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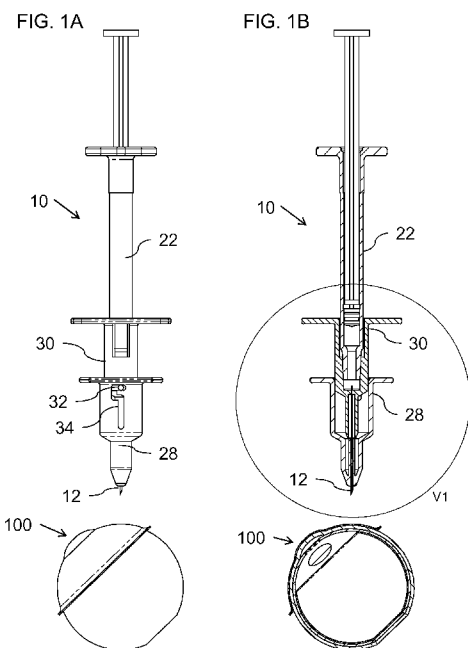
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(54) Title: MINIATURE IMPLANTED DRUG DELIVERY DEVICES AND INSERTER SYSTEMS FOR INTRODUCING SUCH DEVICES



(57) Abstract: A system (10, 200) and corresponding method for introducing a drug delivery device (18, 218) through at least part of a biological barrier starts with the drug delivery device (18, 218) deployed within a channel of a hollow needle (12, 212). The hollow needle is inserted into the biological barrier and the drug delivery device is pushed forward by a suitable plunger. The drug delivery device is preferably anchored to the biological barrier through a radially expanding retention arrangement (36, 36', 36a), and is preferably filled with a liquid drug after deployment via a filling needle (24, 222) extending within the channel of the hollow needle (12, 212).

Miniature Implanted Drug Delivery Devices and Inserter Systems for Introducing Such Devices

FIELD AND BACKGROUND OF THE INVENTION

The present invention relates to miniature drug delivery devices and, in particular, it concerns drug delivery devices with particularly low flow rates, and inserter systems for introducing such devices into the body.

It is known to provide an implantable device which delivers a drug slowly over a period of time. This approach avoids problems of patient compliance, and provides particular advantages where delivery of a drug to a specific target location allows use of much lower overall dosage than would be required for systemic delivery, possibly avoiding undesirable side effects.

In some cases, slow drug delivery is achieved by providing a drug is dispersed in a matrix of resorbable material and is gradually released as the matrix breaks down in the body. Examples of this approach may be found in US Patents Nos. 4351337 and 4450150 to Sidman. This approach typically does not achieve highly uniform drug release rates, and is not suitable for drugs which must be delivered in a liquid form or which have high diffusion rates through the matrix materials.

Examples of implantable devices for delivery of liquid drugs include, but are not limited to, US Patents Nos. 5163920, 4428397, 4820273, 5061242, 5993414, 6183461 and 5836935.

Certain potential applications of such devices impose particularly demanding conditions. For example, ocular applications pose a challenge as to how to anchor an implanted device so that it does not drift within the internal cavity of the eye. Furthermore, the procedure for introducing and anchoring a highly miniature drug delivery device is difficult to perform reliably and safely.

There is therefore a need for inserter systems for introducing and anchoring miniature drug delivery devices into a body structure such as the eye.

SUMMARY OF THE INVENTION

The present invention is a system and method for introducing miniature drug delivery devices. The invention also provides various structures for such miniature drug delivery devices.

5 According to the teachings of an embodiment of the present invention there is provided, a system for introducing a drug delivery device, the system comprising: (a) a hollow needle having a central channel and a tip; (b) a drug delivery device comprising a reservoir and a proximal filling port deployed to allow introduction of a volume of a liquid drug into the reservoir to be released over a period of time, the
10 drug delivery device being deployed within the central channel of the hollow needle; (c) a plunger displaceable so as to push the drug delivery device along the central channel and beyond the tip of the hollow needle; and (d) a liquid injection device including a filling needle extending within the central channel of the hollow needle for engaging the filling port so as to allow filling of the reservoir after at least part of
15 the reservoir has been advanced beyond the hollow needle, such that, when the tip of the hollow needle is inserted into or through a biological barrier of a body and the plunger is displaced towards the tip, the drug delivery device is advanced beyond the hollow needle for deployment within the body and filling by the liquid injection device.

20 According to a further feature of an embodiment of the present invention, the reservoir is an inflatable reservoir sized for deployment within the central channel when substantially empty and inflatable by filling with a liquid drug to assume a deployed size greater than dimensions of the central channel.

 According to a further feature of an embodiment of the present invention, the
25 filling needle is extends through at least part of the plunger such that the filling needle engages the filling port prior to advancing of the drug delivery device.

 According to a further feature of an embodiment of the present invention, the drug delivery device deployed within the hollow needle, the plunger, the filling needle and the liquid injection device pre-filled with a quantity of a liquid drug, are

preassembled into a single integrated delivery system for implanting and filling the drug delivery device within the body.

According to a further feature of an embodiment of the present invention, the tip of the hollow needle is a beveled tip terminating at a point, so as to facilitate
5 introduction of the drug delivery device into a biological barrier without formation of a prior incision.

According to a further feature of an embodiment of the present invention, there is also provided an abutment surface surrounding at least part of the hollow needle so as to define a depth of penetration, and wherein the plunger has a
10 predefined fully-advanced position, such that, after penetration of the tip into or through the biological barrier and advancing of the plunger, the drug delivery device extends to a predefined depth into the body.

According to a further feature of an embodiment of the present invention, the drug delivery device includes a radially expandable retention configuration
15 configured to anchor the drug delivery device within a layer, or between layers, of the biological barrier.

According to a further feature of an embodiment of the present invention, the radially expandable retention configuration includes an expander element resiliently biased to a size greater than the central channel and temporarily compressed for
20 insertion into the central channel.

According to a further feature of an embodiment of the present invention, the radially expandable retention configuration includes a flexible sleeve deployed around an external surface of the drug delivery device while within the hollow needle and configured to become axially compressed and radially expanded during
25 advancing of the drug delivery device from the hollow needle.

There is also provided according to an embodiment of the present invention, a method for introducing a drug delivery device, the method comprising the steps of: (a) inserting the drug delivery device via a channel of a hollow needle through at least part of at least one layer of a biological barrier; and (b) subsequent to the

inserting, filling the drug delivery device with a quantity of liquid drug via a filling needle extending within the channel of the hollow needle.

There is also provided according to an embodiment of the present invention, a system for introducing a drug delivery device, the system comprising: (a) a hollow
5 needle having a central channel and a tip; (b) a drug delivery device comprising a reservoir for receiving a volume of a liquid drug to be released over a period of time, the drug delivery device being deployed within the central channel of the hollow
10 needle; and (c) a plunger displaceable so as to push the drug delivery device along the central channel and beyond the tip of the hollow needle, wherein the drug delivery device includes a radially expandable retention configuration configured to anchor the drug delivery device within a layer, or between layers, of the biological
15 barrier such that, when the tip of the hollow needle is inserted into or through a biological barrier of a body and the plunger is displaced towards the tip, the drug delivery device is advanced beyond the hollow needle and becomes anchored by the radially expandable retention configuration within a layer, or between layers, of the biological barrier.

According to a further feature of an embodiment of the present invention, the radially expandable retention configuration includes an expander element resiliently biased to a size greater than the central channel and temporarily compressed for
20 insertion into the central channel.

According to a further feature of an embodiment of the present invention, the radially expandable retention configuration includes a flexible sleeve deployed around an external surface of the drug delivery device while within the hollow needle and configured to become axially compressed and radially expanded during
25 advancing of the drug delivery device from the hollow needle.

There is also provided according to an embodiment of the present invention, a method for introducing a drug delivery device, the method comprising the steps of: (a) inserting the drug delivery device through at least part of at least one layer of a biological barrier; and (b) anchoring the drug delivery device within a layer, or

between layers, of the biological barrier, wherein the steps of inserting and anchoring are performed sequentially by use of a single deployment system.

According to a further feature of an embodiment of the present invention, subsequent to the inserting, the drug delivery device is filled with a quantity of liquid drug, wherein the step of filling is additionally performed by use of the single deployment system.

There is also provided according to an embodiment of the present invention, a system for introducing a drug delivery device, the system comprising: (a) a hollow needle having a central channel and a beveled tip terminating at a point; (b) a drug delivery device comprising a reservoir for receiving a volume of a liquid drug to be released over a period of time, the drug delivery device being deployed within the central channel of the hollow needle; and (c) a plunger displaceable so as to push the drug delivery device along the central channel and beyond the tip of the hollow needle, such that, when the tip of the hollow needle is inserted into or through a biological barrier of a body and the plunger is displaced towards the tip, the drug delivery device is advanced beyond the hollow needle for deployment within the body.

BRIEF DESCRIPTION OF THE DRAWINGS

The invention is herein described, by way of example only, with reference to the accompanying drawings, wherein:

FIG. 1A is a side view of a system, constructed and operative according to an embodiment of the present invention, for introducing a drug delivery device through a biological barrier, illustrated here schematically as the surface of a human or animal eye;

FIG. 1B is a longitudinal cross-sectional view taken through the view of FIG. 1A;

FIG. 2 is an enlarged view of the region of FIG. 1B designated V1;

FIGS. 3A-3C are enlarged views of a region of FIG. 1A showing an initial state and two successive states during preparation of the system for use;

FIGS. 4A and 4B are enlarged views of the region of FIG. 2 designated V2, and showing the relative positions of a filling needle and the drug delivery device corresponding to the states of FIGS. 3A and 3C, respectively;

FIGS. 5A-5C show a preferred sequence for insertion of a hollow needle of the delivery device into or through at least one layer of the biological barrier;

FIGS. 6A and 6B are views similar to FIGS. 3A-3C showing a further two successive states during advancing of the drug delivery device into or through at least one layer of the biological barrier;

FIG. 7A is a view similar to FIG. 1B corresponding to the state of FIG. 6B;

FIG. 7B is an enlarged view of the region of FIG. 7A designated V11;

FIG. 8A is an enlarged view of the region of FIG. 7B designated V12, showing the deployed drug delivery device extending into the body;

FIG. 8B is an enlarged view of the region of FIG. 8A designated V13, showing a first implementation of a radially expandable retention configuration partially deployed to anchor the drug delivery device between layers of the biological barrier;

FIG. 8C is a view similar to FIG. 8A after operation of a filling syringe to fill an inflatable reservoir of the drug delivery device;

FIG. 8D is a view similar to FIG. 8C after withdrawal of the deployment system, leaving the drug delivery device anchored between layers of the biological barrier and extending into the body for slow release drug delivery;

FIG. 8E is a view similar to FIG. 8D illustrating a variant implementation of an anchoring configuration for anchoring the drug delivery device of the above embodiment in the layers of the biological barrier;

FIG. 8F is a view similar to FIG. 8D illustrating a further variant implementation of an anchoring configuration for anchoring the drug delivery device of the above embodiment in the layers of the biological barrier;

FIGS. 9A and 9B are views similar to FIGS. 4A and 4B, respectively, illustrating a variant implementation of the drug delivery device with an alternative implementation of the radially expandable retention configuration;

FIGS. 10A-10D are views similar to FIGS. 8A-8D, respectively, for the
5 variant implementation of FIGS. 9A and 9B;

FIG. 10E is an enlarged view of the region of FIG. 10D designated V19, illustrating the deployed state of the alternative implementation of the radially expandable retention configuration;

FIGS. 11A-11D are schematic longitudinal cross-sectional views taken
10 through a system, constructed and operative according to an embodiment of the present invention, for introducing a drug delivery device through a biological barrier, showing a sequence of states during filling of a delivery system with a liquid drug in preparation for deployment of the drug delivery device;

FIGS. 12A-12E are schematic longitudinal cross-sectional views taken
15 through the system of FIGS. 11A-11D, showing a sequence of states during deployment and filling of the drug delivery device;

FIG. 13 is a schematic representation of a device according to an aspect of the present invention for refilling a drug delivery device according to the present invention; and

20 FIGS. 14A and 14B are schematic longitudinal cross-sectional views illustrating a retrieval system for retrieving a drug delivery device according to the present invention, the retrieval system being shown in an insertion state and a retrieval state, respectively.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

25 The present invention is, according to a first aspect, a system and method for introducing a drug delivery device into a body. This first aspect of the invention may be implemented with a range of drug delivery devices, particularly those having a reservoir for storing a quantity of liquid drug for slow release. Other aspects of the

present invention relate to devices and methods for refilling and retrieving the corresponding drug delivery devices.

The principles and operation of systems, methods and devices according to the present invention may be better understood with reference to the drawings and
5 the accompanying description.

Referring now to the drawings, FIGS. 1A-8F illustrate the structure and operation of a first exemplary embodiment of a system, generally designated **10**, and a corresponding method for introducing a drug delivery device through at least one layer of a biological barrier into a body, illustrated here with reference to a non-
10 limiting example of the human eye **100**.

In general terms, system **10** includes a hollow needle **12**, having a central channel **14** and a tip **16**, and a drug delivery device **18** comprising a reservoir for receiving a volume of a liquid drug to be released over a period of time. Drug delivery device **18** is small relative to the overall dimensions of system **10**, and is
15 best seen in FIGS. 4A, 4B and 8A-8D. Drug delivery device **18** is initially deployed within central channel **14** of hollow needle **12**. A plunger **20** is displaceable within central channel **14** so as to push the drug delivery device beyond the tip of the hollow needle. When tip **16** of hollow needle **12** is inserted into or through a biological barrier of a body and plunger **20** is displaced towards tip **16**, drug delivery device **18**
20 is advanced beyond the hollow needle for deployment within the body, as shown in FIGS. 7A-8A.

In certain particularly preferred embodiments, the reservoir of drug delivery device **18** is an inflatable reservoir sized for deployment within central channel **14** when substantially empty, and inflatable by filling with a liquid drug to assume a
25 deployed size greater than dimensions of the central channel. Thus, when empty, drug delivery device **18** has lateral dimensions less than the inner diameter of channel **14**, as shown in FIGS. 4A & 8A. After filling, as shown in FIGS. 8C and 8D, the dimensions are increased such that at least two, and typically all three, dimensions of drug delivery device are greater than the inner diameter of channel **14**.

The aforementioned approach of introducing the drug delivery device in a collapsed state and subsequently filling it facilitates introduction of the device via a much smaller needle canula than would otherwise be possible. Thus, certain preferred embodiments of the present invention employ a hollow needle **12** with
5 internal diameter less than 2 millimeters. In certain particularly delicate applications such as ocular application, it may be preferred to employ internal diameters of less than 1 millimeter, and most preferably less than 0.5 millimeter. The deployed volume of the drug delivery device after filling is typically at least 10 times greater than the empty volume during deployment, thereby facilitating controlled delivery of
10 relatively large quantities of drug.

According to certain particularly preferred embodiments of the present invention, introduction of the substantially empty drug delivery device and filling of the device are performed sequentially using a single deployment system, as illustrated here. Thus, in the preferred example illustrated here, system **10** includes a
15 liquid injection device, such as a filling syringe **22** (visible in FIG. 1A), linked to a filling needle **24** (best seen in FIG. 4A). Filling needle **24** extends within central channel **14** for engaging a filling port **26** of drug delivery device **18**, thereby allowing filling of the reservoir after at least part of the reservoir has been advanced beyond the hollow needle.

20 In the particularly preferred non-limiting example illustrated herein, filling needle **24** is integrated with plunger **20** so that the filling needle engages the filling port prior to advancing of the drug delivery device. In this case, according to certain preferred implementations, the engagement of filling needle **24** with filling port **26** is performed as a preparatory step prior to insertion of hollow needle **12** into the
25 biological barrier. This is illustrated here with reference to the external views of FIGS. 3A-3C, which show guided retraction of a distal tip portion **28** of system **10** relative to a main block **30** of the deployment device. In the implementation shown here, the length of this initial movement is limited by a pin-and-slot inter-engagement, shown here as a pin **32** of main block **30** which is engaged in a shaped

slot **34** formed in distal tip portion **28**. The resulting engagement of filling needle **24** with filling port **26** is shown in FIG. 4B.

In a particularly preferred implementation, drug delivery device **18** is introduced and filled sequentially by a single integrated delivery system. Thus, the
5 delivery system includes hollow needle **12**, plunger **20**, filling needle **24** and filling syringe **22**, pre-filled with a quantity of a liquid drug, as illustrated in FIG. 1B.

Turning now to additional features of the preferred exemplary embodiment of FIGS. 1A-8F, needle tip **16** is illustrated in the drawings as being a beveled tip which provides a sharp point. Such an embodiment allows the entire procedure, from initial
10 penetration of the tissue through deployment and filling of the drug delivery device, to be performed using the system of the invention without prior preparation of the insertion site. It should be noted however that alternative implementations may employ a flat-ended or otherwise blunt hollow needle **12**, and rely upon a separately made prior incision, as is known in various other procedures, particularly in the field
15 of ocular procedures where a self-sealing flap may in some cases be cut more easily with a flat cutting tool.

In the case of a pointed needle tip, a preferred sequence for insertion of needle tip **16** is illustrated schematically in FIGS. 5A-5C. According to this sequence, initial penetration of needle tip **16** is performed while the device is significantly inclined
20 relative to the local perpendicular to the tissue surface so that the needle tip penetrates through the outer tissue layers as an oblique angle (FIG. 5B). The device is then straightened to an upright position relative to the local tissue surface (FIG. 5C) prior to completion of the drug delivery device insertion. Without in any way limiting the scope of the invention, the aforementioned angular motion is believed to
25 generate relative lateral displacement between the outer layers of the layered tissue. As a result, after removal of the inserter device, elastic return of the tissue to its original position tends to result in the penetration openings being out of alignment, thereby providing a self-sealing function.

Referring to FIG. 2, distal tip portion **28** preferably provides an abutment surface **28a** surrounding at least part of hollow needle **12** so as to define a depth of penetration of the needle. Additionally, plunger **20** has a predefined fully-advanced position. In the example illustrated here, plunger **20** is integrated with main block **30**,
5 which also features a forward projection **30a** configured to enter a corresponding recess **28b** in the rear of distal tip portion **28**. Forward projection **30a** is here shown as a cylindrical tube and recess **28b** as a corresponding cylindrical recess. These structures define a fully advanced position of main block **30**, and hence also of plunger **20**, relative to distal tip portion **28**, and also provides concentric guiding
10 surfaces that help to maintain alignment of the components during use. The combined effect of the depth-limiting abutment surface **28a** and the predefined range of motion of plunger **20** is to predefine an insertion depth to which drug delivery device **18** extends into the body once deployed, as illustrated in FIG. 7B.

According to another aspect of certain preferred embodiments of the present
15 invention, drug delivery device **18** includes a radially expandable retention configuration **36** configured to anchor the drug delivery device within a layer, or between layers, of the biological barrier. According to a first implementation of this feature, as illustrated in FIGS. 8B and 8D, radially expandable retention configuration **36** includes an expander element resiliently biased to a size greater
20 than the central channel and temporarily compressed for insertion into the central channel. The expander element is typically a C-shaped resilient partial-ring which is squeezed closed or otherwise compressed for insertion within central channel **14**, and which returns resiliently to an oversize diameter as it emerges from tip **16**.

In the example of ocular deployment as exemplified in the drawings, the
25 structure of the biological barrier of eye **100** is made up of a number of different layers, here designated schematically as layers **102**, **104** and **106**, corresponding to the conjunctiva, sclera and choroid, respectively. In this case, it is particularly preferred that the length of hollow needle **12** projecting from abutment surface **28a** is chosen to correspond roughly to the depth of the boundary between two of the layers

104 and **106**, so that expandable retention configuration **36** becomes lodged between layers **104** and **106** when deployed. Radially expandable retention configuration **36** then serves to anchor one end of drug delivery device **18** between the layers after removal of the delivery system, as shown FIG. 8D.

5 FIG. 8E shows a variant implementation in which the depth of penetration is chosen such that radially expandable retention configuration lodges itself within the thickness of the sclera layer **104**, thereby relying on the relatively high structural strength of the sclera for positive anchoring of the device. FIG. 8F shows a further variant in which two radially expandable retention elements **36** and **36'** are spaced
10 apart to provide two-location anchoring, preferably corresponding to the inner and outer surfaces of the sclera **104**.

Referring parenthetically to FIGS. 9A-10E, these illustrate an alternative implementation of a radially expandable retention element in which the retention element includes a flexible sleeve **36a**, best seen in FIGS. 10B and 10E. Flexible
15 sleeve **36a** is deployed around an external surface of the drug delivery device while within the hollow needle, and is made from sufficiently loose and flexible material that it tends to become caught on surrounding tissue as it emerges from hollow
needle **12**, thereby becoming axially compressed and radially expanded (i.e., forming an outward fold or “kink”) during advancing of the drug delivery device from the
20 hollow needle. Hollow needle **12** is inserted through layer **102** and at least part of layer **104**, so that flexible sleeve **36a** does not interact with those layers as device **18** is advanced into the tissue. When flexible sleeve **36a** starts to leave needle **12** and encounters the tissue layer **106**, the sleeve becomes caught on the tissue and slides along the outside of the drug delivery device. The bunching up of the sleeve material
25 forms the aforementioned outward fold which tends to be caught between the layers of the biological barrier and provide the aforementioned anchoring effect. In all other respects, the device of FIGS. 9A-10E may be implemented in a system and method similar to those described in FIGS. 1A-8D.

Although shown here in the context of a multi-layer biological barrier, it should be noted that similar structures may be used to anchor the device of the present invention relative to a single layer barrier, or relative to other body structures, all as will be clear to a person having ordinary skill in the art according to the particular intended application.

Regarding drug delivery device **18** itself, this aspect of the present invention may be implemented with a wide range of drug delivery devices that are based on an inflatable bladder-type reservoir for storing and slowly releasing a quantity of a liquid drug composition or the like. A number of non-limiting examples of suitable drug delivery devices may be found in co-pending international application publication no. WO 2011/101833 filed February 22, 2011, and related US patent application no. 13/430,730, filed 27 March, 2012, which do not constitute prior art. These examples regulate flow rates for release of the liquid drug by employing fine channels formed in facing surfaces between two parts of the drug delivery device, preferably also provide pressure-responsive regulation of the flow rate, or by use of a flow path passing through a porous block which may be biodegradable.

The operation of system **10** will now be clearly understood. In preparation for use, distal tip portion **28** is first retracted towards main block **30** to engage filling needle **24** with filling port **26**, as illustrated in FIGS. 3A-3C and 4A-4B. Needle tip **16** is then introduced as illustrated in FIGS. 5A-5C, or via a preformed incision, and main block **30** is advanced as shown in FIGS. 6A-6B to deploy drug delivery device within the body, as shown in FIG. 7A-8A. During introduction of the device, the radially expandable retention configuration **36** or **36a** is at least partially deployed (FIGS. 8B and 10B). The piston of filling syringe **22** is then advanced so as to fill the reservoir of drug delivery device **18** (FIGS. 8C and 10C), and the deployment system is withdrawn, leaving the deployed and operating drug delivery device **18** anchored between layers of the biological barrier as shown in FIGS. 8D and 10D.

Turning now to FIGS. 11A-12E, there is shown a variant embodiment of a system, generally designated **200** for introducing a drug delivery device through at

least one layer of a biological barrier into a body, such as the human eye. System **200** is generally analogous in structure and function to system **10** described above.

Thus, in general terms, similar to system **10**, system **200** includes a hollow needle **212**, having a central channel **214** and a tip **216**, and a drug delivery device **218** which includes a reservoir for receiving a volume of a liquid drug to be released over a period of time. Drug delivery device **218** is small relative to the overall dimensions of system **200** and is initially deployed within central channel **214** of hollow needle **212**.

One distinguishing feature of system **200** is that sequential deployment and filling of drug delivery device **218** are effected by a single continuous motion of a manually operated actuator **220**, thereby simplifying operation of the system, as will now be detailed.

As seen in FIG. 11A, system **200** includes a filling needle **222** which extends through a hollow spacer **224** to a penetrating tip **226** which is initially embedded in a seal **228**. The opposite end of filling needle **222** passes through, and is integrated with, a first piston **230** which can slide in sealing contact with the inner bore of a syringe body **234**. A second piston **232**, also in sealing contact with the inner bore of syringe body **234**, is initially in close contact with first piston **230**.

Prior to use, system **200** is prepared by introduction of the liquid drug to a storage volume defined within syringe body **234** between pistons **230** and **232**. A non-limiting exemplary implementation of this preparatory process is illustrated in FIGS. 11A-11D. In the implementation shown here, an adapter **236** includes alignment features **238** for centering an adapter needle **240** within the bore of syringe body **234** such that, as the adapter is inserted as shown in FIGS. 11A and 11B, the tip of adapter needle **240** penetrates an elastomer septum of second piston **232** to reach a small recess between the two pistons. The required quantity of drug is then introduced via adapter **236**, typically by attachment of a standard syringe (not shown) to the rear of the adapter. Introduction of the liquid drug forces the two pistons apart, advancing first piston **230** to the position illustrated in FIG. 11C to make space for

the drug which occupies volume **244**. This motion of piston **230** causes a corresponding advancing of filling needle **222** so that the penetrating tip **226** emerges from seal **228** and penetrates through the septum (not shown) of drug delivery device **218**. Drug delivery device **218** is prevented from moving at this stage by the presence
5 of a protective cap **242** which includes an elastomer stopper **246**. Adapter **236** is then removed (FIG. 11D), leaving system **200** ready for use.

For deployment of the drug delivery device through a biological membrane, a manually operated actuator **220**, shown here in a preferred embodiment in the form of a plunger, is attached to second piston **232**, as illustrated in FIG. 12A, and
10 protective cap **242** is removed, leaving the system as illustrated in FIG. 12B. Tip **216** of hollow needle **212** is then inserted into the biological barrier, preferably in a manner similar to that described above with reference to FIGS. 5A-5C.

Manual pressure is then applied to plunger **220** to advance it along the body of syringe **234**. Pressure applied by the plunger to piston **232** applies pressure to the
15 liquid drug which in turn applies pressure to piston **230**. This pressure advances piston **230**, which presses against spacer **224**, thereby advancing spacer **224**, seal **228** and drug delivery device **218** along hollow needle **212** until drug delivery device **218** is deployed in the desired position beyond the tip of the deployment system (FIG. 12C). Thus plunger **220**, piston **232**, liquid-filled volume **244**, piston **230**, spacer **224**
20 and seal **228** all function together as a compound plunger with which filling needle **222** is integrated. Until this point, drug delivery device **218** was prevented from inflating due to the walls of hollow needle **212** surrounding it. After leaving the confines of hollow needle **212**, and with piston **230** reaching a stop position at the end of the bore of syringe **234**, further displacement of plunger **220** and piston **232**
25 forces the liquid drug from volume **244** along filling needle **222**, thereby inflating the reservoir of drug delivery device **218**, as shown in FIG. 12D. Thus, both insertion of the drug delivery device and filling of the drug delivery device with liquid drug are both achieved sequentially by the advancing of plunger **220** unidirectionally along its

range of motion. The deployment system **200** is then withdrawn, leaving drug delivery device **218** in position for slow delivery of the drug over a period of time.

It should be noted that system **200** has been shown here schematically, and has only been detailed to an extent necessary to appreciate the distinctive features of this embodiment, while numerous other features that are similar to features of the embodiment described above have not been detailed here. For example, drug delivery device **218** preferably features one or more radially expandable retention elements, such as is described above with reference to FIGS. 8A-8D, 8E, 8F or 10A-10E. Similarly, the distal end of syringe **234** which surrounds hollow needle **212** preferably defines abutment surfaces for defining the extent of penetration of hollow needle **212**. Except where explicitly stated or self-evident otherwise, it should be understood that features described herein with reference to one embodiment may also be implemented in the context of other embodiments of the present invention.

Turning now to FIG. 13, as mentioned earlier, certain particularly preferred embodiments of the present invention provide an option for refilling of the drug delivery device deployed according to the present invention in situ. Given the very small dimensions of the drug delivery device according to certain implementations of the present invention, there is a need to ensure that a refilling needle has been correctly inserted through the refilling septum of the drug delivery device, and remains correctly inserted throughout the refilling process. There is also a need to assess the quantity of the drug present in the reservoir so as to ensure sufficient drug is provided without dangerously over-filling the device. For this purpose, the present invention provides a refilling device, generally designated **300**, constructed and operative according to a further feature of an embodiment of the present invention. Refilling device **300** is shown here schematically as a syringe type device including a syringe body **302** and a plunger **304** for delivering refill drug to the drug delivery device via a refill needle **306**. Additionally, device **300** includes a pressure measurement arrangement **308** deployed for measuring the fluid pressure within the syringe during operation, and a processing system **310** having a processor, a data

storage device and a display, connected so as to receive the fluid pressure measurements. Monitoring of the pressure after insertion of the device and prior to injection of its contents provides a reliable verification that the needle has correctly reached the elevated-pressure inner volume of the drug storage reservoir. A known and pre-calibrated profile mapping pressure to reservoir content may also facilitate evaluation of the degree of filling of the reservoir during the refilling process. Processing system **310** preferably stores a look-up table or parametrically defined model, preferably empirically derived, indicating the current reservoir contents as a function of the measured pressure within the syringe under zero-flow conditions.

Processing system thus preferably translates measured pressure into a visually displayed indication of the current reservoir content. The processor preferably also provides a warning function if the sensed pressure drops below the expected value, indicating that the refill needle **306** may not be correctly inserted into the drug delivery device or may have become dislodged. Most preferably, the refilling device **300** also features a safety valve, deployed for example as part of plunger **304**, which ensures release of the contained drug outside the patient in the event that the pressure exceeds a maximum allowed threshold value which might lead to a risk of damaging the drug delivery device.

Turning now to FIGS. 14A and 14B, there is shown a preferred embodiment of a retrieval device, generally designated **400**, for retrieving the drug delivery devices of the present invention. Retrieval device **400** preferably features a fine needle **402**, similar to the refilling needles described above, which is sufficiently fine to be inserted through the refilling port of the drug delivery device. Inside needle **402** is a resilient wire **404** pre-formed into a hook or otherwise curled form, but maintained in a straightened configuration by the small diameter inner lumen of needle **402**. An actuator **406** is manually deployable to selectively advance resilient wire **404** so as to extend beyond needle **402**.

In use, needle **402** is introduced into the drug delivery device via the refilling port while resilient wire **404** is in its straightened state. Wire **404** is then advanced

within the device, forming its hooked or otherwise curled form within the device, preferably so as to become anchored on or tangled around internal features of the drug delivery device. The retrieval device can then be removed, drawing after it the drug delivery device.

5 It will now be appreciated that various embodiments of the present invention provides a number of significant advantages over existing options in the field of implantable devices for delivering liquid medications. In particular, certain embodiments of the present invention provide one or more of the following functions:

- 10 • Insertion of the drug delivery device through at least part of at least one layer of a biological barrier and anchoring of the drug delivery device within a layer, or between layers, of the biological barrier, where both the inserting and anchoring are performed sequentially by use of a single deployment system.
- 15 • Insertion of the drug delivery device through at least part of at least one layer of a biological barrier and subsequent filling of the drug delivery device with a quantity of liquid drug, wherein both the inserting and filling are performed sequentially by use of a single deployment system.
- 20 • Introduction of the drug delivery device by a straightforward injection technique via a needle of diameter smaller than the dimensions of the liquid reservoir after filling.
- Introduction of the drug delivery device by a deployment device which predefines the depth of insertion, thereby facilitating consistent results without requiring unusual skill in the deployment process.

25 It will be appreciated that the above descriptions are intended only to serve as examples, and that many other embodiments are possible within the scope of the present invention as defined in the appended claims.

WHAT IS CLAIMED IS:

1. A system for introducing a drug delivery device, the system comprising:

- (a) a hollow needle having a central channel and a tip;
- (b) a drug delivery device comprising a reservoir and a proximal filling port deployed to allow introduction of a volume of a liquid drug into said reservoir to be released over a period of time, said drug delivery device being deployed within said central channel of said hollow needle;
- (c) a plunger displaceable so as to push said drug delivery device along said central channel and beyond said tip of said hollow needle; and
- (d) a liquid injection device including a filling needle extending within said central channel of said hollow needle for engaging said filling port so as to allow filling of said reservoir after at least part of said reservoir has been advanced beyond said hollow needle,

such that, when said tip of said hollow needle is inserted into or through a biological barrier of a body and said plunger is displaced towards said tip, said drug delivery device is advanced beyond said hollow needle for deployment within the body and filling by said liquid injection device.

2. The system of claim 1, wherein said reservoir is an inflatable reservoir sized for deployment within said central channel when substantially empty and inflatable by filling with a liquid drug to assume a deployed size greater than dimensions of said central channel.

3. The system of claim 1, wherein said filling needle is extends through at least part of said plunger such that said filling needle engages said filling port prior to advancing of said drug delivery device.

4. The system of claim 1, wherein said drug delivery device deployed within said hollow needle, said plunger, said filling needle and said liquid injection device pre-filled with a quantity of a liquid drug, are preassembled into a single integrated delivery system for implanting and filling said drug delivery device within the body.

5. The system of claim 1, wherein said tip of said hollow needle is a beveled tip terminating at a point, so as to facilitate introduction of said drug delivery device into a biological barrier without formation of a prior incision.

6. The system of claim 1, further comprising an abutment surface surrounding at least part of said hollow needle so as to define a depth of penetration, and wherein said plunger has a predefined fully-advanced position, such that, after penetration of said tip into or through the biological barrier and advancing of said plunger, said drug delivery device extends to a predefined depth into the body.

7. The system of claim 1, wherein said drug delivery device includes a radially expandable retention configuration configured to anchor said drug delivery device within a layer, or between layers, of the biological barrier.

8. The system of claim 7, wherein said radially expandable retention configuration includes an expander element resiliently biased to a size greater than said central channel and temporarily compressed for insertion into said central channel.

9. The system of claim 7, wherein said radially expandable retention configuration includes a flexible sleeve deployed around an external surface of said drug delivery device while within said hollow needle and configured to become axially compressed and radially expanded during advancing of said drug delivery device from said hollow needle.

10. A method for introducing a drug delivery device, the method comprising the steps of:

- (a) inserting the drug delivery device via a channel of a hollow needle through at least part of at least one layer of a biological barrier; and
- (b) subsequent to said inserting, filling the drug delivery device with a quantity of liquid drug via a filling needle extending within the channel of the hollow needle.

11. A system for introducing a drug delivery device, the system comprising:

- (a) a hollow needle having a central channel and a tip;
- (b) a drug delivery device comprising a reservoir for receiving a volume of a liquid drug to be released over a period of time, said drug delivery device being deployed within said central channel of said hollow needle; and
- (c) a plunger displaceable so as to push said drug delivery device along said central channel and beyond said tip of said hollow needle,

wherein said drug delivery device includes a radially expandable retention configuration configured to anchor said drug delivery device within a layer, or between layers, of the biological barrier such that, when said tip of said hollow needle is inserted into or through a biological barrier of a body and said plunger is displaced towards said tip, said drug delivery device is advanced beyond said hollow needle and becomes anchored by said radially expandable retention configuration within a layer, or between layers, of the biological barrier.

12. The system of claim 11, wherein said radially expandable retention configuration includes an expander element resiliently biased to a size greater than said central channel and temporarily compressed for insertion into said central channel.

13. The system of claim 11, wherein said radially expandable retention configuration includes a flexible sleeve deployed around an external surface of said drug delivery device while within said hollow needle and configured to become axially compressed and radially expanded during advancing of said drug delivery device from said hollow needle.

14. A method for introducing a drug delivery device, the method comprising the steps of:

- (a) inserting the drug delivery device through at least part of at least one layer of a biological barrier; and
- (b) anchoring the drug delivery device within a layer, or between layers, of the biological barrier,

wherein said steps of inserting and anchoring are performed sequentially by use of a single deployment system.

15. The method of claim 14, further comprising, subsequent to said inserting, filling the drug delivery device with a quantity of liquid drug, wherein said step of filling is additionally performed by use of the single deployment system.

16. A system for introducing a drug delivery device, the system comprising:

- (a) a hollow needle having a central channel and a beveled tip terminating at a point;
- (b) a drug delivery device comprising a reservoir for receiving a volume of a liquid drug to be released over a period of time, said drug delivery device being deployed within said central channel of said hollow needle; and
- (c) a plunger displaceable so as to push said drug delivery device along said central channel and beyond said tip of said hollow needle,

such that, when said tip of said hollow needle is inserted into or through a biological barrier of a body and said plunger is displaced towards said tip, said drug delivery device is advanced beyond said hollow needle for deployment within the body.

FIG. 1A

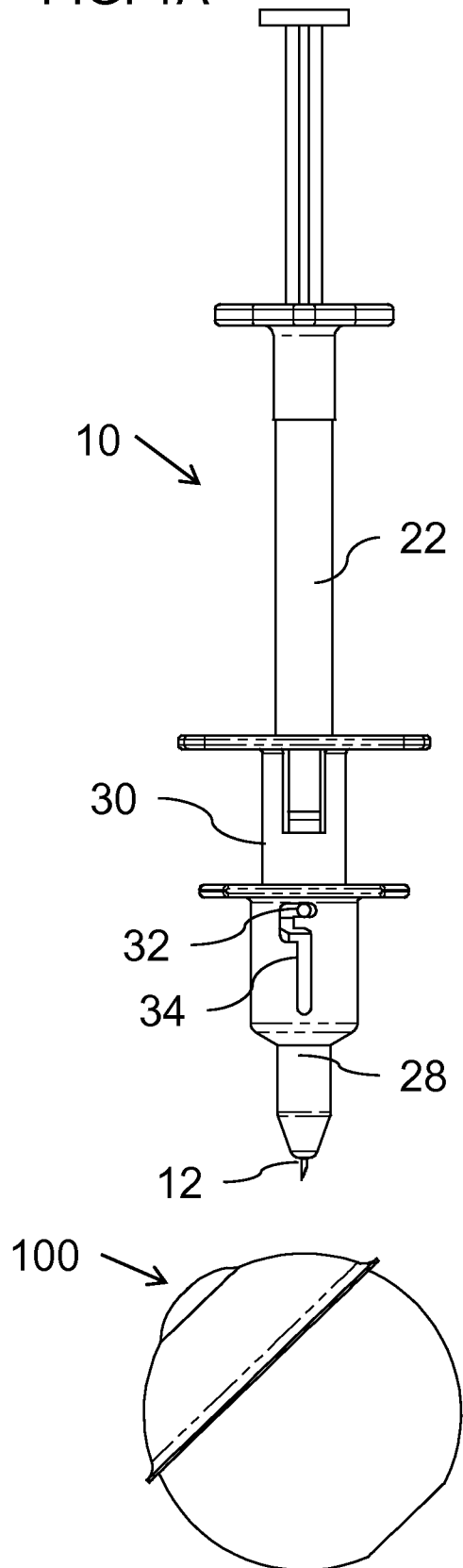


FIG. 1B

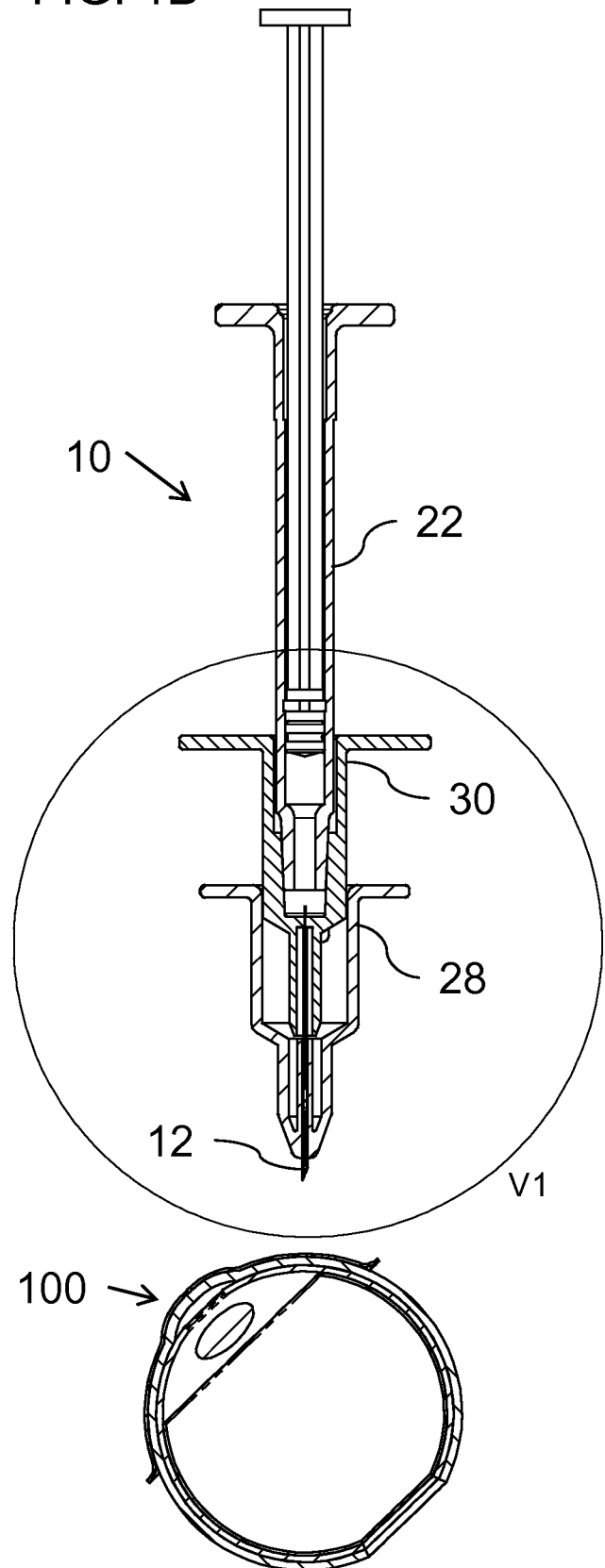


FIG. 2

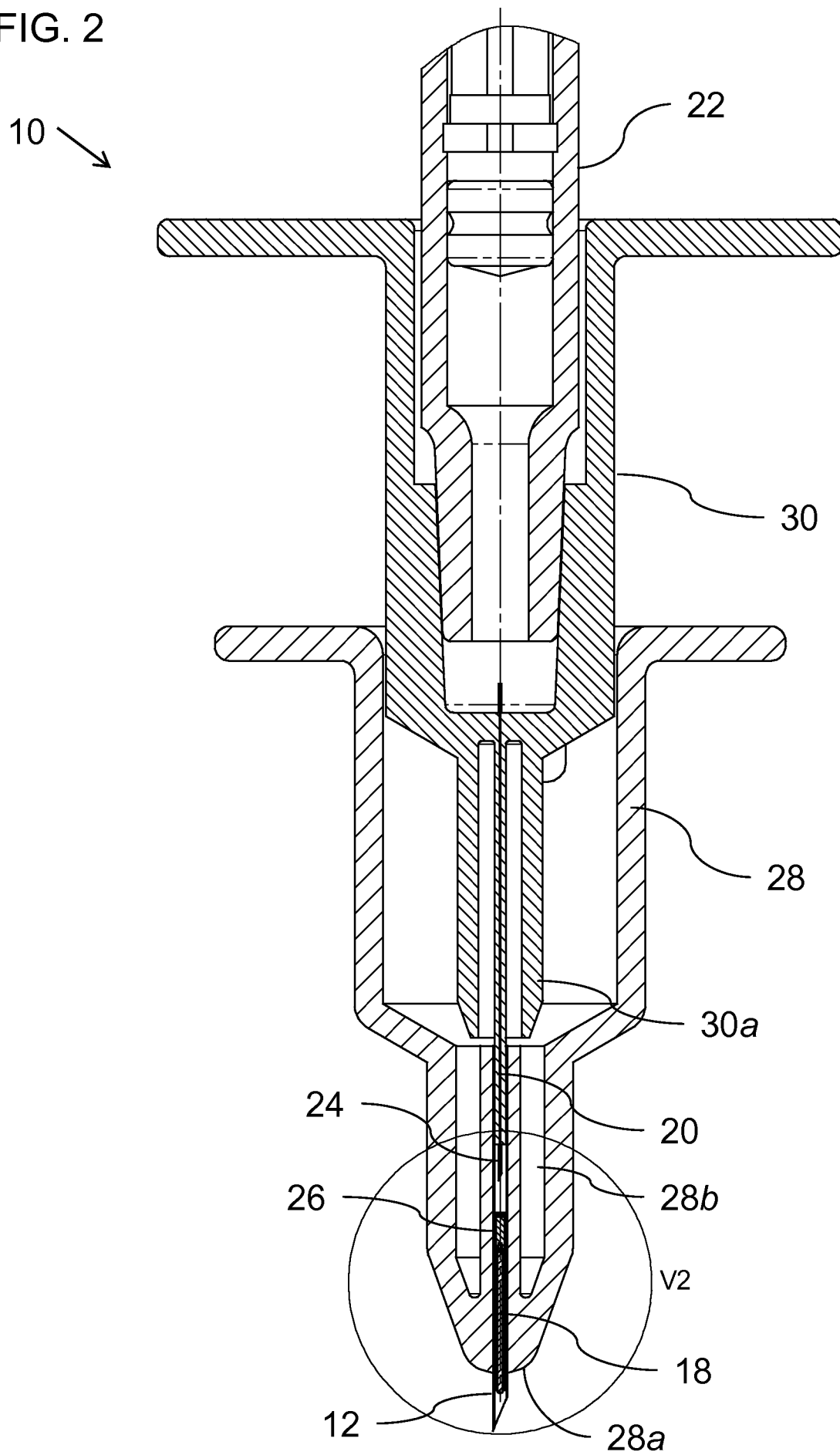


FIG. 3A

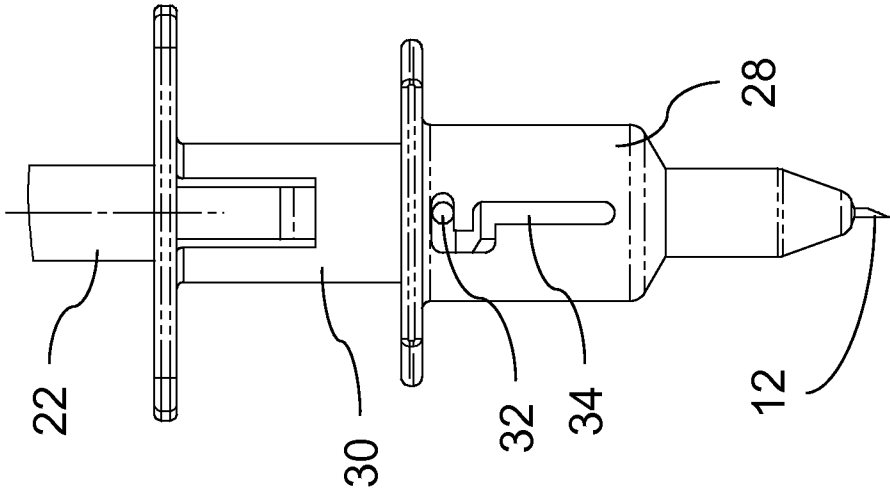


FIG. 3B

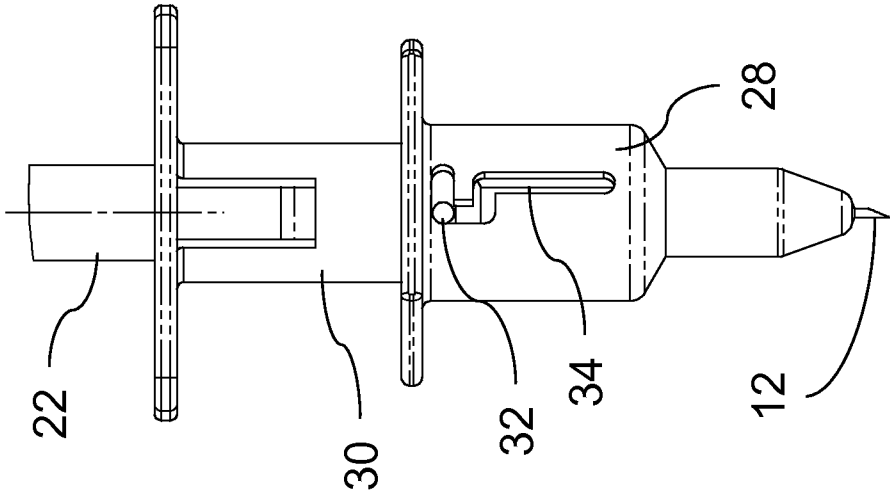


FIG. 3C

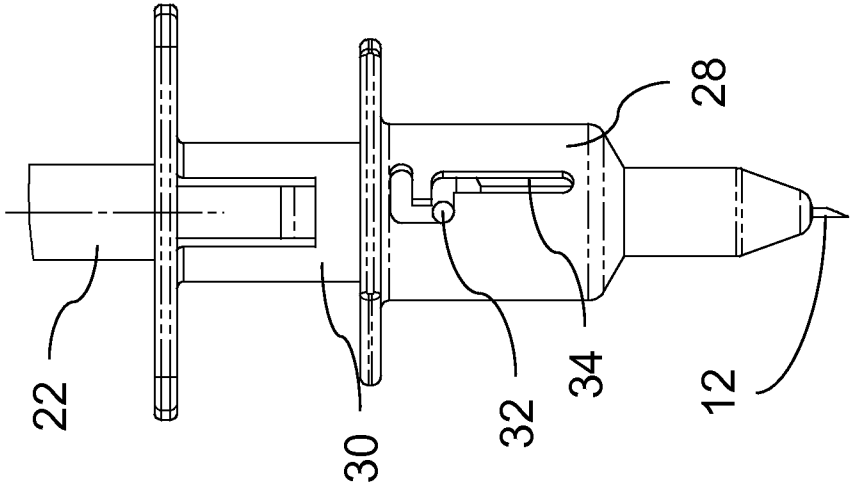


FIG. 4B

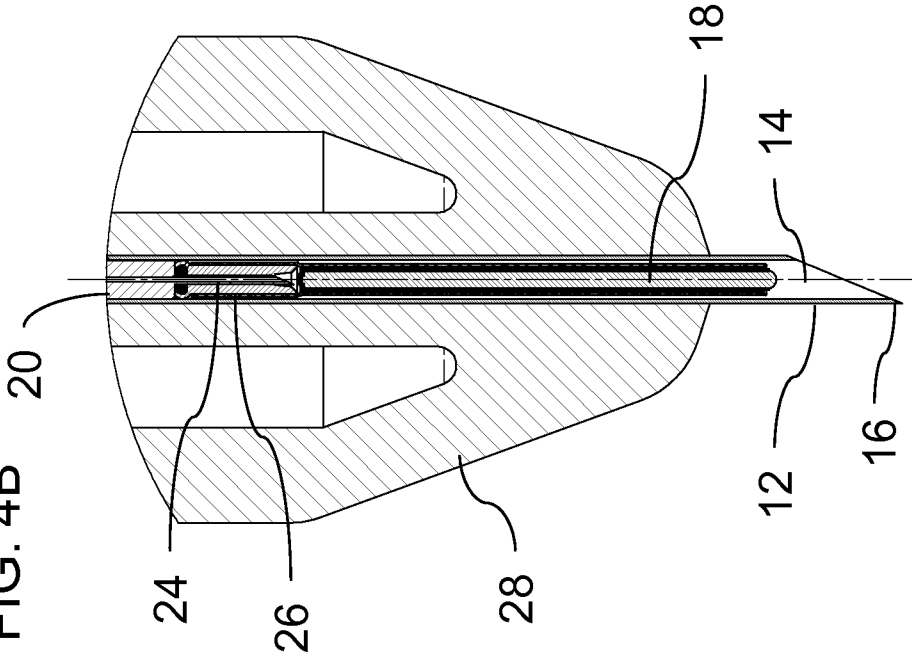


FIG. 4A

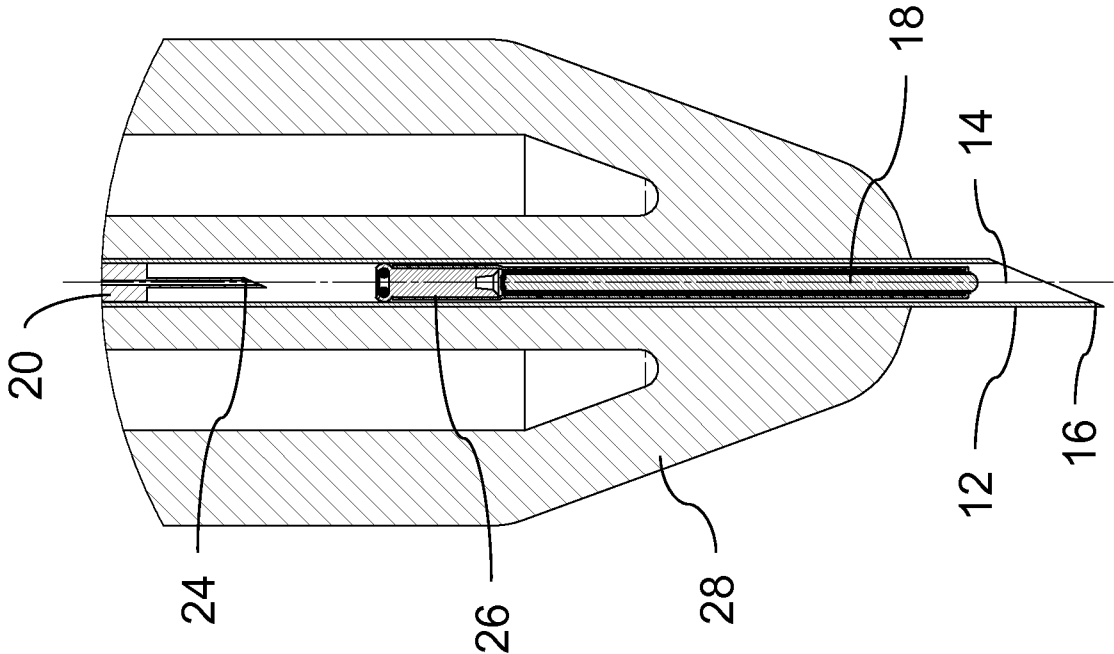


FIG. 5C

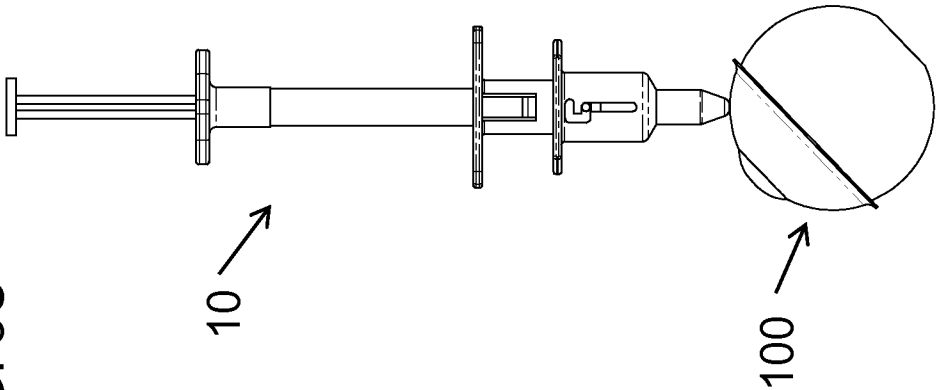


FIG. 5B

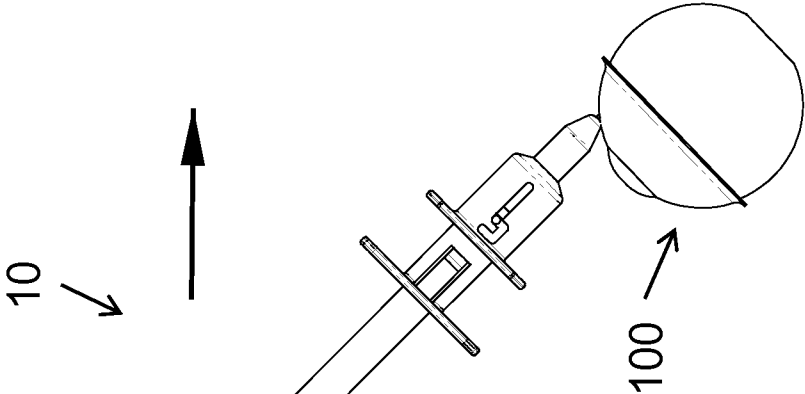


FIG. 5A

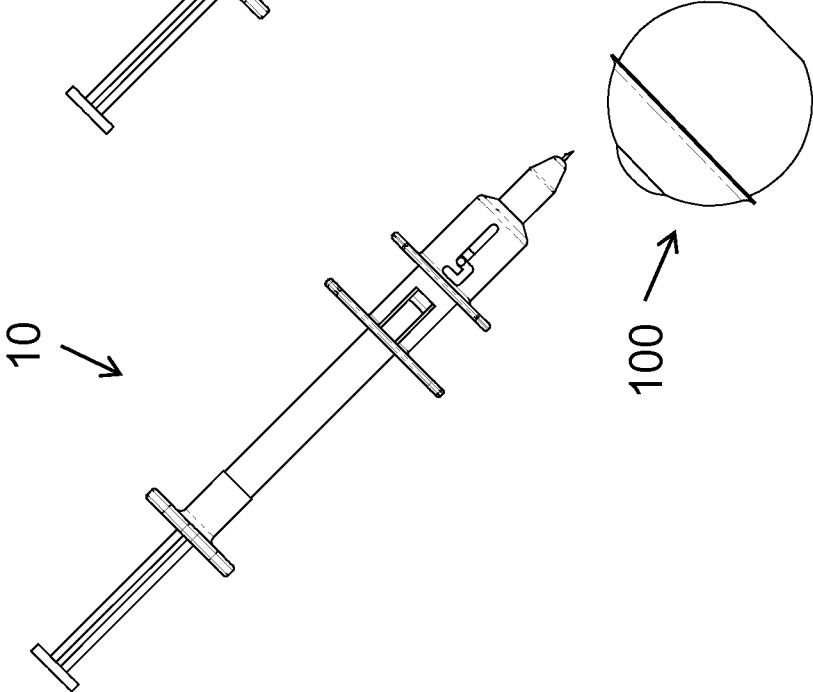


FIG. 6B

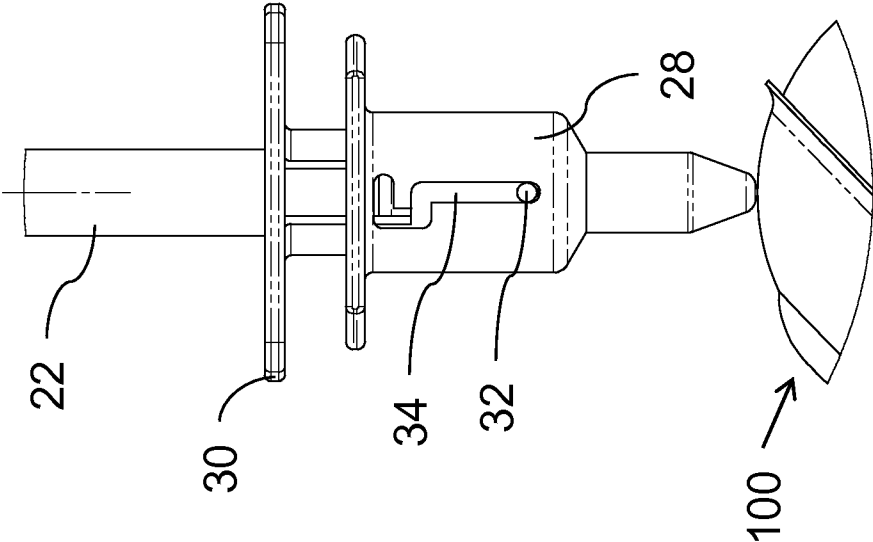
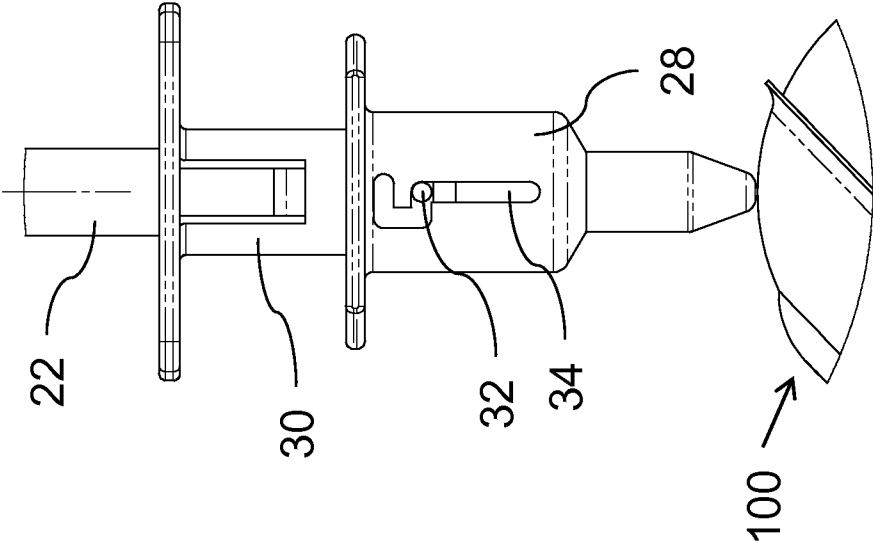
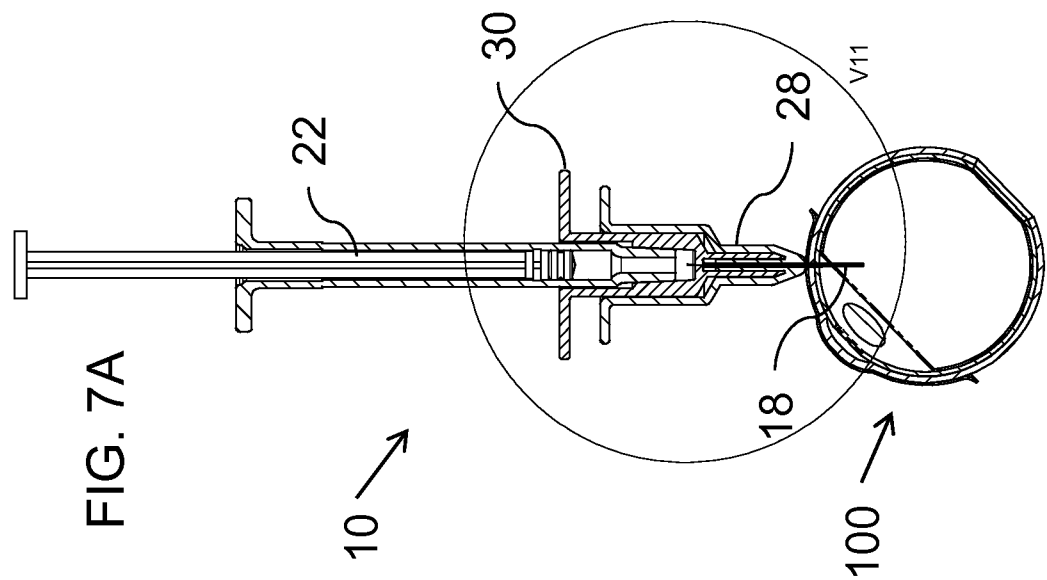
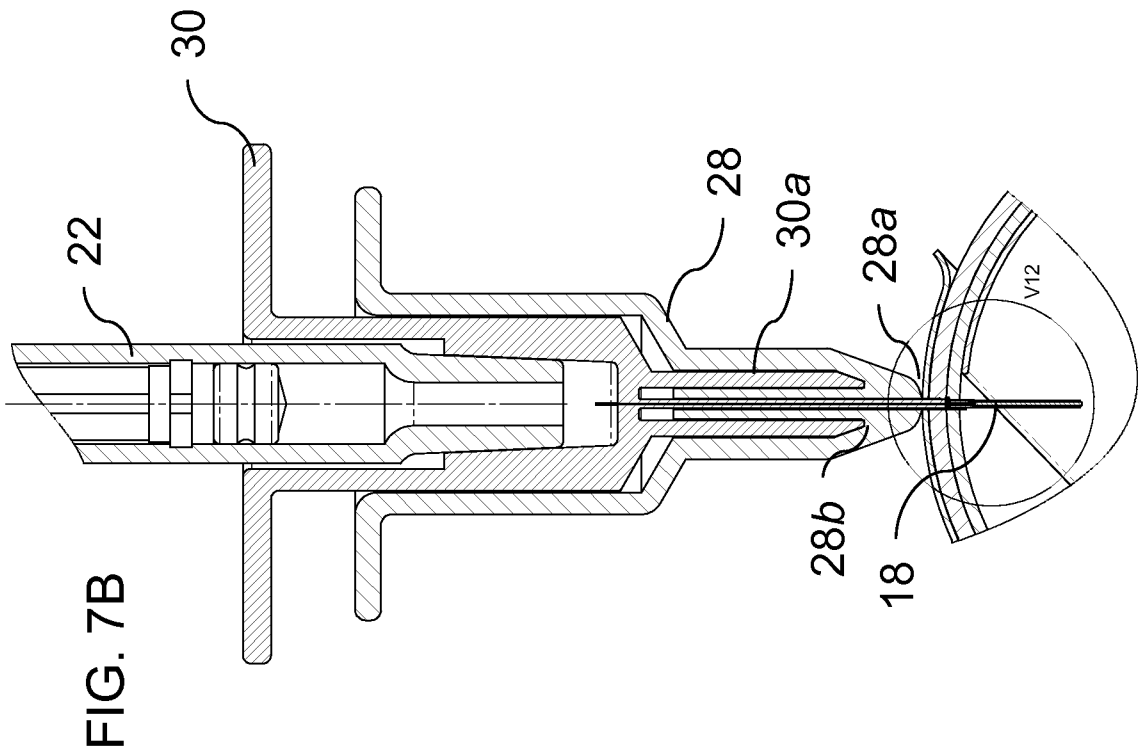


FIG. 6A





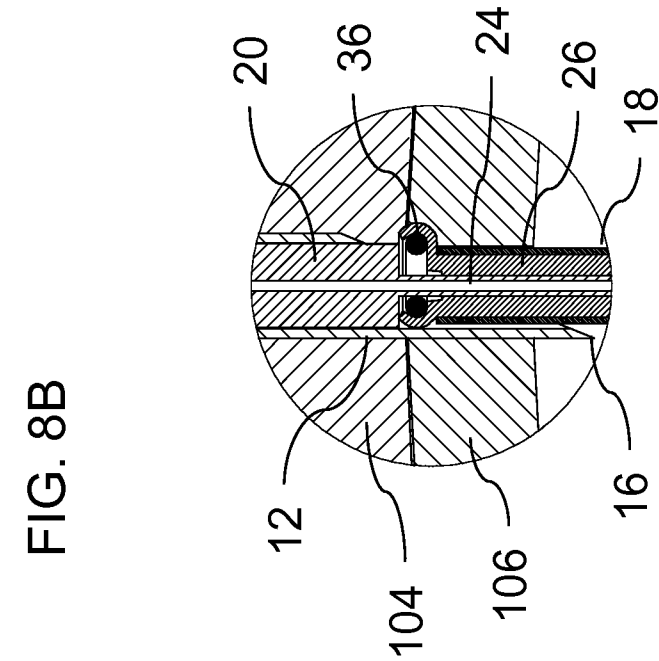
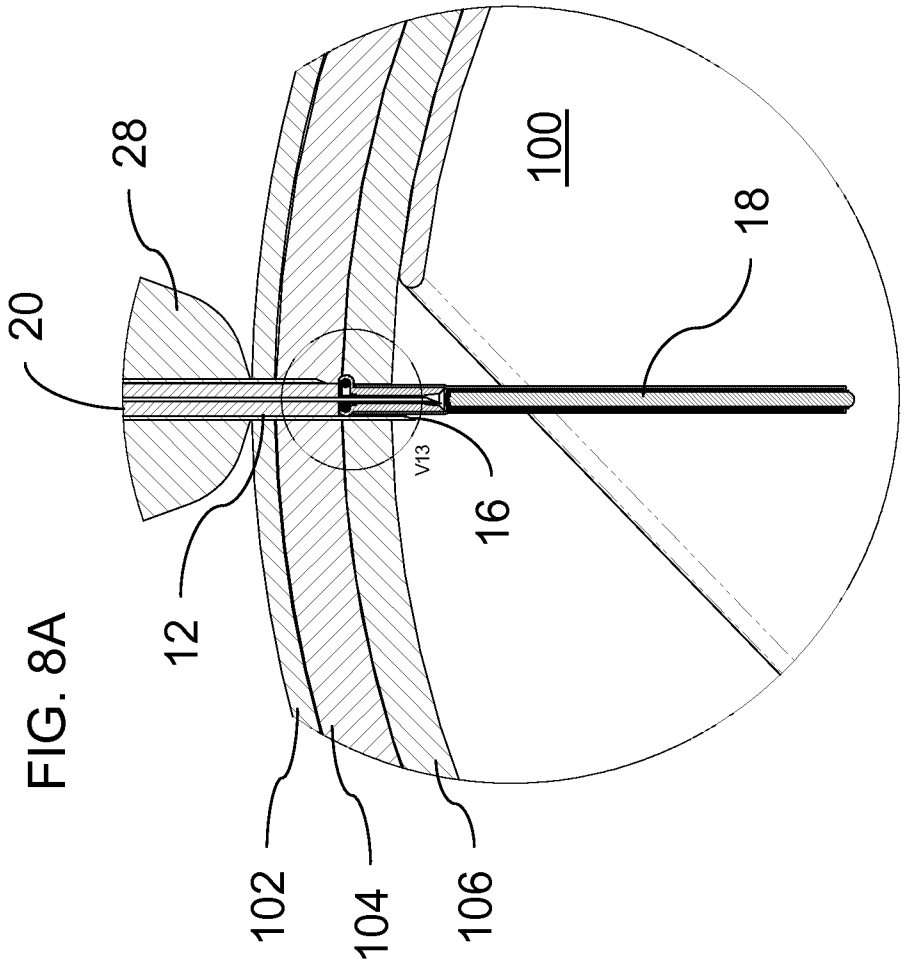


FIG. 8D

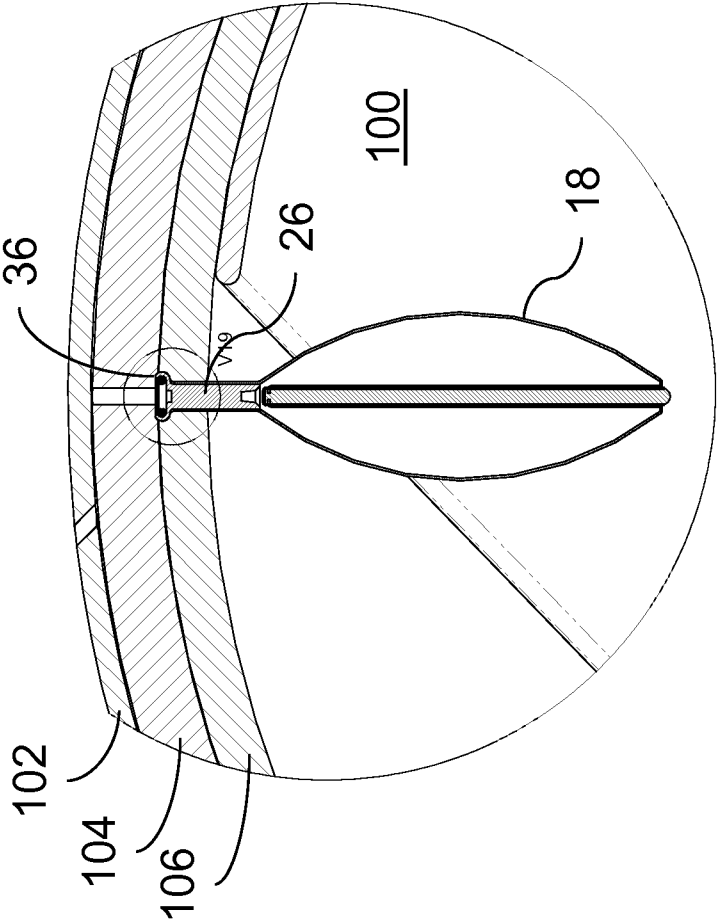
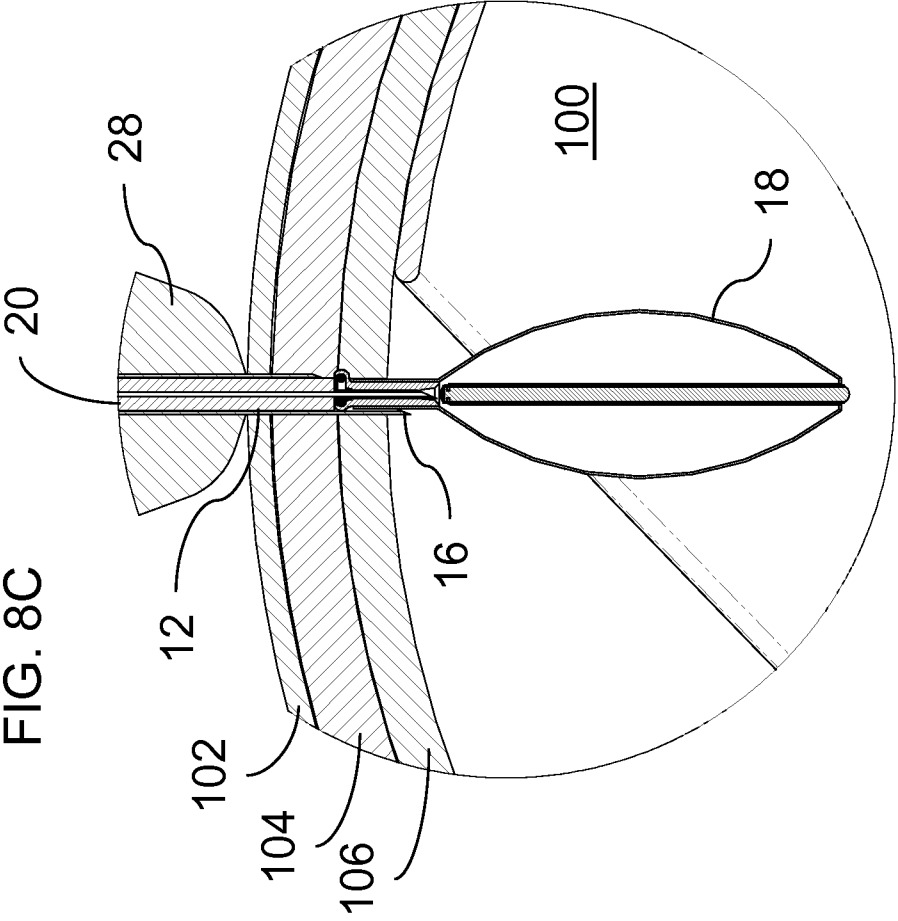


FIG. 8C



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FIG. 8F

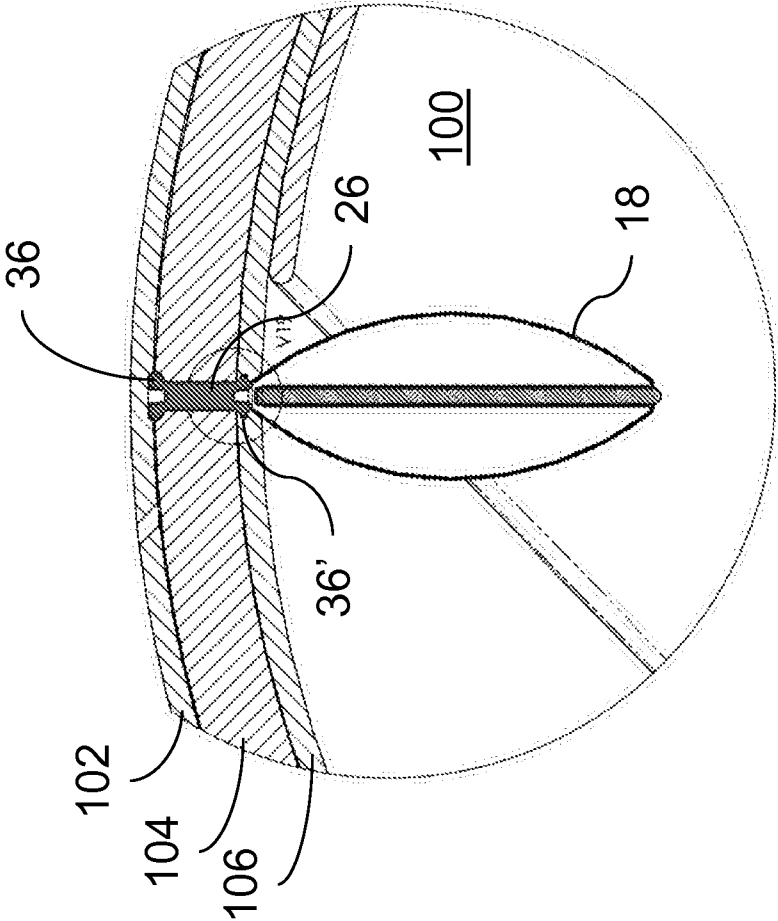
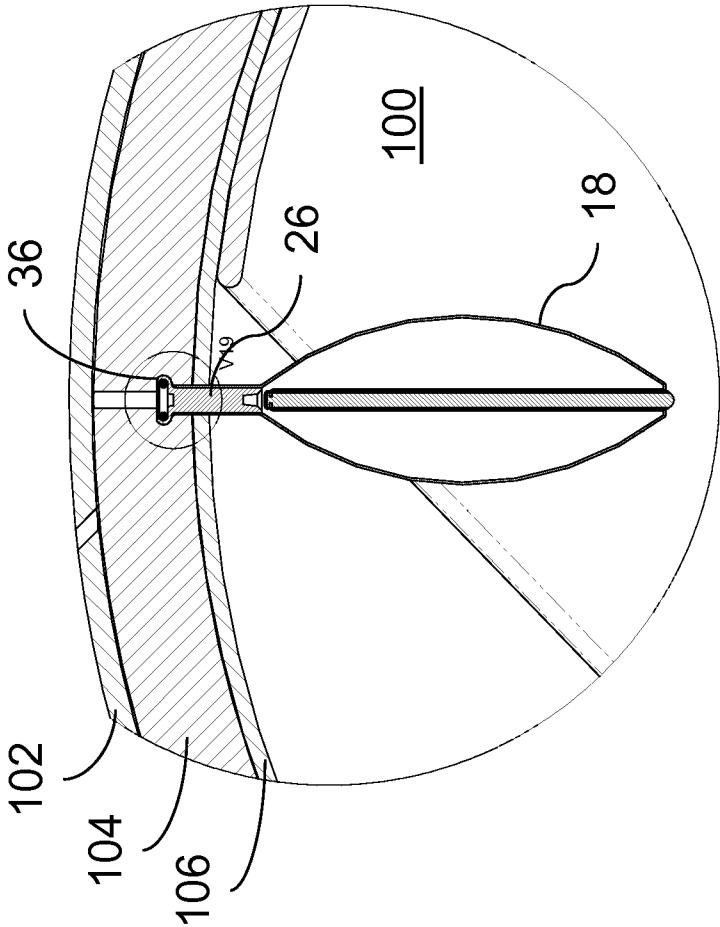


FIG. 8E



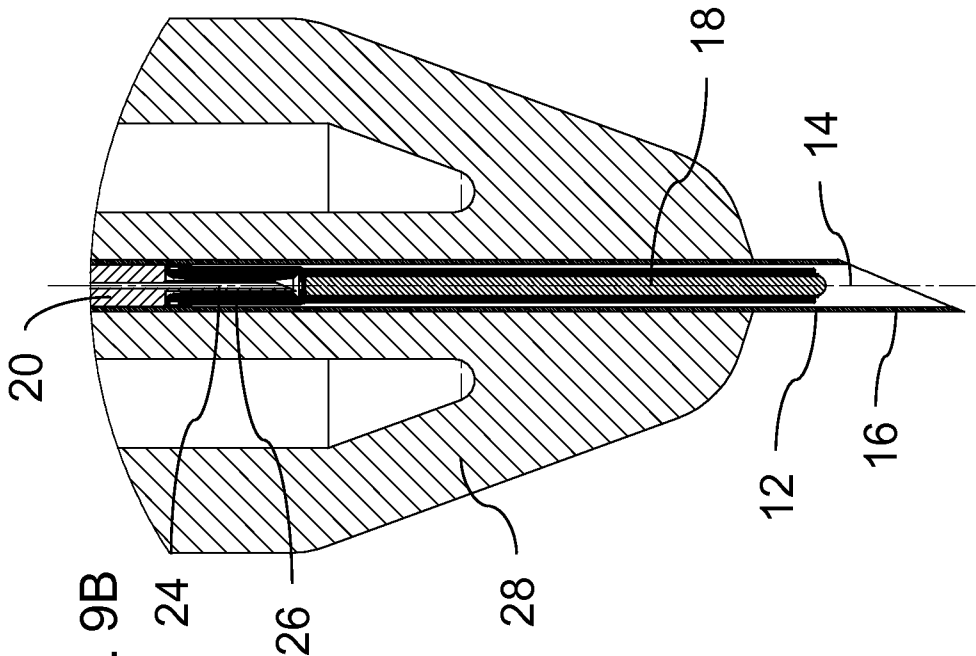


FIG. 9B

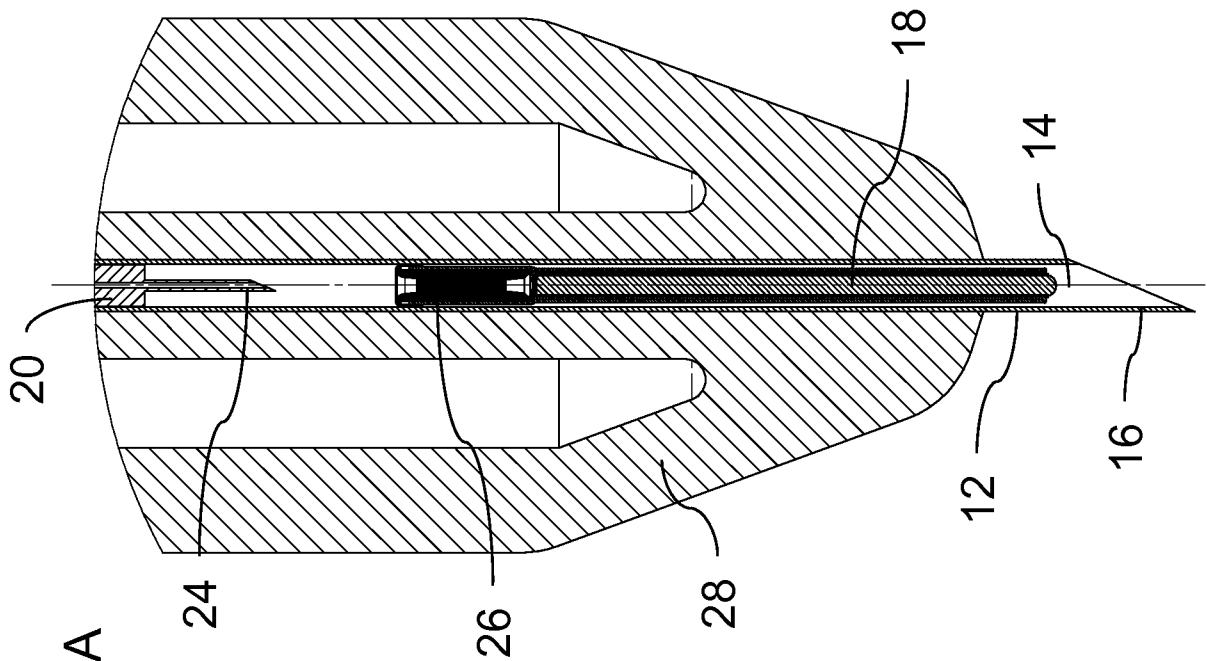


FIG. 9A

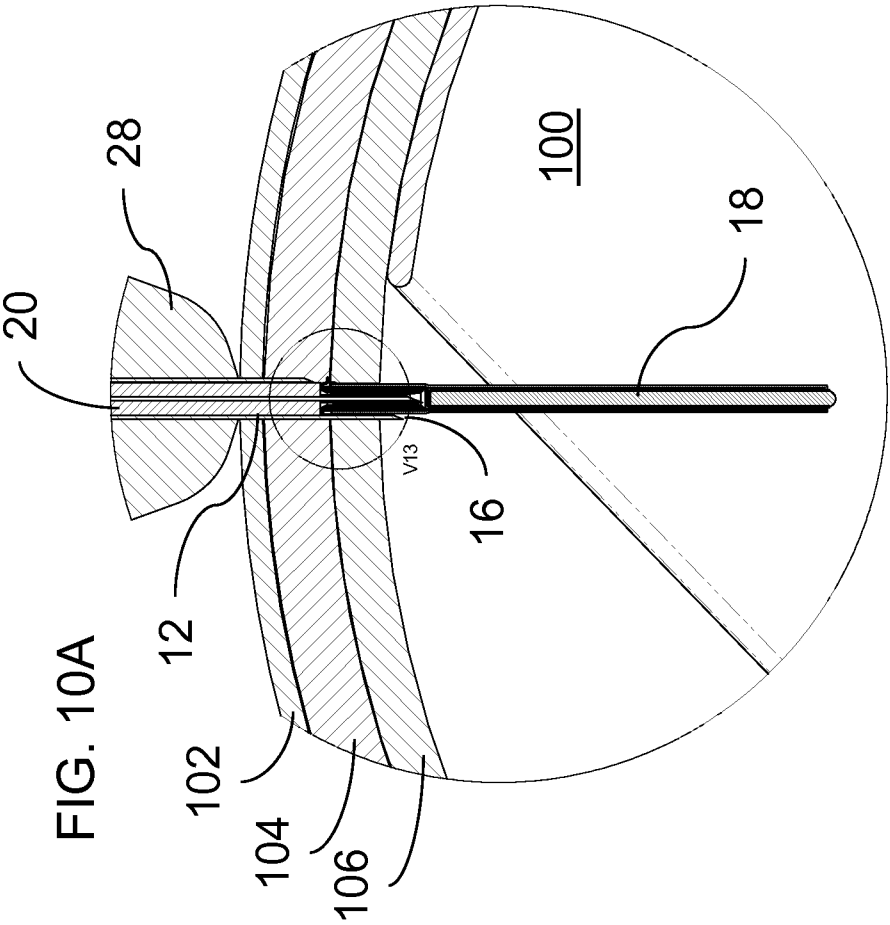


FIG. 10B

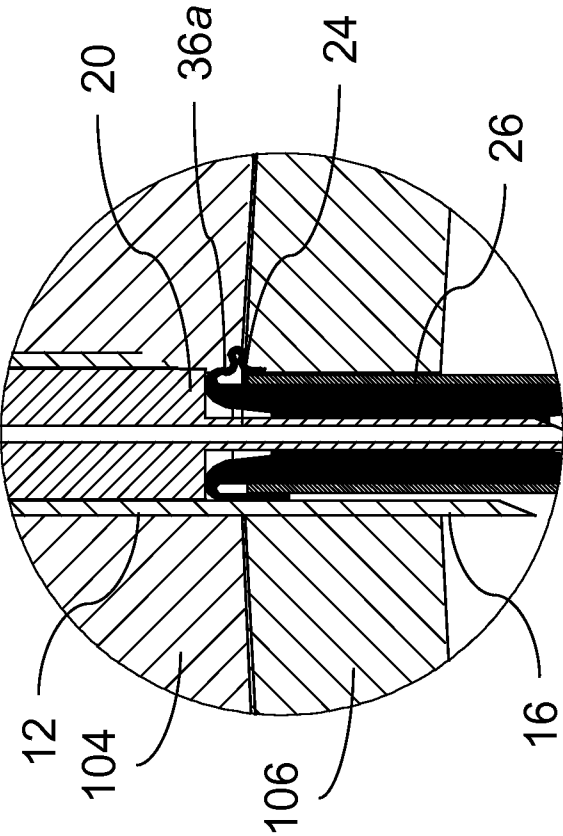


FIG. 10D

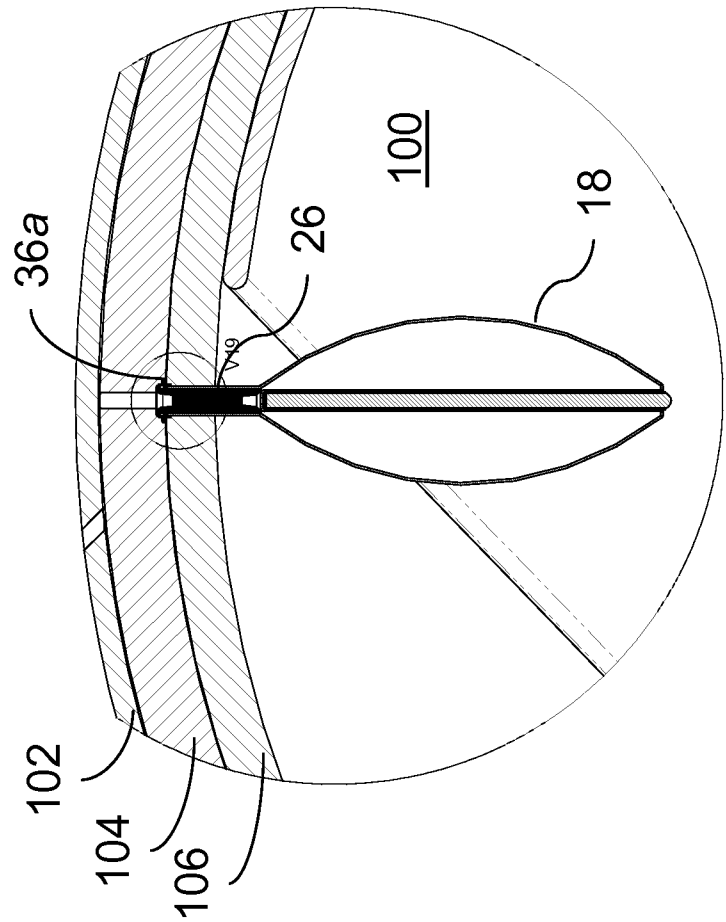
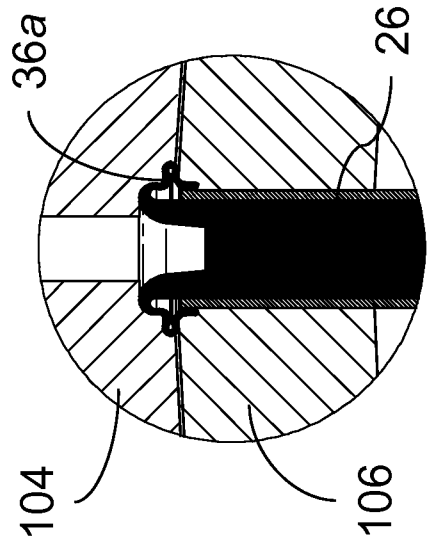
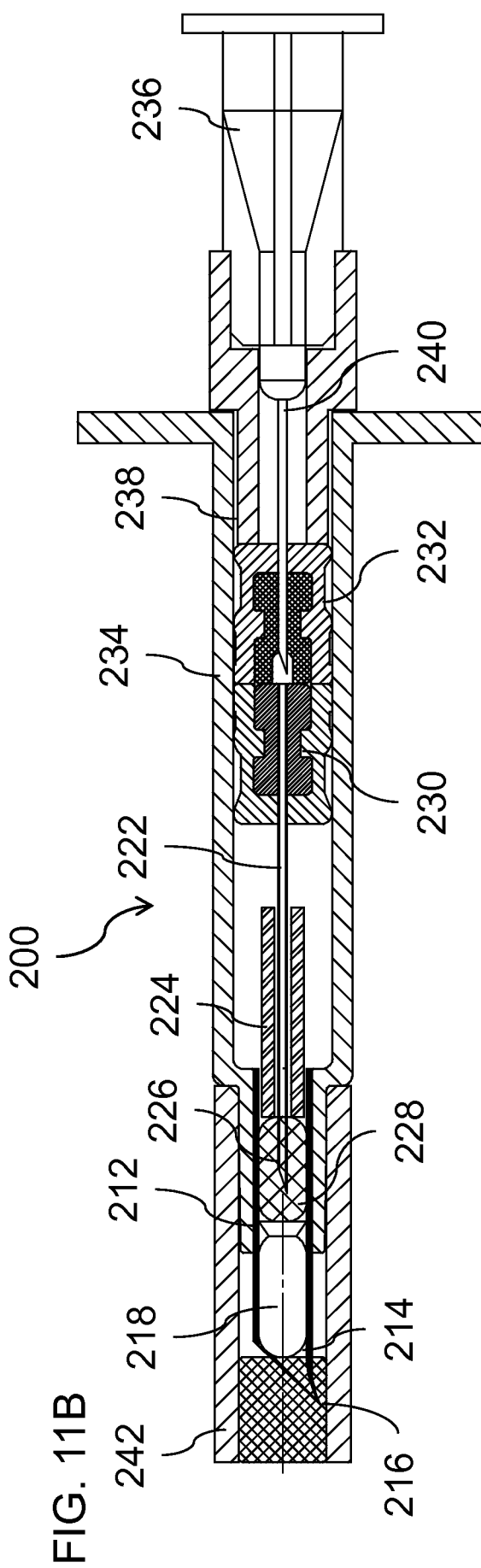
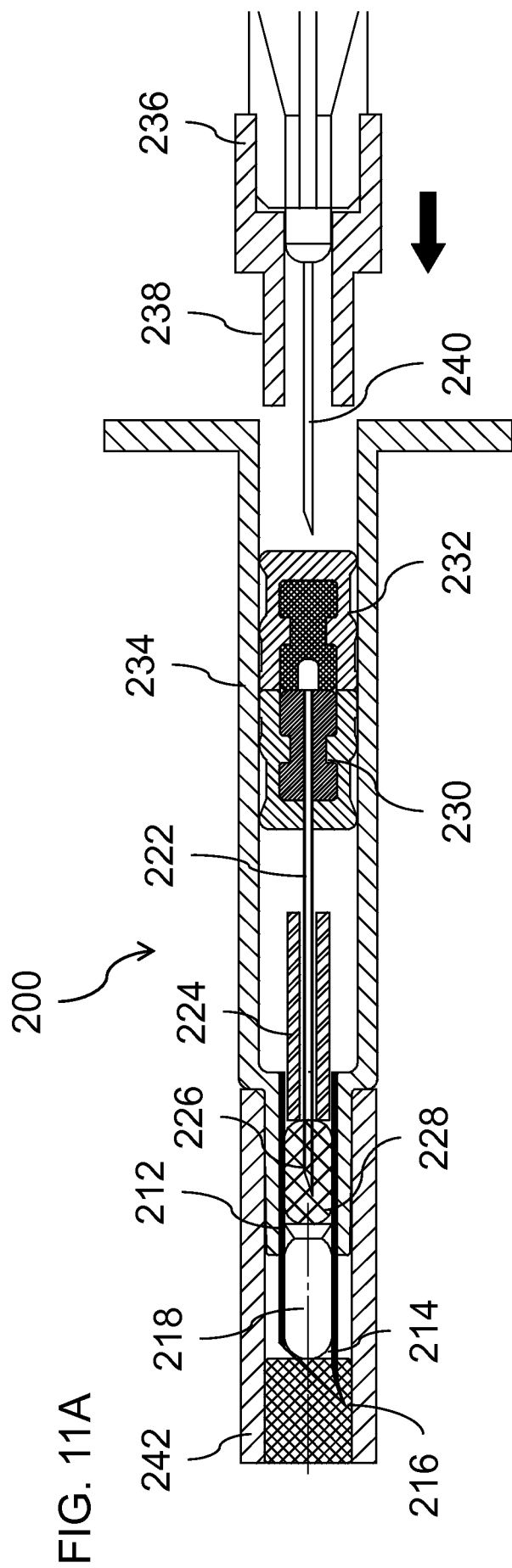
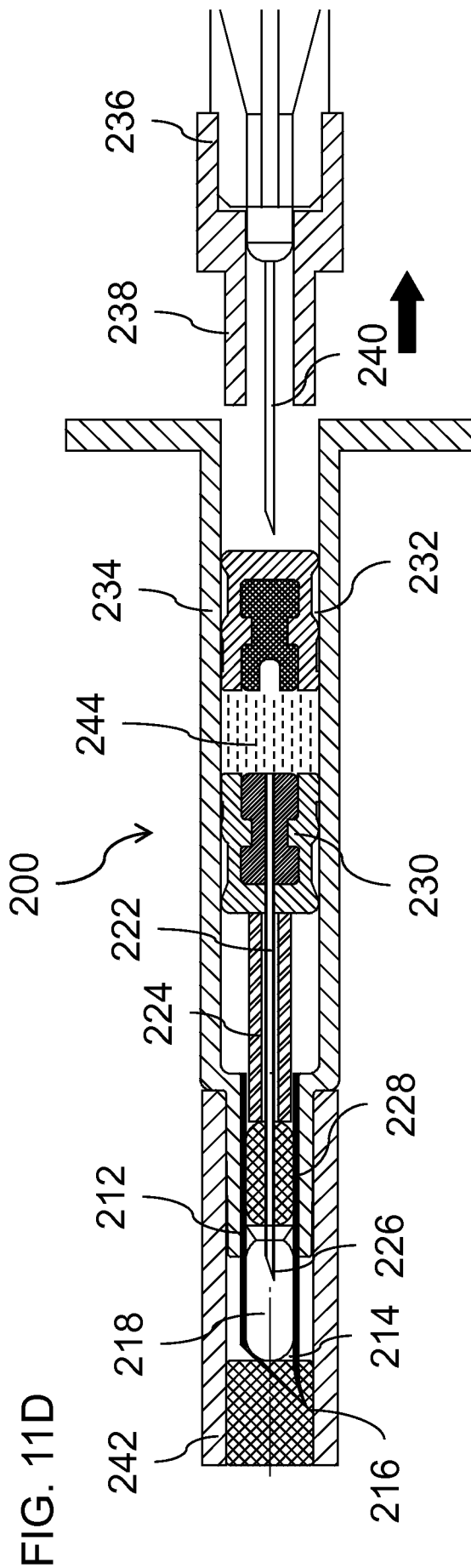
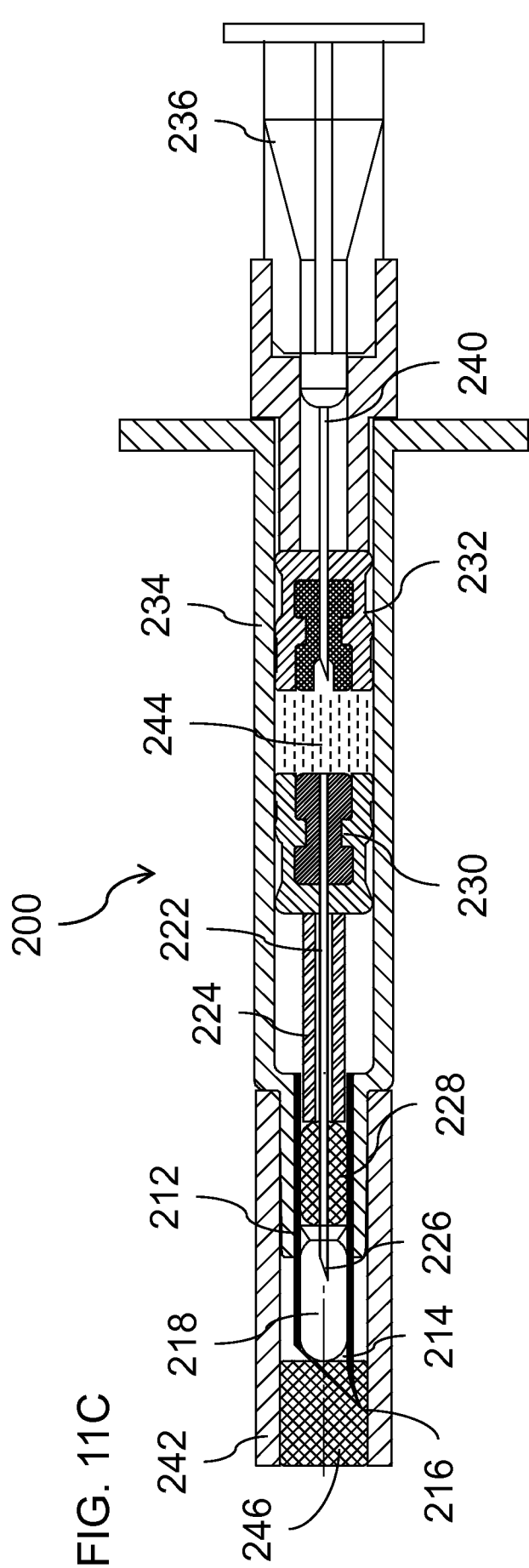
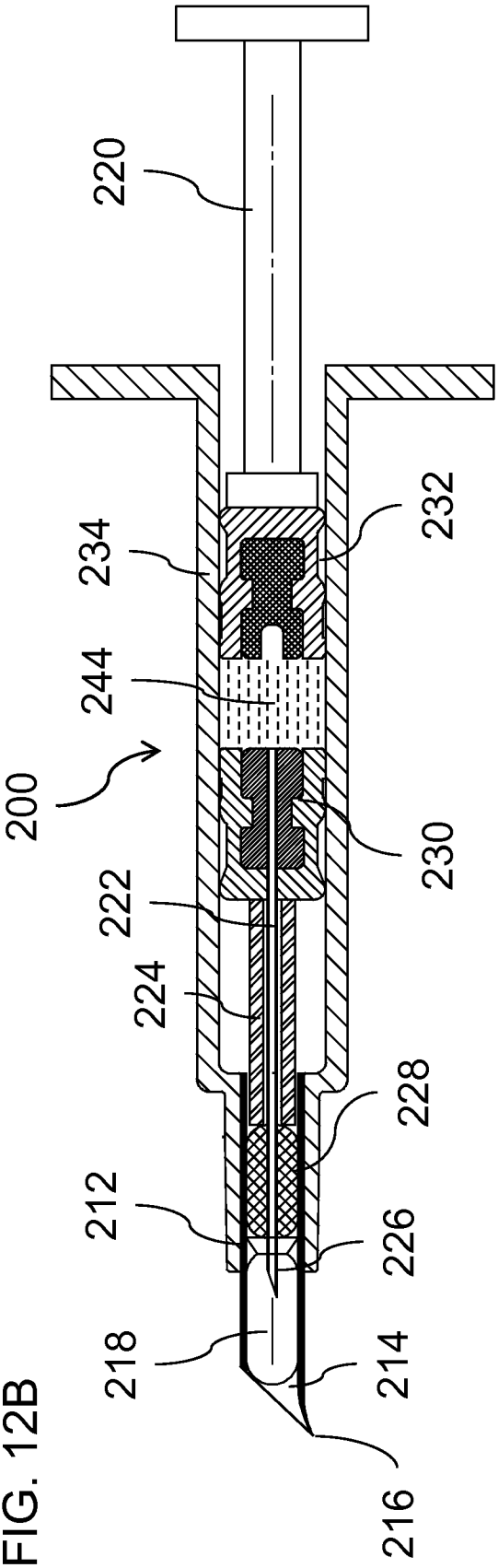
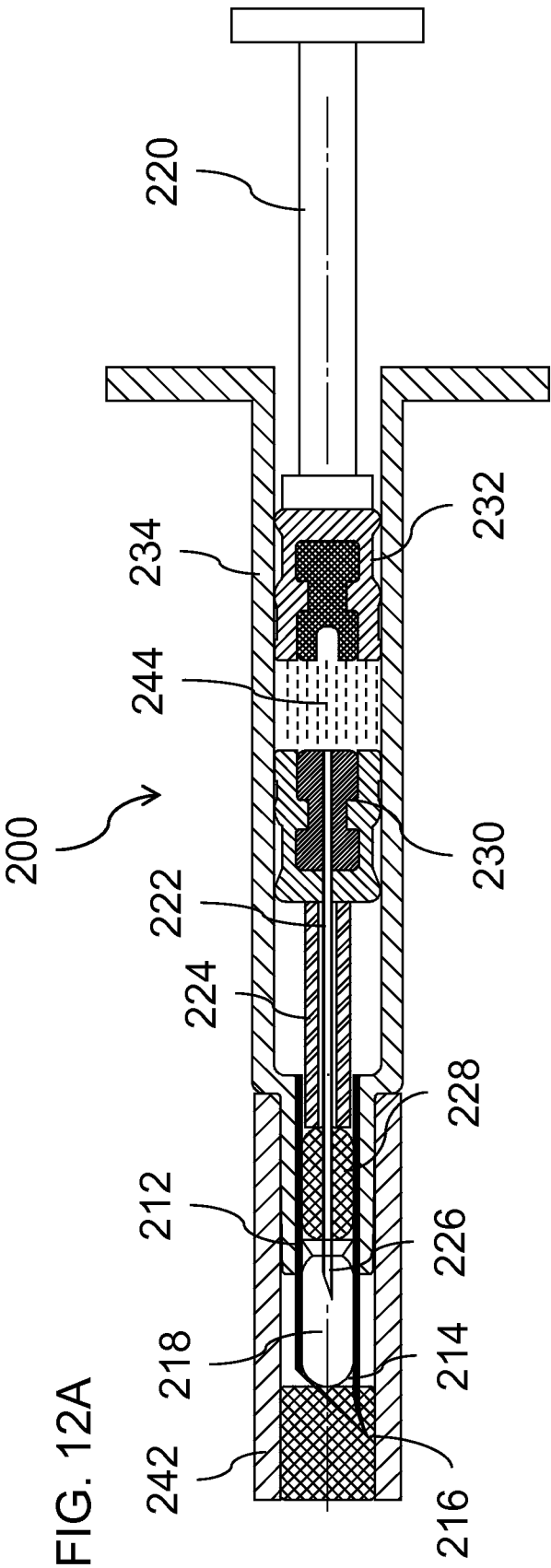


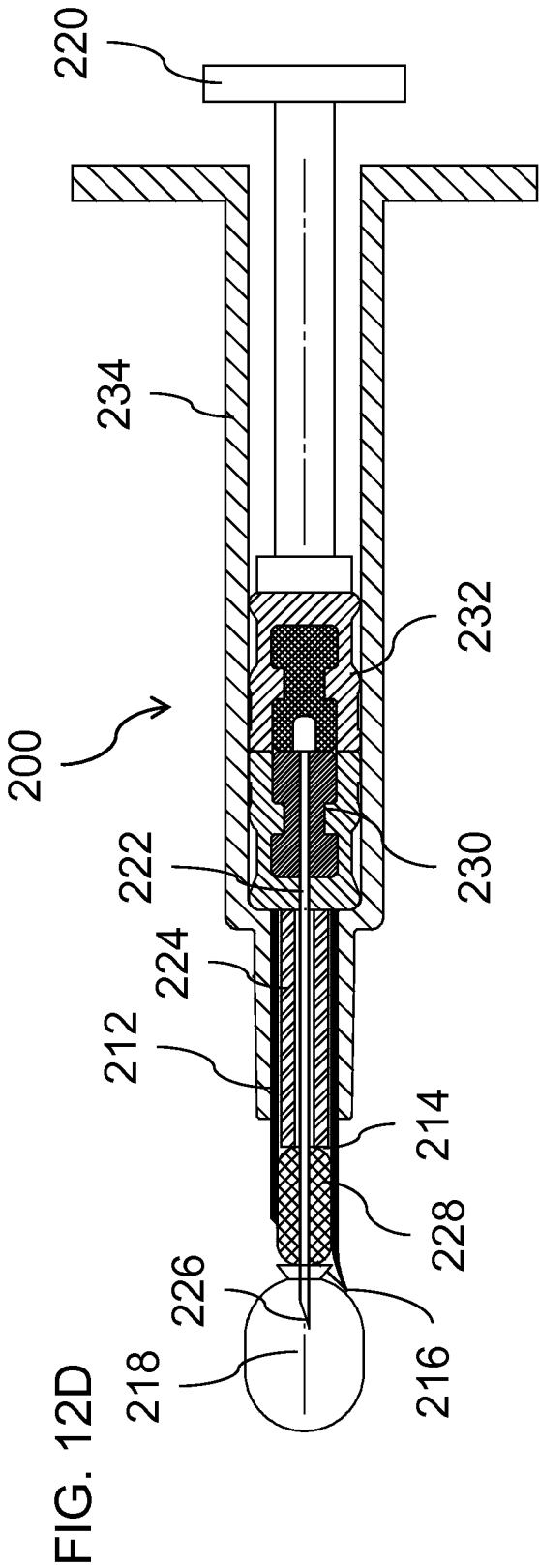
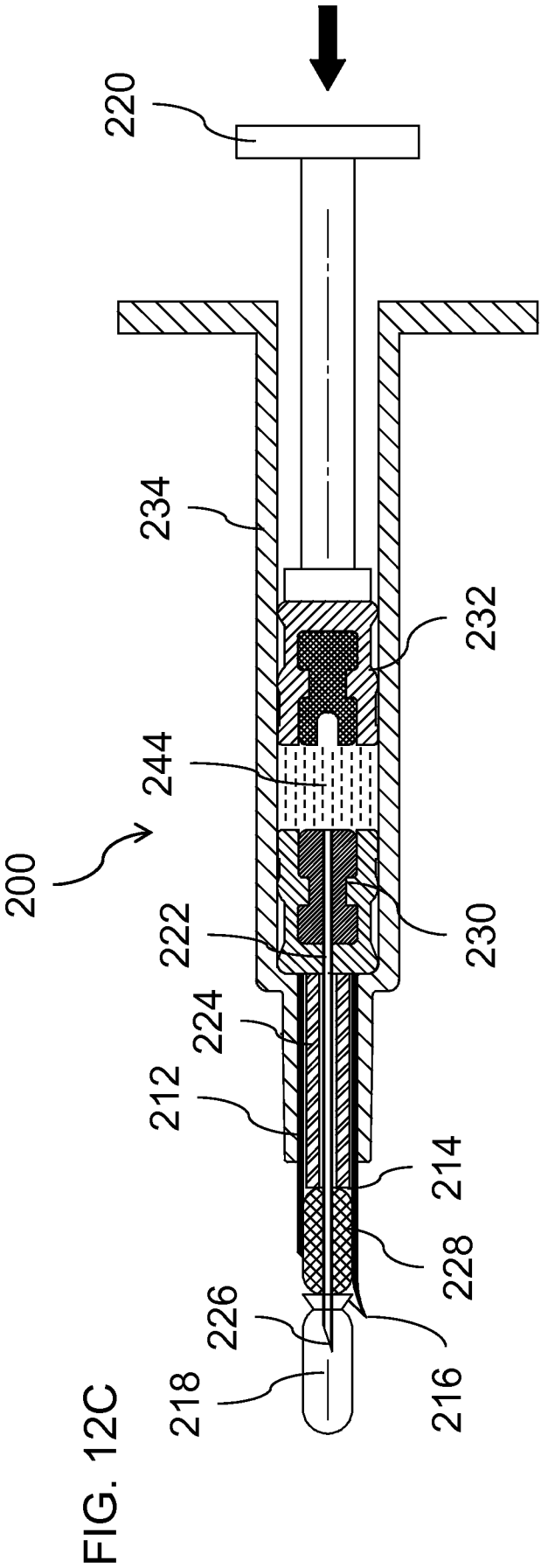
FIG. 10E

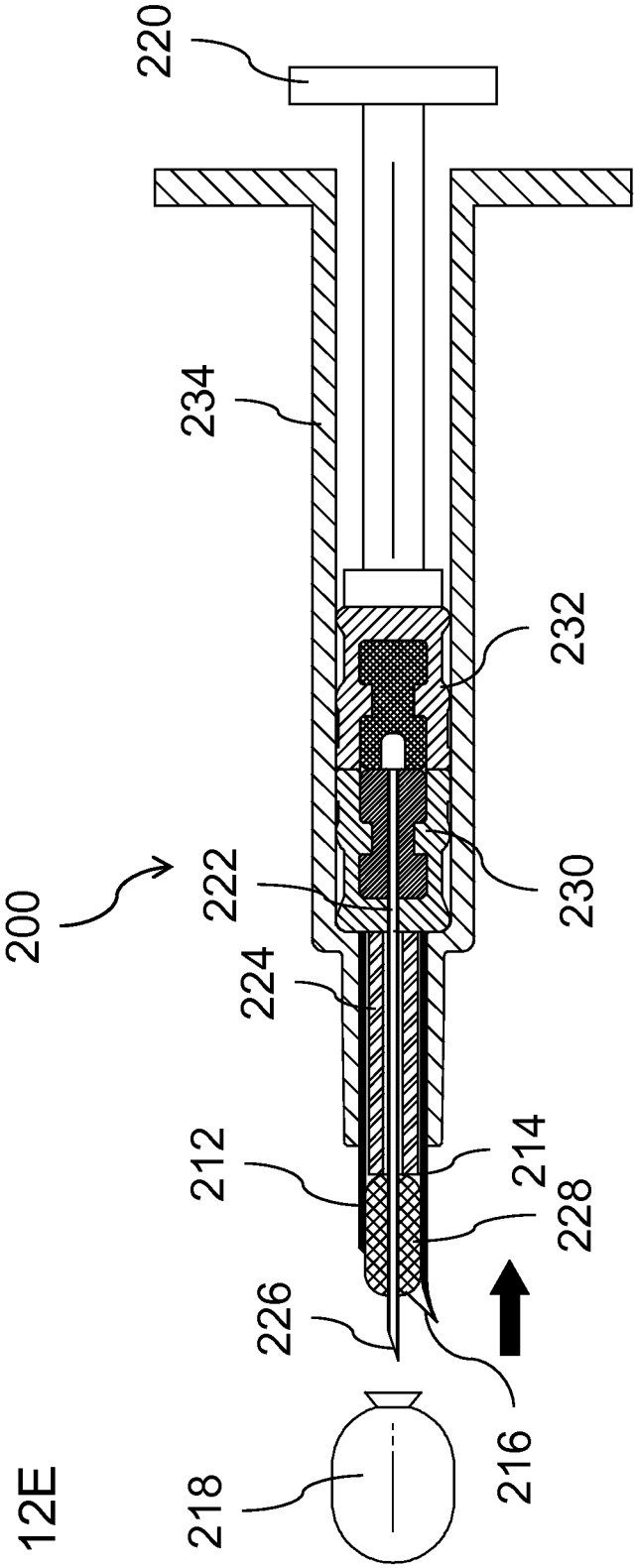












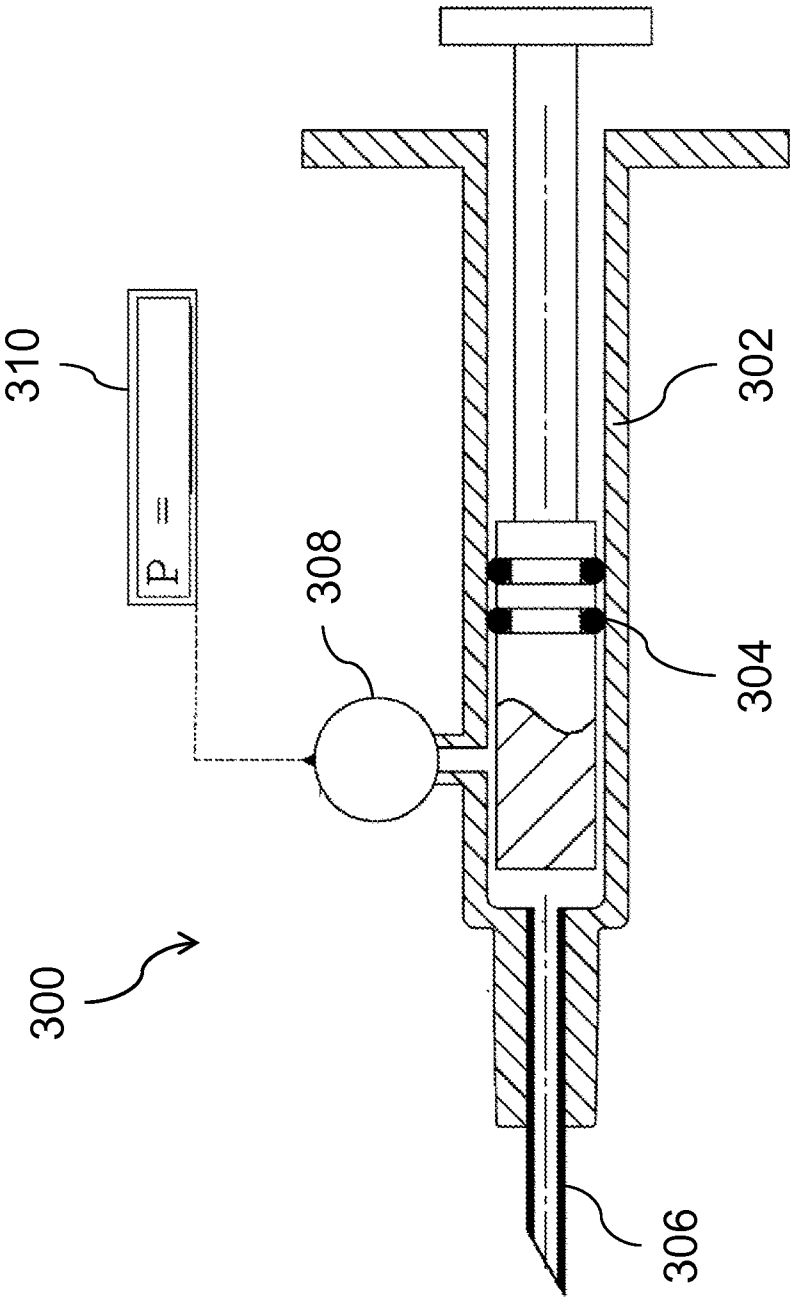


FIG. 13

