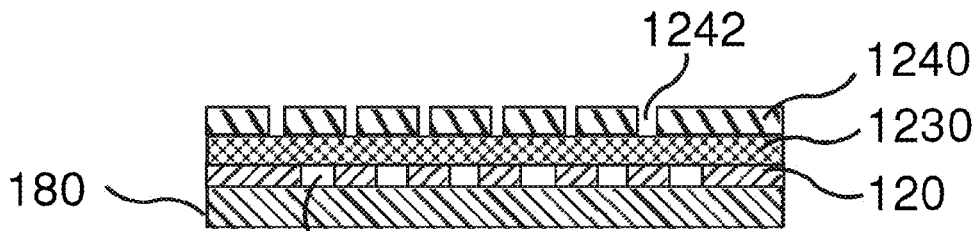


FIG. 12H

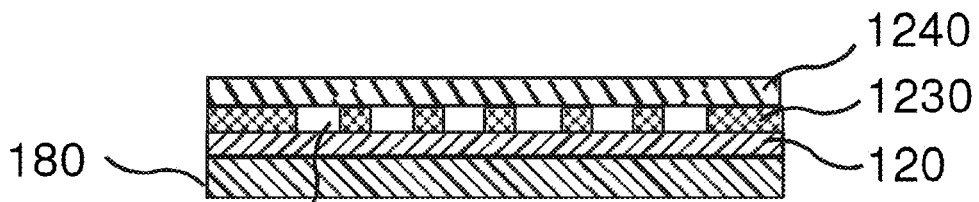


FIG. 12I

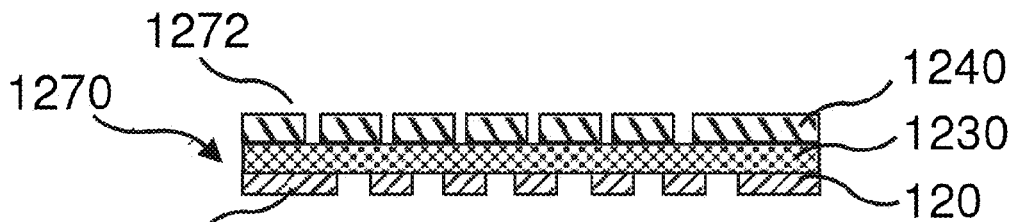


FIG. 12J

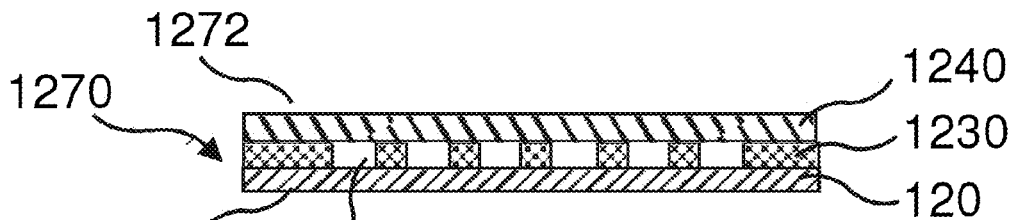


FIG. 12K

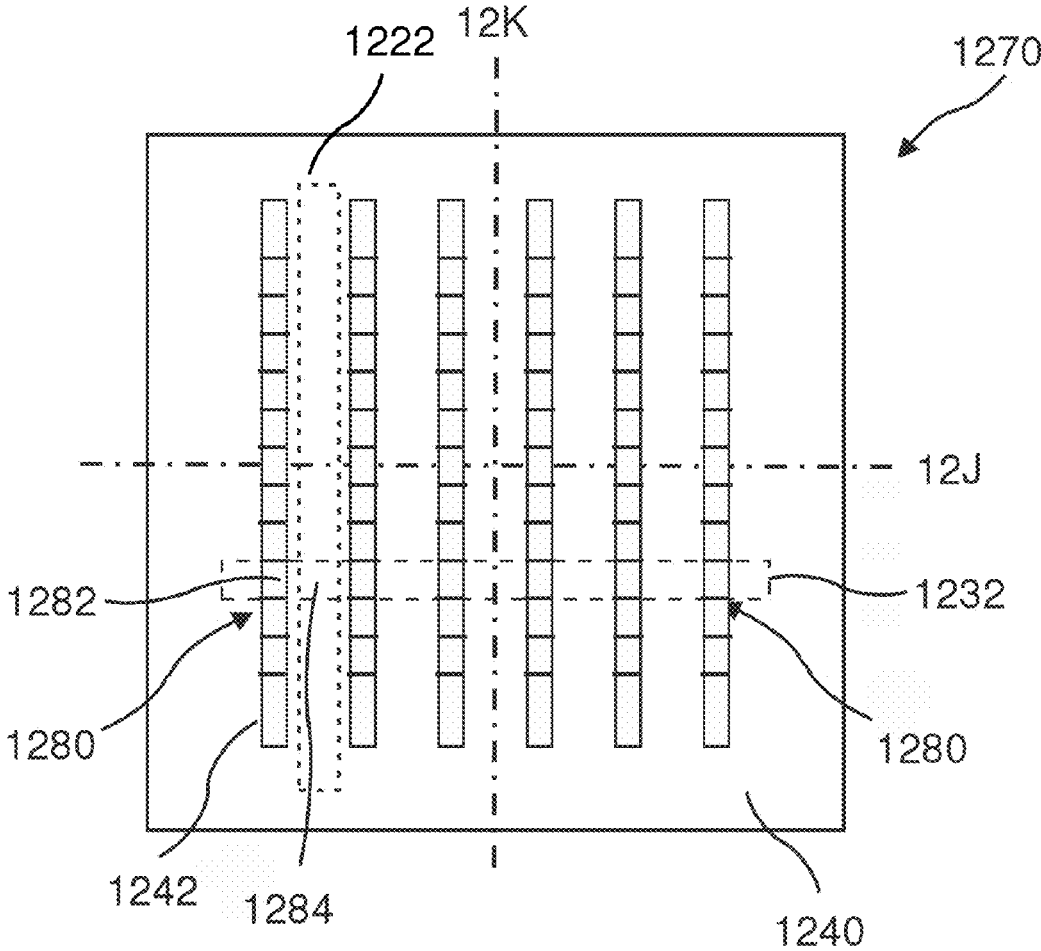


FIG. 12L

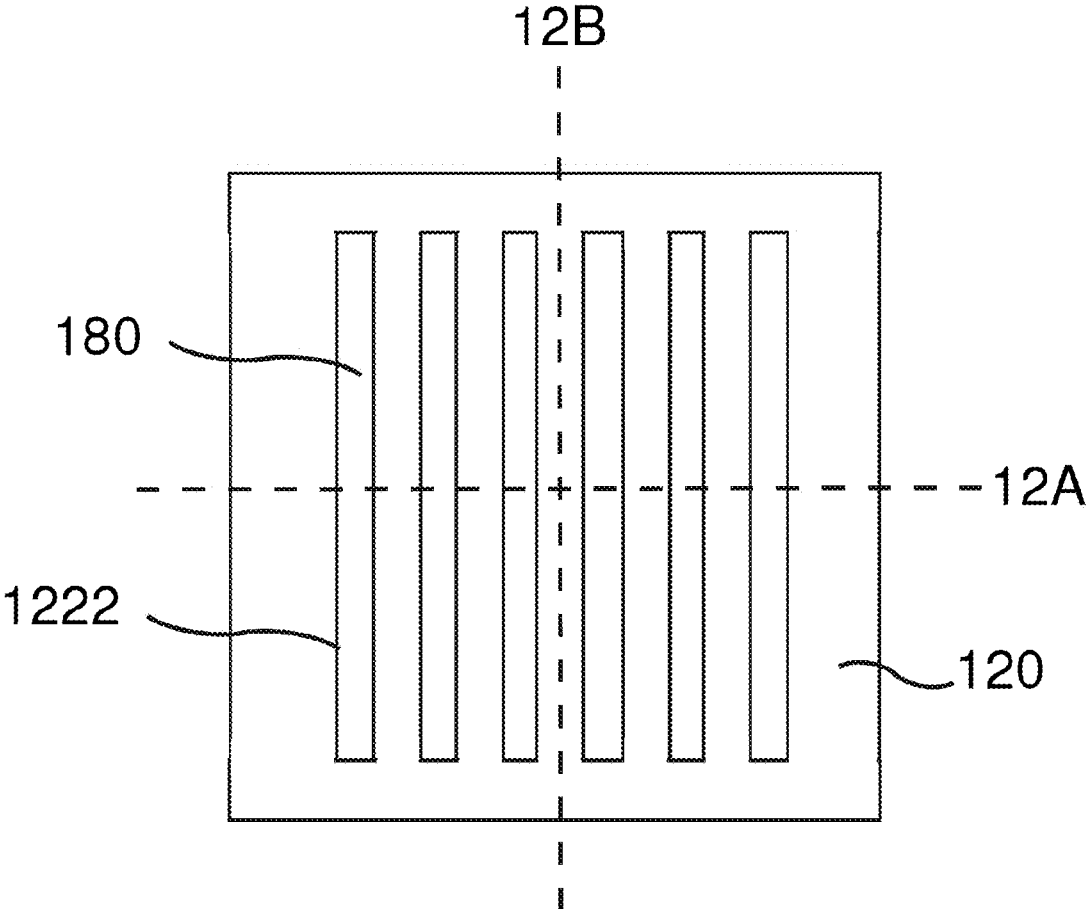


FIG. 13A

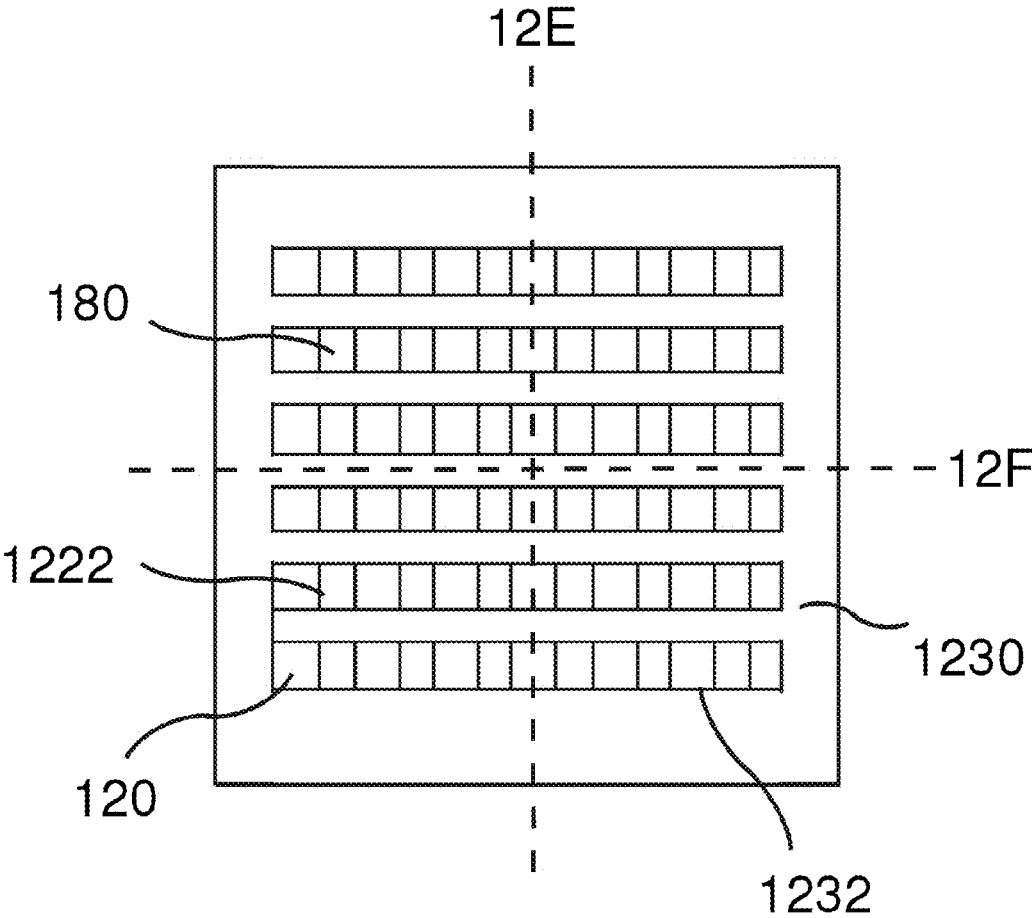


FIG. 13B

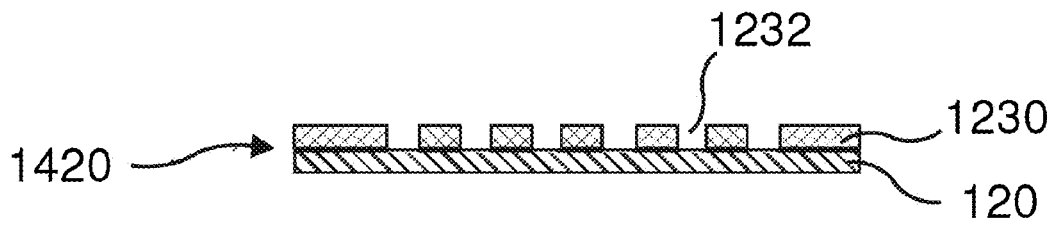


FIG. 14A

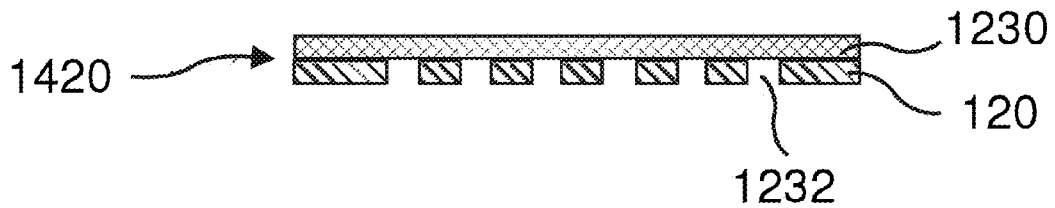


FIG. 14B

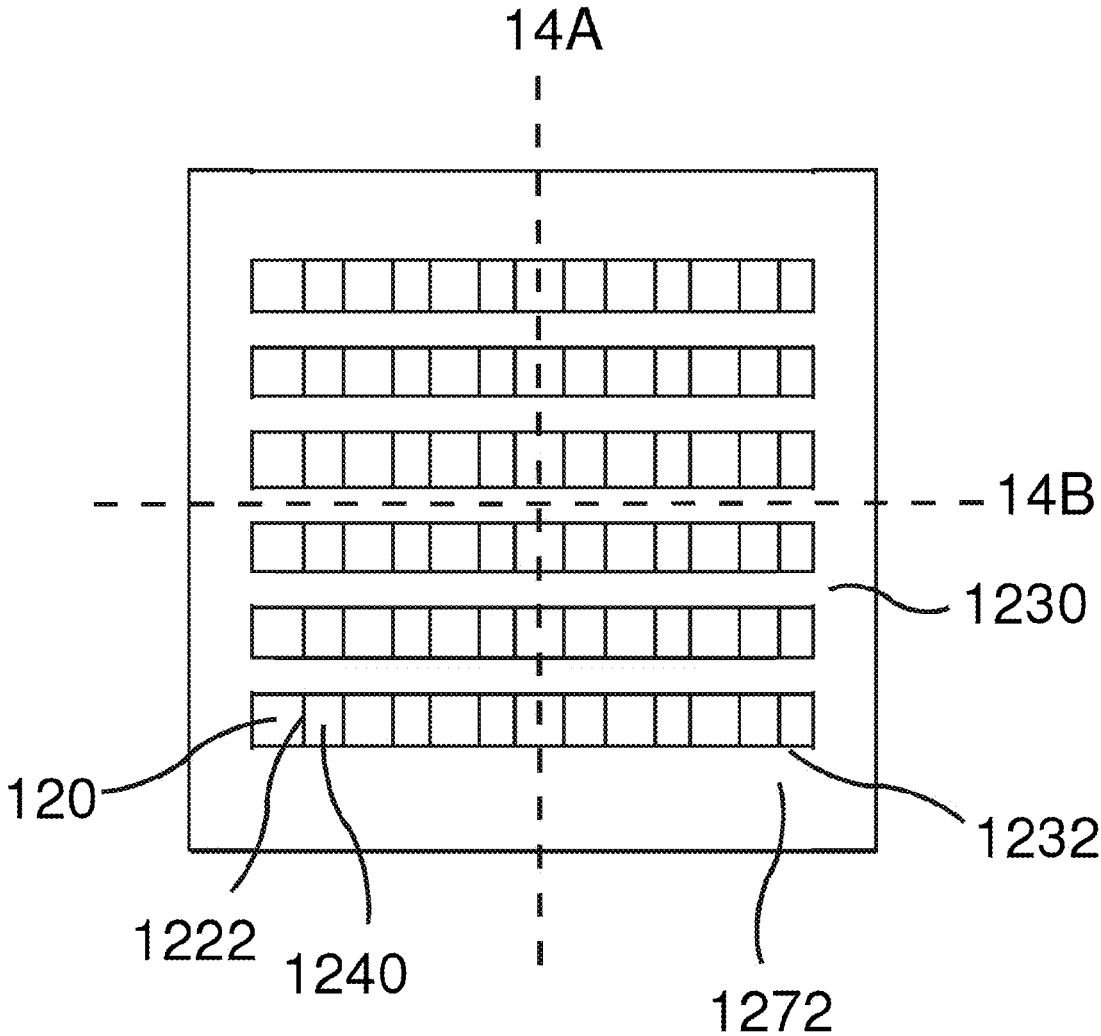


FIG. 14C

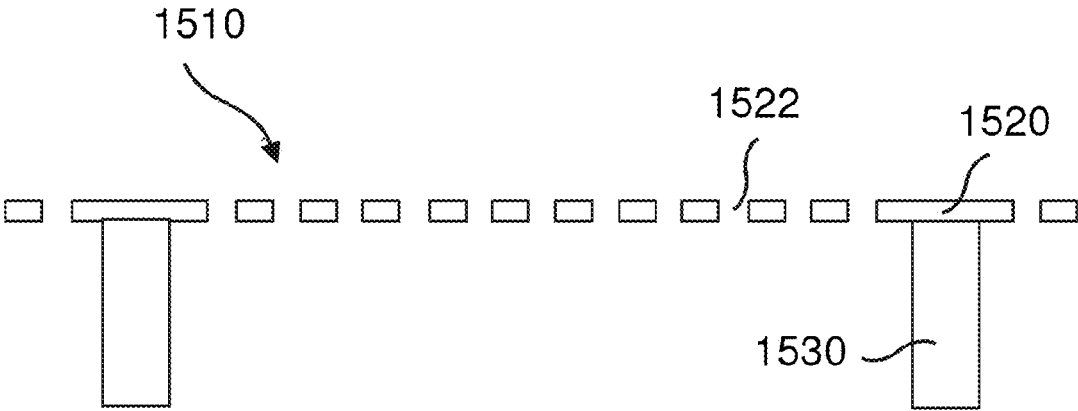


FIG. 15

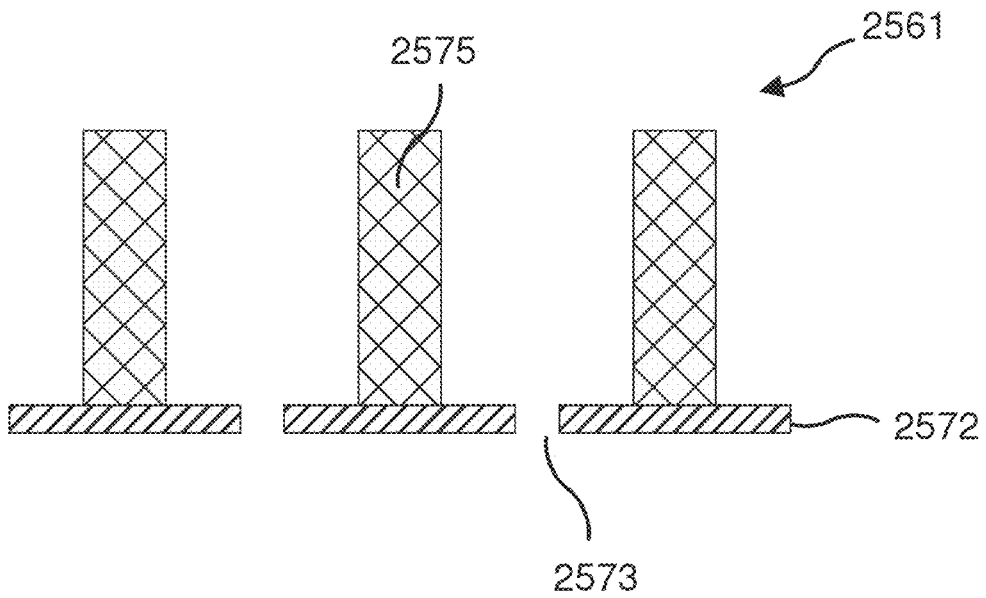


FIG. 16A

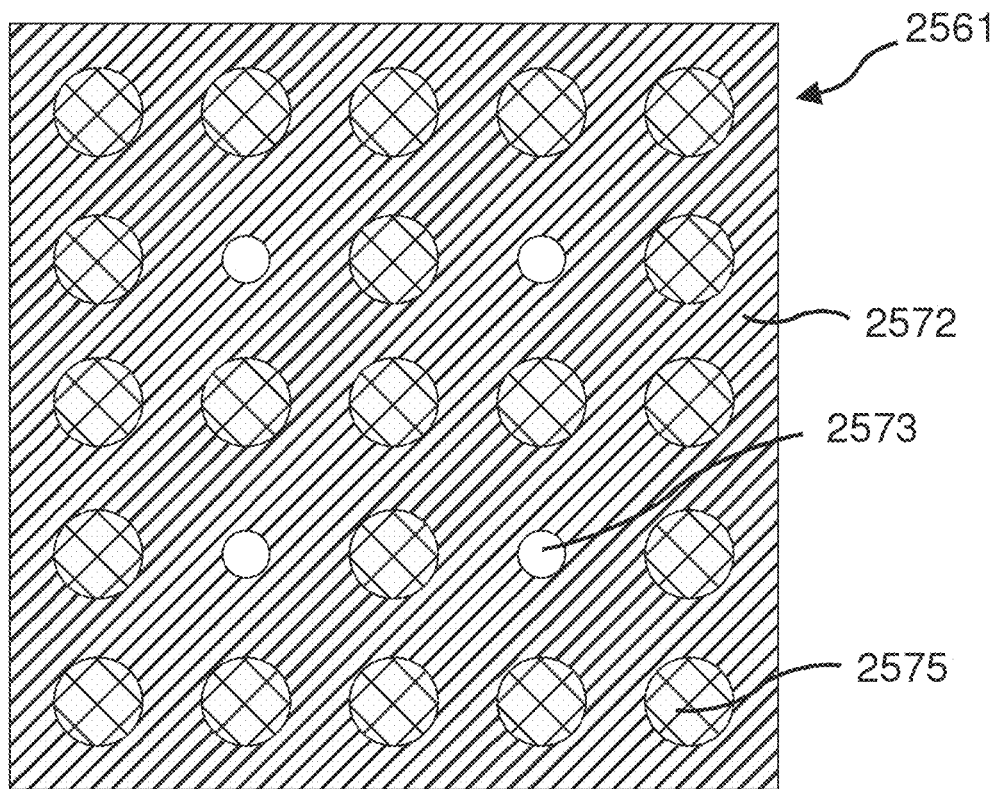


FIG. 16B

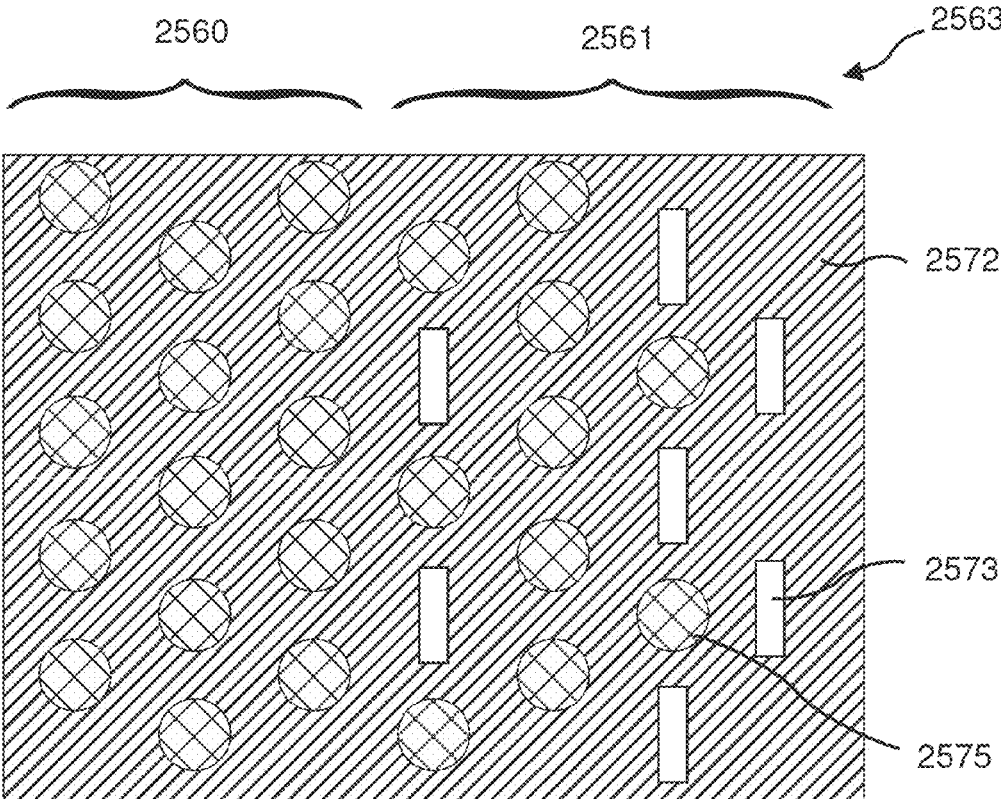


FIG. 16C

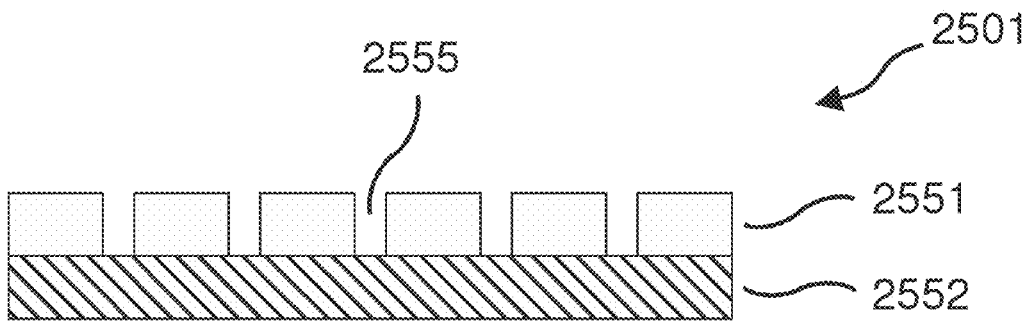


FIG. 16D

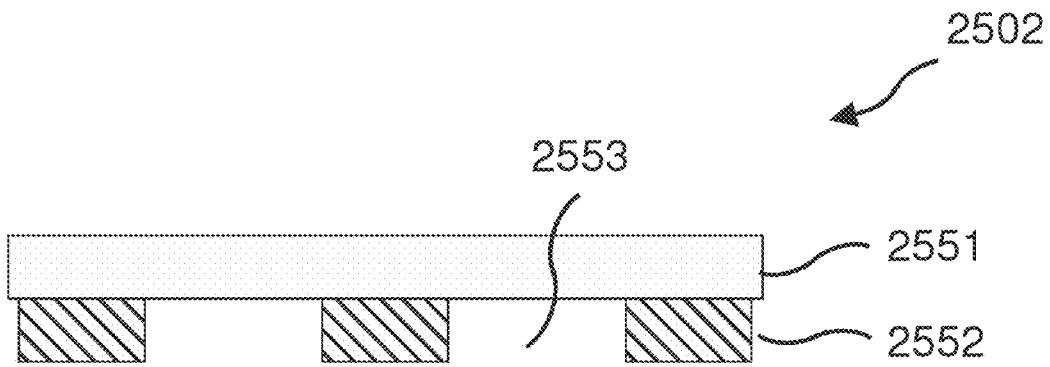


FIG. 16E

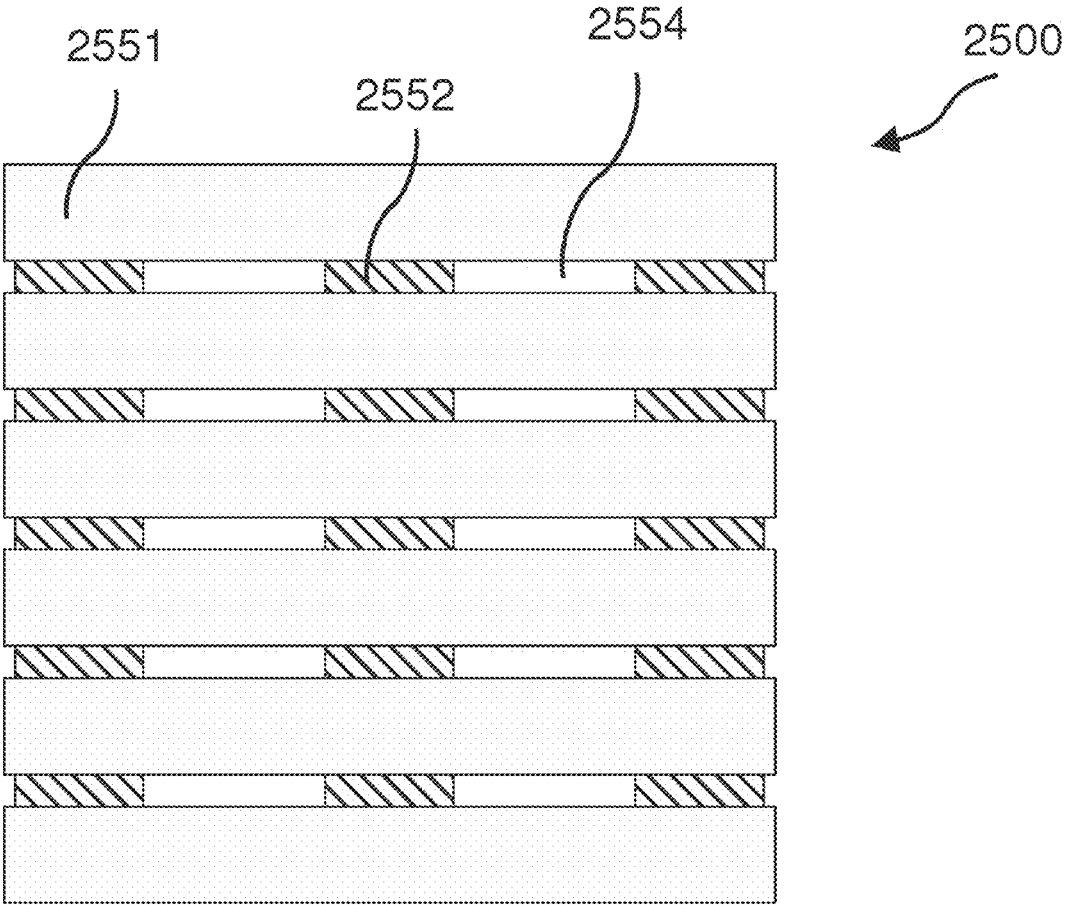
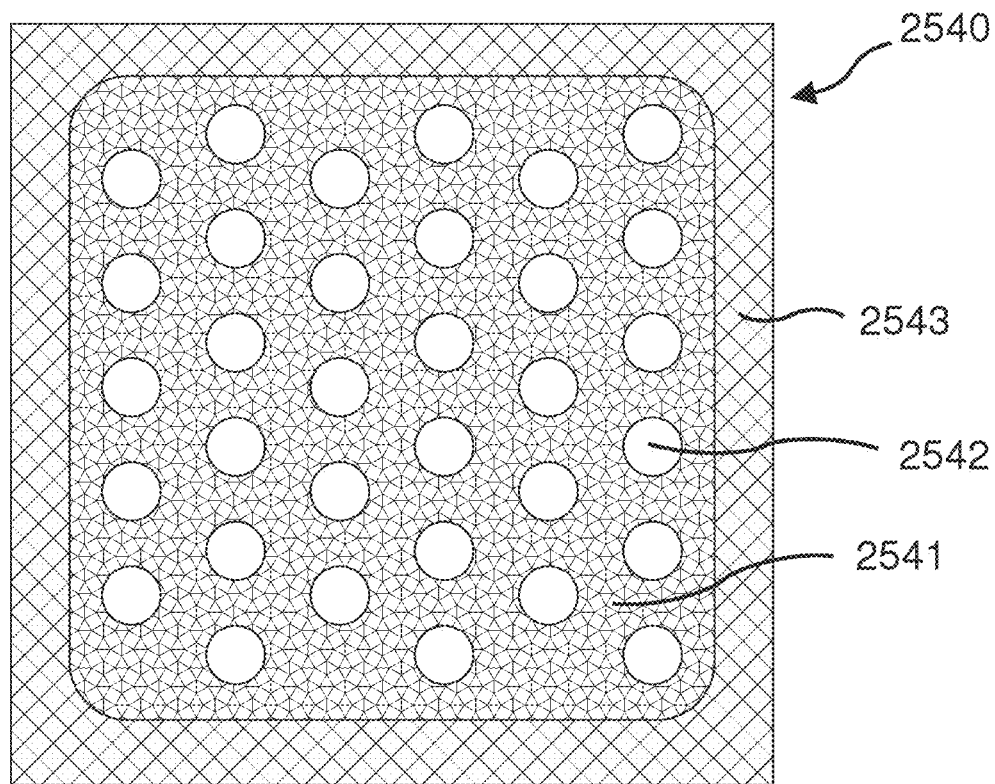
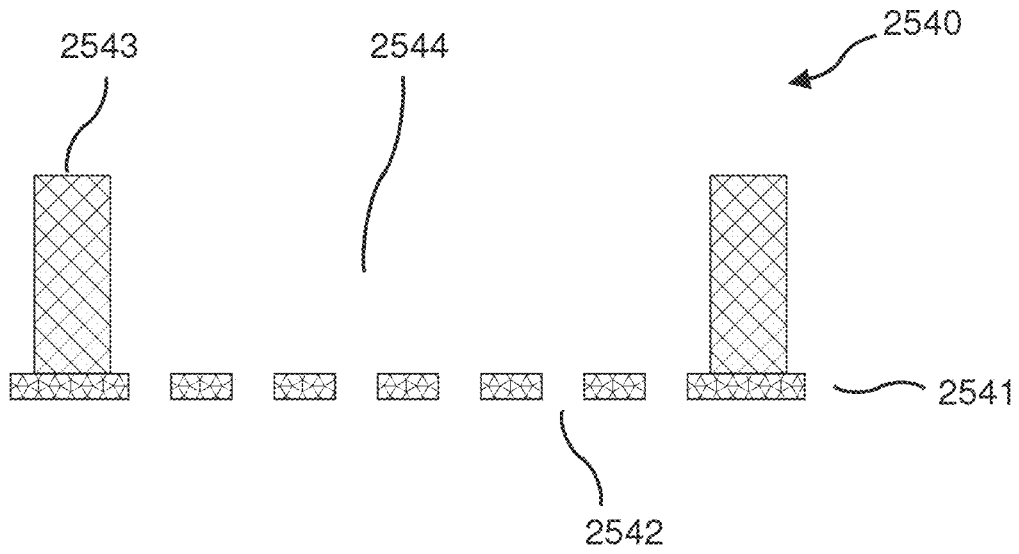


FIG. 16F



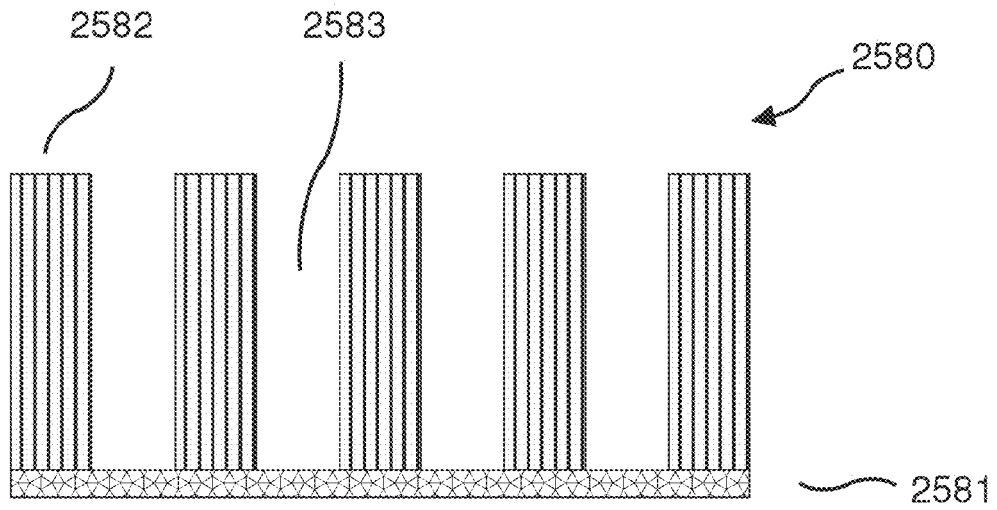


FIG. 16I

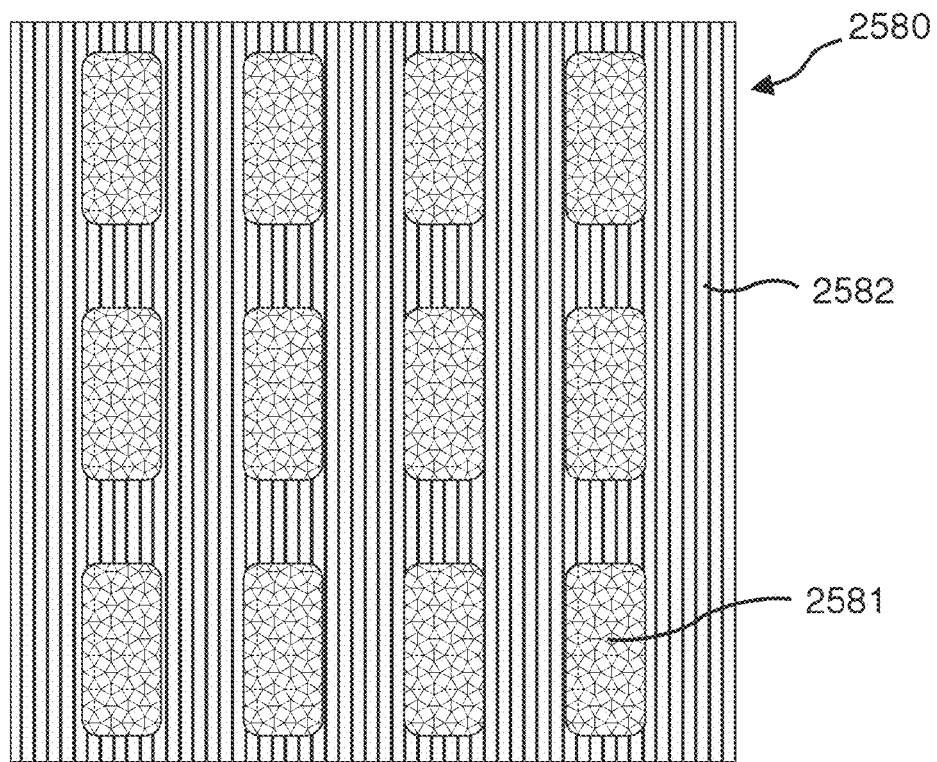


FIG. 16J

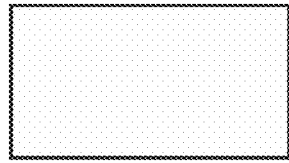


FIG. 17A

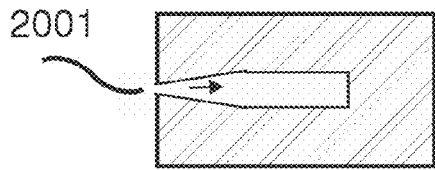


FIG. 17B

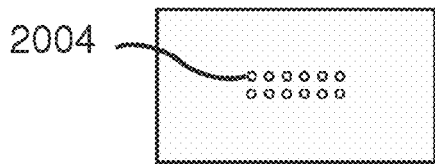


FIG. 17C

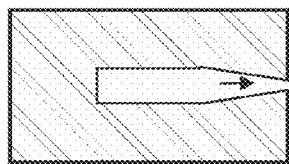


FIG. 17D

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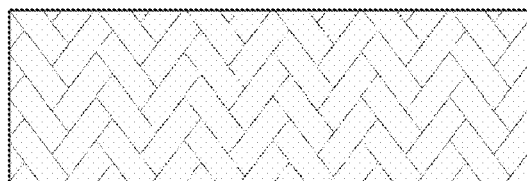


FIG. 17E



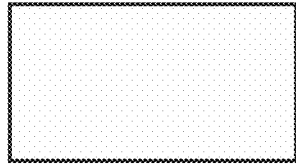


FIG. 18A

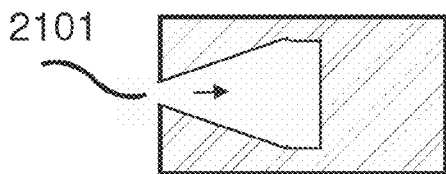
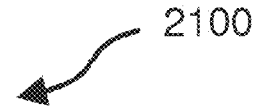


FIG. 18B

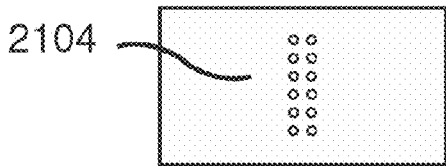


FIG. 18C

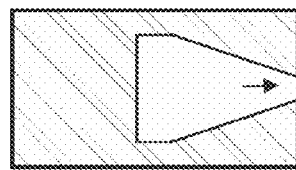


FIG. 18D

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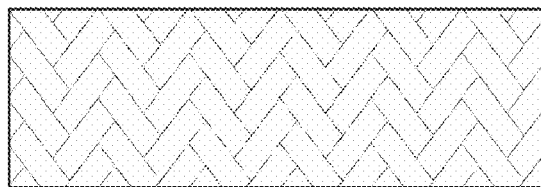
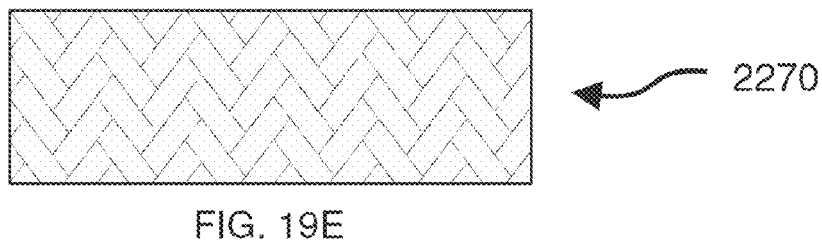
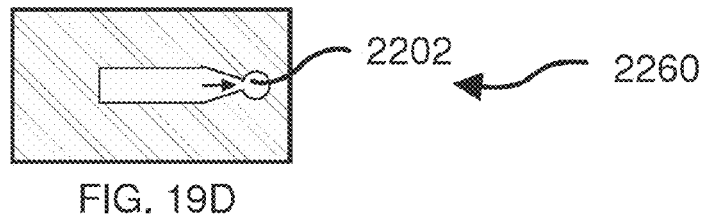
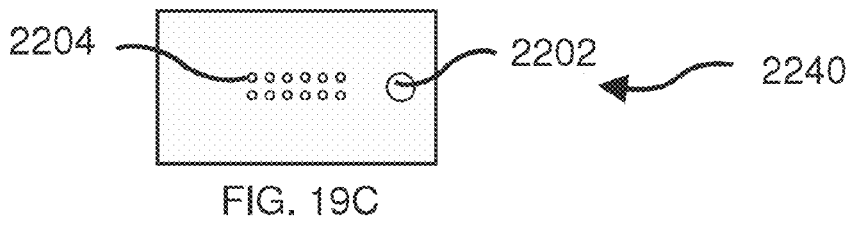
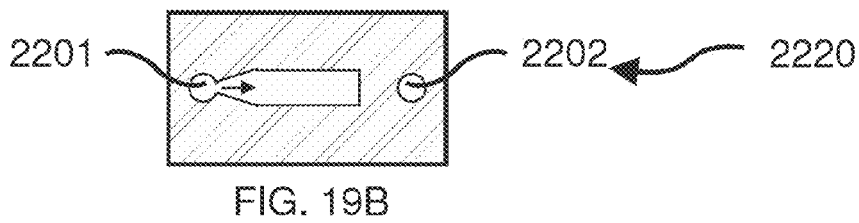
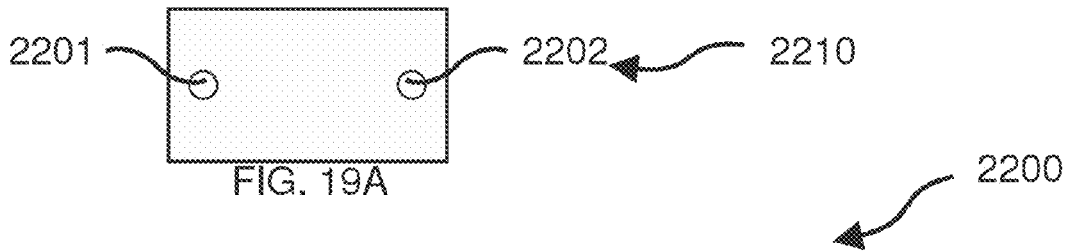


FIG. 18E





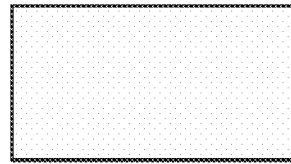


FIG. 20A

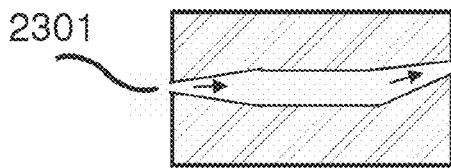
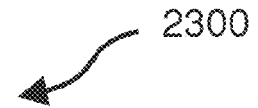
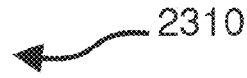


FIG. 20B

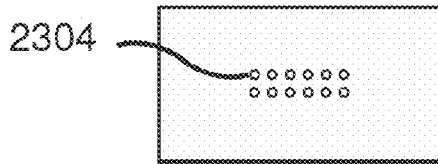


FIG. 20C

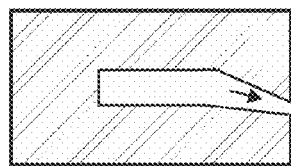


FIG. 20D

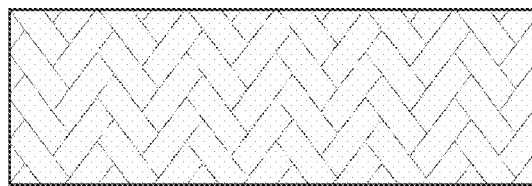
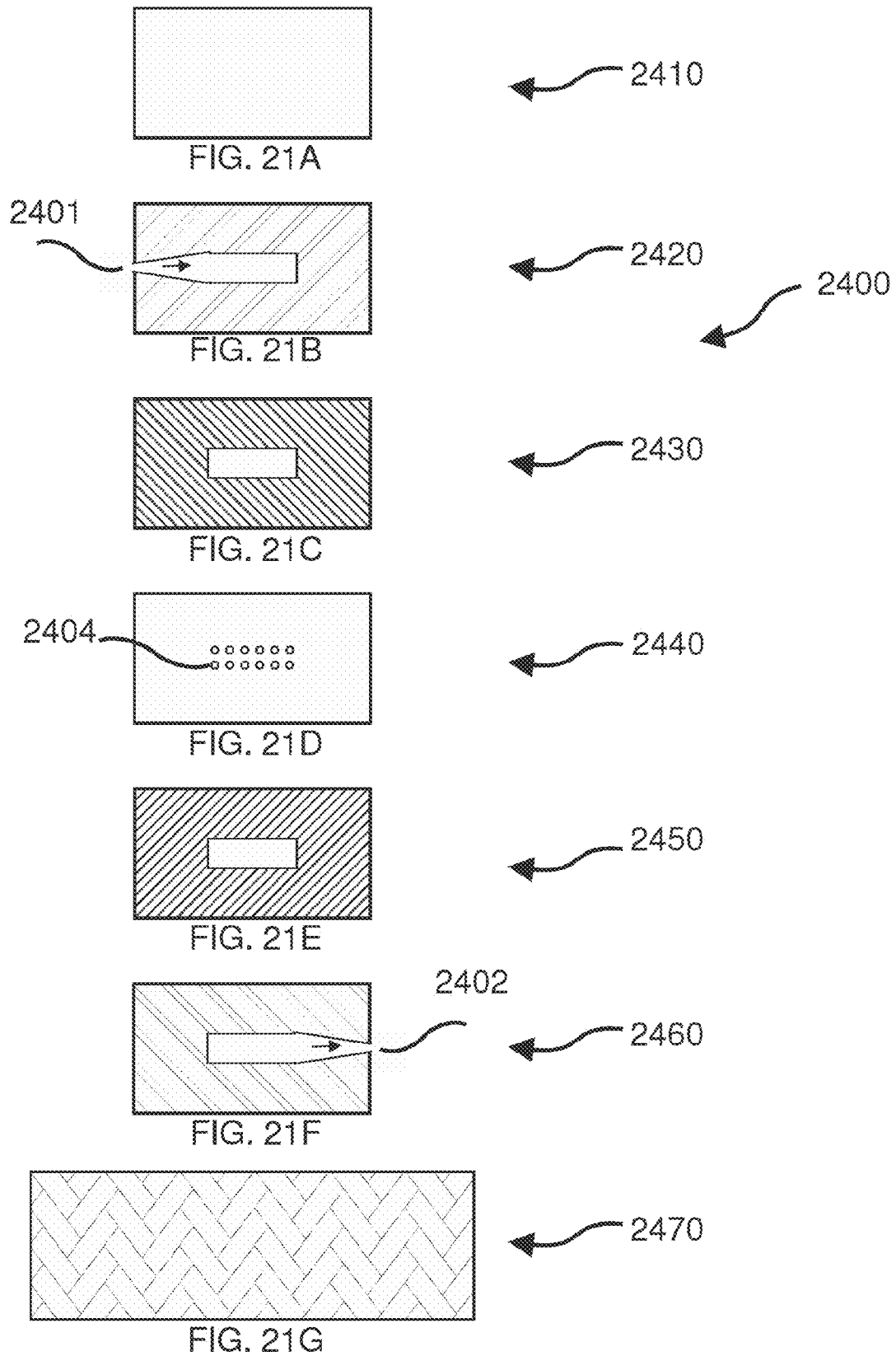


FIG. 20E





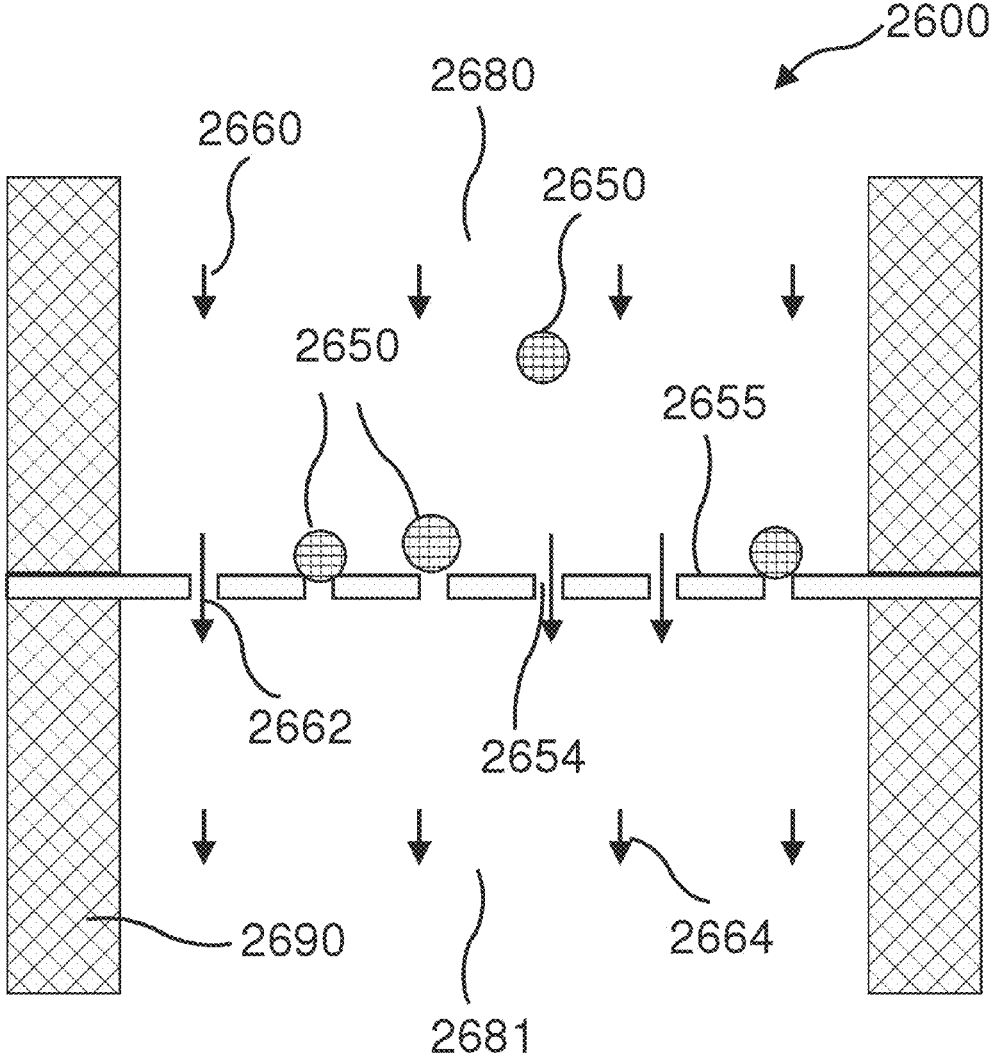


FIG. 22

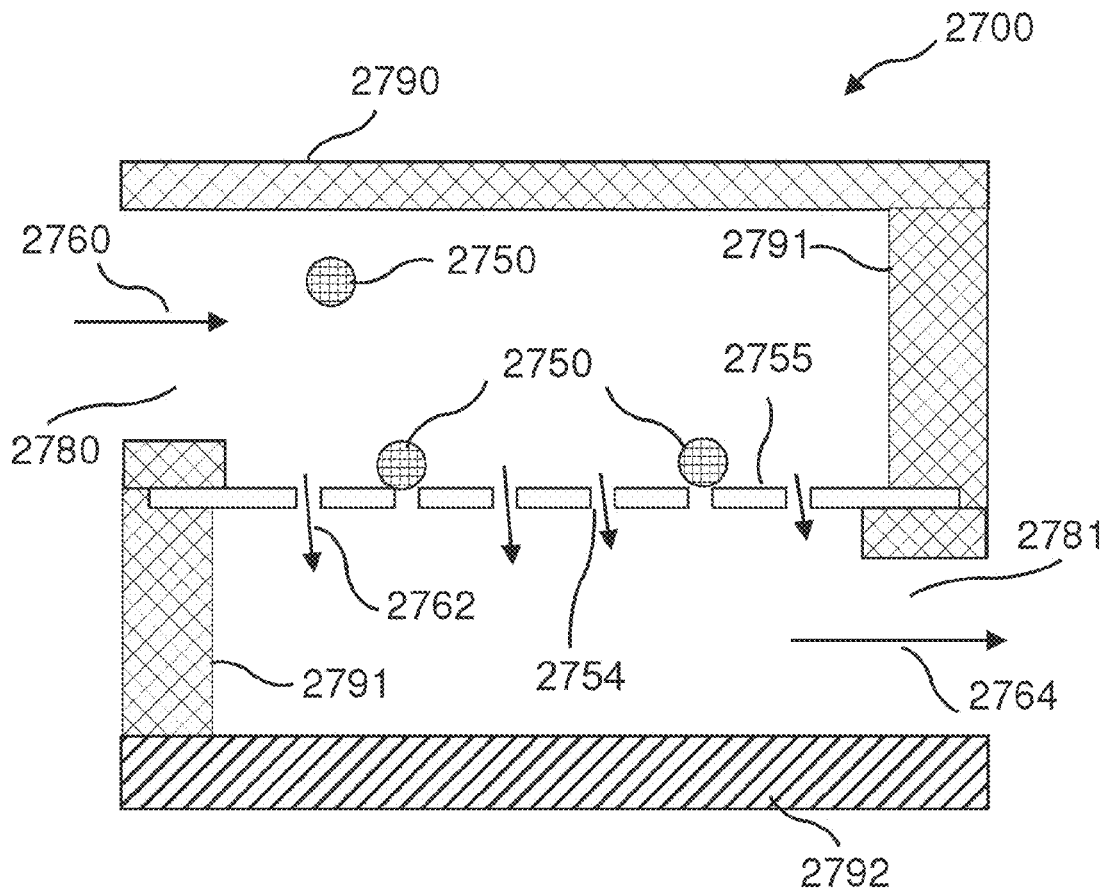


FIG. 23

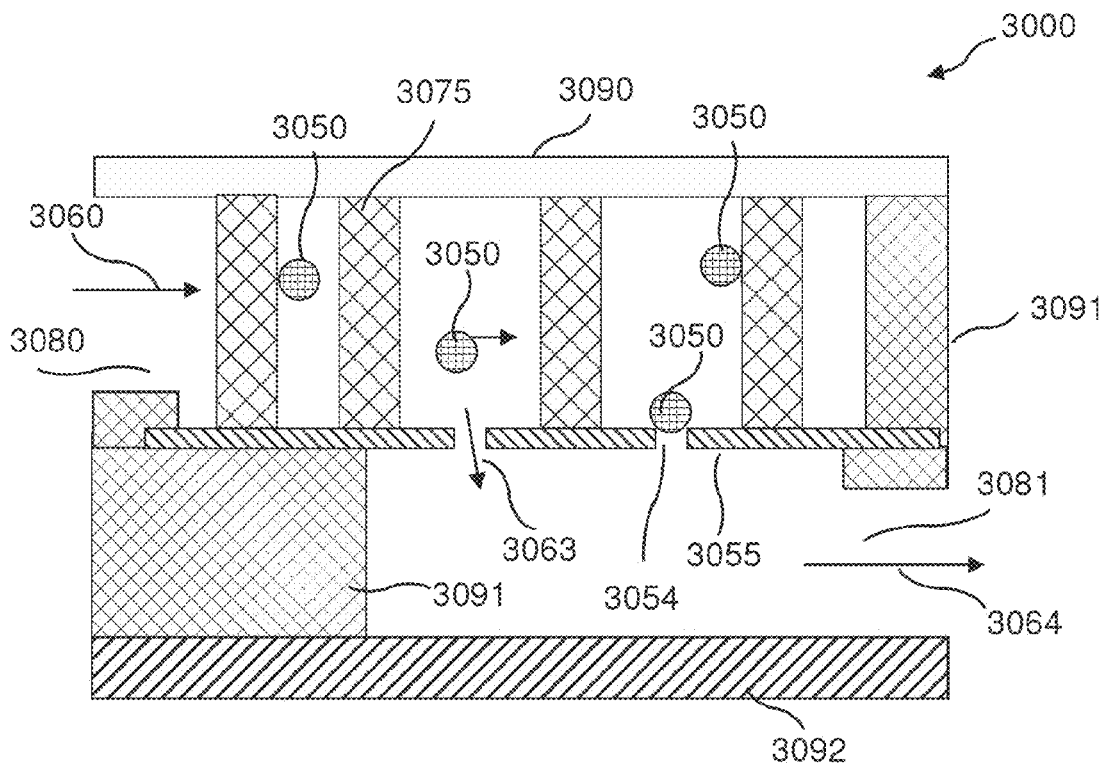


FIG. 24

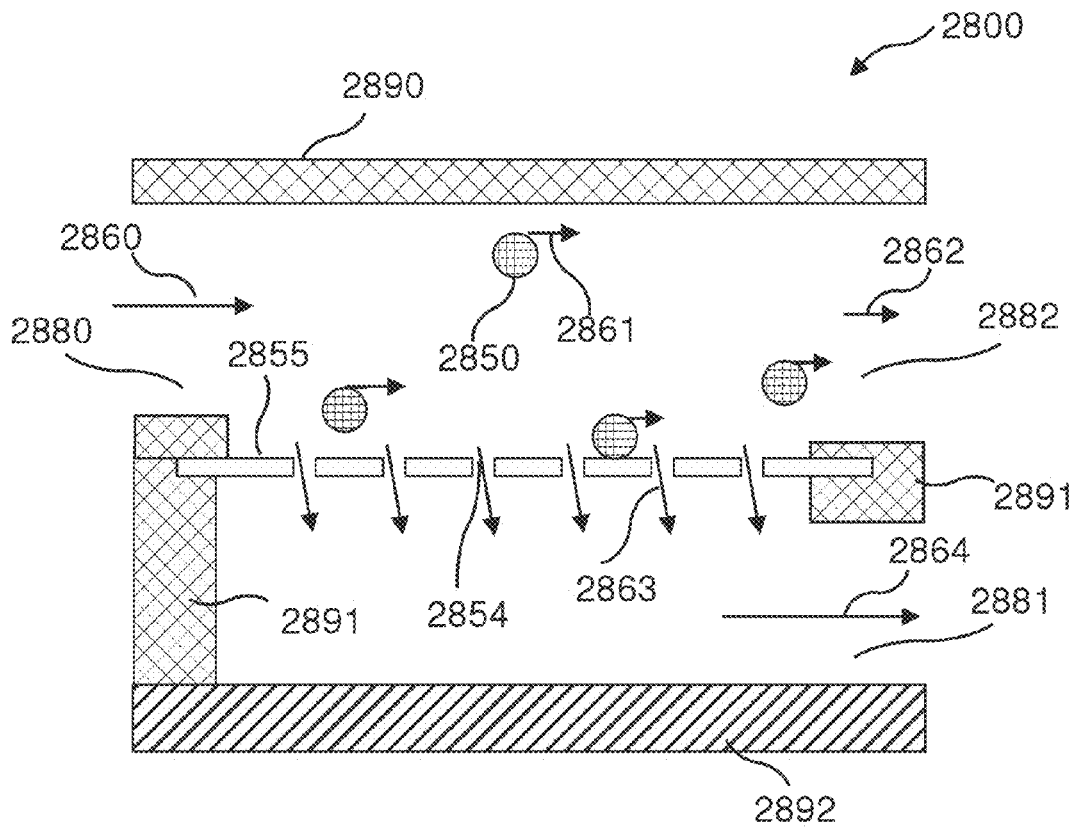


FIG. 25

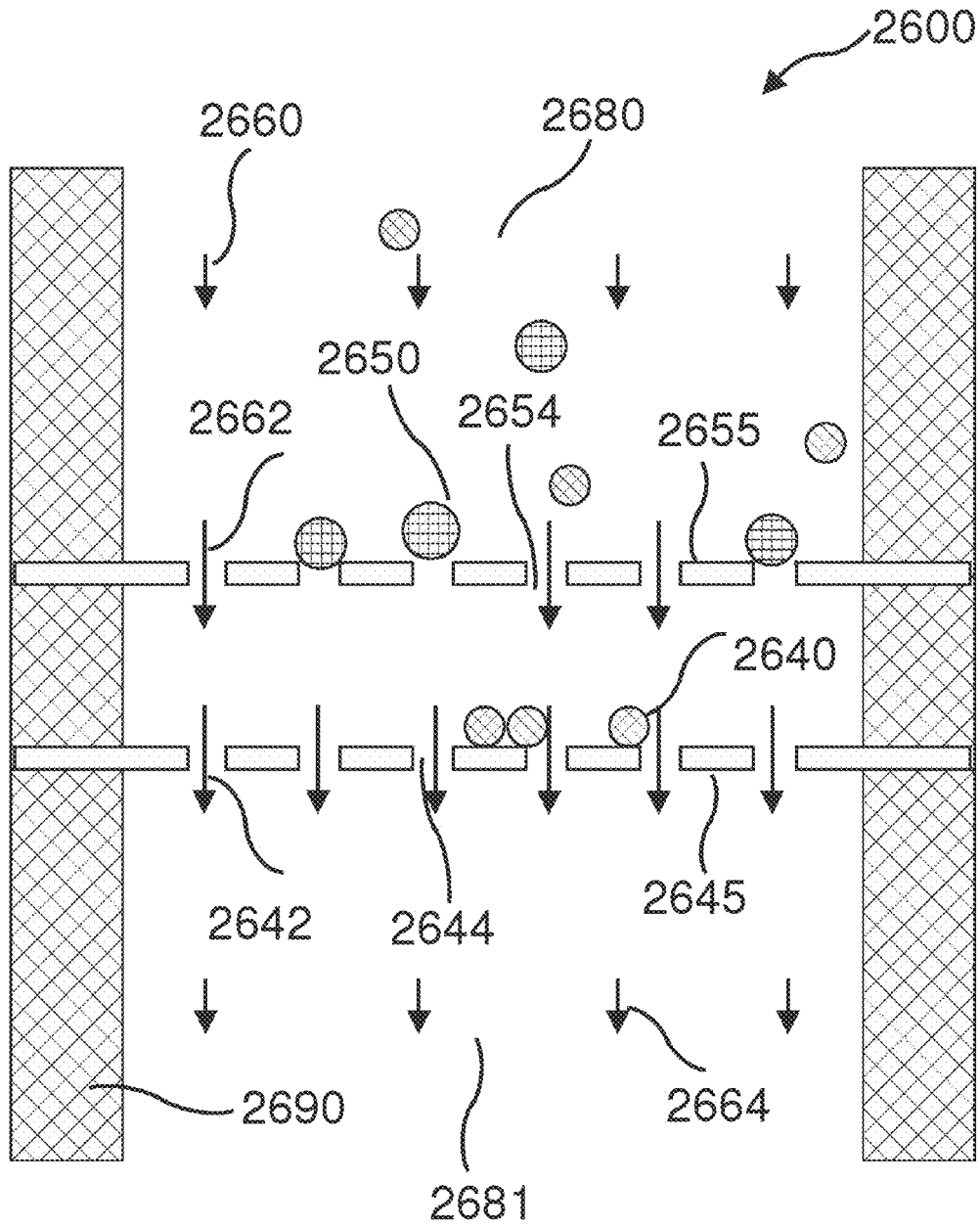


FIG. 26

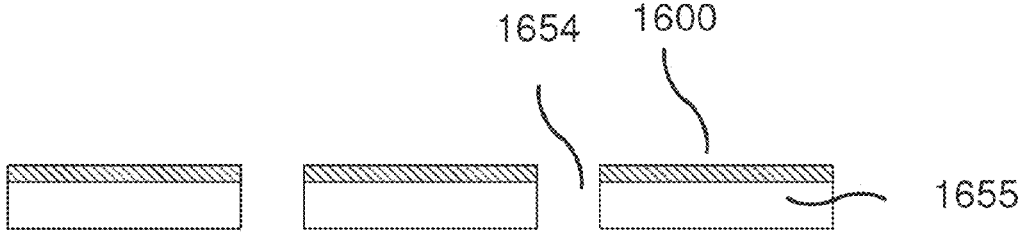


FIG. 27

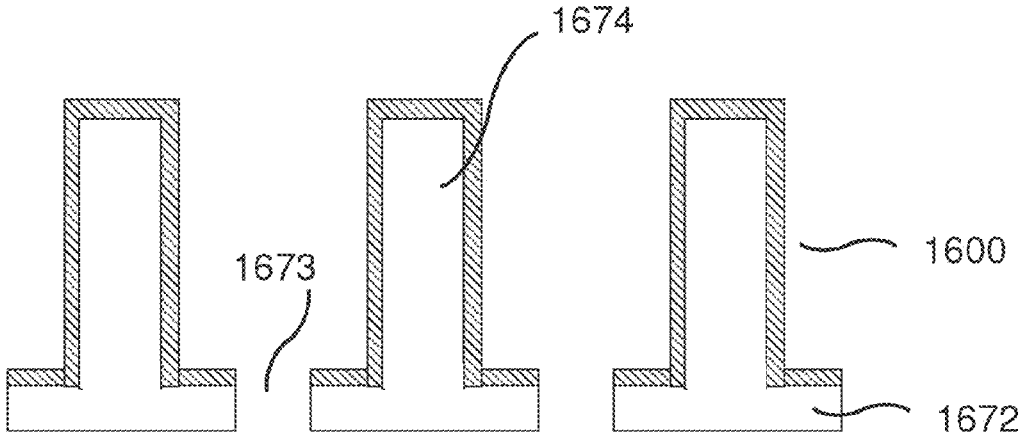


FIG. 28

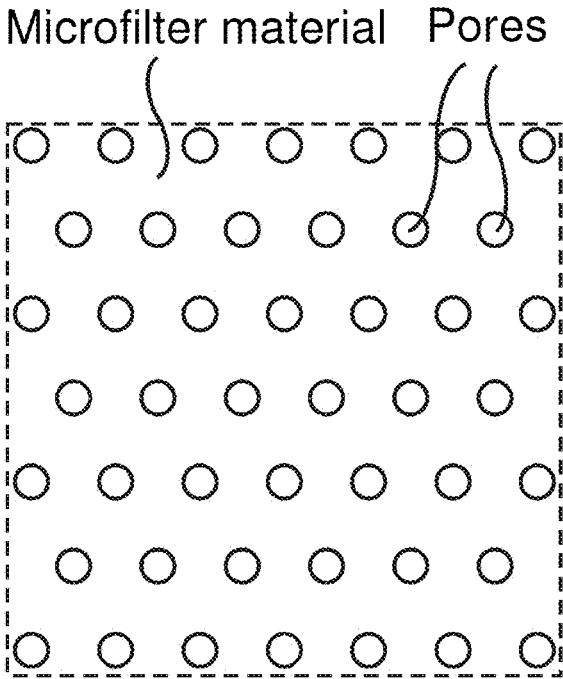


FIG. 29

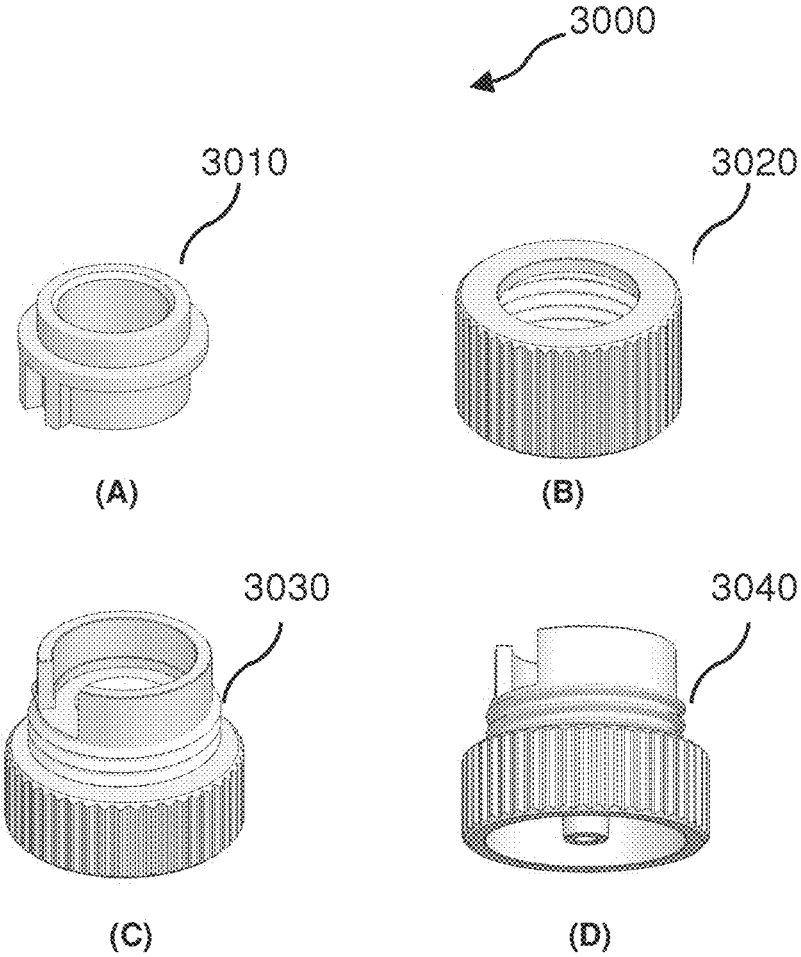


FIG. 30

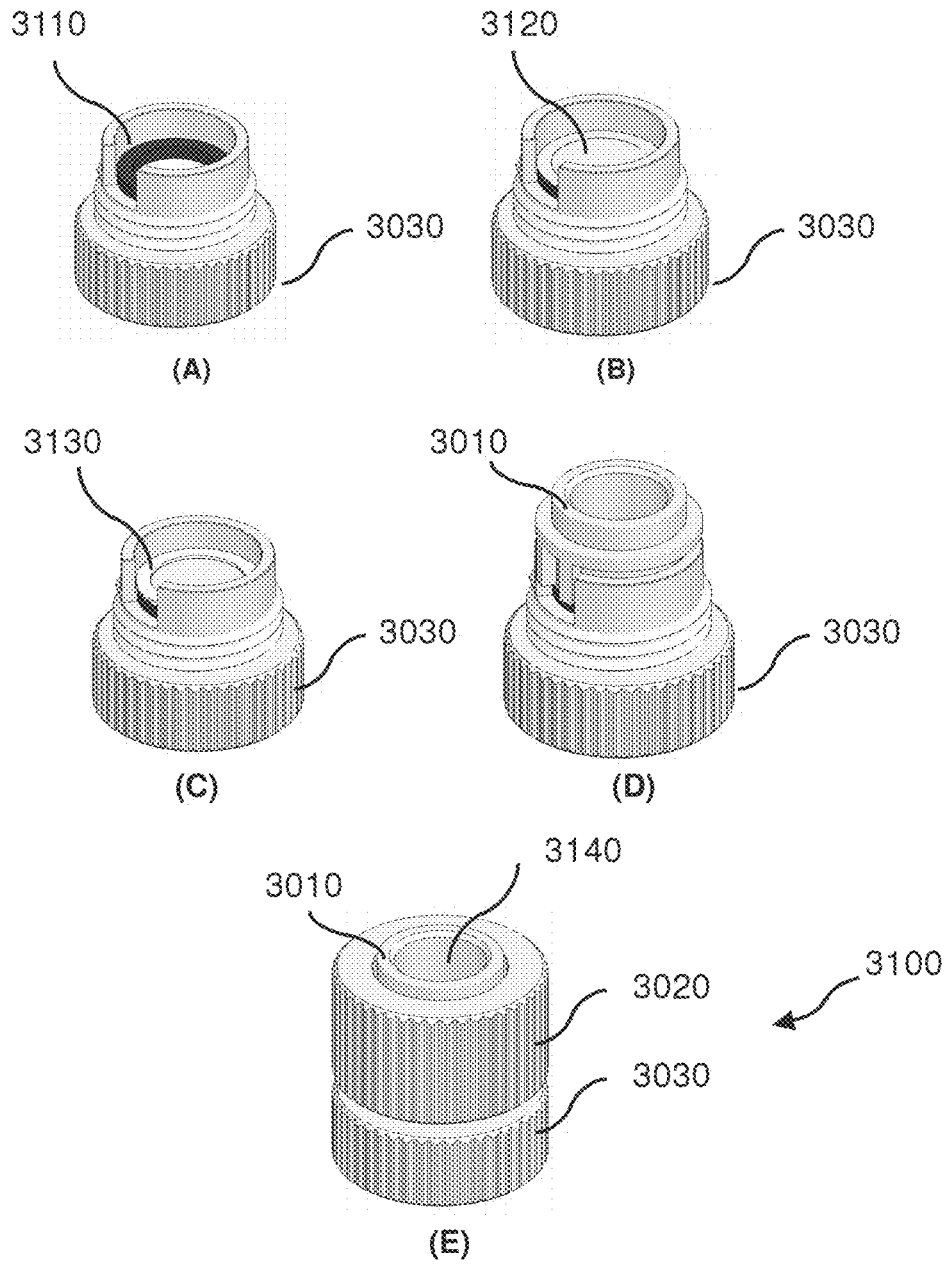


FIG. 31

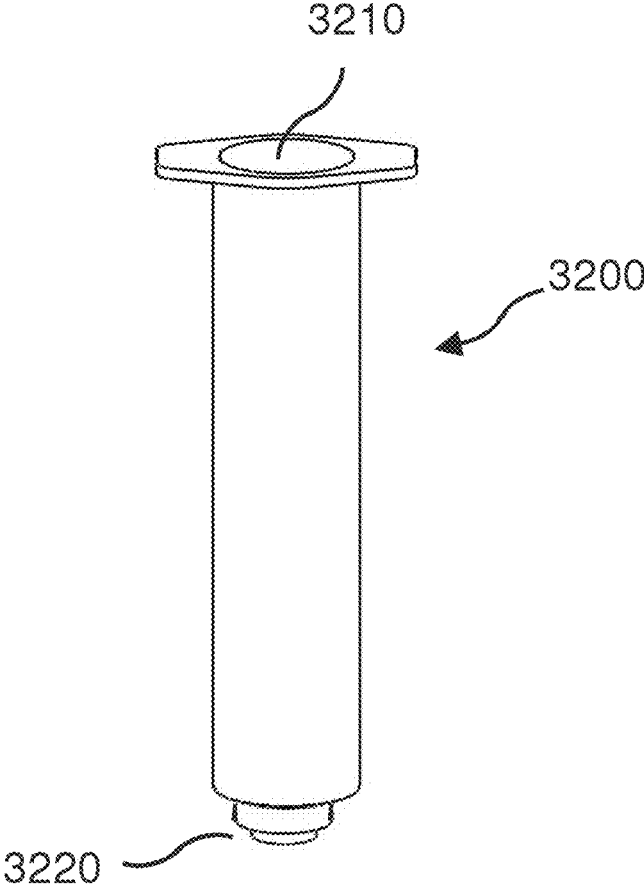


FIG. 32A

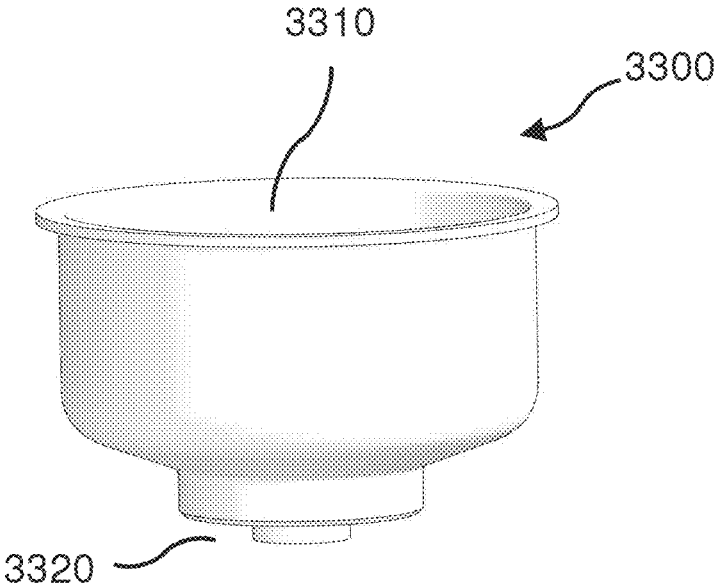


FIG. 32B

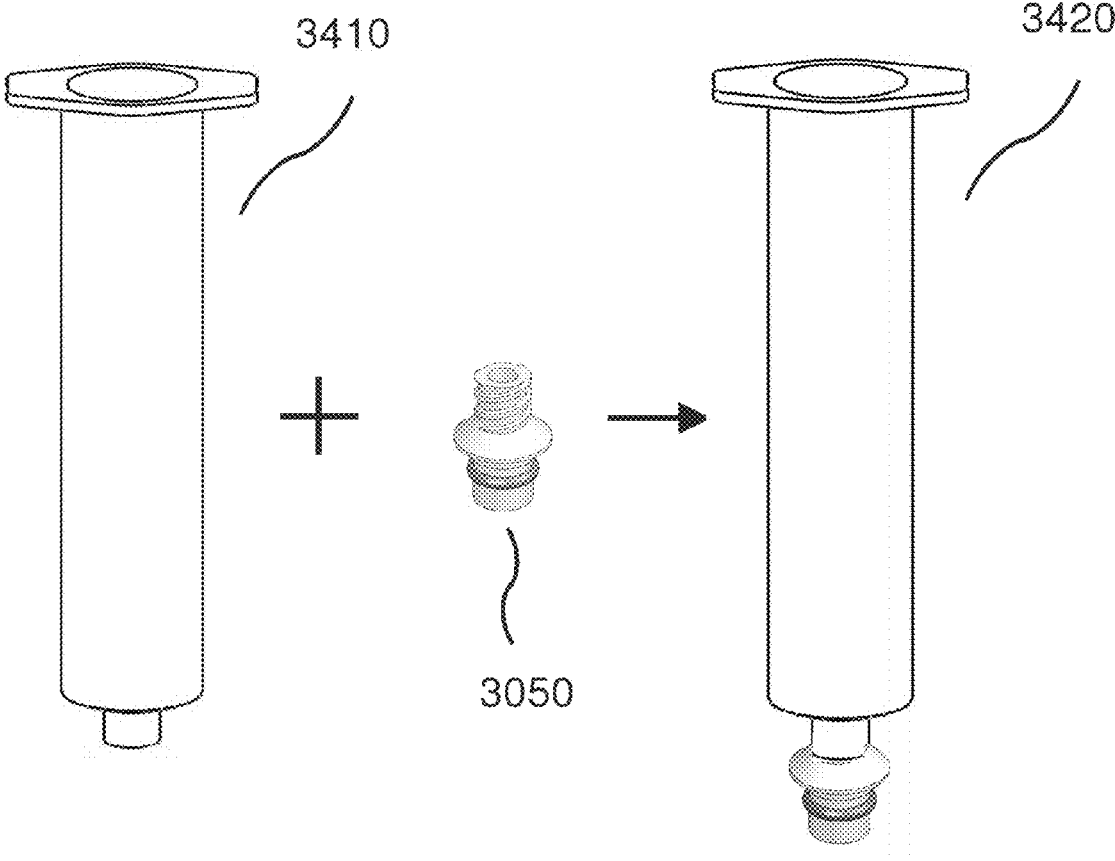


FIG. 33

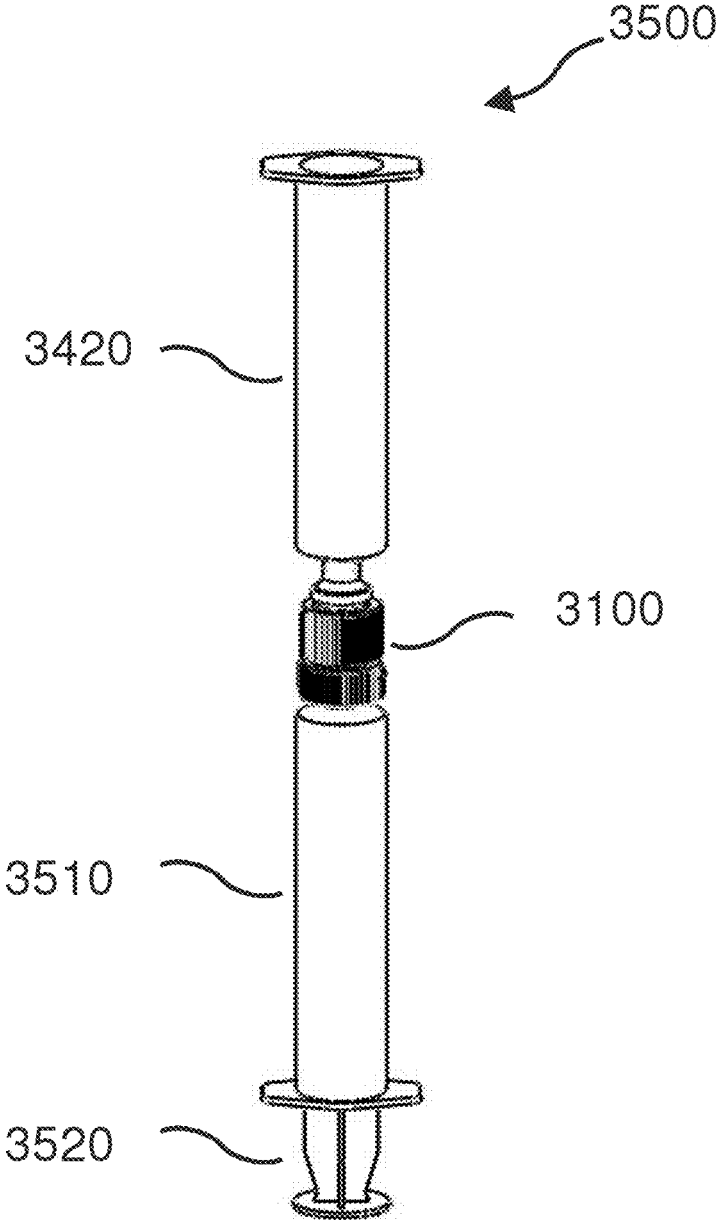


FIG. 34

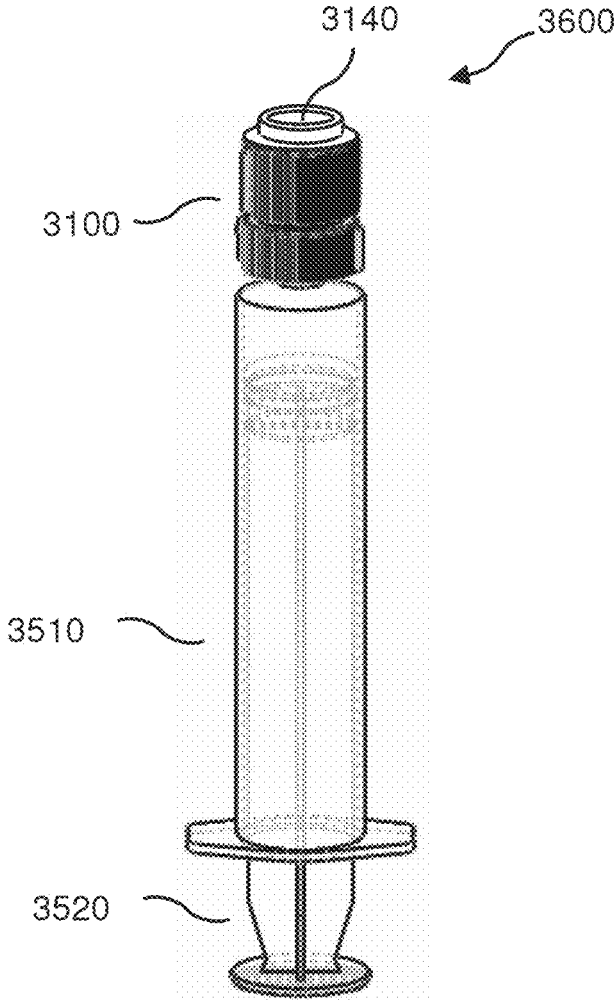


FIG. 35

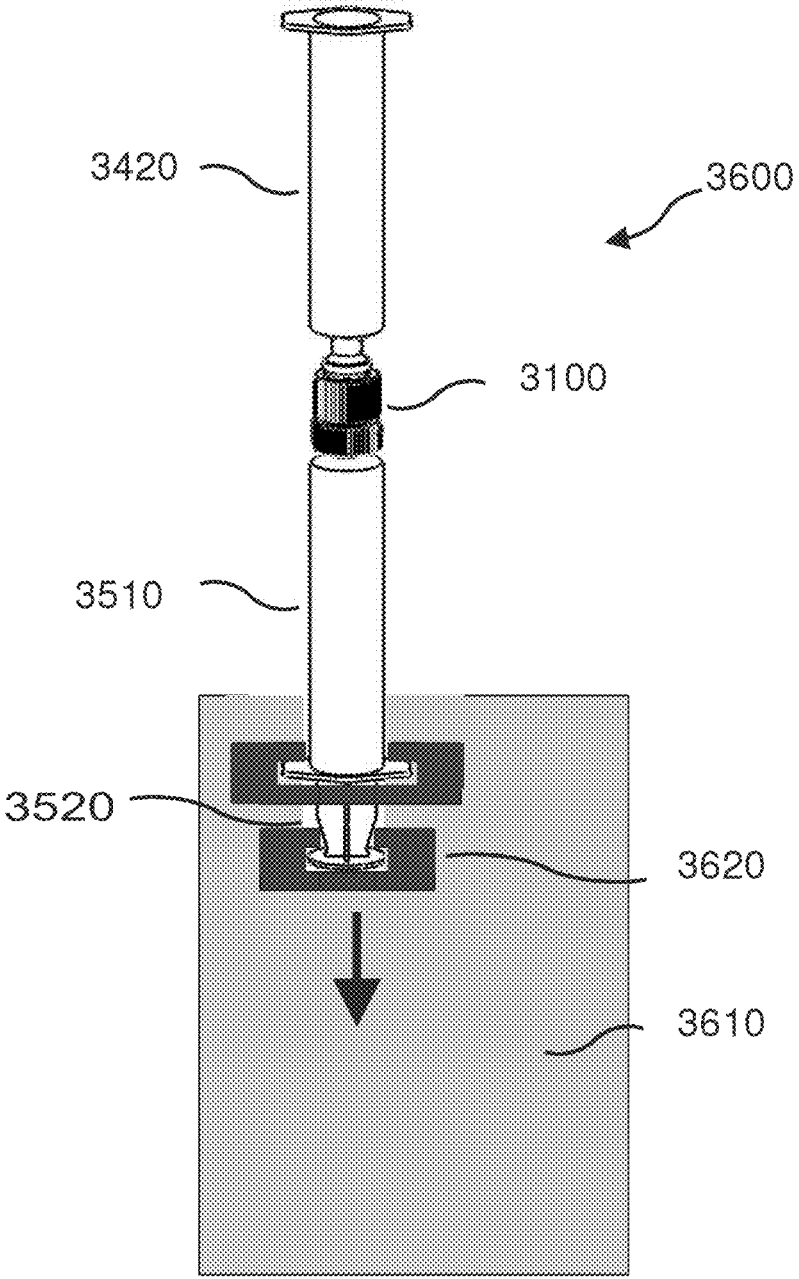


FIG. 36

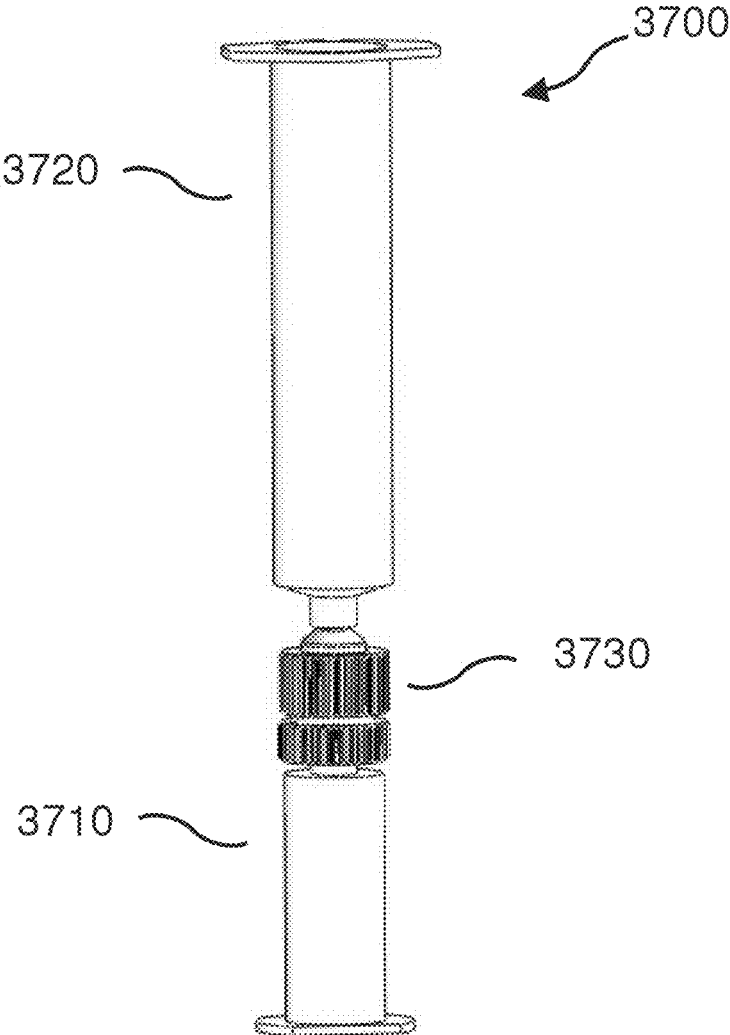


FIG. 37A

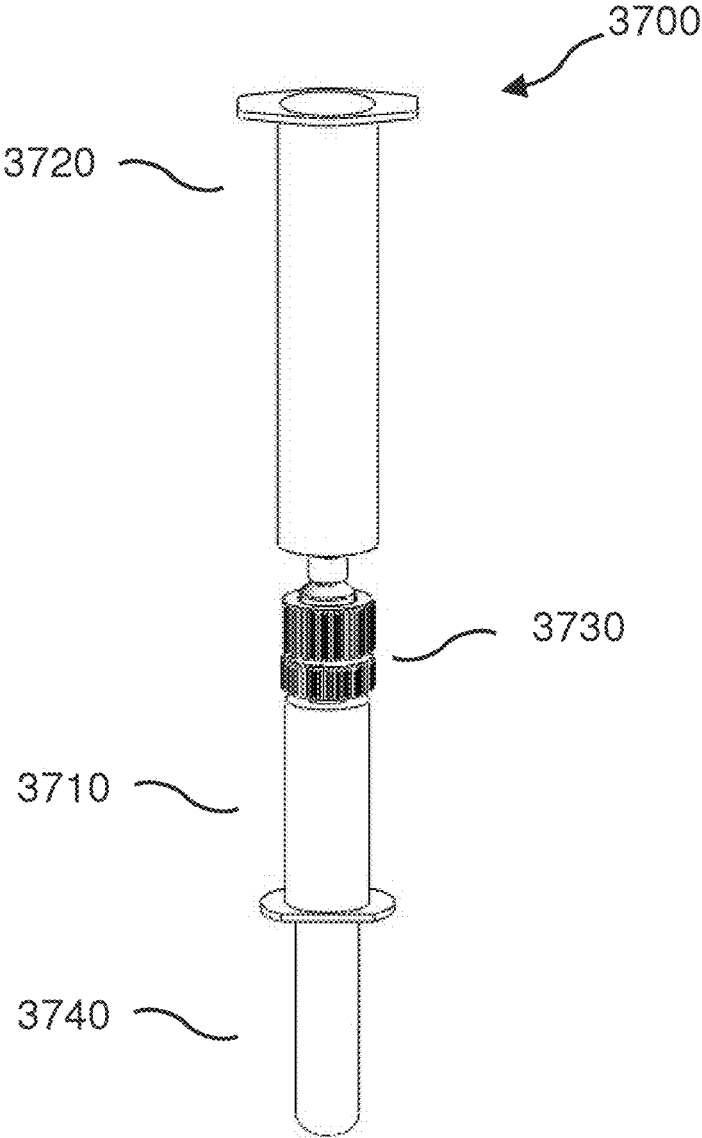


FIG. 37B

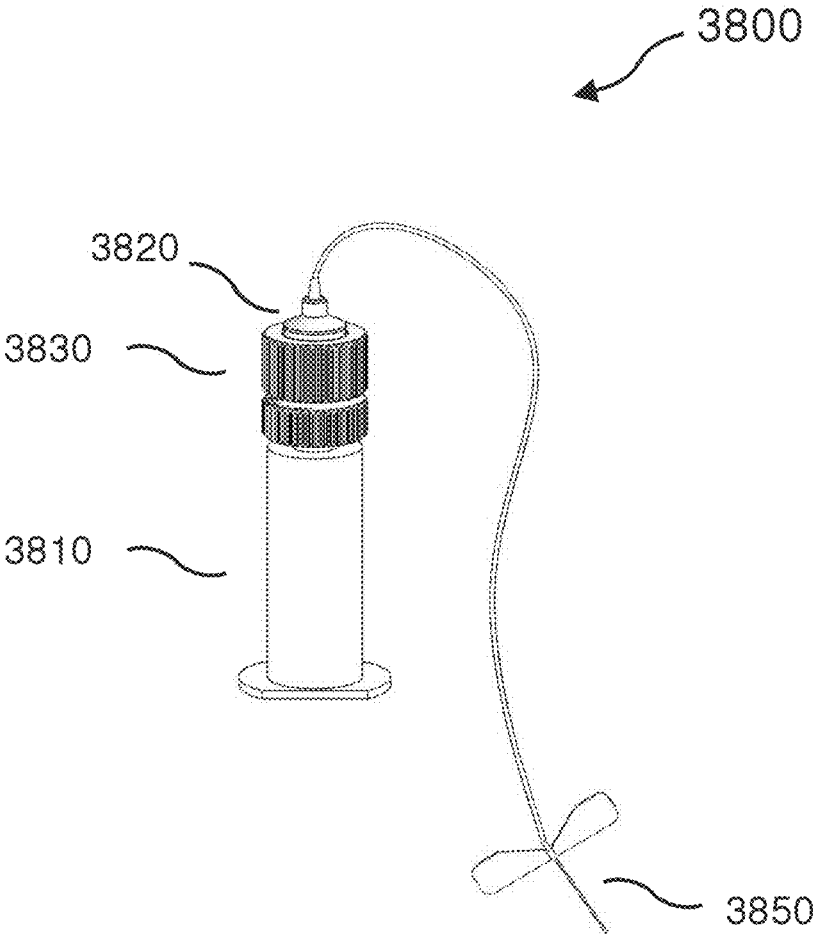


FIG. 38A

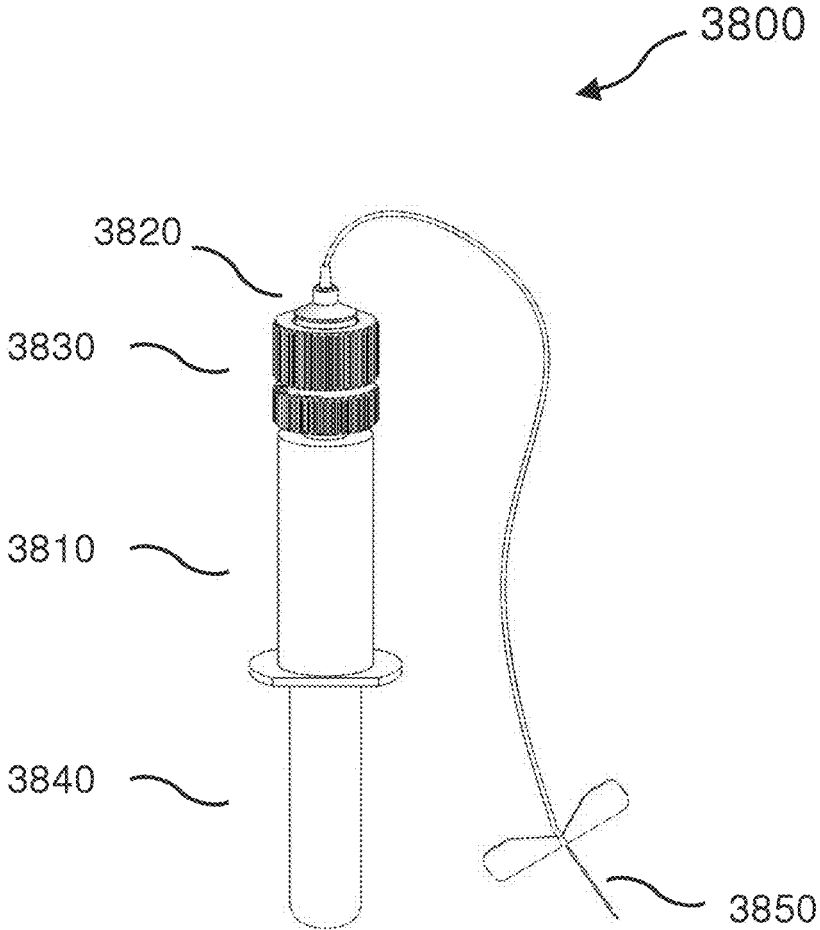


FIG. 38B

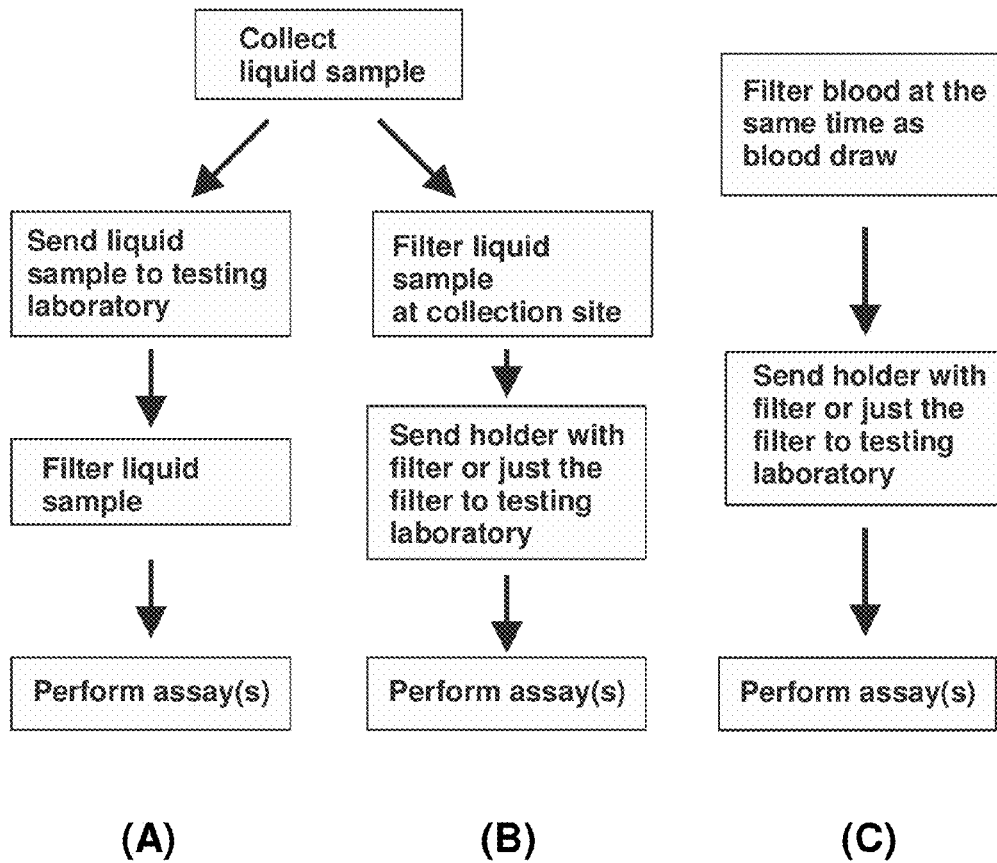


FIG. 39

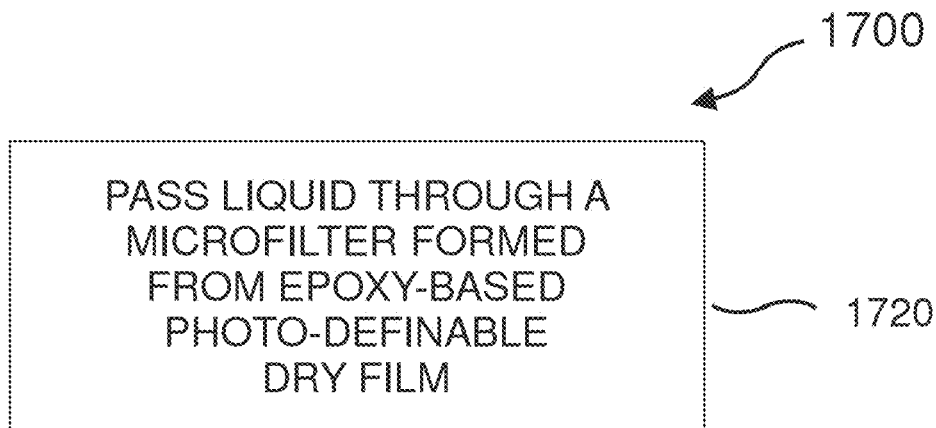


FIG. 40A

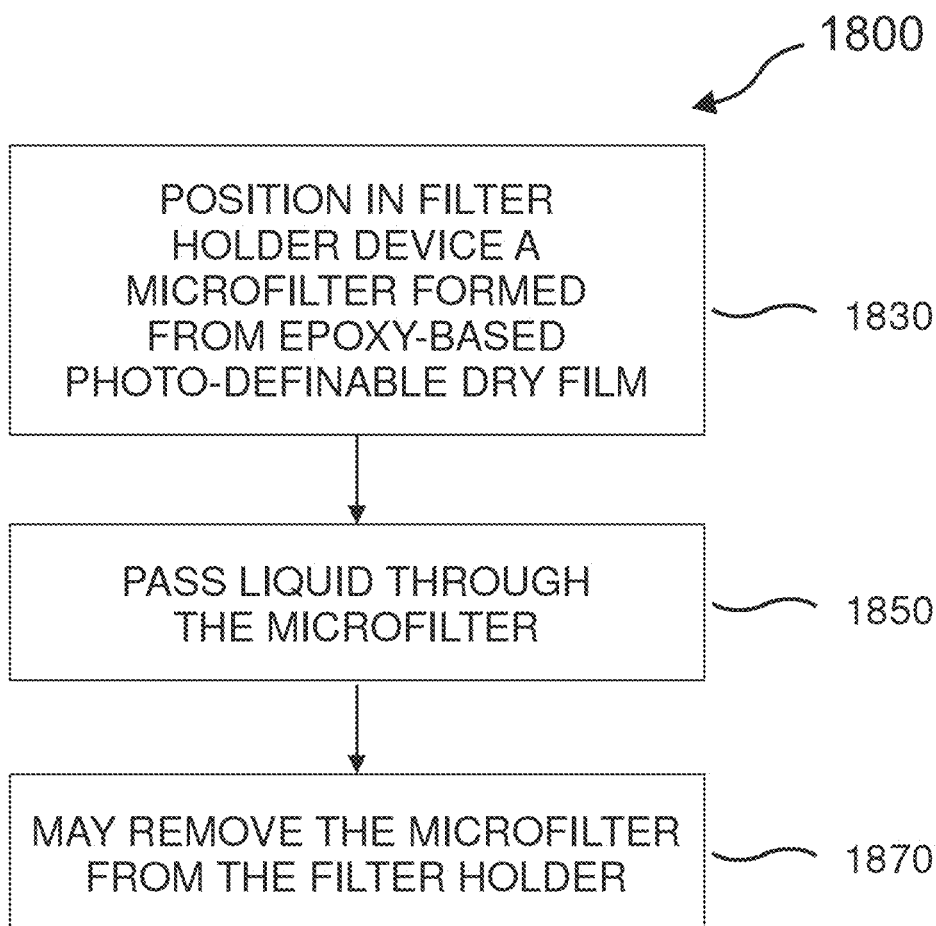


FIG. 40B

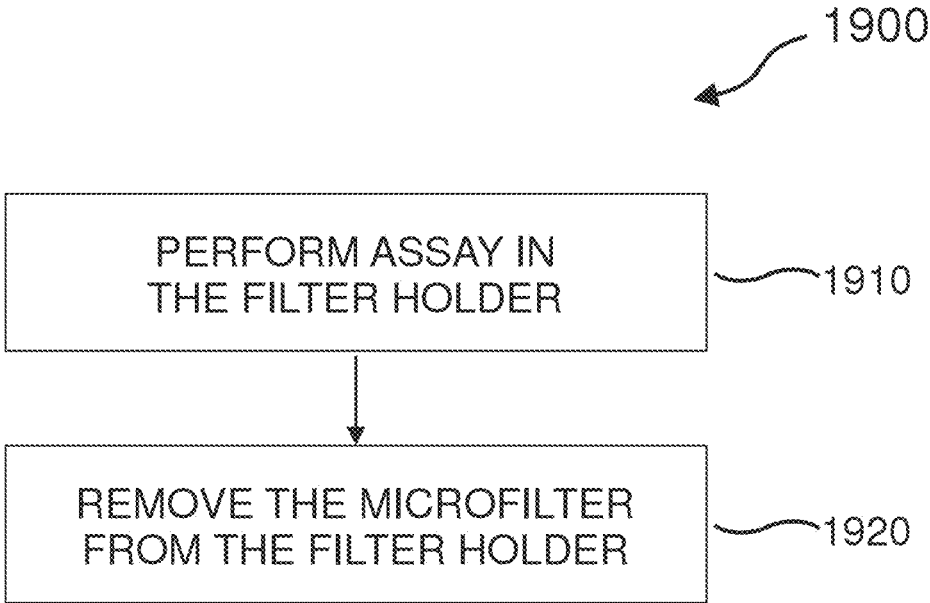


FIG. 40C

**POLYMER MICROFILTRATION DEVICES,
METHODS OF MANUFACTURING THE
SAME AND THE USES OF THE
MICROFILTRATION DEVICES**

**CROSS-REFERENCE TO RELATED
APPLICATIONS**

[0001] This is a continuation in part of prior application titled, "POLYMER MICROFILTRATION DEVICES, METHODS OF MANUFACTURING THE SAME AND THE USES OF THE MICROFILTRATION DEVICES", PCT/US12/66390 filed on Nov. 21, 2012, which claims priority from and incorporates by reference the entire contents of three prior U.S. Provisional Patent Applications (1) "POLYMER MICROFILTRATION DEVICES, METHODS OF MANUFACTURING THE SAME AND THE USES OF THE MICROFILTRATION DEVICES" No. 61/562,404 filed Nov. 21, 2011, (2) "POLYMER MICROFILTRATION DEVICES, METHODS OF MANUFACTURING THE SAME AND THE USES OF THE MICROFILTRATION DEVICES" No. 61/618,641 filed Mar. 30, 2012, (3) "POLYMER MICROFILTRATION DEVICES, METHODS OF MANUFACTURING THE SAME AND THE USES OF THE MICROFILTRATION DEVICES" No. 61/654,636 filed Jun. 1, 2012, and which is a continuation in part of prior application titled, "POLYMER MICROFILTERS AND METHODS OF MANUFACTURING THE SAME", PCT/US11/30966 filed on Apr. 1, 2011, the contents of all of which are hereby incorporated by reference herein.

[0002] This application also claims priority to prior application titled, "POLYMER MICROFILTRATION DEVICES, METHODS OF MANUFACTURING THE SAME AND THE USES OF THE MICROFILTRATION DEVICES", U.S. Provisional Patent Application No. 61/618,641 filed Mar. 30, 2012, the contents of which are hereby incorporated by reference herein.

[0003] This application also claims priority to prior application titled, "POLYMER MICROFILTRATION DEVICES, METHODS OF MANUFACTURING THE SAME AND THE USES OF THE MICROFILTRATION DEVICES", U.S. Provisional Patent Application No. 61/654,636 filed Jun. 1, 2012, the contents of which are hereby incorporated by reference herein.

BACKGROUND

[0004] 1. Field of the Invention

[0005] The present invention relates generally to microfiltration devices, that contain polymer microfilters, methods of manufacturing the same, methods to use the microfiltration device, and applications of the devices.

[0006] 2. Related Art

[0007] Some medical conditions may be diagnosed by detecting the presence of certain types of cells in bodily fluid. In particular, cells indicative or characteristic of certain medical conditions may be larger and/or less flexible than other cells found in certain bodily fluids. Accordingly, by collecting such larger and/or less flexible cells from a liquid sample of a bodily fluid, it may be possible to diagnose a medical condition based on the cells collected.

[0008] Cells larger and/or less flexible than other cells present in a bodily fluid may be collected by filtering the bodily fluid. For example, targeted cells indicative of a condition may be collected by passing a bodily fluid through a

filter having openings that are too small for the target cells to pass through, but large enough for other cells to pass through. Once collected, any number of analyses of the target cells may be performed. Such analyses may include, for example, identifying, counting, characterizing, and/or culturing the collected cells.

[0009] It is desirable for microfilters to have precise pore dimensions, not break during use, and not be autofluorescent for fluorescent microscope imaging. Conventionally, microfilters are installed in a microfiltration device and liquid samples processed to collect cells based on size on the microfilter.

SUMMARY

[0010] In one aspect of the present invention, a microfilter is disclosed. According to an exemplary embodiment, a microfilter comprises a polymer layer formed from epoxy-based photo-definable dry film, and a plurality of apertures each extending through the polymer layer.

[0011] In another aspect of the present invention, multi-layer microfilters are disclosed. According to an exemplary embodiment, a microfilter comprises a first polymer layer formed from epoxy-based photo-definable dry film and having a first aperture extending therethrough, and a second polymer layer formed from epoxy-based photo-definable dry film and having a second aperture extending therethrough, wherein the first and second apertures at least partially define a non-linear passage extending through the first and second layers.

[0012] In yet another aspect of the present invention, a method of manufacturing a microfilter is disclosed. According to an exemplary embodiment, a method comprises providing a first layer of epoxy-based photo-definable dry film disposed on a substrate, exposing the first layer to energy through a mask to form a pattern, defined by the mask, in the first layer of dry film, forming, from the exposed first layer of dry film, a polymer layer having a plurality of apertures extending therethrough, the plurality of apertures having a distribution defined by the pattern, and removing the polymer layer from the substrate.

[0013] In yet another aspect of the present invention, a method of forming a multi-layer microfilter is disclosed. According to an exemplary embodiment, a method comprises forming a first polymer layer comprising a plurality of first apertures from a first layer of epoxy-based photo-definable dry film disposed on a substrate, laminating a second layer of epoxy-based photo-definable dry film on the first polymer layer, and forming a second polymer layer comprising a plurality of second apertures from the second layer of dry film.

[0014] In one aspect of the present invention, filter holders to hold the microfilter are disclosed. According to an exemplary embodiment, filter holders are designed to keep the filter flat and fixed in place, allowing the implementation of various assay steps with ease.

[0015] In yet another aspect of the present invention, a method of using a microfilter is disclosed. According to an exemplary embodiment, a method comprises passing a liquid through a plurality of apertures of a microfilter formed from an epoxy-based photo-definable dry film, wherein the microfilter has sufficient strength and flexibility to filter the liquid, and wherein the apertures are sized to allow passage of a first type of bodily fluid cell and to prevent passage of a second type of bodily fluid cell. An exemplary implementation includes application of a negative pressure using either

vacuum or filter holder connect to syringe(s), which can be performed manually, semi-manually using a syringe pump, or automated.

[0016] In an exemplary implementation, filters can be coated with analyte recognition reagents to capture cells of interest from the body fluids.

[0017] In yet another exemplary implementations of embodiments of the present invention, methods to collect liquid sample and shipping the filtered liquid sample using the microfilter inside the filter holder are also described.

[0018] In yet other exemplary implementations of embodiments of the present invention, methods of using the microfilters to perform assays are disclosed. Exemplary embodiment of the present invention provide methods to isolate cells, methods to recover the cells from the microfilter by backwashing using the filter holder, and methods to perform assay inside the filter holder.

[0019] In yet another aspect of the present invention, exemplary embodiments provide microfiltration chips comprising microfilters.

[0020] In yet another aspect of the present invention, exemplary embodiments provide methods to perform filtration comprising using the microfilter and microfiltration chips.

[0021] In yet another aspect of the present invention, exemplary medical applications of the assays are disclosed.

BRIEF DESCRIPTION OF THE DRAWINGS

[0022] A more complete appreciation of the present invention and many of the attendant advantages thereof will be readily obtained as the same becomes better understood by reference to the following detailed description when considered in connection with the accompanying drawings, wherein:

[0023] FIGS. 1A-1E are cross-sectional views illustrating multiple stages in a process for manufacturing a microfilter in accordance with exemplary embodiments of the present invention;

[0024] FIG. 2A is a flowchart illustrating a process for manufacturing a microfilter in accordance with exemplary embodiments of the present invention;

[0025] FIG. 2B is a flowchart illustrating a process for forming a microfilter from an exposed dry film in the process illustrated in FIG. 2A, in accordance with exemplary embodiments of the present invention;

[0026] FIGS. 3A-3E are cross-sectional views illustrating multiple stages in a process for manufacturing a microfilter 120 in accordance with exemplary embodiments of the present invention;

[0027] FIGS. 4A-4D are cross-sectional views illustrating multiple stages in a process for manufacturing a microfilter in accordance with exemplary embodiments of the present invention;

[0028] FIGS. 5A-5D are cross-sectional views illustrating multiple stages in a process for manufacturing a plurality of microfilters from a plurality of layers of epoxy-based photo-definable dry film in accordance with exemplary embodiments of the present invention;

[0029] FIGS. 6A and 6B are cross-sectional views illustrating multiple stages in a process for attaching dry film structures to a support using an electrostatic chuck apparatus in a process for forming microfilters in accordance with exemplary embodiments of the present invention;

[0030] FIGS. 7A and 7B are cross-sectional views illustrating multiple stages in a process for manufacturing microfil-

ters from a roll of a dry film structure in accordance with exemplary embodiments of the present invention;

[0031] FIGS. 8A and 8B are cross-sectional views illustrating multiple stages in a process for manufacturing microfilters from a plurality of rolls of dry film structures in accordance with exemplary embodiments of the present invention;

[0032] FIGS. 9A-9D are partial top views illustrating various microfilters aperture distributions in accordance with exemplary embodiments of the present invention;

[0033] FIGS. 10A-10E are cross-sectional views illustrating microfilters having various thicknesses and various aperture shapes, sizes and distributions in accordance with exemplary embodiments of the present invention;

[0034] FIG. 11 is a flow chart illustrating a process 1300 for manufacturing a multi-layer microfilter in accordance with exemplary embodiments of the present invention;

[0035] FIGS. 12A-12K are cross-sectional views illustrating multiple stages in a process for manufacturing a multi-layer microfilter in accordance with exemplary embodiments of the present invention;

[0036] FIG. 12L is a top view of multi-layer microfilter in accordance with exemplary embodiments of the present invention;

[0037] FIGS. 13A and 13B are top views illustrating multiple stages in the process of FIG. 11;

[0038] FIGS. 14A and 14B are cross-sectional views of a multi-layer microfilter 1420 in accordance with exemplary embodiments of the present invention.

[0039] FIG. 14C is a top view of a multi-layer microfilter 1420 of FIGS. 14A and 14B;

[0040] FIG. 15 is a cross-sectional view of a microfiltration structure including a microfilter and a support structure in accordance with exemplary embodiments of the present invention.

[0041] FIG. 16A is a cross-sectional view of a microfiltration structure including pores in the microfilter and posts above the microfilter in accordance with exemplary embodiments of the present invention.

[0042] FIG. 16B is a top view of a microfiltration structure including pores in the microfilter and posts above the microfilter in accordance with exemplary embodiments of the present invention.

[0043] FIG. 16C is a top view of a microfiltration structure including rectangular pores in the microfilter and posts above the microfilter in accordance with exemplary embodiments of the present invention.

[0044] FIGS. 16D and 16E are side views of another microfilter according to an exemplary embodiment of the present invention made by two layers of material with different patterns in each layer.

[0045] FIG. 16F is the top view of the microfilter according to an exemplary embodiment of the present invention made by two layers of material with different patterns in each layer.

[0046] FIGS. 16G and 16H are side views of another microfilter according to an exemplary embodiment of the present invention made by two layers of material with different patterns in each layer.

[0047] FIGS. 16I and 16J are side views of a device according to an exemplary embodiment of the present invention with many wells suitable for many assays made by two layers of material with opening on the top layer and solid bottom layer.

[0048] FIGS. 17A-17E are top views of a device according to an exemplary implementation of an embodiment of the

present invention where different layers when stacked together form a microfiltration chip with one inlet and one outlet.

[0049] FIGS. 18A-18E are top views of a device according to another exemplary implementation of an embodiment of the present invention where different layers when stacked together form another microfiltration chip with one inlet and one outlet.

[0050] FIGS. 19A-19E are top views of a device according to yet another exemplary implementation of an embodiment of the present invention where different layers when stacked together form another microfiltration chip with one inlet and one outlet.

[0051] FIGS. 20A-20E are top views of a device according to yet another exemplary implementation of an embodiment of the present invention where different layers when stacked together form another microfiltration chip with one inlet and two outlets.

[0052] FIGS. 21A-21G are top views of a device according to yet another exemplary implementation of an embodiment of the present invention where different layers when stacked together form another microfiltration chip with one inlet and one outlet.

[0053] FIG. 22 shows an exemplary implementation of an embodiment of the present invention where a microfilter in a filter holder capturing large particulates from the liquid as it flows through the microfilter.

[0054] FIG. 23 shows a cross-sectional view of an exemplary implementation of an embodiment of the present invention including the microfiltration chip with one inlet and one outlet. The microfilter in microfiltration chip capturing large particulates from the liquid as it flows through the microfilter.

[0055] FIG. 24 shows a cross-sectional view of another exemplary implementation of an embodiment of the present invention including the microfiltration chip with one inlet and one outlet, where the microfilter has posts disposed above the pores. The microfilter in microfiltration chip capturing large particulates from the liquid as it flows through the microfilter.

[0056] FIG. 25 shows cross-sectional view of an exemplary implementation of an embodiment of the present invention including the microfiltration chip with one inlet and two outlets.

[0057] FIG. 26 shows an exemplary implementation of an embodiment of the present invention including two microfilters in a filter holder capturing large particulates from the liquid as it flows through the microfilter on the first microfilter and collect smaller particulated from the liquid on the second microfilter.

[0058] FIG. 27 is a cross-sectional views of coated planar microfilters in accordance with exemplary embodiments of the present invention;

[0059] FIG. 28 is a cross-sectional views of coated structured microfilters in accordance with exemplary embodiments of the present invention;

[0060] FIG. 29 illustrate an example of lithographically fabricated microfilter according to an exemplary embodiment of the present invention using epoxy-based photo-definable dry film with the pore geometry and locations defined by an optical mask;

[0061] FIGS. 30A-30D illustrates the components of a filter holder in accordance with exemplary embodiments of the present invention;

[0062] FIGS. 31A-31E illustrates the assembly process of the microfilter in the filter holder illustrated in FIGS. 30A-30D in accordance with exemplary embodiments of the present invention;

[0063] FIGS. 32A-32B illustrate input container options for input liquid sample and reagents to be attached to the assembled sample holder illustrated in FIG. 31E in accordance with exemplary embodiments of the present invention;

[0064] FIG. 33 is an illustration of forming an input container according to an exemplary embodiment of the present invention to be attached to the assembled sample holder illustrated in FIG. 31E using a syringe with Luer lock without the plunger and an adaptor in accordance with exemplary embodiments of the present invention;

[0065] FIG. 34 illustration of the filtration system according to an exemplary embodiment of the present invention to perform the assays incorporating a microfilter inside the filter holder in accordance with exemplary embodiments of the present invention;

[0066] FIG. 35 illustrates the filtration system according to an exemplary embodiment of the present invention to perform some steps of assays with the input container removed in accordance with exemplary embodiments of the present invention;

[0067] FIG. 36 illustrates the filtration system according to an exemplary embodiment of the present invention using filtration system illustrated in FIG. 34, where the outlet syringe is pulled by a syringe pump.

[0068] FIGS. 37A-37B shows the configuration according to an exemplary embodiment of the present invention using vacutainer holder to draw the liquid sample from the input sample holder through the filter to a vacutainer.

[0069] FIGS. 38A-38B shows the configuration according to an exemplary embodiment of the present invention using vacutainer holder to draw the liquid sample from the patient through the filter to a vacutainer. This can be used, for example, to filter circulating tumor cells at time of blood drawn in accordance with embodiments of the present invention;

[0070] FIGS. 39A-39B are flowcharts illustrating the options in procedures of collecting liquid samples and testing liquid samples in accordance with exemplary embodiments of the present invention;

[0071] FIG. 39C is a flowchart illustrating the options to filter rare cells when collecting blood using vacutainers in accordance with exemplary embodiments of the present invention;

[0072] FIG. 40A is a flowchart illustrating a filtration process using a microfilter in accordance with exemplary embodiments of the present invention; and

[0073] FIG. 40B is a flowchart illustrating a filtration process using a microfilter in accordance with exemplary embodiments of the present invention.

[0074] FIG. 40C is a flowchart illustrating the process of performing some types of the assays of the cells collected on the microfilter in the filter holder in accordance with exemplary embodiments of the present invention;

DETAILED DESCRIPTION OF EXEMPLARY EMBODIMENTS

[0075] Referring now to the drawings, wherein like reference numerals designate identical or corresponding parts throughout the several views, embodiments of the present invention are shown in schematic detail.

[0076] The matters defined in the description such as a detailed construction and elements are nothing but the ones provided to assist in a comprehensive understanding of the invention. Accordingly, those of ordinary skill in the art will recognize that various changes and modifications of the embodiments described herein can be made without departing from the scope and spirit of the invention. Also, well-known functions or constructions are omitted for clarity and conciseness. Some exemplary embodiments of the present invention are described below in the context of commercial applications. Such exemplary implementations are not intended to limit the scope of the present invention, which is defined in the appended claims

[0077] Aspects of the present invention are generally directed to a microfilter comprising a polymer layer formed from an epoxy-based photo-definable dry film. The microfilter includes a plurality of apertures each extending through the polymer layer. In certain embodiments, the microfilter may be formed by exposing the dry film to energy through a mask and developing the exposed dry film. In some embodiments, the dry film may be exposed to energy in the form of ultraviolet (UV) light. In other embodiments, the dry film may be exposed to energy in the form of X-rays. In certain embodiments, the polymer layer has sufficient strength and flexibility to filter liquid. In some embodiments, the apertures are sized to allow passage of a first type of bodily fluid cell and to prevent passage of a second type of bodily fluid cell.

[0078] Specifically, in certain embodiments, the microfilter may be used to perform assays on bodily fluids. In some embodiments, the microfilter may be used to isolate and detect large rare cells from a bodily fluid. In certain embodiments, the microfilter may be used to collect circulating tumor cells (CTCs) from peripheral blood from cancer patients passed through the microfilter. In certain embodiments, the microfilter may be used to collect circulating endothelial cells, fetal cells and other large cells from the blood and body fluids. In certain embodiments, the microfilter may be used to collect large cells from processed tissue samples, such as bone marrows. In some embodiments, cells collected using the microfilter may be used in downstream processes such as cell identification, enumeration, characterization, culturing, etc.

[0079] More specifically, in certain embodiments, multiple layers of epoxy-based photo-definable dry film may be exposed to energy simultaneously for scaled production of microfilters. In some embodiments, a stack of epoxy-based photo-definable dry film layers is provided, and all of the dry film layers in the stack are exposed to energy simultaneously. In some embodiments, a dry film structure including epoxy-based photo-definable dry film disposed on a substrate is provided in the form of a roll. In such embodiments, a portion of the structure may be unrolled for exposure of the dry film to energy. In certain embodiments, portions of a plurality of rolls may be exposed to energy simultaneously.

[0080] FIGS. 1A-1E are cross-sectional views illustrating multiple stages in a process for manufacturing a microfilter 120 in accordance with embodiments of the present invention. FIG. 2A is a flowchart illustrating a process 200 for manufacturing a microfilter in accordance with embodiments of the present invention. The exemplary process of FIG. 2A will be described below with reference to FIGS. 1A-1E. Other embodiments of the process illustrated in FIG. 2A will be described below with reference to FIGS. 3A-8B.

[0081] At block 220 of FIG. 2A, a layer of epoxy-based photo-definable dry film 100 (which may be referred to herein as “dry film 100”) disposed on a substrate 180 is provided. In some embodiments, dry film 100 is laminated on substrate 180 at block 220. In certain embodiments, a silicon wafer is coated with a thin layer of metallic material, such as copper, and dry film 100 is laminated on the metallic material at block 220. In other embodiments, a dry film 100 with a substrate 180 already attached may be obtained and provided at block 220. As used herein, an “epoxy-based photo-definable” substance refers to a substance including or formed from a photo-definable epoxy resin, such as a polyfunctional epoxy resin, bisphenol A epoxy resin, epoxidized polyfunctional bisphenol A formaldehyde novolac resin, etc. Examples of photo-definable epoxy resins may be found in U.S. Pat. Nos. 7,449,280, 6,716,568, 6,391,523, and 6,558,868, the contents of which are hereby incorporated by reference herein. Examples of photo-definable epoxy resins may also be found in U.S. Patent Publication Nos. 2010/0068648 and 2010/0068649, the contents of which are hereby incorporated by reference herein. As used herein, an “epoxy-based photo-definable dry film” is a dry film including or formed from an epoxy-based photo-definable substance. Examples of epoxy-based photo-definable dry films that may be used in accordance with embodiments of the present invention may be found in U.S. Pat. Nos. 7,449,280, 6,391,523, and 6,558,868, and U.S. Patent Publication Nos. 2010/0068648 and 2010/0068649. Formulations of epoxy-based photo-definable dry films are not limited to those described here.

[0082] The liquid resist form of the epoxy-based photo-definable dry films before they are coated on substrate can be spin coated on substrate and dried to obtain the dry film on the substrate.

[0083] In certain embodiments, substrate 180 is a thin copper foil. In some embodiments, smooth substrates are preferable because irregularities in the surface of the substrate to which the dry film is laminated are transferred to a surface of the dry film. In some embodiments, a thin copper film is preferred as a substrate so that the substrate may be removed in a relatively short amount of time. In other embodiments, substrate 180 may be a silicon wafer, a polyimide film such as Kapton, or any other suitable material.

[0084] As shown in FIGS. 1B and 1C, dry film 100 is exposed to energy through a mask 199 to form an exposed dry film 110 at block 240. In the embodiment illustrated in FIGS. 1B and 1C, dry film 100 is exposed to energy in the form of ultraviolet (UV) light through an optical mask 199 having a mask portion 197 that is transparent to UV light and a mask pattern 198 formed from a thin film of material that is opaque to UV light. In alternative embodiments, dry film 100 may be exposed to X-rays through an X-ray mask at block 240, instead of being exposed to UV light through optical mask 199.

[0085] In the embodiment illustrated in FIGS. 1A-1E, epoxy-based photo-definable dry film 100 is a negative resist. As used herein, a “negative resist” is a photo-definable substance that becomes polymerized when exposed to certain kinds of energy, such as UV light or X-rays. Examples of negative resist epoxy-based photo-definable dry films that may be used in accordance with embodiments of the present invention.

[0086] As shown in FIG. 1C, the portions of exposed dry film 110 that were exposed to UV light through mask 199 become polymerized, leaving portions 116 that are not poly-

merized. The polymerized and non-polymerized portions of exposed dry film **100** form a pattern **118** that is defined by optical mask **199**. More particularly, pattern **118** of exposed dry film **110** is defined by a pattern **198** of optical mask **199**, where pattern **198** is formed by material opaque to UV light. In certain embodiments, pattern **198** may be formed by a thin film of material opaque to UV light, such as a thin film of chromium.

[0087] In alternative embodiments, a positive epoxy-based photo-definable dry film may be used instead of a negative dry film. In such embodiments, the process for forming a microfilter from the positive dry film is similar to the process for forming a microfilter described in relation to FIGS. **1A-2A**, except that a different mask may be used, as described below in relation to FIGS. **4A-4D**. As used herein, a “positive resist” is a photo-definable substance in which polymeric bonds are broken when the substance is exposed to certain kinds of energy, such as UV light or X-rays. In certain embodiments, the positive resist may be a resist based on polydimethylglutarimide (such as PMGI, LOR available from MicroChem), an acetate and xylene free resist (such as an S1800® series resist available from Shipley Corp.), or another type of positive resist. For resist layers that are greater than a few microns in thickness, negative resists are generally much more sensitive than positive resists. Most polymer resists belong to the category of positive resist films. Examples of dry film positive resists that may be used include polymethylmethacrylate (PMMA), and a synthetic polymer of methyl methacrylate. Other examples of positive resists are acrylics, polyimide, polyesters, such as polyethylene terephthalate (PET) (MYLAR™), etc. In certain embodiments, a microfilter may be formed from a photo-definable dry film that is not epoxy based, in accordance with embodiments of the present invention. In such embodiments, the dry film may be a positive or a negative resist. In other embodiments, a microfilter may be formed from a photo-definable liquid resist, rather than a dry film. In such embodiments, the photo-definable liquid resist may be a positive resist or a negative resist. In certain embodiments, the photo-definable liquid resist may be liquid polyimide. In such embodiments, the photo-definable liquid polyimide may be positive resist or negative resist. The liquid resist is spin coated on the substrate and dried to form the dry film on the substrate.

[0088] At block **260**, a microfilter **120** having a plurality of apertures **122** extending through the microfilter is formed from exposed dry film **110**. In certain embodiments, microfilter **120** includes a polymer layer formed from epoxy-based photo-definable dry film and a plurality of apertures extending through the polymer layer. In each of the embodiments of the present invention described herein, a microfilter includes one or more polymer layers and one or more apertures extending through each of the one or more polymer layers. Also, as used herein, an “aperture” refers to any type of passage, pore, trench, gap, hole, etc., that extends between outer surfaces of a layer or other structure. In the embodiment illustrated in FIGS. **1A-1E**, apertures **122** are pores **122**.

[0089] In the embodiment illustrated in FIGS. **1A-1E**, exposed dry film **110** is developed to remove non-polymerized portions **116** to form microfilter **120** having pores **122**. In certain embodiments, exposed dry film **110** is developed by applying a developer to dry film **110** to dissolve non-polymerized portions **116**. In some embodiments, the developer is an aqueous solution that dissolves non-polymerized portions **116** when exposed dry film **110** is submerged in the devel-

oper. At block **280**, microfilter **120** having pores **122** is removed from substrate **180** to form a free-standing microfilter **120**, as shown in FIG. **1E**. In some embodiments, a microfilter is a structure of one or more polymer layers including one or more apertures extending between outer surfaces of the structure, wherein the structure has sufficient strength and flexibility to filter a liquid passed through the one or more apertures. In certain embodiments, a microfilter may include apertures having dimensions small enough to prevent one or more types of bodily fluid cells from passing through the apertures when a bodily fluid or liquid containing a bodily fluid is passed through the filter, wherein the dimensions of the apertures are also small enough to prevent one or more other types of bodily fluid cells from passing through the filter. As used herein, “bodily fluid cell” refers to any cell that may be found in a bodily fluid of a patient, such as red or white blood cells, large rare cells, such as CTCs and fetal cells, circulating endothelial cells, etc. One can also obtain tumor cells from bone marrow after process to remove bone fragments and result in liquid form. In some embodiments, a microfilter includes apertures sized to allow passage of a significant number of red blood cells and to prevent passage of a significant number of CTCs. In certain embodiments, a microfilter formed from one or more layers of epoxy-based photo-definable dry film may be a polymeric microfilter.

[0090] Various embodiments of the process illustrated in FIG. **2A** will be described below in relation to FIGS. **2B-8B**. As noted above, in certain embodiments, substrate **180** may be copper foil. In such embodiments, copper substrate **180** may be removed from microfilter **120** using nitric acid, ferric chloride or another well-known reagent in one variation of block **280**. In certain embodiments, the reagent may be used to etch away copper substrate **180** in order to remove it from microfilter **120**. In other embodiments, substrate **180** may be another type of metallic foil, such as aluminum, and may be removed at block **280** by well-known methods.

[0091] FIG. **2B** is a flowchart illustrating a process for forming a microfilter from an exposed dry film at block **260** of FIG. **2A** in accordance with embodiments of the present invention. In certain embodiments, the process **260** for forming the microfilter comprises forming, from the exposed dry film, a polymer layer comprising a plurality of apertures. In the embodiment of FIG. **2B**, a post bake process is performed on exposed dry film **110** disposed on substrate **180** at block **262**. In certain embodiments, the post bake process includes exposing dry film **110** to a relatively high temperature to post bake dry film **110**. At block **264**, dry film **110** is developed by applying a developer to dry film **110**, as described above in relation to FIGS. **1A-1E**. At block **266**, a hard bake process is performed on the developed dry film **110**. In certain embodiments, the hard bake process includes exposing dry film **110** to a relatively high temperature. In some embodiments of the process illustrated in FIG. **2B**, the hard bake process of block **266** may be omitted. In such embodiments, microfilter **120** is formed by post baking the exposed dry film **110** at block **262**, and developing dry film **110** at block **264**. The processes described above in relation to FIG. **2B** may be used with any of the embodiments described herein. Additionally, in any of the embodiments of the present invention described here, the process for forming a polymer layer of a microfilter from an epoxy-based photo-definable dry film may include exposing the dry film to energy, performing a post-bake process, developing the exposed dry film, and/or post baking the exposed dry film, as described above.

[0092] FIGS. 3A-3E are cross-sectional views illustrating multiple stages in a process for manufacturing microfilter 120 in accordance with embodiments of the present invention. In the embodiment illustrated in FIGS. 3A-3E, the substrate is a polyimide film 181. At block 220, a layer of epoxy-based photo-definable dry film 100 disposed on polyimide film 181 is provided with a separator 182 disposed between a portion of dry film 100 and polyimide film 181. In the embodiment shown in FIG. 3A, a separator 182 is disposed between a portion of dry film 100 and polyimide film 181 along an edge of dry film 100. In certain embodiments, separator 182 may be disposed along one or more edges of dry film 100, or at other locations between dry film 100 and polyimide film 181. Separator 182 may be formed from a polyimide film (such as a KAPTON film) or from any other suitable material that can be laminated to dry film 100 and withstand the temperature of a hard bake process.

[0093] As shown in FIGS. 3B-3C, dry film 100 is exposed to energy, and a microfilter 120 is formed from exposed dry film 110 at blocks 240 and 260, as described above in relation to FIGS. 1B-1D. In certain embodiments, microfilter 120 includes a polymer layer comprising a plurality of apertures there through. In the embodiment illustrated in FIGS. 3A-3E, microfilter 120 is removed from polyimide film 181 by grasping an exposed end of separator 182 and using separator 182 to peel microfilter 120 from polyimide film 181. After dry film 100 is removed from polyimide layer 181, as shown in FIG. 3D, separator 182 is removed from dry film 100 to obtain a free-standing microfilter 120, as shown in FIG. 3E. Removing microfilter 120 from layer 181, and removing separator 182 from microfilter 120, are two steps performed in one variation of block 280, in accordance with embodiments of the present invention.

[0094] In alternative embodiments of the process illustrated in FIG. 2A, a liquid resist may be used instead of dry film 100. In such embodiments, a substrate is coated with a thin layer of a metallic substance and an epoxy-based liquid photoresist is spin coated onto the metallic substance to provide a layer of an epoxy-based photo-definable substance on a substrate, in one variation of block 210. In certain embodiments, the substrate may be a silicon wafer, the metallic substance may be copper, and the liquid photoresist may be an epoxy-based photo-definable liquid. In some embodiments, the epoxy-based photo-definable liquid is a liquid negative resist, such as SU-8. The layer of the epoxy-based photo-definable substance is exposed to energy, and a microfilter 120 is formed from the exposed layer at blocks 240 and 260, as described above in relation to FIGS. 1B-1D. In one variation of block 280, microfilter 120 is released from the substrate by etching away the metallic substance via conventional processes as described above. In certain embodiments, the liquid resist may be a liquid negative resist, such as SU-8 or KMPR® available from MicroChem Corp.

[0095] In other alternative embodiments of the process illustrated in FIG. 2A, a liquid negative resist may be used instead of dry film 100, and a positive resist between the negative resist and the substrate may be used as a release layer. In such embodiments, to provide a layer of an epoxy-based photo-definable substance on a substrate in one variation of block 210, a liquid positive resist is spin coated on a substrate, the positive resist is exposed to energy (such as UV light) at the appropriate dose for the thickness of the coating, and a liquid, epoxy-based negative resist is spin coated on the positive resist. In certain embodiments, the positive resist

may be exposed to energy without the use of a mask. The layer of the epoxy-based photo-definable substance is exposed to energy, and a microfilter 120 is formed from the exposed layer at blocks 240 and 260, as described above in relation to FIGS. 1B-1D. In such embodiments, in one variation of block 280, microfilter 120 is released from the substrate by developing the positive resist. In certain embodiments, the same developer may be used to develop both the positive and the negative resists. In other embodiments, one developer may be used to form the pores in microfilter 120, and another developer may be used to release the microfilter from the substrate. In alternative embodiments, a dry film positive resist may be used as the release layer instead of a liquid positive resist. Examples of dry film positive resists that may be used include polymethylmethacrylate (PMMA), and a synthetic polymer of methyl methacrylate.

[0096] In other embodiments, a negative epoxy-based photo-definable dry film 100 may be used in combination with a positive resist release layer. Such embodiments are similar to the embodiments described above utilizing a positive resist release layer, except that a layer of the negative dry film 100 may be laminated on the spin coated positive resist at block 210, rather than spin coating a liquid negative resist.

[0097] FIGS. 4A-4D are cross-sectional views illustrating multiple stages in a process for manufacturing a microfilter 420 in accordance with embodiments of the present invention. In the embodiment illustrated in FIGS. 4A-4D, a layer of positive, epoxy-based photo-definable dry film 400 (which may be referred to herein as "dry film 400") disposed on substrate 180 is provided at block 220. In certain embodiments, positive dry film 400 is laminated on substrate 180 at block 220. As shown in FIG. 4B, positive dry film 400 is exposed to energy at block 240, as described above in relation to FIGS. 1B and 1C, except that, rather than exposed portions of dry film 400 becoming polymerized, polymeric bonds of dry film 400 are broken at portions 416, which are exposed to energy (e.g., UV light) through mask 499. A pattern 418 of exposed portions 416 and unexposed portions is formed in exposed dry film 410. As shown in FIG. 4B, mask 499 includes a transparent portion 497 and an opaque portion 498. As described above, mask 199 of FIG. 1B is used with a negative resist and is configured to cover portions of dry film 100 where pores will be formed in dry film 100. In the embodiment illustrated in FIGS. 4A-4D, opaque portions 498 of mask 499 are configured to cover all portions of positive dry film 400 except the locations where apertures will be formed, allowing UV light to pass through mask 499 to positive dry film 400 at locations where apertures are to be formed in positive dry film 400.

[0098] In the embodiment illustrated in FIGS. 4A-4D, a microfilter 420 is formed from exposed dry film 410 in one variation of block 260, by developing dry film 410 using a developer that dissolves the portions 416 of dry film 400 where the polymeric bonds were broken. At block 280, microfilter 420 having apertures 422 is removed from substrate 180 to form a free-standing polymeric microfilter 420, as shown in FIG. 4D. In some embodiments, microfilter 420 includes a polymer layer formed from epoxy-based photo-definable dry film and includes a plurality of apertures extending through the polymer layer. In certain embodiments, the apertures 422 are pores 422. In some embodiments, at block 280, microfilter 420 may be released from the substrate by developing the positive resist, as described above.

[0099] FIGS. 5A-5D are cross-sectional views illustrating multiple stages in a process for manufacturing a plurality of microfilters from a plurality of layers of epoxy-based photo-definable dry film in accordance with embodiments of the present invention. In certain embodiments of the process illustrated in FIG. 2A, a plurality of dry film structures 501, each including a layer of epoxy-based photo-definable dry film 500 (which may be referred to herein as "dry film 500") disposed on a substrate 580, are provided at block 220. In the embodiment illustrated in FIGS. 5A-5D, dry films 500 disposed on substrates 580 are provided at block 220 by stacking structures 501 on a support 590, as illustrated in FIG. 5A.

[0100] In certain embodiments, as illustrated in FIG. 5A, dry films 500 in the stack of structures 501 are simultaneously exposed to energy in the form of X-rays through an X-ray mask 599 at block 240 of FIG. 2A. In some embodiments, the penetration of X-rays is much deeper than UV light. Unlike UV light, X-rays do not diverge within a material having a thickness of less than 5 mm, even for features significantly smaller than one micron. In some embodiments, X-ray lithography may be typically performed on a beamline of a synchrotron. In addition, X-ray lithography can be used for both negative and positive resists. In the embodiment illustrated in FIGS. 5A-5D, dry films 500 are each negative resists. In other embodiments, dry films 500 may be positive resists. In such embodiments, a mask configured to form apertures in a positive resist, as described above in relation to mask 499 of FIG. 4B, may be used. Additionally, in embodiments in which dry films 500 are positive resists, dry films 500 may be attached to support 290 and stacked directly on one another, without each being disposed on a respective substrate.

[0101] As shown in FIG. 5B, the portions of each dry film 500 exposed to X-rays through mask 599 become polymerized, leaving portions 516 of dry film 500 that are not polymerized. The polymerized and non-polymerized portions of each dry film 500 form a pattern 518 that is defined by pattern 598 of optical mask 599. In some embodiments, mask 599 includes an X-ray transparent portion 597, and a pattern 598 configured to substantially block X-rays. In certain embodiments, pattern 598 is formed from gold. In addition, in some embodiments, X-ray transparent portion 597 may be a thin graphite sheet or a silicon wafer. In the embodiment illustrated in FIGS. 5A-5D, each of substrates 580 transmits most of the X-ray energy applied to it. In certain embodiments, substrates 580 are formed from a metallic foil. In such embodiments, when the foil is sufficiently thin, each substrate 580 will transmit most of the X-ray energy applied to it. In some embodiments, the number of structures 501 that may be stacked and then exposed simultaneously is based on the reduction in the dose of the X-rays caused by the X-rays passing through the metallic foil.

[0102] In certain embodiments, in one variation of block 260, the plurality of exposed dry films 510 are developed to form a plurality of microfilters 520 each having apertures 522 in a manner similar to that described above in relation to FIGS. 1A-1E. In certain embodiments, apertures 522 are pores 522. In some embodiments, the process illustrated in FIGS. 5B and 5C may be performed in one variation of block 260. In such embodiments, structures 501 are separated from one another, as shown in FIG. 5B, and a post bake procedure is performed on exposed dry films 510 disposed on respective substrates 580 in one variation of block 262. In certain embodiments, each of exposed dry films 510 is developed, as described above, in one variation of block 264 to form pores

522 in each of dry films 500, as shown in FIG. 5C. In some embodiments, a hard bake procedure is performed on dry films 510 disposed on respective substrates 580, in one variation of block 266, to form microfilters 520 having pores 522. In other embodiments, the hard bake procedure may be omitted. In certain embodiments, in one variation of block 280, substrates 580 are chemically removed from microfilters 520, as described above, to obtain free-standing microfilters 520 having pores 522, as shown in FIG. 5D. In certain embodiments, each of microfilters 520 is a polymer layer including apertures 522.

[0103] As described above, in certain embodiments, each of substrates 580 may be formed from a metallic foil. In alternative embodiments, substrate 580 may be a polymer based substrate that transmits most of the X-rays applied to it, and which has a melting point higher than a post bake temperature for dry film 500. For example, in certain embodiments, substrate 580 may be formed from a positive resist. In such embodiments, substrate 580 may be exposed to energy, such as UV light or X-rays, sufficient to break polymeric bonds in the positive resist such that substrate 580 may be removed chemically by a developer solution at block 280 of FIG. 2A. In other embodiments, substrate 580 may be a polyimide film and may be removed at block 280 by peeling the polyimide substrate 580 from microfilter 520.

[0104] In alternative embodiments, layers of epoxy-based photo-definable dry film 500 (which may be referred to as "dry films 500") may be stacked and simultaneously exposed without each of the layers being disposed on a respective substrate. In such embodiments, in one variation of block 220, dry films 500 are stacked on a support 590 without substrates disposed between adjacent dry films 500. The stacked dry films 500 are exposed in one variation of block 240. In some embodiments, the process illustrated in FIG. 2B may be performed at block 260. In such embodiments, the exposed dry films 510 may be separated and placed on separate substrates on which exposed dry films 510 undergo a post bake process in one variation of block 262. In such embodiments, the substrates used are able to withstand the post bake temperature and can be dissolved by water or one or more chemicals. While attached to respective substrates, exposed dry films 510 are developed at block 264 and may undergo a hard bake procedure at block 266. At block 280, the substrates 580 are removed from the microfilters 520 formed from exposed dry films 510.

[0105] In certain embodiments, structures 501 may be attached to support 590 using adhesive, a clamp, or any other suitable mechanism or method. In some embodiments, structures 501 are held to a support by an electrostatic chuck. FIGS. 6A and 6B are cross-sectional views illustrating multiple stages in a process for attaching dry film structures 501 to a support using an electrostatic chuck apparatus 600 in a process for forming microfilters in accordance with embodiments of the present invention. In the embodiment illustrated in FIGS. 6A and 6B, a plurality of dry film structures 501, each including a layer of epoxy-based photo-definable dry film 500 disposed on a substrate 580, are provided stacked on a support 690, as shown in FIG. 6A. As shown in FIG. 6A, support 690 includes a water cooling frame 692 with a duct 693, an insulator 664 disposed on frame 692, and a conductive layer 662 disposed on insulator 664. Additionally, a transparent conductive layer 660 is placed on the stack of structures 501 such that the stack of structures 501 is disposed between conductive layers 660 and 662, as shown in FIG. 6A.

Also as shown in FIG. 6A, a circuit connecting the conductive layers is open so that a voltage 665 of zero is applied to the conductive layers.

[10106] As shown in FIG. 6B, closing the circuit between the conductive layers and applying a non-zero voltage 665 between conductive layers 660 and 662 causes apparatus 600 to press together structures 501 between conductive layers 660 and 662. With structures 501 pressed together by apparatus 600, X-rays may be applied to dry films 500 through X-ray mask 599, as described above in relation to FIGS. 5A-5D. In certain embodiments, stacking structures 501 on apparatus 600 and pressing together structures 501 using apparatus 600, as described above in relation to FIGS. 6A and 6B, may be performed in one variation of block 220 of FIG. 2A. While the use of an electrostatic chuck apparatus has been described above in relation to forming microfilters from dry films each disposed on a respective substrate, in alternative embodiments, an electrostatic chuck may similarly be used to press together a stack of free standing polymeric films, such as free standing dry films, that are not each attached to a respective substrate. Such free standing dry films may be stacked and pressed together during a process for forming a plurality of microfilters from the plurality of dry films.

[10107] FIGS. 7A and 7B are cross-sectional views illustrating multiple stages in a process for manufacturing microfilters from a roll of a dry film structure in accordance with embodiments of the present invention. In the embodiment illustrated in FIGS. 7A and 7B, a dry film structure 785 is provided in the form of a roll 702 of the dry film structure 785. Dry film structure 785 includes a layer of epoxy-based photo-definable dry film 700 (which may be referred to herein as "dry film 700") disposed on a removable substrate 782. In some embodiments, substrate 782 may be a chemically dissolvable metallic foil. In certain embodiments, the metallic foil may include aluminum or copper, which may be etched away as described above. In the embodiment illustrated in FIGS. 7A and 7B, a portion of roll 702 is disposed on a roller 774 and another portion is disposed on a roller 775. A working portion 787 of dry film structure 785 extends between rollers 774 and 775 and is held substantially flat by rollers 770 for exposure to energy through mask 199.

[10108] In certain embodiments of the process illustrated in FIG. 2A, a layer of epoxy-based photo-definable dry film 700 disposed on a substrate 782 is provided, in one variation of block 220, by unrolling a portion of dry film structure 785 from roll 702 and advancing the portion of structure 785 in the direction of arrow 772 to provide working portion 787 of structure 785 between support 791 and mask 199. In some embodiments, the working portion 787 provided at block 220 includes a portion of dry film 700 that has not been patterned by exposure to energy through a mask. In certain embodiments, support 791 and mask 199 are moved away from structure 785 when structure 785 is advanced.

[10109] In the embodiment illustrated in FIGS. 7A and 7B, mask 199 and support 791 are moved adjacent to structure 785, and dry film 700 is exposed to energy through mask 199, as shown in FIG. 7B, in one variation of block 240 of FIG. 2A. In the embodiment shown in FIGS. 7A and 7B, mask 199 is an optical mask and the energy is UV light, although a different type of energy may be used along with a different mask, as described above. In the embodiment illustrated in FIGS. 7A and 7B, dry film 700 is a negative resist. In other embodiments, dry film 700 may be a positive resist. In such embodiments, a mask configured to form pores in a positive resist, as

described above in relation to mask 499 of FIG. 4B, may be used. Exposing dry film 700 to energy through the mask forms a pattern in dry film 700, as described above in relation to other embodiments. In certain embodiments, support 791 presses against structure 785, as shown in FIG. 7B, to stretch dry film 700 for the exposure process to thereby provide additional tension and stability to dry film 700 during the exposure process. In some embodiments, after exposing dry film 700, structure 785 may be advanced again as described above, to provide a new working portion 787 that has not yet been exposed at block 220, and the new working portion 787 may be exposed at block 240, as described above. In certain embodiments, this process of advancing structure 785 and exposing dry film 700 may be continuously repeated. In some embodiments, the process may be repeated until most or all portions of dry film 700 have undergone an exposure process.

[10110] In some embodiments, a microfilter having apertures is formed from an exposed portion of dry film 700 by developing the exposed portion, as described above in relation to other embodiments, in one variation of block 260. In such embodiments, the exposed portion of dry film 700 may be developed before it is rolled onto roller 775, or may be developed after all desired portions of dry film 700 have been exposed. In some embodiments, the process illustrated in FIG. 2B may be performed at block 260. In such embodiments, the exposed portion of dry film 700 may be advanced through an oven for a post bake procedure at block 262, the exposed portion of dry film 700 may be developed at block 264, and then undergo a hard bake procedure at block 266. In other embodiments, the procedures at blocks 262, 264 and 266 may be performed after all desired portions of dry film 700 have been exposed. In certain embodiments, the hard bake procedure may be omitted.

[10111] In some embodiments, after developing the exposed dry film 700, substrate 782 is removed at block 280, as described above in relation to other embodiments. In certain embodiments, after removing substrate 782, individual microfilters are diced from the roll of dry film from which the microfilters were formed. In certain embodiments, forming microfilters from a dry film provided as a roll may simplify the manufacture of microfilters in accordance with embodiments of the present invention, and may allow automation of the manufacturing process.

[10112] FIGS. 8A and 8B are cross-sectional views illustrating multiple stages in a process for manufacturing microfilters from a plurality of rolls of dry film structures in accordance with embodiments of the present invention. The embodiment illustrated in FIGS. 8A and 8B is similar to the embodiment illustrated in FIGS. 7A and 7B, except that layers of epoxy-based photo-definable dry film 700 of multiple rolls 702 are exposed to energy simultaneously. In such embodiments, a plurality of layers of epoxy-based photo-definable dry film 700 each disposed on a respective substrate 782 are provided, in one variation of block 220, by advancing the structure 785 of each roll 702 in the direction of arrow 772 to provide a stack 887 of structures 785 between support 891 and mask 899. In the embodiment illustrated in FIGS. 8A and 8B, mask 899 and support 891 are moved adjacent to the stack 887, and the portions of dry films 700 in stack 887 disposed between mask 899 and support 891 are exposed to energy simultaneously through mask 899, as shown in FIG. 8B, in one variation of block 240 of FIG. 2A. In the embodiment illustrated in FIGS. 8A and 8B, dry films 700 are each negative resists. In other embodiments, dry films 700 may be

positive resists. In such embodiments, a mask configured to form apertures in a positive resist, as described above in relation to mask 499 of FIG. 4B, may be used. Further processes for forming microfilters from dry films 700 are similar to the processes described above in relation to the embodiment illustrated in FIGS. 7A and 7B.

[0113] In the embodiment illustrated in FIGS. 8A and 8B, support 891 is disposed on a water cooling frame 692 including a duct 693. In addition, as shown in FIG. 8B, working portions 887 of structure 785 may be securely held in place between support 891 and mask 899 by a clamp 860. In alternative embodiments, stack 887 may be held secure using an electrostatic chuck, as described above in relation to other embodiments. In some embodiments, the number of dry films 700 exposed simultaneously may be determined based on the precision yielded when exposing a stack of a particular number of films. In certain embodiments, forming microfilters from a plurality of dry films provided as a plurality of rolls, as described above, may simplify the manufacture of microfilters and/or facilitate high volume production of microfilters.

[0114] In embodiments in which non-epoxy-based dry films are used, the dry films may be provided in roll form without a substrate. In such embodiments, each roll 702 includes only the dry film and not any substrate. In embodiments in which epoxy-based dry films are used, each roll 702 may include an additional cover layer on dry film 700. In such embodiments, substrate 782 is disposed on a first side of dry film 700 and the cover layer on the opposite side of dry film 700. In certain embodiments, lithography-based microfabrication in accordance with embodiments of the present invention may enable efficient mass production of highly uniform precision microfilters. In certain embodiments, fabricating microfilters in accordance with embodiments of the present invention may yield increased porosity and pore uniformity in the microfilters produced.

[0115] FIGS. 9A-9D are partial top views illustrating various microfilters aperture distributions in accordance with embodiments of the present invention. In certain embodiments, microfilters having different aperture sizes, shapes and distributions may be provided. In some embodiments, certain combinations of aperture size, shape and distribution may be more advantageous than others for a particular application of a microfilter. For example, for the microfiltration of rare cells, such as circulating tumor cells and fetal cells in blood, a microfilter having round pores each having a diameter of 7-8 microns may be preferable in certain embodiments. In some applications, a microfilter having round pores each with a diameter of 7-8 microns can trap the rare cells while retaining a very small percentage of blood cells.

[0116] In the embodiments illustrated in FIGS. 9A and 9B, microfilters 910 and 912 each have a uniform distribution of pores 920 and 922, respectively. Additionally, pores 920 are uniform in size, as are pores 922. In the embodiment illustrated in FIG. 9C, microfilter 914 includes uniform rectangular pores 924, distributed over microfilter 914 in several groupings of pores 924. In the embodiment illustrated in FIG. 9D, microfilter 916 a plurality of pores 926 of a first size, and a plurality of pores 928 of a second size. In other embodiments, any or pores 920, 922, 924, 926 and 928 may be any other type of aperture. Any of microfilters 910, 912, 914 and 916 may be manufactured using any of the microfilter manufacturing processes described above in accordance with embodiments of the present invention. Additionally, any of the microfilter manufacturing processes described above in

accordance with embodiments of the present invention may be used to form apertures of a plurality of different cross-sectional shapes. For example, in certain embodiments, apertures may be formed which have the cross-sectional shape of a circle, triangle, square, rectangle, ellipse, oval, trapezoid, parallelogram, etc.

[0117] Microfilters having various thicknesses and various aperture shapes, sizes and distributions may be manufactured in accordance with embodiments of the present invention described herein. FIGS. 10A-10E are cross-sectional views illustrating microfilters having various thicknesses and various aperture shapes, sizes and distributions in accordance with embodiments of the present invention. FIGS. 10A and 10B show microfilters 1010 and 1012, each formed via one of the processes described above in accordance with embodiments of the present invention. Microfilter 1010 includes a plurality of pores 1020, each having a width 1040. Microfilter has a thickness 1030 substantially perpendicular to the width 1040 of pores 1020. In the embodiment illustrated in FIG. 10A, thickness 1030 is not significantly larger than width 1040. In certain embodiments, it is preferable that the thickness of the microfilter is on the same order as the width of one or more pores of the microfilter in order to reduce the pressure required to pass a liquid sample through the pores. In some applications, if the thickness of a microfilter is significantly greater than the width of some or all of the pores, a much larger amount of pressure may be applied to the microfilter to pass a liquid sample through a microfilter than if the microfilter has a thickness on the same order as some or all of the pores. Passing the liquid sample through the filter with a relatively large amount of pressure may distort the shape of one or more pores, or risk breaking the microfilter.

[0118] For example, for the microfiltration of rare cells, such as CTCs and fetal cells in blood, a microfilter having a thickness of 8-14 microns may be preferable in certain embodiments. In certain embodiments, a microfilter for such an application may have pores each having a diameter of 7-8 microns and a thickness of 8-14 microns. In other embodiments, a microfilter for such applications may include a rectangular aperture having a width of between 5-7 microns and a length greater than 7 microns, wherein the length and width of the aperture are both substantially perpendicular to the thickness of the microfilter. In certain embodiments, the rectangular aperture may be an elongate trench. In certain embodiments, it may be preferred that the width of the apertures in the microfilter are near in size to the thickness of the microfilter. In some embodiments, the thickness of the microfilter is less than ten times the width of some or all the pores. In other embodiments, the thickness of the microfilter is within 10 microns of the width of some or all of the pores. Microfilters formed in accordance with embodiments of the present invention may be used in applications other than capturing circulating tumor cells from blood. In some embodiments, the desired aperture geometry, aperture dimensions, aperture distribution, microfilter materials, microfilter thickness, microfilter size, etc., may vary for different applications. In some embodiments, desired aperture geometry, dimensions, and distribution may be provided by using an appropriate mask, such as an optical or X-ray mask. In certain embodiments, a consideration for microfilters is strength of the material from which the microfilter is made to prevent breakage of the filter material or distortion of the aperture shape.

[0119] As shown in FIG. 10B, microfilter 1012 has a plurality of pores 1022 each having a width 1042. Microfilter 1012 also has a thickness 1032 that is substantially perpendicular to the width 1042 of pores 1022. The thickness 1032 of microfilter 1012 is greater than the thickness of microfilter 1010. In the embodiment illustrated in FIG. 10B, pores 1022 are uniform in size and are each substantially perpendicular to a first surface 1050 and a second surface 1052 of microfilter 1012. In the embodiment illustrated in FIG. 10C, microfilter 1014 has pores 1024 with a first width 1044 and pores 1026 with a second width 1046 that is smaller than the first width 1044. Microfilter 1014 also has a thickness 1034. In the embodiment illustrated in FIG. 10D, microfilter 1016 has pores 1028 with non-uniform cross-sectional shapes. Each pore 1028 has a first opening in a first surface 1054 of microfilter 1016 and a second opening in a second surface 1056 of microfilter 1016. As shown in FIG. 10D, the width 1048 of pore 1028 at the first surface 1054 is greater than the width 1049 of pore 1028 at the second surface 1056. Microfilter 1016 also has a thickness 1036. In the embodiment illustrated in FIG. 10E, microfilter 1018 has pores 1029 with non-uniform cross-sectional shapes. Each pore 1029 has a first opening 1045 in a first surface 1053 of microfilter 1018 and a second opening 1047 in a second surface 1057 of microfilter 1018. As shown in FIG. 10E, the width 1045 of pore 1029 at the first surface 1053 is smaller than the width 1047 of pore 1029 at the second surface 1057. Microfilter 1018 also has a thickness 1038.

[0120] FIGS. 12A-12K are cross-sectional views illustrating multiple stages in a process for manufacturing a multi-layer microfilter 1270 in accordance with embodiments of the present invention. FIG. 12L is a top view of multi-layer microfilter 1270 in accordance with embodiments of the present invention. FIG. 11 is a flow chart illustrating a process 1100 for manufacturing a multi-layer microfilter 1420 in accordance with embodiments of the present invention. FIGS. 14A and 14B are cross-sectional views of a multi-layer microfilter 1420 in accordance with embodiments of the present invention. FIG. 14C is a top view of a multi-layer microfilter 1420 of FIGS. 14A and 14B. The exemplary process of FIG. 11 will be described below with reference to FIGS. 12A-12F and FIGS. 14A-14C. FIGS. 13A and 13B are top views illustrating multiple stages in the process of FIG. 11.

[0121] At block 1120 of FIG. 11, a first microfilter 120 is formed on a substrate 180 from a layer of epoxy-based photo-definable dry film. In the embodiment illustrated in FIGS. 12A-12L, first microfilter 120 may be formed on substrate 180 by a process similar to the process 200 of FIG. 2A, described above, omitting the removal of microfilter 120 from substrate 180 at block 280. In certain embodiments, microfilter 120 comprises a polymer layer including a plurality of apertures. In certain embodiments of process 1100, a mask with a pattern configured for forming a plurality of elongate trenches in the dry film 100 may be used instead of mask 199 with pattern 198 configured for forming a plurality of pores in dry film 100. In such embodiments, the mask may have a pattern including elongate strips of metal so that corresponding elongate trenches may be formed in dry film 100 when dry film 100 is exposed through the mask.

[0122] FIG. 13A is a top view of microfilter 120 formed at block 1120 in accordance with embodiments of the present invention. As shown in FIG. 13A, microfilter 120 includes a plurality of elongate trenches 1222 and is disposed on a

substrate 180 exposed through trenches 1222. FIG. 12A is a cross-sectional view of microfilter 120 taken along line 12A of FIG. 13A and FIG. 12B is a cross-sectional view of microfilter 120 taken along line 12B of FIG. 13A. As shown, line 12B is perpendicular to line 12A.

[0123] At block 1140 of FIG. 11, a layer of epoxy-based photo-definable dry film 1210 (which may be referred to herein as “dry film 1210”) is laminated on microfilter 120, as shown in FIG. 12C. In certain embodiments, dry film 1210 is capable of bridging over features formed in the surface on which it is laminated. In such embodiments, dry film 1210 does not significantly fill trenches 1222 when laminated on microfilter 120. At block 1160, a second microfilter 1230 is formed from the layer of epoxy-based photo-definable dry film 1210, as described below. In certain embodiments, microfilter 1230 comprises a polymer layer including a plurality of apertures. As shown in FIG. 12D, dry film 1210 is exposed to energy through a mask 1290 to form an exposed dry film 1212 having a pattern 1218 of polymerized portions and non-polymerized portions 1216, as described above in relation to block 240 of FIG. 2A. In the embodiment illustrated in FIGS. 12A-12L, dry film 1210 is a negative resist. In other embodiments, dry film 1210 may be a positive resist and a different mask configured for use with a positive resist may be used. In the embodiment illustrated in FIGS. 12A-12L, dry film 1210 is exposed to energy in the form of ultraviolet (UV) light through an optical mask 1290 having a mask portion 1295 that is transparent to UV light and a mask pattern 1293 including a plurality of elongate strips that are opaque to UV light. In alternative embodiments, dry film 1210 may be exposed to X-rays through an X-ray mask instead of being exposed to UV light through optical mask 1290.

[0124] In the embodiment illustrated in FIGS. 12A-12L, a polymeric microfilter 1230 having a plurality of trenches 1232 extending through microfilter 1230 is formed from exposed dry film 1212, as described above in relation to block 260 of FIGS. 2A and 2B, in one variation of block 1160. FIG. 13B is a top view of first and second microfilters 120 and 1230 in accordance with embodiments of the present invention. As shown in FIG. 13B, microfilter 1230 includes a plurality of elongate trenches 1232 and is disposed on first microfilter 120, which is exposed through trenches 1232. As shown in FIG. 13B, trenches 1232 of microfilter 1230 are formed substantially perpendicular to trenches 1222 of microfilter 120. FIG. 12E is a cross-sectional view of microfilters 120 and 1230 taken along line 12E of FIG. 13B, and FIG. 12F is a cross-sectional view of microfilters 120 and 1230 taken along line 12F of FIG. 13B. As shown, line 12F is perpendicular to line 12E. The thickness of each layer can be different.

[0125] In certain embodiments, after forming second microfilter 1230, substrate 180 may be removed from microfilter 120, as described above in relation to block 280 of FIG. 2A, to form a multi-layer microfilter 1420 shown in FIGS. 14A-14C. In the embodiment illustrated in FIGS. 14A-14C, multi-layer microfilter 1420 includes second microfilter 1230 disposed on first microfilter 120. As shown in FIG. 14C, multi-layer microfilter 1420 includes apertures 1240 extending through multi-layer microfilter 1420 where trenches 1222 and 1232 intersect. In certain embodiments, microfilter 1240 comprises a polymer layer including a plurality of apertures. FIG. 14A is a cross-sectional view of microfilter 1420 taken along line 14A of FIG. 14C, and FIG. 14B is a cross-sectional

view of microfilter 1420 taken along line 14B of FIG. 14C. As shown, line 14B is perpendicular to line 14A. The thickness of each layer can be different.

[0126] In certain embodiments, as an alternative to forming microfilter 1420, a microfilter 1270 having a non-linear passage 1280 may be formed, as illustrated in FIGS. 12G-12L. In such embodiments, multi-layer microfilter 1270, shown in FIG. 12L, is formed by forming a third microfilter 1240 on first and second microfilters 120 and 1230 and removing substrate 180. Additionally, in such embodiments, after forming second microfilter 1230 on first microfilter 120 and substrate 180, as shown in FIG. 12F, a layer of epoxy-based photo-definable dry film 1215 is laminated on second microfilter 1230, as shown in FIG. 12G. Subsequently, third microfilter 1240 is formed from dry film 1215, as described above in relation to the formation of second microfilter 1230 and the processes of blocks 240 and 260 of FIGS. 2A and 2B. As shown in FIG. 12H, third microfilter 1240 includes a plurality of elongate trenches 1242 that are substantially perpendicular to trenches 1232 and substantially parallel with trenches 1222. Additionally, in certain embodiments, trenches 1242 are offset from trenches 1222 such that trenches 1242 are not directly above trenches 1222, as shown in FIG. 12H.

[0127] In certain embodiments, after forming third microfilter 1240, substrate 180 may be removed from microfilter 120, as described above in relation to block 280 of FIG. 2A, to form multi-layer microfilter 1270. FIG. 12L is a top view of multi-layer microfilter 1270. FIG. 12J is a cross-sectional view of multi-layer microfilter 1270 taken along line 12J of FIG. 12L, and FIG. 12K is a cross-sectional view of multi-layer microfilter 1270 taken along line 12K of FIG. 12L. As shown, line 12K is perpendicular to line 12J.

[0128] As shown in FIG. 12L, multi-layer microfilter 1270 includes non-linear passages 1280 extending through each of microfilters 1240, 1230, and 120 so as to extend from a first surface 1272 to a second surface 1274 (see FIG. 12J) of multi-layer microfilter 1270. In certain embodiments, each non-linear passage 1280 is defined by a first aperture 1282 at an intersection of trenches 1242 and 1232, a second aperture 1284 at an intersection of trenches 1232 and 1222, and a portion of trench 1232 connecting the first and second apertures. In embodiments in which a multi-layer microfilter 1270 has one or more non-linear passages 1280, the filtration path is longer than if multi-layer microfilter 1270 included only linear apertures. For clarity, only selected trenches 1232 and 1222 and selected non-linear passages are illustrated in FIG. 12L. In some embodiments, each non-linear aperture 1280 is interconnected with many other non-linear passages 1280 via trenches 1232.

[0129] In certain embodiments of the multi-layer microfilter 1270, the respective thicknesses of microfilters 120, 1230 and 1240 can be the same or different, the trenches of a microfilter may or may not all have the same size and/or shape, the trenches of different microfilters of the multi-layer microfilter may or may not all have the same size and/or shape. In some embodiments, elongate trenches 1242 may have a width of 5-7 microns and a length greater than 7 microns, wherein the length and the width are both perpendicular to the thickness of the microfilter. Alternatively or in addition, the trenches of thicknesses of microfilters may be non-linear, and the trenches of adjacent microfilters may be oriented at an angle other than 90 degrees with respect to one another. Alternatively or in addition, one or more of microfilters 120, 1230 and 1240 may include pores like any of the

pores illustrated in FIGS. 9A-9D instead of trenches, and, in some embodiments, multi-layer microfilter 1270 may include more than three microfilters disposed on one another. Additionally, in certain embodiments, each of microfilters 120, 1230 and 1240 may be formed from the same type of epoxy-based photo-definable dry film.

[0130] FIG. 15 is a cross-sectional view of a microfiltration structure including a microfilter and a support structure in accordance with embodiments of the present invention. In the embodiment illustrated in FIG. 15, microfiltration structure 1510 includes a microfilter 1520 having pores 1522 and disposed on a support structure 1530 configured to provide structural strength to microfilter 1520. In certain embodiments, support structure 1530 may be integrated with microfilter 1520. In some embodiments, support structure 1530 is a grid support structure. In certain embodiments, microfiltration structure 1510 may be formed by a process similar to the process described above in relation to FIGS. 12A-12F and FIGS. 14A-14C. In such embodiments, microfilter 1520 and support structure 1530 are each formed from a layer of epoxy-based photo-definable dry film and patterned using an appropriate mask. In such embodiments, microfilter 1520 is formed on support structure 1530 or support structure 1530 is formed on microfilter 1520.

[0131] The microfilter can be a combination of pores with other structure elements above or below the layer that form the pores to form many filtration devices. A few examples of two layers or three layers structures with and without pores are illustrated in FIGS. 16A-16J. These devices have applications for isolating cells from body fluids and for biological assays.

[0132] FIG. 16A is an exemplary embodiments of a two layered microfilter, showing a side view of microfilter 2561 containing posts 2575 on microfilter base 2572 with pores 2573. FIG. 16B shows the top view of posts 2575 on microfilter base 2572 with pores 2573. The pore can be circular, square, rectangle, etc. The posts can also of a variety of shapes such as circles, squares, rectangles, etc. The arrangement of the posts and the density can be regular, varied spatially or random as long as the posts are not closer than 30 μm and the posts don't cover the pores. The pores can have various shapes, sizes and distributions. The posts can also have various shapes, sizes and distributions.

[0133] FIG. 16C is another exemplary embodiments of a two layered microfilter, showing the top view of a microfilter 2563 where pore is rectangular shaped 2573, but not distributed everywhere on the base support 2572, and posts 2575 is also not distributed everywhere. The pores can have various shapes, sizes and distributions. The posts can also have various shapes, sizes and distributions.

[0134] FIG. 16I is an exemplary embodiments of a two layered device 2580, showing the side view of bottom layer 2581 without pores. The top layer 2582 forms wells 2583. FIG. 16J shows the top view. This well structure can be fabricated using the same procedure described in FIGS. 1A-1D followed by procedure described in FIGS. 12A-12F except no mask is needed for FIG. 1B.

[0135] Pore dimensions depends on the needs of specific applications. For example, pore diameter of 7-8 microns is the commonly preferred for circulating tumor cells from human blood, while circulating tumor cells originated in mice are smaller than in humans, so smaller pores are preferred for mice studies. For rectangular pores, width of 5-7 μm are

preferred for collecting circulating tumor cells from human blood and the length is not critical as long as the filters don't deform during filtration.

[0136] FIGS. 16D-16F is another exemplary embodiment of a two layered microfilter with different patterns in each layer of the material according to an exemplary implementation. FIG. 16D showing the first side view 2501 of a two layer microfilter with the top layer 2551 and bottom layer 2552. The top layer are of strips with slot opening 2555. FIG. 16E is the second side view 2502 of the two layer microfilter turned 90 degrees. The bottom layer 2552 has open slots 2553. FIG. 16F shows the top view of the microfilter 2500. The cross strips forms effective rectangular pores 2554.

[0137] In an advantageous exemplary implementation, dimensions of the top channel 2555 between two top strips of 2551 are 5-7 μm for isolation of tumor cells. Small width of 10-20 μm for 2551 would allow high filtration rate, but a wide range of width for 2551 are functional. The width of 2552 can vary, but for high filtration rate, the width may be 5-20 μm . The gap between two strips of 2552 that form the pore 2553 can also vary. For high filtration rate, the gap can be from 10-60 μm . The thickness of 2551 about 10 μm may be preferable. In an exemplary implementation, thickness of bottom layer 2552 can advantageously be approximately the same as the gap between two strips of 2552.

[0138] FIGS. 16G-16H are side view and top view of an exemplary device 2540 consisting of microfilter 1541 with pores 2542 at the bottom of wells 2544. The structure shown in FIG. 16H is repeated over the whole microfilter area. An application of the device 2540 is to filter the cells and followed by culture of the cells in the wells 2544. Again the pores can have a variety of shapes and sizes. The size, shape and depth of the well 2544 can also be varied as appropriate for different uses. The density of the wells 2583 can also be varied.

[0139] An exemplary application of devices 2500, 2540, 2561 and 2563 includes rare cell isolation where the mechanism of isolation of the cells is based on size.

[0140] FIGS. 16I-16J are side view and top view of an exemplary device 1580 consisting of wells 2583 formed by 2582 with solid bottom 2581. These can be used for a variety of biological assays. The size, shape and depth of the wells 2583 can be varied. The density of the wells 2583 can also be varied.

[0141] Various materials for forming devices described in FIGS. 16A-16J above can include epoxy-based photo-definable dry films and other types of photo-definable dry films. The methods to fabricate those structures using photo-definable dry films are described above and in, for example, PCT/US11/20966.

Microfiltration Chips

[0142] The steps of filtration of large rare cells from body fluids on the microfilter or concentrated in a reduce volume with reduced contaminants can be performed with minimal human intervention if the microfilter can be installed in a microfiltration device. This microfiltration device contains at least one inlet, at least one outlet, and at least one microfilter. An exemplary microfiltration device having such components can be referred to as a "microfiltration chip". The term "microfiltration chip" is used herein simply as a non-limiting reference label for consistency of description and to facilitate understanding of exemplary implementations. Applications and procedures described herein which reference "microfil-

ters" are also applicable to and may employ microfiltration chips according to exemplary embodiments of the present invention.

[0143] FIGS. 17A-17E illustrate top views of an exemplary implementation where five layers when stacked form a microfiltration chip according to an exemplary embodiment of the present invention. The top layer 2010 is attached to 2020, with inlet 2001 and opening to the microfilter layer 2040 with pores 2004. Layer 2060 with opening for the pores 2004 is below the microfilter 2040 with outlet 2002. Finally, the layer 2060 is attached to a base layer 2070.

[0144] Layers 2010 and 2020 may be formed at the same time with same material. Layers 2060 and 2070 may be formed at the same time with the same material. Layers 2010 and 2070 may be formed with material that is transparent for optical radiation. Layers 2010 and 2070 may be glass. Layers 2020 and 2060 may be photo-definable dry films. These are exemplary implementations according to embodiments of the present invention.

[0145] In another exemplary implementation, if the cells and particles captured on the microfiltration chip are to be analyzed using a microscope, at least the top cover 2010 should be clear and thin, such as a microscope slide cover slip. In yet another exemplary implementation, if a fluorescence analysis is required, the top cover 2010 and the bottom layer 2070 both should be essentially non-fluorescent. In yet another exemplary implementation, bottom layer 2070 can be clear or opaque. In yet another exemplary implementation, base layer 2070 can be the same size as the top four layers, 2010, 2020, 2040, and 2060, or it can be any other size. In yet another exemplary implementation, a size for layer 2070 can be the size of glass slide 25 mm \times 75 mm. In yet another exemplary implementation, the thickness of layer 2020 and 2060 may be $>50 \mu\text{m}$. The dimensions of the inlet 2001, the outlet 2002 may be varied.

[0146] The microfilter can have a few variations without departing from the scope of the embodiments of the present invention. In an exemplary embodiment of the present invention illustrated in FIGS. 18A-18E, the microfilter pores have a rectangular pattern where the short side of the rectangle is in the direction of flow, while in another exemplary embodiment of the present invention illustrated in FIGS. 17A-17E, the long side of the rectangular microfilter pore pattern is in the direction of the flow. The microfilter pores may have a square pattern or other geometric patterns within the scope of the embodiments of the present invention.

[0147] The position of the inlet and outlet can be varied without departing from the scope of the embodiments of the present invention. An exemplary variation is shown in FIGS. 19A-19E, where the inlet and outlet are both on the top. Other inlet and outlet positions, geometry and implementation are also possible within the scope of the embodiments of the present invention.

[0148] FIGS. 20A-20E show an example of an implementation according to an embodiment of the present invention where there are two openings 2301 and 2302 on the same layer 2320. They can be one inlet and one outlet or both inlets. The inlet and outlet can be switched during different times of the assay.

[0149] FIGS. 20A-20E also show another implementation according to an embodiment of the present invention where the inlet 2301 and outlets 2320 and 2360 can be on the sides of the microfiltration chip.

[0150] FIGS. 21A-21G show examples of yet other implementations according to embodiments of the present invention where additional layers 2430 and 2450 can be included without effecting the filtration process. In yet further exemplary implementations, additional layer similar to 2430 can be included between 2410 and 2410, and additional layer similar to 2450 can be included between 2460 and 2470.

[0151] Layers 2420 and 2430, 2410, 2420, and 2430, 2450 and 2460, or 2450, 2460, and 2470 may be formed at the same time, or with same material. Layers 2410 and 2470 may be formed with material that is transparent for optical radiation. Layers 2010 and 2070 may be glass. Layers 2420, 2430, 2450 and 2060 may be photo-definable dry films. These are yet other exemplary implementations according to embodiments of the present invention.

[0152] According to further exemplary embodiments of the present invention, microfilter in any of the microfiltration chips can be a single layer with pores penetrating from the top to the bottom. The pores can be circular, square, rectangle, or any shape that can be obtained by photo lithography that is of useful application.

[0153] According to still further exemplary embodiments of the present invention, microfilter in any of the microfiltration chips can also be double layered. An example of two layered filter with rectangular opening is shown in FIGS. 13A-13B. Other examples of double layered microfilters are shown in FIGS. 15 and 16. The microfilter in any of the microfiltration chip can also be triple layer as shown in FIGS. 14A-14C.

Methods to Perform Microfiltration

[0154] FIG. 22 illustrates a device according to an exemplary embodiment of the present invention that can implement the installation of a microfilter for filtration showing the flow of liquid containing large particulates 2650 and small particulates (not shown) through the pores 2654 in the microfilter 2655 held in a structure 2690 formed either by the microfilter holder or microfiltration chip. According to an exemplary implementation, liquid enters through the input 2680, and exits through the outlet 2681. The liquid flow 2660 enters the microfiltration chip through opening 2680. The liquid flow 2662 through the microfilter 2655. The liquid flow 2681 exits the device through opening 2681. The large particulates 2650 bigger than the pores 2654 are retained on the microfilter 2655.

[0155] FIG. 23 illustrates a device according to an exemplary embodiment of the present invention that can implement the installation of a microfilter for filtration showing the flow of liquid containing large particulates 2750 and small particulates (not shown) through the pores 2754 in the microfilter 2755 in a microfiltration chip. According to an exemplary implementation, liquid enters through the input 2780, and exits through the outlet 2781. The liquid flow 2760 enters the microfiltration chip through opening 2780. The liquid flow through the microfilter 2755. The liquid flow 2764 exits the microfiltration chip through opening 2781. The large particulates 2750 bigger than the pores 2754 are retained on the microfilter 2755.

[0156] FIG. 24 illustrates a device that can implement the installation of a microfilter for filtration showing the flow of liquid containing large particulates 3050 and small particulates (not shown) through the pores 3054 in the microfilter 3055 held in a structure 3091 formed by the microfiltration chip. According to an exemplary implementation, liquid

enters through the input 3080, and exits through the outlet 3081. The liquid flow 3060 enter the microfluidic chip through opening 3080. The liquid flow through the microfilter 3055 with posts 3075. The liquid flow 3064 exits the microfiltration chip through opening 3081. The large particulates 3050 bigger than the pores 3054 are retained on the microfilter 3055 or captured on the posts 3075.

[0157] FIG. 25 illustrates a device according to an exemplary embodiment of the present invention that can implement the installation of a microfilter for filtration showing the flow of liquid containing large particulates 2850 and small particulates (not shown) through the pores 2854 in the microfilter 2855 in a microfiltration chip. According to an exemplary implementation, liquid enters through the input 2880, and exits through the outlet 2881 and 2882. The liquid flow 2860 enters the microfiltration chip through opening 2880. Most of the liquid flow 2863 through the microfilter 2855. The liquid flow 2864 exits the microfiltration chip through opening 2881. The large particulates 2850 bigger than the pores 2854 are retained above the microfilter 2855, and they flow out the opening 2882 on the same side of the microfilter as the input opening 2860.

[0158] According to exemplary embodiments of the present invention microfilter in any of the microfiltration devices shown in FIGS. 22-25 can be a single layer with pores penetrating from the top to the bottom. The pores can be circular, square, rectangle, or any shape that can be obtained by photo lithography that is of useful application.

[0159] According to exemplary embodiments of the present invention microfilter in any of the microfiltration device shown in FIGS. 22-25 can also be double layered. An example of two layered filter with rectangular opening is shown in FIGS. 13A-13B. Other examples of double layered microfilters are shown in FIGS. 15 and 16. The microfilter in any of the microfiltration chip can also be triple layer as shown in FIGS. 14A-14C.

[0160] FIG. 26 illustrates a device according to an exemplary embodiment of the present invention that can implement the installation of more than one microfilter for filtration showing the flow of liquid containing large particulates 2650, small particles of special characteristics 2640 and other small particulates (not shown) through the pores 2654 in the microfilter 2655 and pores 2644 in the microfilter 2645 held in a structure 2690 formed either by the device. According to an exemplary implementation, liquid enters through the input 2680, and exits through the outlet 2681. The liquid flow 2660 enters the microfiltration chip through opening 2680. The liquid flow 2662 through the first microfilter 2655, and the liquid flow 2642 through the second microfilter 2645. The liquid flow 2664 finally exits the device through opening 2681. The large particulates 2650 bigger than the pores 2654 are retained on the microfilter 2655 by size seclusion. The second microfilter 2645 can have a smaller pore size to capture particulates 2640 again based on size seclusion. The second microfilter 2645 can also be coated with a material that recognize markers on the desired particulates 2640 in the liquid that are smaller than the pore sizes or deformable to squeeze through the pores in the first microfilter. The types of coatings and methods of coating are described in sections below. These particles 2640 are captured on the second microfilter 2645.

[0161] In FIG. 26, the first microfilter 2655 may also be coated with the same or different material as the second microfilter 2645 or have no coating. The number of microfil-

ters placed sequentially are not limited to two. The geometry of the microfilters in the same device can be different.

[0162] According to an exemplary implementation, more than one microfilter can also be installed in any of devices shown in FIGS. 23-25. According to another exemplary implementation, the microfilters can be of different geometries with different properties.

[0163] According to exemplary embodiments of the present invention, in any of the microfiltration devices, FIGS. 22-26, a prefilter with large pores may be used to eliminate large particulates in the liquid.

Coating of the Microfilters

[0164] In certain embodiments, surface functionalization of a polymeric microfilter may provide a surface of the microfilter with surface properties desired for a particular application of the microfilter. Some materials can be directly disposed on the microfilter surfaces. Other times, the microfilter surfaces need to be treated. In one embodiment, the surface of the polymeric microfilter can be functionalized by performing a plasma treatment on the surface of the microfilter to activate the surface to enable chemical compounds and/or organic materials to attach to the surface. In some embodiments, another surface modification technique is to coat the microfilter with a thin layer of a metallic substance.

[0165] FIGS. 27 and 28 are cross-sectional views of coated microfilters in accordance with embodiments of the present invention. In the embodiment illustrated in FIG. 27, microfilter 1655 includes a coating 1600 on surface of one layer microfilter 1655. Coating 1600 can also be disposed on multi-layered microfilters. Coating 1600 can also be disposed on microfilters produced by other methods and by other materials. In FIG. 28, the multi-layer device is formed from epoxy-based photo-definable dry film, as described above in relation to FIGS. 16A-16J. In the embodiment illustrated in FIG. 28, microfilter 1672 includes a coating 1600 on surface of the top structure and flat portions of the microfilter 1672. Coating 1600 can also be disposed on microfilter structures produced by other methods and by other materials. Coating 1600 can also be disposed on the surface of structure shown in FIGS. 16I-16J even without pores.

[0166] In certain embodiments, coating 1600 may be formed from a metallic substance, a nanoparticle colloidal substance, chemical compound, or an organic substance. In such embodiments, these surface coatings can be used to attach analyte recognition elements, DNA, aptamers, surface blocking reagents, etc. In other embodiments, coating 1600 may include analyte recognition elements, DNA, aptamers, surface blocking reagents, etc. In certain embodiments, the surface coatings may be used to attach, for example, macromolecules such as polypeptides, nucleic acids, carbohydrates and lipids. Examples of polypeptides that may be used as analyte recognition elements include, for example, an antibody, an antigen target for an antibody analyte, a receptor (including a cell receptor), a binding protein, a ligand, or other affinity reagent to the target analyte. Examples of nucleic acids that may be used as analyte recognition elements include, for example, DNA, cDNA, or RNA of any length that allows sufficient binding specificity. In such embodiments, both polynucleotides and oligonucleotides can be used as analyte recognition elements. In other embodiments, gangliosides, aptamers, ribozymes, enzymes, antibiotics or other chemical compounds may be used as analyte recognition elements. In certain embodiments, analyte rec-

ognition coatings or elements may include, for example, biological particles such as a cell, a cell fragment, a virus, a bacteriophage or tissue. In some embodiments, analyte recognition coatings or elements may include, BSA, fetal bovin serum (FBS), selectins including P-selectins, E-selectins, L-selectins, nanoparticles, nanotubes, halloysites, dendrimers, chemical linkers or other chemical moieties that can be attached to a microfilter and which exhibit selective binding activity toward a target analyte.

[0167] In some embodiments, coating 1600 may be formed from a metallic substance including gold, nickel, etc. In certain embodiments, coating 1600 includes gold coated on chromium. In some embodiments, it may be preferable to form coating 1600 from gold, as certain chemical compounds and organic materials readily attach to gold. In other embodiments, coating 1600 may be formed from carbon nanotubes. In the embodiment illustrated in FIGS. 27-28, coating 1600 is disposed on one surface of microfilters. In other embodiments, one or more surfaces of microfilters may be coated with coating 1600. In some embodiments, microfilters may be completely coated with coating 1600. Coating 1600 may be disposed on one or more surfaces of any of the microfilters described herein in accordance with embodiments of the present invention, including multi-layer microfilters. In certain embodiments, coating 1600 may be disposed on one or more surfaces of multi-layer microfilter 1620.

[0168] In some embodiments, examples of chemical compounds and organic materials that may be useful for assays when deposited on the surface of a microfilter include are self-assembled monolayers with a range of functionality including amine, carboxyl, hydroxyl, epoxy, aldehyde, and polyethylene glycol (PEG) groups. These compounds and materials may be deposited on the surface of a microfilter using silane chemistry with solution immersion or vapor deposition. In certain embodiments, for example, grafting PEG-triethoxysilane onto an oxidized polymer renders the surfaces hydrophilic in a controlled manner. In other embodiments, a surface of a polymeric microfilter can be functionalized with avidin, biotin, protein A, protein G, antibodies, etc.

[0169] In certain embodiments, coating a surface of a microfilter with a metallic substance may provide other benefits in addition to facilitating the attachment of chemical compounds and/or organic materials. In some embodiments, for example, a layer of a metal metallic substance, having an appropriate thickness, can block transmission of light through the microfilter. In certain embodiments, a thickness sufficient to block the transmission of light is about 40 nm. In other embodiments, this thickness may vary depending on the substance used. Additionally, metallic substances are generally electrically conductive. In some embodiments, when the metallic substance is electrically conductive, the coating may reduce or eliminate charging of the surface of the microfilter. In alternative embodiments, a microfilter may be coated with a thin layer of PARYLENE. In other embodiments, a microfilter may be coated with a thin layer of fluorinated ethylene propylene (FEP), polytetrafluoroethylene (PTFE), perfluoroalkoxy (PFA), or another similar material. In such embodiments, coating a microfilter with one of these materials can reduce nonspecific binding; however, the fluorescent nature of these materials may make them disadvantageous when a microfilter is to be analyzed via microscope imaging.

[0170] In certain embodiments, single layer or multi-layered microfiltration devices can be coated with an antibody

against surface markers on the CTCs to further improve the collection of live CTCs. In such embodiments, capture efficiency of live CTCs may be improved. In some embodiments, useful surface markers may include antibodies against EpCAM, HER2, EGFR, KRAS, Vimentin, and MUC-1, but not limited to these surface markers, P-selectin, E-selectin, other selectins, ligands, aptamers, etc. In embodiments in which the microfilter is coated with cell surface recognition elements, the microfiltration can capture CTCs through size exclusion and surface markers simultaneously.

Filter Holder and Filtration Devices

[0171] The steps of filtration of large rare cells from body fluids on the microfilter or concentrated in a reduce volume with reduced contaminants can be performed with minimal human intervention if the microfilter can be installed in a microfiltration device. The present invention details a method to use microfilters using filter holders.

[0172] Microfilters made from epoxy-based photo-definable dry films, as shown in FIG. 29, can be used in a variety of ways to analyze rare cells such as CTCs from body fluids. Rare cells can be collected on the epoxy-based photo-definable dry film microfilters and other precision microfilters. Following the rare cells collection, rare cells can be analyzed using the following methods:

[0173] Enzymatic assays in rare cells such as CTCs can be test to evaluate viability and enzymatic activities;

[0174] Histopathological staining to look at cell morphology;

[0175] Immunofluorescence staining to look for expression of biomarkers to identify cancer subtype, or to determine biomarker mutation, or cell type. These information can be used to determine cancer therapy and monitor treatment; immunofluorescent staining can be used to perform enumeration of CTCs, based on DAPI (positive), Cytokeratins (CKs) (positive), such as CK8, 18, 19 and others, and CD45 (negative) cells, and count the number per ml of blood to monitor cancer treatment and recurrence;

[0176] Fluorescent in situ hybridization (FISH) to identify gene amplification, number of copies of biomarker genes, gene translocation, to identify cancer subtypes to determine therapy and monitor treatment; mRNA FISH can also be performed to determine the marker is over-expressing.

[0177] Nucleic acid assays for mRNA, microRNA and genomic DNA biomarkers, and gene mutations to identify cancer subtypes to determine therapy and monitor treatment;

[0178] Sequencing the genes to identify gene mutations, amplification and translocations to identify cancer subtypes to determine therapy and monitor treatment; and

[0179] Culture rare cells, such as CTCs, to increase the number of cells. These cells can then be used to perform the assays described in bullets above. In addition, viable cells can be used to determine the effect of drugs on the cancer cells to determine therapy.

[0180] Culture rare cells on microfilters coated with analyte/markers recognition elements for analytes/markers that can be secreted by the CTCs. Assay similar to EPISPOT can be performed using the microfilter.

[0181] Some of those assay methods can be combined or performed sequentially on rare cells collected on epoxy-based photo-definable dry films. For example, rare cells col-

lected on epoxy-based photo-definable dry films can be performed sequentially to obtain different information from the same cells. Some examples are: (1) Perform enumeration and for a cancer surface biomarker, followed by FISH assay and finally histopathological staining, (2) perform immunofluorescent staining to count the number of rare cells and at the same time determine over-expression of biomarker(s), mutated biomarker(s), cell type, and/or other information, followed by histopathological staining, (3) perform FISH to identify a marker followed by histopathological staining.

[0182] According to exemplary embodiments of the present invention, filtration devices to hold the filter and perform filtration are designed to have the following features:

[0183] The filter holder to hold the filter flat without causing twisting.

[0184] This filter holder has a large opening in the inlet unit. This is to allow easy access of the reagents to microfilter surface and visual access to the microfilter, where the cells are collected, and allow visualization of the filter from above.

[0185] The filter holder will allow performance of at least some types of assay in the filter holder.

[0186] The filter holder will allow different attachment of input sample holder above the filter holder, and syringe, stopper on the filter flask connected to a vacuum pump or vacutainer holder below.

[0187] According to an exemplary embodiment of the present invention, a filter holder allows for (i) filtration of the liquid sample through the microfilter, (ii) ability to perform assays with the microfilter in the holder, (iii) easy access to the microfilter in the filter holder, (iv) easy installation of the microfilter into the filter holder, (v) easy removal of the microfilter from the filter holder, and (vi) filter holder material capable of tolerating temperature required for the assays, and (vii) filter holder material capable of tolerating chemicals used in the assays.

[0188] FIGS. 30A-30C show three components that forms a filter holder according to an exemplary embodiment of the present invention, including: an inlet unit **3010**, a nut **3020** and an outlet unit **3030** (top view FIG. 30C and the bottom view FIG. 30D). In an exemplary implementation a syringe or vacutainer holder with a Lure-Lock can be attached to the outlet unit of the filter holder. In an exemplary implementation an output unit can include a cut out on the top ring, for example, to allow access to the microfilter to allow easy removal of microfilters from the filter holder.

[0189] FIGS. 31A-31E show the assembly of the filter holder according to an exemplary embodiment of the present invention where a gasket **3110** is placed inside the outlet unit as shown in FIG. 31A. A round microfilter **3120** with the appropriate diameter for the filter holder is placed above the gasket, FIG. 31B. A second gasket **2130** is place above the microfilter **3120**, FIG. 21C. The inlet unit **3010** is placed into the outlet unit **3030** by properly aligning the two components as shown in FIG. 21D. Finally the nut **3020** is installed to tighten the filter holder system to prevent leakage of the liquid sample around the microfilter inside the holder. While one gasket may be sufficient, a plurality of gaskets may be used, for example to prevent leakage, as needed. The gasket can be of different material or design. In an exemplary filter holder design, a microfilter installed in a filter holder system will remain flat, experience compression to form a tight seal, but not experience any twisting force.

[0190] There are a lot of potential variations of the filter holders in keeping with the scope of exemplary embodiments of the present invention, including without limitation:

[0191] The filter holder can have different sizes to accommodate different microfilter sizes. The common microfilter sizes are 0.5 inch (13 mm) and 1 inch diameters, but are not limited to these dimensions.

[0192] The filter outlet unit can have more than one opening. The opening is to allow easy removal of microfilter from filter holder and prevent the microfilter from becoming twisted by fixing in place the inserted inlet unit.

[0193] The shape and dimensions of different component of the filter holder can vary. One of the features of the filter holder concept according to exemplary embodiments of the present invention is pressing down on the inlet unit to prevent leakage. In an exemplary embodiment, a nut is used to hold down the inlet unit.

[0194] The inlet unit can also be held down by changing the nut to a snap-on part.

[0195] The outlet 3030 can have a female connector or a male connector.

[0196] The filter holder can be designed to accommodate one gasket under the microfilter or two gaskets, one above and one below the microfilter.

[0197] A filter support structure can be added to the inside of output units of the filter holders. Support structure may not be needed for epoxy-based photo-definable dry film microfilters because the material is sufficiently strong.

[0198] The inlet unit 3010 can be combined with inlet adaptor 3050, in FIG. 33, into one piece.

[0199] Exemplary embodiments of filter holders described in this application are applicable to most filters and microfilters. Certain exemplary embodiments can be particularly suitable to very thin and strong microfilters such as epoxy-based photo-definable dry film microfilters.

[0200] FIGS. 32A-32B show examples of sample input containers for holding the liquid sample and wash buffers. FIG. 32A shows an inlet container 3200 with the shape similar to a syringe with sample inlet opening 3210 and connection to the filter holder 3220. FIG. 32B shows an inlet container 3300 with wider opening 3310 and connection to the filter holder 3320 to allow easy access to the microfilter. FIGS. 32A-32B show examples of sample input container made as one piece.

[0201] FIG. 33 shows an example of a sample input container 3420 that can also be constructed out of an off the shelf syringe 3410 without a plunger and an inlet adaptor 3050 with a male Lure-Lock. The inlet adaptor can have various designs as long as it does not leak and it can be easily removed from the filter holder. The shape at the bottom of the sample input containers 3200, 3300, and the inlet adaptor 3050 should have a shape to facilitate tight fit into the inlet unit 3010 on the top of the assembled filter holder 3100 in FIG. 31E. Alternatively, or in combination, an O-ring can be provided to facilitate the tight fit into the inlet unit 3010.

[0202] The filtration system 3500 can assume the configuration shown in FIG. 34. In the example of this drawing, the input container 3420, constructed as shown in FIG. 33, is connect to the top of the assembled filter holder 3100 from FIG. 31E, and the bottom of the assembled filter holder 3100 shown in FIG. 31E is connected to a waste syringe 3510 with plunger 3520.

[0203] FIG. 35 shows a filtration system with just the filter holder 3100 and the waste syringe 3510. The opening 3140 allows performance of many assay steps directly on the microfilter.

[0204] The filtration can be performed manually by drawing the plunger, but manual operation may not provide consistent speed. The filtration system 3600 using syringe pump 3610 in FIG. 36 can provide more consistent speed for pulling the plunger 3520 by a pusher block 3620. The syringe pump can have just infusion or both infusion and withdrawn functions.

[0205] FIG. 37A shows another exemplary method for drawing blood through the microfilter by connecting a vacutainer holder 3710 to the outlet of the filter holder 3730. When a vacutainer 3740 is inserted into the vacutainer holder 3710, shown in FIG. 37B, the vacuum in the vacutainer will draw the liquid sample into the vacutainer and the microfilter will collect the rare cells.

[0206] FIG. 38A shows a filtration device 3800 that can perform filtration at the time of getting blood draw. The filtration device combines a filter holder 3830 with the inlet adaptor 3820 and vacutainer holder 3810. A needle 3850 with female Luer-Loc can be attached to the inlet adaptor 3820. FIG. 38B shows a vacutainer 3840 inserted into the vacutainer holder 3810.

Sample Collection and Performing Assays Using the Filtration Devices

[0207] The logistics of collecting the sample and shipping the sample to clinical laboratories to perform the rare cell assays according to exemplary embodiments of the present invention are described below with reference to FIGS. 39A-39C.

[0208] FIG. 39A illustrates an example of a method to obtain samples and shipping to clinical laboratories for testing including: (i) collecting body fluid, (ii) sending sample to testing laboratory, (iii) filtering rare cells out of body fluids, and (iv) performing an assay.

[0209] Because cells can degrade and blood might form clots and time required for shipping can be 24-48 hours, FIG. 39B describes an alternative method for obtaining samples, and for testing in clinical laboratories including: (i) collecting body fluid, (ii) filtering the rare cells from the body fluid at the collection site, (iii) sending filter holder with microfilter or just the microfilter with captured rare cells to testing laboratory, and (iv) performing the assay in the testing laboratory.

[0210] For blood samples, it is also possible to filter the rare cells from blood at the time of drawing the blood using the filtration device 3800 shown in FIG. 39C. After performing a washing step, the filter holder with microfilter or just eh microfilter with captured rare cells can be sent to testing laboratory. Assays will be performed in the testing laboratory.

Microfilter Applications

[0211] In exemplary embodiments of the present invention described above, microfilters can be formed from epoxy-based photo-definable dry film having a thickness between 1-500 μm . In certain embodiments, such microfilters can be formed using UV light to expose the dry film (i.e., using UV lithography). In some embodiments, X-rays (i.e., X-ray lithography) may be preferred for exposing relatively thick dry films, for simultaneously exposing multiple stacked dry films, or for exposing resists that require a relatively high

dose. In certain embodiments, relatively thick microfilters may provide more structural strength than thinner microfilters, but may also utilize higher pressure during filtration.

[0212] As noted above, in certain exemplary embodiments, epoxy-based photo-definable dry films are a preferred material from which to form microfilters in accordance with embodiments of the present invention. In some embodiments, properties of epoxy-based photo-definable dry film that make it a suitable material from which to form microfilters for medical diagnostic applications are that it is photo-definable by UV light, it is clear, it has a high tensile strength of 75 Mpa, it can be laminated to itself, it can be directly coated on a substrate, and it has no auto-fluorescence in the visible wavelengths. In addition, while the processes described above in accordance with embodiments of the present invention may be used to form microfilters, the processes described above may also be used to manufacture other kinds of free-standing patterned polymeric films.

[0213] Microfilters formed in accordance with exemplary embodiments of the present invention have many possible applications. In some embodiments, exemplary applications for such microfilters include medical applications, water filtration applications, beer and wine filtration applications, pathogen detection applications, etc.

[0214] FIG. 40A is a flowchart illustrating a filtration process 1700 using a microfilter in accordance with exemplary embodiments of the present invention. At block 1720 of FIG. 40A, a liquid may be passed through a microfilter formed from a layer of epoxy-based photo-definable dry film, in accordance with any one of the embodiments described above, having a plurality of apertures. In certain embodiments, the liquid may be pushed through the microfilter. In other embodiments, the liquid may be drawn through the microfilter. The draw can be produced by a syringe or by vacuum. In some embodiments, the liquid may be passed back and forth through the microfilter one or more times. In some embodiments, the particulates retained on the microfilters may be backwashed using appropriate liquid.

[0215] In certain exemplary embodiments, the process illustrated in FIG. 40A may be used to perform an assay using the microfilter. In certain embodiments, the process may be used to filter cells, such as CTCs, from a solution including a patient's bodily fluid. This can be accomplished using vacutainer as shown in FIG. 38A-38B, or by placing the filter on a support above a vacuum pump.

[0216] FIG. 40B is a flowchart illustrating a filtration process 1800 using a microfilter in accordance with exemplary embodiments of the present invention. At block 1830 of FIG. 40B, a microfilter formed from a layer of epoxy-based photo-definable dry film, in accordance with any one of the embodiments described above, is positioned in a filter holder. In certain embodiments, the filter holder includes an inlet, an outlet, and securely holds the microfilter around the edges of the filter. In some embodiments, a liquid may be input into the filter holder through the inlet. At block 1850, the liquid is passed through the microfilter. In certain embodiments, the liquid is a bodily fluid or a solution including a bodily fluid. In certain embodiments, the liquid is drawn through the microfilter by applying negative pressure at the outlet of the filter holder such that all or substantially all of the liquid is drawn through the pores of the microfilter. In other embodiments, the liquid is pushed through the microfilter. At block 1870, the

microfilter may also be removed from the filter holder. For microscope imaging, the microfilter may be placed on a glass slide.

[0217] FIG. 40C is a flowchart 1900 illustrating examples for performing some types of assays in the filter holder 1910. After the assay, the filter can be removed from the filter holder 1920 for analysis.

[0218] Enzymatic activity assay, histopathology staining (colorimetric staining), and immunofluorescent staining for applications such as determination of biomarker expression, EPISPOT and enumeration, can follow the flow chart steps in FIG. 40C after step shown in FIG. 40B. These assays can be performed in a filter holder as described. These assays can also be performed on glass slide, or plate after removing the microfilter from the filter holder follow the flow chart shown in FIG. 40A or 40B.

[0219] Nucleic acid assays, sequencing, fluorescence in situ hybridization (FISH), mRNA in situ hybridization, and culture require removing the microfilter with captured rare cells from the filter holder and perform the assays in each of their own appropriate ways as shown in flow chart of FIG. 40A or 40B.

[0220] In the exemplary flow chart of FIG. 40B, the protocol to isolate cells is as follows using the syringe pump:

- [0221] 1. Assemble the microfilter in the filter holder as shown in FIGS. 31A-31E;
 - [0222] 2. Mount a waste syringe 3510 to a syringe pump 3610;
 - [0223] 3. Attach the assembled filter holder 3100 on to the waste syringe 3510;
 - [0224] 4. Attach the inlet container 3420 (FIG. 32A, 32B or 33) on top of the filter holder 3100, and the complete assemble looks like FIG. 36;
 - [0225] 5. Place liquid sample into the input container 3420;
 - [0226] 6. Using negative pressure, suck the liquid sample through the microfilter into the waste syringe.
 - [0227] 7. To perform wash step, place wash buffer into the input container 3420, and suck the wash buffer through the microfilter into the outlet syringe. Perform wash a few times.
 - [0228] 8. Remove the input container 3420 and the filtration system looks like FIG. 35.
 - [0229] 9. Open the filter holder to retrieve the microfilter.
- [0230] The syringe pump can be operated manually or can be automated.

[0231] The filtration can be performed manually. To perform the assay manually, skip step 2. At the start, complete assembled system looks like FIG. 34.

[0232] The filter holder 3100 can also be placed on a stopper on a filter flask connect to a vacuum pump with the inlet container 3420. The liquid sample and the wash buffer can be placed into the inlet container 3420. The vacuum pump turned on to draw the liquid through the filter. If the inlet container is not used, the liquid sample and the wash buffer can be pipette into the reaction well 3010 while the vacuum is on.

[0233] The filtration can be performed using the vacutainer systems 3700 as shown in FIG. 37A. The steps are:

- [0234] 1. Assemble the microfilter in the filter holder as shown in FIGS. 31A-31E;
- [0235] 2. Mount a vacutainer holder to the outlet of the filter holder 3730 as shown in FIG. 37A.

- [0236] 3. Attach the inlet container 3720 (FIG. 32A, 32B or 33) on top of the filter holder 3730, and the complete assemble looks like FIG. 37A;
- [0237] 4. Place liquid sample into the input container 3720;
- [0238] 5. Insert vacutainer 3740 into the vacutainer holder 3710, as shown in FIG. 37B.
- [0239] 6. To perform washing step, place wash buffer into the input container 3720.
- [0240] 7. Install a fresh vacutainer 3740 into the vacutainer holder 3710 to perform the wash.
- [0241] 8. Further processing of the cells collect on the microfilter as needed, or open the filter holder to retrieve the microfilter.
- [0242] The filtration can be performed at the time of blood draw using the vacutainer systems 3800 as shown in FIG. 38A. The steps are:
- [0243] 1. Assemble the microfilter in the filter holder as shown in FIGS. 31A-31E;
- [0244] 2. Mount a vacutainer holder to the outlet of the filter holder as shown in FIG. 38A.
- [0245] 3. Attach a needle 3850 with female Luer-Loc to the inlet adaptor 3820.
- [0246] 4. Insert needle into the vein of the patient.
- [0247] 5. Insert vacutainer 3840 into the vacutainer holder 3810, as shown in FIG. 38B.
- [0248] 6. Remove vacutainer 3840 after draw.
- [0249] 7. Remove the needle from the patient's vein.
- [0250] 8. Remove the needle from the filter holder, and install an input container 3720 as shown in FIG. 37A.
- [0251] 9. To perform the washing step, place wash buffer into the input container 3720.
- [0252] 10. Install a fresh vacutainer 3740 into the vacutainer holder 3710 to perform the wash, as shown in FIG. 37B or insert the syringe into the outlet of the filter holder as shown in FIG. 33.
- [0253] 11. Further processing of the cells collect on the microfilter as needed, or open the filter holder to retrieve the microfilter.
- [0254] The filter with cells attached can be used for various assays and analysis.
- [0255] FIG. 40C shows that many assay steps can be performed in the filter holder after the steps in the flow chart of FIGS. 40A-40B. The details of the assay steps can vary depending on the assay. This is performed using the configuration shown in FIG. 35. The steps will vary depending on the assay. The general steps are described below.
- [0256] 1. Incubation: Place reagents into the reservoir 3140 above the filter and incubate. The incubation of with reagents above the microfilter in the filter holder is possible because the epoxy-based photo-definable dry film microfilter is hydrophobic. Reagents will not leak through the microfilter.
- [0257] 2. Wash: For small amount of wash buffers after the incubation, wash buffer can be placed directly into the reservoir 3140 and sucked out by negative pressure. This can be repeated. When larger volumes of washing buffers are required, place the input container back on the filter holder to form the system shown in FIG. 34 and perform washing step by sucking out the wash buffer placed into the input container.
- [0258] 3. Anytime the outlet syringe becomes full, replace with a new outlet syringe.

- [0259] 4. When the assay is completed, unassembled the filter holder and take out the microfilter and place it on a glass slide, plate reader or other appropriate device for image analysis.

[0260] Nucleic acid assays and sequencing are suited for performing the assay from the lysed cells. The conceptual protocol follows the steps of collecting the cells on the microfilter. After taking the microfilter out of the filter holder, place the microfilter containing rare cells in an Appendorf centrifugation tube with lysis buffer. The rest of the steps for the nucleic acid assays are the same as common samples.

[0261] For culturing the rare cells, the conceptual protocol follows the steps of collecting the cells on the microfilter. After taking the microfilter out of the filter holder, place the microfilter containing rare cells in culture medium. Cells can also be backwashed out of the filter into culture medium. In some situations the rare cells need to be removed from the microfilter and inject into an animal such as mice.

Analysis of Microfiltered Cells

[0262] In some embodiments, the particulates retained on the microfilter may then be subject to processing and or analysis to analyze any cells or other materials, substances, etc. collected by the microfilter. The analysis may be performed in the filter holder device or microfiltration chip. The analysis may also be performed outside the filter holder after the microfilter is removed from the filter holder. Exemplary applications of this process in accordance with embodiments of the present invention will be described below.

[0263] In certain embodiments, a microfilter formed from a layer of epoxy-based photo-definable dry film in accordance with any one of the embodiments described above may be used for medical diagnostics and/or prognostics. In certain embodiments, the microfilter may be used to collect certain types of cells from bodily fluids based on cell size. In some embodiments, the microfilter can be used for isolating and detecting rare cells from a biological sample containing other types of cells. In some embodiments, the microfilter can be used to filter a fluid sample, and the collected cells can be used in a downstream processes such as cell identification, enumeration (cell counting), characterization of the collected cells, culturing the collected cells, separating the cells into individual cells or groups of cells, or use the of cells in other ways. The final enriched target cells can be subjected to a variety of characterization and manipulations, such as staining, immunofluorescence of markers, cell counting, DNA, mRNA, microRNA analysis, fluorescence in-situ hybridization (FISH), immunohistochemistry, flow cytometry, immunocytochemistry, image analysis, enzymatic assays, gene expression profiling analysis, sequencing, efficacy tests of therapeutics, culturing of enriched cells, and therapeutic use of enriched rare cells. In addition, depleted plasma protein and white blood cells can be optionally recovered and subjected to other analysis, such as inflammation studies, gene expression profiling, etc.

[0264] In certain embodiments, the microfilter can be held in a filter holder for medical diagnostics and/or prognostics. In some embodiments, the filter holder may include a built in support for the microfilter. In certain embodiments, the filter holder may have gasket above and below the filter. In some embodiments, the microfilter may be used to collect circulating tumor cells (CTCs) in blood. In such embodiments, a blood sample, typically in the range of 1-10 ml, is taken from a patient. The blood sample is then drawn through the micro-

filter by applying negative pressure, such as a sucking force. In certain embodiments, the blood is pulled through the microfilter via an outlet. In some embodiments, passing the blood through the filter by pushing can cause cell rupture except at very low pressure or low speed.

[0265] In certain embodiments, most cells having a dimension larger than the width of one or more pores of the microfilter are retained. Most white blood cells are deformable and can pass through pores having a smaller width than a dimension of the white blood cell. In certain embodiments, nearly no red blood cells are retained on the microfilter. In some embodiments, the microfilter may include pores 7-8 μm in diameter for enriching circulating tumor cells and fetal cells; however, the microfilter pore size and shape can be varied for these applications as well.

[0266] In some embodiments, CTCs collected by the microfilter can be enumerated on the microfilter. In one experiment conducted to determine the capture efficiency of a microfilter, tumor cell lines were used. The microfilter used to demonstrate filtration efficiency was a microfilter having a 7-8 micron diameter pores separated by 20 microns and arranged within a 9 mm diameter area. The microfilter was placed into a filter holder. A prestained MCF-7 cell line was spiked into 7.5 ml of whole blood. To capture live CTCs, the blood was diluted 1:1 with a buffer solution. One exemplary buffer solution is phosphate buffered saline (PBS). Another exemplary buffer solution is a mild fixation buffer to make the CTCs slightly more rigid. The liquid sample was drawn through the microfilter using negative pressure at approximately 10 ml/min. Afterwards, the filter was washed twice in a buffer solution. The microfilter was removed from the holder and mounted onto a microscope slide to be counted. The recovery rate of live MCF-7 cells was $85\% \pm 3\%$. If the blood is mildly fixed, the capture efficiency of MCF-7 cells increases to $98\% \pm 2\%$.

[0267] In certain embodiments, collected CTCs can be subjected to a variety of analyses and manipulations, such as immunofluorescence, genetic characterization and molecular phenotyping, fluorescence in-situ hybridization (FISH), mRNA FISH, in-situ hybridization (ISH), mRNA ISH, immunohistochemistry (IHC), flow cytometry, immunocytochemistry, colorimetric staining, histopathological staining (for example, hematoxylin and eosin staining), image analysis, epithelial immunospot (EPISPOT), enzymatic assays, gene expression profiling analysis, efficacy tests of therapeutics, culturing of enriched cells, and therapeutic use of enriched rare cells. In addition, in some embodiments, depleted plasma protein and white blood cells can be optionally recovered, and subjected to other analysis such as inflammation studies, gene expression profiling, etc.

[0268] In certain embodiments, CTCs collected from blood can be stained to identify them as potential tumor cells and not blood cells. It is possible to identify the CTCs by morphology using colorimetric staining. Other methods are based on fluorescence staining. Some typical fluorescence staining methods to identify the cells as tumor cells use DAPI to identify the nucleus and use cytokeratins 8, 18 and 19 conjugated to a fluorescent dye to identify them as epithelial cell. Since normal epithelial cells are not found in blood, epithelial cells found in blood are accepted as tumor cells. CD45 antibody is used to identify white blood cells, eliminating the blood cells retained on the microfilter as CTCs. It is commonly accepted that a cell from blood that is DAPI positive, CK 8, 18, and 19 positive and CD45 negative to be a CTC. Other markers, such

as EpCAM, MUC-1, and others) can be included to further provide specificity and increase of fluorescence signal.

[0269] In certain embodiments, captured CTCs are stained to specifically identify the origin of the tumor cells, such as breast, prostate, colon, etc. For example, PSA marker on the cell would identify its origin as prostate. For each cancer, specific markers can be found either on the surface or inside the cells.

[0270] In certain embodiments, captured CTCs can be characterized to determine if it contains specific mutations of the DNA. This can be identified by for DNA, mRNA, and microRNA expression by PCR, by sequencing, or by antibodies, ligands, aptamers and others that recognize the mutated proteins.

[0271] In certain embodiments, captured CTCs can be characterized to determine over-expression of genes. Sometimes the number of copies of a gene is more than it should be for each cell. When there are more copies of the gene, it produces more copies of mRNA, which in turn produces more copies of the protein. The number of copies of the gene can be determined by FISH or by ISH. The amount of mRNA produced can be obtained by PCR, mRNA FISH and mRNA ISH. The amount of proteins produced can be determined by immunofluorescent stains for that protein. Cells over-expression a marker will stain brighter for that marker than normal tissue. One example of over expression is HER-2 in some subtypes of breast cancer.

[0272] In certain embodiments, captured CTCs can be characterized to determine if they are viable cells.

[0273] In certain embodiments, captured CTCs can be determined if they are viable. Viability can be determined by trypan blue staining, or culture.

[0274] In certain embodiments, captured CTCs can be determined if they are stem cells. They can be stained for stem cell marker phenotype ($\text{CD44}^+/\text{CD24}^{\text{low}}$ or $\text{CK19}^+/\text{MUC1}^-$)

[0275] In certain embodiments, viable CTCs may be captured by microfilter coated by analyte recognition elements, such as antibodies, ligands, aptamers, etc. The analytes of interest are to be secreted by the CTC. A secondary analyte recognition element can be used to produce a detectable signal if the analyte is produced by the CTC. The signal may be produced by fluorescent dyes. The concept is similar to EPISPOT.

[0276] In certain embodiments, captured live CTCs can be cultured directly on the microfilter to increase the number of CTCs and to evaluate the characteristics of CTCs. In other embodiments, the CTCs can be backwashed from the microfilter prior to culturing or sorting.

[0277] In other embodiments, a microfilter formed from a layer of epoxy-based photo-definable dry film in accordance with embodiments of the present invention may be used in therapeutic applications in which circulating tumor cells are removed from the blood of cancer patients. Circulating tumor cells are the cause of cancer spreading from the original site to other locations such as brain, lung and liver. Most carcinoma cancer patients die from the metastatic cancer. In certain embodiments, microfiltration using a microfilter formed in accordance with embodiments of the invention is a suitable method for removing circulating tumor cells from the blood stream of a patient because the filtration speed is fast and microfilters retain very little white blood cells and almost no red blood cells when used to filter blood.

[0278] Microfiltration for circulating tumor cells in blood can provide a large array of diagnostic, prognostic and research applications. For collecting circulating tumor cells, previous research reports utilized track etch filters with random pore locations with some overlapping pores and not straight pores, and microfilters with orderly arranged pores produced by reactive ion etching. In certain embodiments of the present invention, microfilters formed from epoxy-based photo-definable dry film and having precisely arranged pores are used to collect circulating tumor cells in blood.

[0279] One exemplary application of a microfilter formed in accordance with embodiments of the present invention is to monitor effectiveness of treatment by counting the number of CTCs collected in the blood. Large number of CTC per ml of blood can indicate short lifespan. The change of CTCs per ml of blood can indicate whether treatment is working or not. If the number of CTCs is decreasing, it indicates the treatment is having an effect. In contrast if the number of CTCs is increasing, it indicates the treatment is ineffective.

[0280] One exemplary application of a microfilter formed in accordance with embodiments of the present invention is using the microfilter for capturing cells and subsequently culturing the cells in the filter holder, or culturing the cells after back flushing the cells off the microfilter. Various drugs can be applied to the cultured CTCs can be used to evaluate drug efficacy to determine the best treatment for the patients.

[0281] One exemplary application of a microfilter formed in accordance with embodiments of the present invention is to determine gene mutation in CTCs to determine the appropriate drug.

[0282] One exemplary application of a microfilter formed in accordance with embodiments of the present invention is to determine over expression of a marker when there is drug to treat tumors with the over expression

[0283] One exemplary application of a microfilter formed in accordance with embodiments of the present invention is to determine if the cancer is returning after remission. If the number of CTCs become more than five from 7.5 ml of blood and the CTC count is increasing in time, then the cancer is returning.

[0284] Another exemplary application of a microfilter manufactured in accordance with embodiments of the present invention is capturing circulating endothelial cells. Endothelial cells in the peripheral blood provides information about various medical conditions.

[0285] Exemplary application of a microfilter manufactured in accordance with embodiments of the present invention is capturing circulating fetal cells in a mother's blood during pregnancy. Such fetal cells may include primitive fetal nucleated red blood cells. Fetal cells circulating in the peripheral blood of pregnant women are a potential target for non-invasive genetic analyses. Fetal cells include nucleated red blood cells and epithelial (trophoblastic) cells, which are 14-60 μm in diameter, larger than peripheral blood leukocytes. Enrichment of circulating fetal cells followed by genetic diagnostic can be used for noninvasive prenatal diagnosis of genetic disorders using PCR analysis of a DNA target or fluorescence in situ hybridization (FISH) analysis of genes.

[0286] Another exemplary application of a microfilter manufactured in accordance with embodiments of the present invention is collecting or enriching stromal cells, mesenchymal cells, endothelial cells, epithelial cells, stem cells, non-hematopoietic cells, etc. from a blood sample, collecting tumor or pathogenic cells in urine, and collecting tumor cells

in spinal and cerebral fluids. Another exemplary application is using the microfilter to collect tumor cells in spinal fluids. Another exemplary application is using the microfilter to capture analytes bound to latex beads or antigen caused particle agglutination whereby the analyte coated bead or agglutinated clusters are captured on the membrane surface.

[0287] Another exemplary application of a microfilter formed in accordance with embodiments of the present invention is for erythrocyte deformability testing. Red blood cells are highly flexible cells that will readily change their shape to pass through pores. In some diseases, such as sickle cell anemia, diabetes, sepsis, and some cardiovascular conditions, the cells become rigid and can no longer pass through small pores. Healthy red cells are typically 7.5 μm and will easily pass through a 3 μm pore membrane, whereas a cell with one of these disease states will not. In the deformability test, a microfilter having 5 μm apertures is used as a screening barrier. A blood sample is applied and the membrane is placed under a constant vacuum. The filtration rate of the cells is then measured, and a decreased rate of filtration suggests decreased deformability.

[0288] Another exemplary application of a microfilter formed in accordance with embodiments of the present invention is leukocyte/Red blood cell separation. Blood cell populations enriched for leukocytes (white blood cells) are often desired for use in research or therapy. Typical sources of leukocytes include whole peripheral blood, leukopheresis or apheresis product, or other less common sources, such as umbilical cord blood. Red blood cells in blood can be lysed. Then the blood is caused to flow through the microfilter with small pores to keep the leukocytes. Another exemplary application is using the microfilter for chemotaxis applications. Membranes are used in the study of white blood cell reactions to toxins, to determine the natural immunity in whole blood. Since immunity is transferable, this assay is used in the development of vaccines and drugs on white blood cells. Another exemplary application is using the microfilter for blood filtration and/or blood transfusion. In such applications, microfilters can be used to remove large emboli, platelet aggregates, and other debris.

[0289] Additionally, arrays of precision micro-pores can be fabricated in rolls of polymer resists in accordance with embodiments of the invention described above. Such arrays may be used for applications for which wafer-sized microfilters are not suitable. Examples of such applications include water filtration, kidney dialysis, etc.

[0290] While various embodiments have been described above, it should be understood that they have been presented by way of example only, and not limitation. It will be apparent to persons skilled in the relevant art that various changes in form and detail can be made therein without departing from the spirit and scope of the present invention. Thus, the breadth and scope of the present invention should not be limited by any of the above-described exemplary embodiments, but should be defined only in accordance with the following claims and their equivalents. The present embodiments are, therefore, to be considered in all respects as illustrative and not restrictive. Additionally, it will be appreciated that any features, components, elements, etc., described above in relation to different exemplary embodiments may be implemented together.

1-26. (canceled)

27. A filtration device comprising:

an inlet layer including a first volume and a first inlet layer opening to said first volume;

a filter structure including a polymer filter layer having a first filter surface and a second filter surface diametrically opposite to said first filter surface; and an outlet layer including a second volume and a first outlet layer opening to said second volume, wherein said first filter surface is exposed to said first volume,

said second filter surface is exposed to said second filter volume, and

at least a filtered portion of a sample entering said first inlet layer opening, passes from said first volume to said second volume via said filter layer and exits said second volume via said first outlet layer opening.

28. The filtration device of claim **27**, further comprising: a top layer covering at least a portion of said first volume opposite said first surface of said filter layer; and a base layer covering at least a portion of said second volume opposite said second surface of said filter layer.

29. The filtration device of claim **28**, wherein: said top layer includes a first top opening and a second top opening;

said first top opening communicatively connected with said first inlet top layer opening;

said top layer includes a top outlet layer opening communicatively connected with said second top opening;

said filter layer includes a first filter layer opening communicatively connected with said top outlet layer opening; said sample entering said first volume via said first inlet layer opening; and

an unfiltered portion of said sample exits said top outlet opening connected with second top opening in the first volume, and

at least a filtered portion of a sample entering said first inlet layer opening, passes from said first volume to said second volume via said filter layer and exits said second volume via said first outlet layer opening.

30. The filtration device of claim **27**, wherein the polymer filter comprises photo-definable dry film.

31. The filtration device of claim **30**, wherein the photo-definable dry film comprises epoxy-based photo-definable dry film.

32. The filtration device of claim **27**, further comprising a coating of one or more elements on first filter surface of the first polymer layer.

33. The filtration device of claim **32**, wherein said elements include analyte recognition elements.

34. The filtration device of claim **27**, wherein said inlet layer includes a second inlet layer opening to said first volume.

35. The filtration device of claim **27**, wherein the filter structure comprises a plurality of filter layers in a stacked formation.

36. A filtration method comprising: configuring a filtration device comprising:

an inlet layer including a first volume and a first inlet layer opening to said first volume;

a filter structure including a polymer filter layer having a first filter surface and a second filter surface diametrically opposite to said first filter surface; and

an outlet layer including a second volume and a first outlet layer opening to said second volume,

wherein said first filter surface is exposed to said first volume,

said second filter surface is exposed to said second filter volume; and

passing a liquid sample into said first inlet layer opening whereby at least a filtered portion of a sample entering said first inlet layer opening passes through apertures in the filter from said first volume to said second volume via said filter layer and exits said second volume via said first outlet layer opening, and wherein the apertures are sized to essentially allow passage of a first type of component in the fluid and to substantially prevent passage of a second type of component in the fluid.

37. The filtration method of claim **36**, wherein said filtration device further comprises:

a top layer covering at least a portion of said first volume opposite said first surface of said filter layer; and

a base layer covering at least a portion of said second volume opposite said second surface of said filter layer.

38. The filtration method of claim **37**, wherein: said top layer includes a first top opening and a second top opening;

said first top opening communicatively connected with said first inlet top layer opening;

said top layer includes a top outlet layer opening communicatively connected with said second top opening;

said filter layer includes a first filter layer opening communicatively connected with said top outlet layer opening; said sample entering said first volume via said-first inlet layer opening; and

an unfiltered portion of said sample exits said top outlet opening connected with second top opening in the first volume, and

at least a filtered portion of a sample entering said first inlet layer opening, passes from said first volume to said second volume via said filter layer and exits said second volume via said first outlet layer opening.

39. The method of claim **36**, wherein the polymer filter comprises a photo-definable dry film filter.

40. The method of claim **36**, wherein the photo-definable dry film comprises epoxy-based photo-definable dry film.

41. The method of claim **36**, further comprising: performing, on the captured second component, one or more of identification, immunofluorescence staining, enumeration, cell lysis, fluorescence in situ hybridization, mRNA in situ hybridization, other molecular characterizations, immunohistochemistry, histopathological staining, flow cytometry, image analysis, enzymatic assays, gene expression profiling analysis, erythrocyte deformability, white blood cell reactions, efficacy tests of therapeutics, culturing of enriched cells, and therapeutic use of enriched rare cells, and separation from the microfilter.

42. The method of claim **36**, wherein the first filter surface includes a coating of one or more elements on at least one surface of the microfilter.

43. The method of claim **42**, wherein the elements comprise analyte recognition elements configured to capture components in the fluid.

44. The filtration method of claim **36**, wherein the filter structure comprises a plurality of filter layers in a stacked formation.

45. A method of configuring a filtration device comprising: providing an inlet layer including a first volume and a first inlet layer opening to said first volume;

providing a filter structure including a polymer filter layer having a first filter surface and a second filter surface diametrically opposite to said first filter surface; and providing an outlet layer including a second volume and a first outlet layer opening to said second volume, wherein said first filter surface is exposed to said first volume, said second filter surface is exposed to said second filter volume, and at least a filtered portion of a sample entering said first inlet layer opening, passes from said first volume to said second volume via said filter layer and exits said second volume via said first outlet layer opening.

46. The method of claim **45**, further comprising:
providing a top layer covering at least a portion of said first volume opposite said first surface of said filter layer; and providing a base layer covering at least a portion of said second volume opposite said second surface of said filter layer.

47. The method of claim **36**, wherein the liquid comprises a body fluid comprising at least one of the group consisting of blood, urine, cerebral spinal fluid and saliva.

48. The method of claim **36**, wherein the liquid comprises water.

49. The method of claim **36**, wherein the liquid contains disbursed components from bone marrow, tumor tissue, and other tissue.

50. The method of claim **36**, wherein said second component comprises at least one of circulating tumor cells, circulating endothelial cells, stem cells, stromal cells, epithelial cells, non-hematopoietic cells, fetal cells, nucleated red blood cells, trophoblastic cells, macrophages, circulating cancer associated macrophage-like cells, epithelial-mesenchymal transition cells, blood cells, processed tissue sample cells, white blood cells, T-cells, B-cells, and tumor cells from bone marrow.

51. The method of claim **36**, wherein said second component comprises at least one of leukocytes, T-cells and B-cells, and the liquid sample is blood.

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