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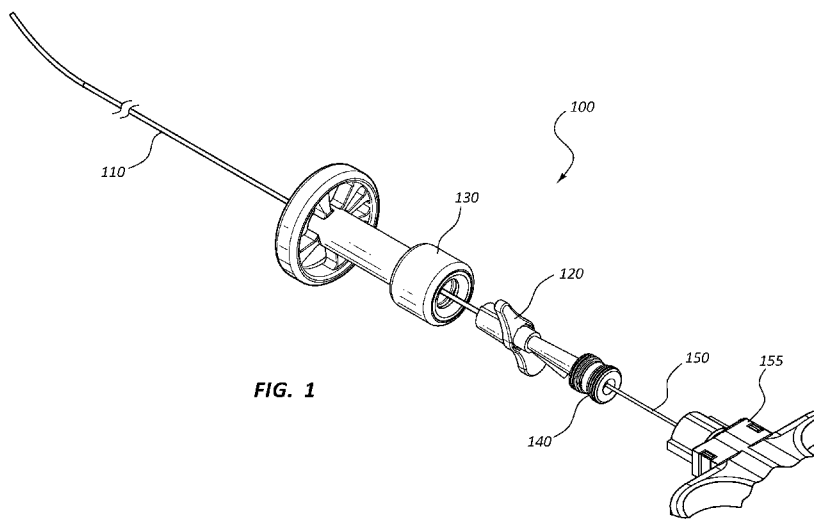


FIG. 1

(57) Abstract: Various components of an interventional therapy assembly are disclosed. One such component is a pointer which may be configured to indicate the disposition of another portion or component of the assembly. A rotary indexing device which may be configured to control or facilitate rotation of other components is also disclosed. Further, various components of an elongate instrument having flexible segments and operative segments are also disclosed. In some embodiments, the elongate instrument may comprise a biopsy assembly and may be used in connection with other components of the disclosed assembly.



## BIOPSY DEVICE AND METHOD OF USE

### RELATED APPLICATIONS

**[0001]** This application claims priority to United States Provisional Application Number 61/739,112 filed on December 19, 2012 and titled "Biopsy Device and Method of Use," which is hereby incorporated by reference in its entirety.

### TECHNICAL FIELD

**[0002]** The present disclosure relates generally to medical devices. More specifically, the present disclosure relates to transvascular devices or other minimally invasive devices. In some embodiments the present disclosure relates to transvascular liver therapy devices, including transjugular liver biopsy devices.

### BRIEF DESCRIPTION OF THE DRAWINGS

**[0003]** The embodiments disclosed herein will become more fully apparent from the following description and appended claims, taken in conjunction with the accompanying drawings. The drawings depict only typical embodiments, which embodiments will be described with additional specificity and detail in connection with the drawings in which:

**[0004]** Figure 1 is a perspective view of a transvascular biopsy assembly in a first configuration.

**[0005]** Figure 2 is a front elevation view of a portion the transvascular biopsy assembly of Figure 1 in a second configuration.

**[0006]** Figure 3 is a cross-sectional view of the portion of the transvascular biopsy assembly of Figure 2 taken through plane 3-3.

**[0007]** Figure 4 is a perspective view of the introducer sheath, pointer, and valve of the assembly of Figure 1.

**[0008]** Figure 5 is an exploded view of the rotary indexer of the assembly of Figure 1.

**[0009]** Figure 6 is a front assembled view of the rotary indexer of Figure 5.

**[0010]** Figure 7 is an exploded view of another embodiment of a rotary indexer.

**[0011]** Figure 8 is a first perspective view of the base of the rotary indexer of Figure 7.

**[0012]** Figure 9 is a second perspective view of the base of the rotary indexer of Figure 7.

- [0013]** Figure 10 is a first perspective view of the rotary member of the rotary indexer of Figure 7.
- [0014]** Figure 11 is a second perspective view of the rotary member of the rotary indexer of Figure 7.
- [0015]** Figure 12 is a cross-sectional view of the rotary member and base of the rotary indexer of Figure 7.
- [0016]** Figure 13 is an exploded view of an embodiment of a valve.
- [0017]** Figure 14 is a perspective view of another embodiment of a valve.
- [0018]** Figure 15 is an exploded view of the valve of Figure 14.
- [0019]** Figure 16 is a cross-sectional view of a portion of the valve of Figure 14 with an elongate device disposed therein.
- [0020]** Figure 17 is a perspective view of another embodiment of a valve.
- [0021]** Figure 18 is an exploded view of the valve of Figure 17.
- [0022]** Figure 19 is a cross-sectional view of a portion of the valve of Figure 17 with an elongate device disposed therein.
- [0023]** Figure 20 is a side view of the biopsy assembly of the assembly of Figure 1.
- [0024]** Figure 21 is an exploded perspective view of the stylet and cannula of the biopsy assembly of Figure 20.
- [0025]** Figure 22 is a side view of the stylet of the biopsy assembly of Figure 20.
- [0026]** Figure 23 is an enlarged view of a portion of the stylet of Figure 22, taken around line 23-23.
- [0027]** Figure 24 is a side view of the cannula of the biopsy assembly of Figure 20.
- [0028]** Figure 25 is an enlarged view of a portion of the cannula of Figure 24, taken around line 25-25.
- [0029]** Figure 26 is a side view of another embodiment of a flexible segment of a component of a biopsy assembly.
- [0030]** Figure 27 is a side view of another embodiment of a flexible segment of a component of a biopsy assembly.
- [0031]** Figure 28 is an enlarged view of a portion of the flexible segment of Figure 27, taken around line 28.
- [0032]** Figure 29 is a side view of another embodiment of a flexible segment of a component of a biopsy assembly.

- [0033]** Figure 30 is an enlarged view of a portion of the flexible segment of Figure 29, taken around line 30-30.
- [0034]** Figure 31A is an enlarged view of a portion of the flexible segment of Figure 30, taken around line 31A.
- [0035]** Figure 31B is an enlarged view of another embodiment of a portion of a flexible segment, analogous to the view of Figure 31A.
- [0036]** Figure 31C is an enlarged view of a portion of Figure 31B.
- [0037]** Figure 32 is a top view of a portion of an operative segment of another embodiment of a portion of a biopsy assembly.
- [0038]** Figure 33 is a side view of a portion of the operative segment of Figure 32.
- [0039]** Figure 34 is an enlarged view of the portion of the operative segment of Figure 32, taken around line 34-34.
- [0040]** Figure 35A is a side view of a portion of a biopsy assembly in a first configuration.
- [0041]** Figure 35B is a side view of the biopsy assembly of Figure 35A in a second configuration.
- [0042]** Figure 35C is a side view of the biopsy assembly of Figure 35A in a third configuration.
- [0043]** Figure 36 is a top view of a portion of another embodiment of a biopsy assembly.
- [0044]** Figure 37 is an enlarged view of a portion of the biopsy assembly of Figure 36, taken around line 37-37.
- [0045]** Figure 38 is a side view of a portion of the stylet of the biopsy assembly of Figure 36.
- [0046]** Figure 39 is a side view of a portion of the cannula of the biopsy assembly of Figure 36.
- [0047]** Figure 40 is a side view of the outer tubular member of the biopsy assembly of Figure 36.
- [0048]** Figure 41 is a side view of a portion of the biopsy assembly of Figure 36.
- [0049]** Figure 42 is a cross-sectional view of the portion of the biopsy assembly of Figure 41, taken through plane 42-42.
- [0050]** Figure 43 is a cross-sectional view of the portion of the biopsy assembly of Figure 41, taken through plane 43-43.

**[0051]** Figure 44 is an enlarged view of a portion of the cross-sectional view of Figure 42, taken around line 44-44.

**[0052]** Figure 45 is an enlarged view of a portion of the cross-sectional view of Figure 44, taken around line 45-45.

**[0053]** Figure 46A is a side view of a portion of the biopsy assembly of Figure 36 in a first configuration.

**[0054]** Figure 46B is a side view of a portion of the biopsy assembly of Figure 36 in a second configuration.

**[0055]** Figure 46C is a side view of a portion of the biopsy assembly of Figure 36 in a third configuration.

**[0056]** Figure 46D is a side view of a portion of the biopsy assembly of Figure 36 in a fourth configuration.

**[0057]** Figure 46E is a side view of a portion of the biopsy assembly of Figure 36 in a fifth configuration.

#### DETAILED DESCRIPTION

**[0058]** Various therapies and procedures may be performed through transvascular or other minimally invasive techniques. For example, one or more instruments may be introduced into the vasculature of a patient and advanced to a treatment site, and a therapy may then be performed.

**[0059]** In some embodiments, vascular access techniques may be used to treat the liver. For example, a device may be configured for use in a transjugular liver biopsy procedure; i.e., a procedure directed to retrieving a liver sample through a device introduced at the jugular vein of a patient.

**[0060]** It will be readily understood that the components of the embodiments as generally described and illustrated in the Figures herein could be arranged and designed in a wide variety of different configurations. Thus, the following more detailed description of various embodiments, as represented in the Figures, is not intended to limit the scope of the disclosure, but is merely representative of various embodiments. While the various aspects of the embodiments are presented in drawings, the drawings are not necessarily drawn to scale unless specifically indicated.

**[0061]** The phrases “connected to” and “coupled to” refer to any form of interaction between two or more entities, including mechanical, electrical, magnetic, electromagnetic, fluid, and thermal interaction. Two components may be coupled to

each other even though they are not in direct contact with each other. For example, two components may be coupled to each other through an intermediate component.

**[0062]** The directional terms “proximal” and “distal” are used herein to refer to opposite locations on a medical device. The proximal end of the device is defined as the end of the device closest to the practitioner when the device is in use by the practitioner. The distal end is the end opposite the proximal end, along the longitudinal direction of the device, or the end furthest from the practitioner.

**[0063]** Figure 1 is a perspective view of a transvascular biopsy assembly 100 in a first configuration. The assembly 100 of Figure 1 comprises various components as further detailed below. In other embodiments, any combination of the individual components may comprise an assembly or subassembly for use in connection with a transvascular procedure.

**[0064]** In the embodiment of Figure 1, the assembly 100 comprises an introducer sheath 110 coupled to a pointer 120 adjacent the proximal end of the introducer sheath 110. A delivery lumen (105 of Figure 3) may extend from a proximal end of the valve 140, through the pointer 120, to a distal end of the introducer sheath 110. A biopsy device 150 may be configured to be disposed within the delivery lumen (105 of Figure 3). The assembly may further comprise a rotary indexer 130. These components are disclosed in further detail below. A handle 155 of the biopsy device 150 is also shown in Figure 1.

**[0065]** A biopsy device 150 may be configured to be disposed within the delivery lumen (105 of Figure 3) of the introducer sheath 110, pointer 120, and valve 140. In the configuration of Figure 1, the biopsy device 150 is shown partially disposed within the delivery lumen (105 of Figure 3). In some embodiments the biopsy device 150 may be configured to be delivered through the vasculature of a patient to obtain a tissue sample from the patient. For example, the biopsy device 150 may be configured to obtain a liver sample by introducing the biopsy device 150 into a vessel of the liver, then obtaining a sample. In other embodiments the biopsy device may be configured to obtain a tissue sample from another part of the body. The biopsy device 150 may be configured to obtain a partial core tissue sample, a full core tissue sample, a blood or other fluid sample from a particular region of the body, and so forth. Other treatment devices or assemblies (used in connection with or in place of the biopsy device 150) are within the scope of this disclosure. For example, the assembly 100 may be configured to guide or deliver devices such as stent

deployment tools, snares, balloons, probes, forceps, and so forth. In some embodiments, the assembly 100 may be configured for use in connection with a transjugular intrahepatic portosystemic shunt ("TIPS") procedure. In some such embodiments, the assembly 100 may or may not comprise a biopsy device 150.

**[0066]** Figure 2 is a front elevation view of a portion of the transvascular biopsy assembly 100 of Figure 1 in a second configuration, and Figure 3 is a cross-sectional view of the portion of the transvascular biopsy assembly 100 of Figure 2. Referring to Figures 1-3, the biopsy device 150 may be axially displaceable within the delivery lumen 105 of the assembly 100. In the configuration of Figures 2 and 3, the biopsy device 150 is shown partially advanced within the delivery lumen 105, such that a distal end of the biopsy device 150 is disposed within a portion of the delivery lumen 105 adjacent the rotary indexer 130. The introducer sheath 110, the pointer 120, and the valve 140 may each comprise a lumen: an introducer sheath lumen 115, a pointer lumen 125, and a valve lumen 145. The introducer sheath lumen 115, the pointer lumen 125, and the valve lumen 145 may be aligned such that these lumens collectively comprise the delivery lumen 105 of the assembly 100. In other embodiments or configurations, the assembly 100 may or may not include each of the introducer sheath 110, the pointer 120, and the valve 140. In such embodiments, the delivery lumen 105 may only comprise a subset of the introducer sheath lumen 115, the pointer lumen 125, and the valve lumen 145. For example, in some configurations the assembly 100 may not include a pointer 120. In such instances the valve lumen 145 may be disposed directly adjacent the introducer sheath lumen 115 to form the delivery lumen 105 of the assembly 100. In other embodiments, additional components of the assembly may be disposed such that lumens or openings in these additional components also comprise the delivery lumen 105. As shown in Figure 3, the rotary indexer 130 may also comprise a rotary indexer lumen 135. In the illustrated embodiment, the delivery lumen 105 runs through the rotary indexer lumen 135, though the rotary indexer lumen 135 may not comprise a direct portion of the delivery lumen 105. In other embodiments, these components may be arranged such that a portion of the rotary indexer lumen 135 comprises a portion of the delivery lumen 105.

**[0067]** Further, in the configuration of Figures 2 and 3, the pointer 120 has been axially advanced with respect to the rotary indexer 130 (as compared to the configuration of Figure 1) such that a ridge 128 of the pointer 120 is engaged with a

slot 138 of the rotary indexer 130. This engagement may couple a rotary member 132 of the rotary indexer 130 to the pointer 120 such that rotation of the pointer 120 also rotates the rotary member 132.

**[0068]** Figure 4 illustrates the introducer sheath 110, pointer 120, and valve 140 of the assembly 100 of Figure 1. In some embodiments, the pointer 120 may be fixedly coupled adjacent a proximal end 111 of the introducer sheath 110, including embodiments wherein these components are integrally formed. In other embodiments, the pointer 120 may be removably coupled to the introducer sheath 110. The pointer 120 may be coupled to the introducer sheath 110 such that the pointer 120 and introducer sheath 110 are constrained to rotate together about a longitudinal axis of the components. The introducer sheath 110 may be formed of a single material, comprise a laminate material, and/or comprise a reinforced material. In some embodiments the introducer sheath 110 may be configured with a particular stiffness to facilitate a particular procedure. For example, the introducer sheath 110 may be sufficiently stiff to facilitate advancement of the introducer sheath 110 along the vasculature of a patient.

**[0069]** The introducer sheath 110 may comprise a curved distal tip 112. The magnitude of the curve may be greater or smaller than shown in the illustrated embodiment. The curved distal tip 112 may be configured to facilitate steering of the introducer sheath 110 as the introducer sheath 110 is advanced within a body lumen of a patient. The curved distal tip 112 may also or alternatively be configured to access a particular portion of the patient's body during a therapy. For example, during a transjugular liver biopsy procedure, an introducer sheath 110 may be advanced from an insertion point at a patient's jugular vein, through the vasculature to the patient's liver. The curved distal tip 112 of the introducer sheath 110 may be configured to direct instruments passed through the introducer sheath 110 (e.g., a biopsy device) such that the instruments emerge from the sheath and contact the desired tissue. Rotation of the introducer sheath 110 about its longitudinal axis may be configured to alter or direct the orientation or disposition of the curved distal tip 112 of the introducer sheath 110.

**[0070]** The pointer 120 may be disposed such that its orientation corresponds to the orientation or direction of the curved distal tip 112 of the introducer sheath 110. For example, the curved distal tip 112 of the introducer sheath may be curved substantially within a plane, and a transverse axis of the pointer 120 may lie

substantially within the same plane. Thus, rotation of the introducer sheath 110 (which may be coupled to the pointer 120) may cause the pointer 120 to rotate such that the pointer 120 extends in the same direction as the curved distal tip 112 of the introducer sheath 110. When the introducer sheath 110 is in use, the curved distal tip 112 may be disposed within a patient, and thus not directly observable by a practitioner. The pointer 120, however, may be disposed outside the patient and thereby provide the practitioner with a visual indication of the orientation of the curved distal tip within the patient's body.

**[0071]** An introducer sheath 110 or any other elongate instrument configured to be positioned or otherwise controlled by rotating a curved portion thereof (such as the curved distal tip 112 of introducer sheath 110) may be utilized in a variety of procedures within a patient's body, including transvascular procedures. In some instances, a practitioner may control or observe the rotational position of a curved distal tip 112 through use of a pointer 120 and/or a rotary indexer 130. These components, and their relationship to rotation of another device, are further discussed below.

**[0072]** A pointer analogous to pointer 120 of Figure 4 may be used in connection with any elongate instrument. For example an analogous pointer may be coupled to a drainage catheter, a stent delivery device, a guidewire, or any other device configured to be inserted into a patient's body. Some such devices may be configured for use within the vasculature of a patient, while others may be configured for insertion into other body lumens, or configured for insertion into portions of the body not comprising lumens.

**[0073]** The pointer 120 of Figure 4 further comprises a ridge 128 disposed on a distal portion of the pointer 120. The ridge 128 may be configured to engage another element of an assembly to couple the rotational displacement of the pointer 120 with the rotational displacement of the coupled element. For example, as discussed in connection with Figure 3, the ridge 128 may couple the pointer 120 to a rotary member (132 of Figure 3) of a rotary indexer (130 of Figure 3). In some embodiments the ridge 128 or other engagement member may be disposed on the introducer sheath 110 or some other component of the assembly. The rotary indexer (130 of Figure 3) is discussed in more detail below.

**[0074]** As shown in Figure 4, a valve 140 may be coupled to a proximal end of the pointer 120. The valve 140 may be configured to control flow and access to the

valve lumen 145 and/or the delivery lumen 105 of the assembly. In some embodiments the valve 140 may be directly coupled to the pointer 120, including embodiments wherein the valve 140 and pointer 120 are integrally formed. In other embodiments, the valve 140 may be coupled directly to the introducer sheath 110 and may be integrally formed with the introducer sheath 110. In some embodiments, the valve 140 may be provided in connection with an introducer sheath 110 but no pointer 120. Similarly, an introducer sheath 110 and pointer 120 may be configured for use without a valve 140.

**[0075]** Figure 5 is an exploded view of the rotary indexer 130 of the assembly 100 of Figure 1. Figure 6 is a front assembled view of the rotary indexer 130. The rotary indexer 130 may be configured to control the rotation and/or indicate the rotational position of certain components of the assembly. The rotary indexer 130 may comprise a base 131 configured to receive a rotary member 132. The rotary member 132 may be configured to be rotatable relative to the base 131 when the rotary indexer 130 is assembled. The rotary member 132 may be configured with an engagement feature, such as slot 138, configured to engage other components. Thus, rotation of the rotary member 132 with respect to the base 131 may also rotate an engaged component relative to the base 131.

**[0076]** For example, referring collectively to Figures 1-6, an assembly 100 may be configured such that the pointer 120 comprises a ridge 128 configured to engage the slot 138 of the rotary member 132. In the illustrated embodiment, rotation of the introducer sheath 110 and pointer 120 may thus be coupled to rotation of the rotary member 132 with respect to the base 131. In the illustrated embodiment, the ridge 128 of the pointer 120 is configured to slide into the slot 138 of the rotary member 130 when the components are engaged. In other embodiments, the pointer 120 or some other component, such as the introducer sheath 110, may be fixedly coupled to the rotary member 132. In still further embodiments, the rotary indexer 130 may be configured such that a component such as a medical device is coupled to the base 131 such that rotation of the base 131 with respect to the rotary member 132 (which may be coupled to the pointer 120) rotates the medical device with respect to the pointer 120. In such an arrangement, the relative position of the pointer 120 with respect to the base 131 may still be configured to indicate the rotational position of the medical device.

**[0077]** In some embodiments, the base 131 may comprise visual indicia corresponding to the position of the pointer 120 with respect to the base 131 which indicate the position of the pointer 120 (and therefore the curved distal tip 112 of the introducer sheath 110). For example, numbers, colors, lines, and so forth on the base 131 may align with the pointer 120 and indicate the relative rotation of the curved distal tip 112 of the introducer sheath 110.

**[0078]** The rotary member 132 may also be configured with detents or other catch mechanisms configured to releasably couple the rotary member 132 to the base 131 such that the rotary member 132 cannot easily rotate when the detents or catch mechanisms are engaged. Interaction between the rotary member 132 and the base 131 may thus allow a practitioner to rotatably displace the rotary member 132 (and therefore the pointer 120 and introducer sheath 110 in some embodiments) a certain amount, then release the rotated components. Interaction with the base 131 may be configured to resist unwanted rotational displacement of these components, for example, when the practitioner desires to leave the components in place for a portion of a therapeutic procedure.

**[0079]** In the embodiment of Figures 5 and 6, the rotary member 132 may be coupled to the base 131 by interaction with a biasing element 133 and a retainer 134. The retainer 134 may be coupled to the base 131 and the biasing element 133 configured to bias the rotary member 132 into contact with a portion of the base 131.

**[0080]** Figure 7 is an exploded view of another embodiment of a rotary indexer 230. The embodiment of Figure 7 may include components that resemble components of the embodiment of Figure 5 in some respects. For example, the embodiment of Figure 7 includes a rotary member 232 that may resemble the rotary member 132 of the embodiment of Figure 5. It will be appreciated that all the illustrated embodiments have analogous features and components. Accordingly, like or analogous features are designated with like reference numerals, with the leading digits incremented to "2." Relevant disclosure set forth above regarding similarly identified features thus may not be repeated hereafter. Moreover, specific features of the system and related components shown in Figure 7 may not be shown or identified by a reference numeral in the drawings or specifically discussed in the written description that follows. However, such features may clearly be the same, or substantially the same, as features depicted in other embodiments and/or described with respect to such embodiments. Accordingly, the relevant descriptions of such

features apply equally to the features of the system and related components of Figure 7. Any suitable combination of the features, and variations of the same, described with respect to the system and components illustrated in Figure 5 can be employed with the system and components of Figure 7, and vice versa. This pattern of disclosure applies equally to further embodiments depicted in subsequent figures and described hereafter. Moreover, this pattern of disclosure and numbering applies to any of the components or systems herein disclosed, not simply to various embodiments of rotary indexers. Thus, any component or assembly depicted in any embodiment herein may have analogous features to, or function in an analogous manner to, components or assemblies of other embodiments.

**[0081]** The rotary indexer 230 shown in Figure 7 comprises a base 231, a rotary member 232, a biasing element 233, and a retainer 234. Recessed portions 237 are also shown in a surface of the base 231. When the rotary indexer 230 is assembled, the retainer 234 and biasing element 233 may interact with the rotary member 232 to bias the rotary member 232 into contact with the surface of the base 231 in which the recessed portions 237 are formed. Figures 8-9 are perspective views of the base 231 of the rotary indexer 230, Figures 10-11 are perspective views of the rotary member 232 of the rotary indexer 230, and Figure 12 is a cross-sectional view of the rotary member 232 and base 231.

**[0082]** Referring to Figures 7-12, the rotary indexer 230 may be configured such that detents between the rotary member 232 and the base 231 releasably couple the rotary member 232 to the base 231 such that the rotary member 232 may only rotate with respect to the base 231 when the detents are not engaged. In the illustrated embodiment, the base 231 may include one or more recessed portions 237 in a surface of the base 231. The rotary member 232 may include a protrusion 236 configured to interact with the recessed portions 237. The protrusion 236 and recessed portions 237 may function as detents, tending to constrain rotation of the rotary member 232 with respect to the base 231 when the protrusion 236 is disposed within a recessed portion 237. The biasing element 233 may tend to bias the rotary member 232 such that disengaging the protrusion 236 from a recessed portion 237 may be accomplished by overcoming the force exerted by the biasing element 233 on the rotary member 232 such that the rotary member 232 may move distally with respect to the base 231, allowing the protrusion 236 to disengage and the rotary member 232 to rotate with respect to the base 231.

**[0083]** The recessed portions 237 may be disposed in an arcuate path, creating multiple detents in the path of rotational displacement of the rotary member 232. The detents may be spaced in a variety of configurations; for example the detents could be equally spaced about an arc, could be disposed closer or farther apart along particular segments of the arc, could increase or decrease in spacing in one direction or the other, and so forth. In some embodiments, the detents may be spaced such that they represent a particular displacement of the rotary member 232 and thus, in some embodiments, the curved distal tip (112 of Figure 4) of an introducer sheath or other component coupled to the rotary member 232. In some embodiments the adjacent detents may represent rotational displacement of about 5 to about 25 degrees, of about 10 to about 20 degrees, or of about 15 degrees. In some embodiments there may be from about five to about nine detents disposed in an arc, or more or fewer detents disposed in an arc or complete circle.

**[0084]** The spacing of the detents may be configured for use in connection with a particular therapy. For example, a particular procedure may be configured to obtain multiple tissue samples from differing locations in the body. In one instance, a practitioner may dispose a biopsy device (150 of Figure 1) within a vessel of a patient's liver via transvascular access. The practitioner may then obtain a first tissue sample, rotate the rotary member 232 to the next detent, obtain another tissue sample, and so forth.

**[0085]** The interaction of the rotary member 232 and the base 231 may be configured to provide visual indicia of the rotational position of an instrument coupled to the rotary member 232. For example, as discussed above, a pointer (120 of Figure 1) may be coupled to the rotary member 232 and correspond to marks or other indicia on the base 231. Interaction of the rotary member 232 and base 231 may also be configured to provide audible or tactile indicia of the relative displacement of an instrument. For example, the rotary indexer 230 may be configured such that the device creates an audible or tactile response as the detents between the rotary member 232 and base 231 engage and disengage.

**[0086]** As an analogous component, the rotary indexer 130 of Figures 1-3 and 5-6 may be configured with detents in an analogous manner to those described in connection with the rotary indexer 230 of Figures 7-12. Referring again to Figure 1, the rotary indexer 130 may comprise a base 131, a rotary member 132, a biasing element 133, and a retainer 134 disposed in an analogous manner to that discussed

in connection with the embodiment of Figure 7. Any configuration of detents or catches between the rotary member 132 and the base 131 are within the scope of this disclosure. In some embodiments the detents may be similar to those described in connection with Figures 7-12. For example, the cross-sectional view of Figure 3 illustrates a protrusion 136 and a recessed portion 137. As with the other embodiments, additional recessed portions may be positioned in a rotational path about the base 131.

**[0087]** Other arrangements of detents are within the scope of this disclosure. For example recesses may be formed in an outside diameter of one component with arms or tabs of a second component configured to interact therewith. Further, one component may comprise a plurality of protrusions configured to interact with a plurality of recesses. The components and detents may be arranged to allow full 360-degree rotation of the parts.

**[0088]** Any of the embodiments of rotary indexers or related components described herein may be used in connection with any other component or assembly described herein. Referring again to Figure 1, a rotary indexer 130 may comprise a portion of a medical device assembly, such as transvascular biopsy assembly 100. At various points during a therapy, a practitioner may rotate a portion of the assembly, utilizing the rotary indexer 130 to control the degrees of rotation (e.g., the number of stops the indexer 130 is rotated) or track the amount of rotation (e.g., by observing visual indicia, including the disposition of a pointer 120 with respect to the rotary indexer 130).

**[0089]** One exemplary procedure which may utilize a rotary indexer 130 is a transjugular liver biopsy. A practitioner may advance a liver biopsy device to a desired location within a lumen of the liver. The practitioner may then obtain a first sample, rotate the rotary indexer 130 between one or more stops, and obtain another sample. The rotary indexer 130 may facilitate controlling the spacing of where samples are obtained. A rotary indexer 130 may similarly be used in connection with a drainage catheter or other instrument disposed within a patient's body. For example, a practitioner may wish to rotate a drainage catheter in order to access various locations within the body. A rotary indexer 130 may facilitate control of the spacing of such displacement.

**[0090]** A rotary indexer 130 may also be used in connection with a TIPS procedure. A practitioner may dispose a cannula within a first lumen of the liver,

then attempt to advance the cannula through the liver to a second lumen of the liver, the cannula connecting the two lumens. In order to ascertain whether the cannula has accessed the second lumen, the practitioner may attempt to draw blood back through the cannula; if the tip of the cannula is disposed in the second lumen, blood or other fluids from that lumen should draw through the cannula. In instances where sufficient fluid is not drawn back, the practitioner may determine the cannula is disposed within liver tissue, not within a lumen. The practitioner may then rotate the cannula in order to access a different spot, and attempt to locate the second lumen. A rotary indexer 130 may be configured to indicate rotational positions a practitioner has already tested as well as control rotation of the cannula.

**[0091]** It is within the scope of this disclosure to utilize a rotary indexer 130 with various other procedures or treatments. For example, placement of catheters, balloons, snares, stents, and so forth may each be done during a procedure wherein one or more components are rotated. Rotary indexers 130 may be configured to facilitate or control such rotation in any such procedure.

**[0092]** Figures 13-19 are various embodiments of valves which may be configured for use in connection with a transvascular access assembly such as the transvascular biopsy assembly 100 of Figure 1. The assembly 100 of Figure 1 comprises a valve 140 coupled adjacent a proximal end of the pointer 120. Any of the valves described herein may be used in place of and/or in an analogous manner to the valve 140 of Figure 1. As with the assembly 100 of Figure 1, these valves may be coupled to the assembly at various points, including embodiments wherein a valve is directly coupled to the introducer sheath 110. Luer connectors, threads, or other releasable coupling mechanisms may be used to couple the valves and/or other components.

**[0093]** In the embodiment of Figure 1, a pointer 120 is coupled adjacent an end of the valve 140. The pointer 120 may be configured to be rotatable with respect to the valve 140, or the pointer may be fixedly coupled to the valve 140. In any of the embodiments discussed below, the valves may be releasably or fixedly coupled to a pointer (such as 120 of Figure 1) and/or an introducer sheath (such as 110 of Figure 1) when used in an assembly.

**[0094]** Valves, such as hemostasis valves, may be configured to allow vascular access while still maintaining a degree of hemostasis at the insertion site. For example, the assembly 100 of Figure 1 may be configured such that the introducer

sheath 110 is disposed within the vasculature. The valve 140 may be configured to allow a practitioner to access the delivery lumen (105 of Figure 3) with an elongate device (such as biopsy device 150 of Figure 1) while selectively closing the valve 140 to prevent blood flow through the delivery lumen (105 of Figure 3).

**[0095]** Analogous to the valve 140 of Figure 1, the valves (and related components) described below have analogous reference numerals with the leading digit incremented between embodiments.

**[0096]** Figure 13 is an exploded view of an embodiment of a valve 240. The valve 240 may comprise a seal 242, an actuating component 244, and a valve lumen 245. The valve 240 may be configured such that the practitioner may selectively open and close the seal 242 to selectively allow access to the valve lumen 245. For example, a practitioner may open the seal 242 in order to introduce an instrument into the valve lumen 245 and close the seal 242 in order to prevent bleed back when the therapy allows. In some instances the seal 242 may be configured to seal the valve lumen 245 when no instrument is disposed therein and/or to seal about an instrument that has been introduced into the valve lumen 245.

**[0097]** The valve 240 may be configured with an actuator 244 configured to selectively open and close the seal 242. In some embodiments, displacement of the actuator 244 may be configured to open and/or close the seal 242. The actuator 244 may be configured to toggle between open and closed seal 242 positions, meaning the practitioner may displace the actuator 244 to change the configuration of the seal 242 and release the actuator 244 without the seal 242 changing configurations. In other embodiments the seal 242 or actuator 244 may be biased in either the closed or open position, requiring the practitioner to maintain pressure or contact with the actuator 244 to maintain the non-biased position. In the embodiment of Figure 13, axially displacing the actuator 244 with respect to the seal 242 may be configured to open and/or close the seal 242.

**[0098]** The valve 240 may further comprise a side port 246. The side port 246 may be configured to allow a practitioner to introduce fluid flow (e.g., contrast fluid, saline, etc.) into the valve lumen 245 whether the seal 242 is open or closed. Flow through or pressure within the side port 246 may prevent bleed back through the side port 246.

**[0099]** Figure 14 is a perspective view of another embodiment of a valve 340, Figure 15 is an exploded view thereof, and Figure 16 is a cross-sectional view of a

portion thereof. Figure 16 further includes an elongate device 350 disposed within a valve lumen 345 of the valve 340. The valve 340 of Figures 14-16 comprises a seal 342, an actuator 344, a valve lumen 345, and a side port 346. These components may function analogously to similarly indicated components of the valve 240 of Figure 13.

**[00100]** The actuator 344 of the valve 340 of Figures 14-16 may be a lever, which may be biased in an extended position. Depressing the actuator 344 may open or close the seal 342 depending on the arrangement of internal components of the valve 340. For example, the seal 342 may be compressed in a closed position when the actuator 344 is extended and may open as the actuator 344 is depressed. A mechanism such as a lever actuator 344 may be configured to allow a practitioner to incrementally open or close the seal 342 by incrementally depressing the actuator 344. This may allow a practitioner to open the seal 342 just enough to facilitate displacement of an elongate device 350 within the valve lumen 345 without opening the seal 342 such that there is bleed back around the elongate device 350. The elongate device 350 may comprise, for example, a biopsy device, a catheter, a stent delivery device, or any other transvascular therapy device. In the cross-sectional view of Figure 16, the seal 342 is closed about the elongate device 350 disposed in the valve lumen 345.

**[00101]** Figure 17 is a perspective view of another embodiment of a valve 440, Figure 18 is an exploded view thereof, and Figure 19 is a cross-sectional view of a portion thereof. Figure 19 further includes an elongate device 450 disposed within a valve lumen 445 of the valve 440. The valve 440 of Figures 17-19 comprises a seal 442, an actuator 444, a valve lumen 445, and a side port 446. These components may function analogously to similarly indicated components of the valves 240, 340 of Figures 13-16.

**[00102]** The actuator 444 of the valve 440 of Figures 17-19 may be configured such that axially displacing the actuator 444 in a distal direction with respect to the seal 442 causes a distal portion of the actuator 444 to interact with the seal 442, forcing the seal 442 open. Retracting the actuator 444 in a proximal direction with respect to the seal 442 may remove the actuator 444 from contact with the seal 442, allowing the seal 442 to close. The actuator 444 may be configured to remain in the open or closed configuration whether or not the practitioner is applying a force to the actuator 444, or the actuator 444 may be biased in one position or the other.

**[00103]** The actuator 444 may be configured to provide an audible and/or tactile indication that the seal 442 has been opened and/or closed. For example, the actuator 444 may be configured with ridges 447 configured to interact with other portions of the valve 440. Features such as ridges 447 or other detents may be configured to maintain the position of the actuator 444 (and therefore the seal 442) and/or provide an audible or tactile indication that the seal 442 has been changed from one configuration to another.

**[00104]** The valve 440 of Figure 18 may further comprise a rotary member 443. The rotary member 443 may be configured to interact with threads positioned on another portion of the valve 440 such that the rotational position of the rotary member 443 controls compressive forces acting on a conical member 441. The conical member 441 may be configured such that, when compressed, the conical member 441 tends to engage an elongate instrument disposed within the valve lumen 445. Thus, in some embodiments, compression of the conical member 445 couples such an instrument to the valve 440. This coupling may also act as a secondary seal, in connection with seal 442.

**[00105]** Any of the valves disclosed in any embodiment herein may comprise certain features described in connection with any other embodiment. For example, any of the valves may be configured to produce an audible and/or tactile indication of a change to the configuration of the valve. Other features of each embodiment may be analogously applied to the other embodiments.

**[00106]** Further, any of the valves described herein may be utilized in connection with any of the other components or assemblies described herein. Referring to Figure 1, a valve 140 may be used in connection with a medical device assembly such as transvascular liver biopsy assembly 100. In one exemplary procedure, a practitioner may insert an introducer sheath 110 into a patient with the valve 140 in a closed configuration. The practitioner may then open the valve 140 and insert a biopsy assembly 150 into the introducer sheath 110. Once the biopsy assembly 150 is inserted, the practitioner may then close the valve 140. The assembly may then be rotated (for example, the introducer sheath 110 may be rotated in connection with the rotary indexer 130) or otherwise manipulated into a desired position or configuration. The biopsy assembly 150 may then be used to obtain a tissue sample, with the valve 140 still in the closed configuration. The valve 140 may then be disposed in the open configuration and the biopsy assembly 150 withdrawn. The

valve 140 may again be closed. In other procedures, the configuration of the valve 140 may be other than that described, depending on the therapy or practitioner preference. For example, a practitioner may open the valve 140 when actuating a biopsy assembly 150 to obtain a sample or when manipulating the position of the assembly 100 or any component thereof. Further, a practitioner may advance or retract a component, such as biopsy assembly 150, with the valve 140 in a closed configuration, essentially pushing or pulling the component while in contact with a seal of the valve 140. Moreover, valves configured to toggle between the open and closed configurations without requiring a practitioner to maintain contact with an actuator, or valves biased in either the open or closed position (which change to the non-biased configuration only in response to continued input at an actuator) may each be used in connection with any of the procedures disclosed herein.

**[00107]** Figure 20 is a side view of the biopsy assembly 150 of the assembly 100 of Figure 1. The biopsy assembly 150 may comprise a handle 155 configured to manipulate, fire, or otherwise operate other components of the biopsy assembly 150, such as needles, trocars, stylets, cannulas, and so forth, in order to obtain a tissue sample. Any handle 155 shape, style, or design is within the scope of this disclosure. The biopsy assembly 150 of Figure 20 further comprises a stylet 160 disposed within a cannula 170. The stylet 160 and cannula 170 may be operably coupled to handle 155 such that manipulation or actuation of the handle 155 is configured to advance or retract the stylet 160 and/or cannula 170 in order to obtain a tissue sample. The biopsy assembly 150 of Figure 20, as well as any other biopsy assembly disclosed herein, may further comprise one or more outer sheath members (not shown), which may be disposed around the cannula 170 and/or stylet 160. An outer sheath member may be configured to deliver or protect the operative components and may be fixedly coupled to the handle 155.

**[00108]** Figure 21 is an exploded perspective view of the stylet 160 and cannula 170 of the biopsy assembly 150 of Figure 20. The stylet 160 may comprise a sharp distal end 162 configured to be advanced through tissue. A recessed trough 164 on the stylet 160 may be configured to isolate a tissue sample, which may be severed by the cannula 170. For example, and as further described below, the stylet 160 may be advanced into tissue such that the trough 164 of the stylet 160 extends beyond a distal tip 172 of the cannula 170. Tissue may prolapse into, or otherwise enter, the void formed by the trough 164. Tissue within the trough 164 may then be

severed by the distal tip 172 of the cannula 170 as the cannula is distally advanced with respect to the stylet 160. A distal segment 174 of the cannula 170 may then retain the severed tissue within the trough 164. The stylet 160 and cannula 170 may then be retracted together to remove the severed tissue sample.

**[00109]** Referring to Figure 1 and Figures 20-21, in some instances a biopsy assembly 150 may be configured to traverse an introducer sheath 110 or other path when positioning the distal tip 162 of the stylet 160 adjacent desired tissue within the body. The stylet 160 and cannula 170 may thus comprise elongate proximal segments 161, 171, respectively, configured to advance the distal tips 162, 172 of these components within the body. The proximal segments 161, 171 may operably connect the distal tips 162, 172 to the handle 155 while allowing the distal tips 162, 172 to be offset from the handle 155. The length of the proximal segments 161, 171 may be related to the therapy for which the biopsy assembly 150 is configured for use. For example, the biopsy assembly 150 may be configured for use in connection with a transjugular liver biopsy. In such instances, the proximal segments 161, 171 may be sized such that the handle 155 may be disposed proximally of the insertion site, while the distal tips 162, 172 are disposed adjacent liver tissue to be biopsied.

**[00110]** Referencing Figures 1 and 4, an introducer sheath 110 configured for use in connection with a biopsy assembly 150 may comprise a curved distal tip 112. The biopsy assembly 150 may comprise flexible segments configured to traverse the curvature of the curved distal tip 112.

**[00111]** Referring again to the biopsy assembly 150 of Figures 20-21, the stylet 160 and cannula 170 may each comprise a flexible segment 166, 176 disposed distally of the proximal segments 161, 171 of each component. The flexible segments 166, 176 may be configured to traverse a curved path and/or be disposed in a curved path when the biopsy assembly 150 is in use. Operative segments or segments configured to isolate, sever, or contain tissue samples (such as distal tip 162, trough 164, distal tip 172, and distal segment 174) may be disposed distally of the flexible segments 166, 176. In other embodiments, any of these components may be configured with multiple flexible segments disposed at various locations in order to facilitate use of the biopsy assembly 150 across delivery paths comprising multiple bends or curves.

**[00112]** Figure 22 is a side view of the stylet 160 of the biopsy assembly 150 of Figure 20, and Figure 23 is an enlarged view of a portion of the stylet 160 of Figure

22, taken around line 23-23. Figures 22 and 23 illustrate a portion of the proximal segment 161 of the stylet 160, as well as the flexible segment 166, the trough 164, and the sharp distal tip 162. In some embodiments, the stylet 160 may be generally formed of an elongate member having a generally circular cross-section. Certain features, such as the trough 164, may be formed by removing material from the generally circular member. The flexible segment 166 may be formed by grinding or otherwise removing material such that the flexible segment 166 has a smaller outside diameter than the proximal segment 161. The stylet 160 may comprise tapered or transition zones between the smaller-diameter flexible segment 166 and the proximal segment 161 and/or the operative distal end of the stylet 160. In some embodiments, the flexible segment 166 may have a generally circular cross-section. In other embodiments, the flexible segment 166 may have a cross-section other than a circular cross-section, including in embodiments where the proximal segment 161 has a generally circular cross-section.

**[00113]** Various elongate members, which may be configured to be passed through a sheath, are discussed herein. (For example: stylet 160 of Figures 22-23, cannula 170 of Figures 24 and 25, analogous stylets—such as 760 of Figure 36—and analogous cannula—such as 770 of Figure 36—of other embodiments, and the outer tubular member 780 of Figure 36). Any of these elongate instruments may be configured with a flexible segment. In some such embodiments, the flexible segment may be characterized as more flexible than another portion of the elongate instrument, such as a portion of the instrument disposed proximally or distally of the flexible segment. For example, a flexible segment may be more flexible than an operative segment disposed distally of the flexible segment. In some instances, the apparent bending modulus of the segments may be used to compare relative flexibilities. Further, in some embodiments, the relative flexibility of various segments may be determined by measuring the force required to deflect portions of identical lengths an identical amount. In some instances, the force required to deflect a length of a flexible segment a particular amount is an order of magnitude less than the force required to deflect a stiffer segment (either proximal or distal) having the same length, the same amount.

**[00114]** Figure 24 is a side view of the cannula 170 of the biopsy assembly 150 of Figure 20. Figure 25 is an enlarged view of a portion of the cannula 170 of Figure 24, taken around line 25-25. Figures 24 and 25 illustrate the proximal segment 171

of the cannula 170, as well as the flexible segment 176 and the distal tip 172. In some embodiments, the cannula 170 may comprise a generally hollow member. The flexible segment 176 may be formed by creating a spiral cut 177 in the wall of the hollow member. The spiral cut 177 may extend completely through a wall of the hollow member, from the outside diameter to the inside diameter. The spiral cut 177 may extend in one continuous helix along the flexible segment 176, or may be comprised of multiple cuts extending along certain portions of the flexible segment 176. Spiral cuts described in connection with this or any other embodiment here may be formed by laser cutting, grinding, mechanical cutting, or any other method.

**[00115]** Flexible segments, such as segment 166 of Figure 22 and segment 176 of Figure 25, may be positioned at various points along members of any biopsy device or similar device within the scope of this disclosure. Any disclosure provided herein in connection with the flexible segment of a solid member (such as flexible segment 166 of Figure 22) may be applied to any other elongate solid member of any embodiment of a biopsy device or other medical device. Similarly, any disclosure provided herein in connection with the flexible segment of a hollow member (such as the flexible segment 176 of Figure 25) may be applied to any other elongate hollow member of any embodiment of a biopsy device or other medical device.

**[00116]** Figure 26 is a side view of another embodiment of a flexible segment 276 of a component of a biopsy assembly. As with disclosure relative to other flexible segments, any disclosure provided in connection with this embodiment is applicable to any hollow member of any elongate device. In the embodiment of Figure 26, the flexible segment 276 comprises four spiral cuts 277a, 277b, 277c, 277d extending from an outside diameter to an inside diameter of the flexible segment 276. The spiral cuts 277a, 277b, 277c, 277d may be disposed such that the cuts do not cross each other, or may be disposed to cross at certain points. The cuts 277a, 277b, 277c, 277d may or may not be substantially parallel along the length of the cuts 277a, 277b, 277c, 277d. While the illustrated embodiment comprises four spiral cuts, other embodiments may comprise one, two, three, five, six, seven, eight, or any other number of spiral cuts. Further, certain embodiments may have segments of a different number of cuts at different points along the length of the flexible segment. For example, a flexible segment may have one spiral cut that runs the entire length of the flexible segment and additional spiral cuts that only extend along a portion of the length of the flexible segment.

**[00117]** Figure 27 is a side view of another embodiment of a flexible segment 376 of a component of a biopsy assembly. Figure 28 is an enlarged view of a portion of the flexible segment 376 of Figure 27, taken around line 28. In the illustrated embodiment, the flexible segment 376 comprises one spiral cut 377 extending the entire length of the flexible segment 376. In the illustrated embodiment, the spiral cut 377 is configured with a constant pitch along the entire length of the flexible segment 376. The pitch may be from about four revolutions per inch to about ten revolutions per inch, including about five, six, seven, eight, or nine revolutions per inch or any partial number in the range, such as 7.5 revolutions per inch.

**[00118]** Figure 29 is a side view of another embodiment of a flexible segment 476 of a component of a biopsy assembly. Figure 30 is an enlarged view of a portion of the flexible segment 476 of Figure 29, taken around line 30-30. Figure 31A is an enlarged view of a portion of the flexible segment 476 of Figure 30, taken around line 31A. In the embodiment of Figures 29-31A, a spiral cut 477 in the flexible segment 476 comprises a transition portion 478. In the illustrated embodiment, the pitch of the spiral cut 477 is not constant in the transition portion 478. Rather, it transitions from no pitch (or aligned with a longitudinal axis of the flexible segment 476) to a pitch about the flexible segment 476. Thus, in some embodiments, a spiral cut may begin at an end of the flexible segment 476 with very little or no pitch. The spiral cut 477 may then transition to a constant pitch, continuously vary along the length of the flexible segment 476, or vary for portions and remain constant for portions of the flexible segment 476. A spiral cut 477 with a transition portion 478 having a shallow pitch disposed at an end of the flexible segment 476 may be configured to reduce stress concentrations at the end of the flexible segment 476. A flexible segment may have a transition portion 478 at a distal end of the flexible segment, the proximal portion of the flexible segment, or both. Flexible segments having multiple spiral cuts may also be configured with a transition portion or other segment of non-constant pitch.

**[00119]** The embodiment of Figures 29-31A further comprises struts 479 disposed along the length of the spiral cut 477. The struts 479 may comprise small portions of the member which are not removed to form the spiral cut 477. The struts 479 may be configured to transfer longitudinal forces along the length of the flexible segment 476 while still increasing the flexibility of the flexible segment 476 with a spiral cut 477.

**[00120]** Figure 31B is an enlarged view of another embodiment of a portion of a flexible segment, analogous to the view of Figure 31A, showing another embodiment of a strut. Figure 31C is an enlarged view of a portion of Figure 31B. In the embodiment of Figures 31B and 31C, the strut comprises a socket portion 479a and a ball portion 479b. These components may be disposed to provide a strut structure which may flex along and around the strut itself. In some embodiments, a top portion of the ball portion 479b may be removed, such that the top of the ball portion 479b does not contact the socket portion 479a when the flexible portion is compressed along its axis. Rather, opposing sides of the spiral cut itself (such as 477 of Figure 30) may contact each other to bear a compressive force. This design may prevent the ball portion 479b from being easily deformed in response to compressive forces. Struts that comprise a socket portion 479a and a ball portion 479b may be configured to transfer longitudinal forces along the length of a flexible segment.

**[00121]** Some spiral cuts may be configured with no struts, some may comprise only struts such as that illustrated in Figure 31A, others comprise only struts such as that shown in Figures 31B and 31C, while other embodiments comprise struts of both varieties at various points along a spiral cut.

**[00122]** Figure 32 is a top view of a portion of an operative segment of another embodiment of a portion of a biopsy assembly; Figure 33 is a side view of a portion of the operative segment of Figure 32; and Figure 34 is an enlarged view of the portion of the operative segment of Figure 33, taken around line 34-34. In the illustrated embodiment, features of a sharpened distal tip 572 of a hollow biopsy device member are shown. The sharpened distal tip 572 may be configured to sever a tissue sample disposed within a recessed trough (such as trough 164 of Figure 23). The sharpened distal tip 572 may be configured both to sever the longitudinal length of the tissue sample and to interact with a distal end of a trough (such as trough 164 of Figure 23) to sever the distal end of the sample.

**[00123]** Various angles are illustrated in Figure 34. These angles may vary in various embodiments. For example, angle  $\alpha$  may vary from about 8 degrees to about 11 degrees, angle  $\beta$  from about 33 degrees to about 39 degrees, and angle  $\gamma$  from about 18 degrees to about 21 degrees.

**[00124]** Figures 35A-35C are side views of a portion of a biopsy assembly in a first configuration, a second configuration, and a third configuration, respectively. The

biopsy assembly may be operated by first advancing the device in the configuration of Figure 35A to a position within the body adjacent a site to be biopsied. The device may be advanced through an introducer sheath (such as introducer sheath 110 of Figure 4), including transjugular procedures wherein the introducer sheath is introduced to the vasculature at the jugular vein. In some such procedures, the biopsy device may be positioned adjacent a portion of the liver to be biopsied. In some such embodiments, flexible segments (such as 166 and 176 of Figures 22 and 25) may be disposed in a curved portion of the introducer sheath (such as curved distal tip 112 of Figure 4) when the sample is obtained. The illustrated embodiment comprises a stylet 660 having a distal end 662 and a recessed trough 664. The illustrated embodiment further comprises a cannula 670 which may be disposed over stylet 660. The cannula 670 may comprise a sharpened distal tip 672.

**[00125]** In some embodiments, the sample may be obtained by advancing the stylet 660 with respect to the cannula 670 such that the distal tip 662 of the stylet 660 and the trough 664 extend beyond the distal tip 672 of the cannula 670, as shown in Figure 35B, and into the tissue to be biopsied. Tissue may then prolapse into the recessed trough 664 of the stylet 660. The cannula 670 may then be advanced such that the distal tip 672 of the cannula 670 severs the length of the sample (along the length of the trough 664) and the distal end of the sample by interaction with the distal end of the trough 664, as shown in Figure 35C. In some embodiments the biopsy assembly may be configured with springs or other mechanisms such that the components displace to obtain the sample automatically when the device is actuated.

**[00126]** Figure 36 is a top view of a portion of another embodiment of a biopsy assembly. The biopsy assembly of Figure 36 comprises a stylet 760, a cannula 770, and an outer tubular member 780. In the embodiment of Figure 36, and as further described below, the cannula 770 and outer tubular member 780 may be configured to sever a tissue sample. Any of the disclosure above relating to any other biopsy device, including the positioning and function of flexible segments, may be applicable to biopsy devices with a cannula 770 and outer tubular member 780 such as in Figure 36.

**[00127]** Figure 37 is an enlarged view of a portion of the biopsy assembly of Figure 36, taken around line 37-37. As shown in Figure 37, the stylet 760 may comprise a

sharpened distal tip 762. The cannula 770 may comprise an opening 775 configured to receive a cutting member 785 coupled to the outer tubular member 780.

**[00128]** Figure 38 is a side view of a portion of the stylet 760 of the biopsy assembly of Figure 36. In the illustrated embodiment, the stylet 760 comprises a sharpened distal tip 762, a proximal segment 761, and a flexible segment 766. As with the stylets of other embodiments, the stylet 760 may comprise a generally solid circular member wherein the diameter of the flexible segment 766 is smaller than the diameter of the proximal segment 761. Unlike the stylets of earlier embodiments, the stylet 760 may not comprise a recessed trough, because in the embodiment of Figures 36-46D, the cannula 770 and outer tubular member 780 interact to isolate and sever a tissue sample.

**[00129]** Figure 39 is a side view of a portion of the cannula 770 of the biopsy assembly of Figure 36. The cannula 770 comprises a sharpened distal tip 772, a flexible segment 776, and a proximal segment 771 which may be less flexible than the flexible segment 776. The cannula 770 may comprise a generally hollow member, and the flexible segment 776 may comprise a spiral cut 777 extending between the inside diameter and the outside diameter of the member. Any spiral cut arrangement disclosed herein may be applied to the cannula 770 of this embodiment or any other hollow member of this embodiment, such as the outer tubular member (780 of Figure 40), for example.

**[00130]** The cannula 770 may also comprise an opening 775 and slots 773a, 773b configured to interact with the outer tubular member (780 of Figure 40) as further disclosed below. The slots 773a, 773b, opening 775 and other features of the cannula 770 or other elongate members (such as the tabs 783a, 783b or cutting member 785 of the outer tubular member 780 of Figure 40) are not necessarily drawn to scale for illustrative purposes.

**[00131]** Figure 40 is a side view of the outer tubular member 780 of the biopsy assembly of Figure 36. The outer tubular member 780 may comprise a cutting member 785 and a flexible segment 786 which may comprise a spiral cut 787. A proximal segment (not shown), which may be less flexible than the flexible segment 786, may be disposed proximally of the flexible segment 786. The outer tubular member 780 may also comprise a tabs 783a, 783b configured to interact with a slots (773a, 773b of Figure 39) as further described below.

**[00132]** Figure 41 is a side view of a portion of the biopsy assembly of Figure 36; Figure 42 is a cross-sectional view of the portion of the biopsy assembly of Figure 41, taken through plane 42-42; and Figure 43 is a cross-sectional view of the portion of the biopsy assembly of Figure 41, taken through plane 43-43. The stylet 760, the cannula 770, and the outer tubular member 780 are shown in these views. The stylet 760 may be disposed within the cannula 770, and the outer tubular member 780 may be disposed around the cannula 770. The sharpened distal tip 762 of the stylet 760, the opening 775 of the cannula 770, and the cutting member 785 and tabs 783a, 783b of the outer tubular member 780 are also shown.

**[00133]** Figure 44 is an enlarged view of a portion of the cross-sectional view of Figure 42, taken around line 44-44. Figure 44 illustrates two slots 773a of the cannula 770 as well as two tabs 783a of the outer tubular member 780. The stylet 760 is also shown. Figure 45 is a further enlarged view of a portion of the cross-sectional view of Figure 44, taken around line 45-45, further illustrating a slots 773a and tabs 783a. In the illustrated embodiment, two slots 773a and tabs 783a, disposed at 180 degrees to each other, are shown. In some embodiments, the assemblies may comprise multiple slots and tabs positioned around a circumference of the assembly at the same longitudinal position. In some embodiments they may be equally spaced, while in other embodiments they may be irregularly spaced. Further, additional slots (such as 773b of Figure 39) and tabs (such as 783b of Figure 40) may be positioned at other longitudinal points on the assemblies.

**[00134]** Referencing Figures 38-45, the tabs 783a, 783b may be configured to engage the slots 773a, 773b such that the outer tubular member 780 and cannula 770 are coupled such that they cannot rotate relative to each other at the engagement point. In some assemblies, both the outer tubular member 780 and cannula 770 may be configured with flexible segments which may comprise one or more spiral cuts on each member. These flexible segments may have differing spring constants, and thus the outer tubular member 780 and cannula 770 of an assembly may tend to rotate to different degrees as the assembly is advanced along a path. Thus, the tabs 783a, 783b and slots 773a, 773b may be configured to retain the relative rotational positions of the outer tubular member 780 and cannula 770 as the assembly is advanced along a path. Retention of such alignment may tend to keep other components (such as the cutting member 785 and opening 775 of Figure 46A) aligned while the assembly is in use. Further, the slots 773a, 773b may be

sized such that the cannula 770 and outer tubular member 780 may be axially displaceable relative to each other (for example during actuation of the assembly as described below) while still retaining the rotational positions of these components. In some embodiments the tabs may be disposed on the cannula 770 and configured to interact with slots on the outer tubular member 780.

**[00135]** Figures 46A-46E are side views of a portion of the biopsy assembly of Figure 36 in five configurations. The outer tubular member 780, cutting member 785, cannula 770, distal tip of the cannula 772, opening 775, stylet 760, and distal tip of the stylet 762 are shown in these views.

**[00136]** The configurations of Figures 46A-46E illustrate five positions of the assembly during a procedure. As with any other biopsy assembly disclosed herein, the biopsy assembly of these figures may be used in connection with a variety of procedures, including transvascular procedures. In some embodiments, for example, the biopsy assembly may be advanced through an introducer sheath (such as introducer sheath 110 of Figure 4). The assembly may be configured such that the flexible segments of the outer tubular member 780, cannula 770, and stylet 760 are disposed within a curved portion (such as the curved distal tip 112 of Figure 4) when the biopsy assembly is actuated to obtain a sample. Further, displacement of particular portions of the biopsy assembly may be controlled by a handle (such as handle 155 of Figure 20). In some embodiments, springs or other mechanisms may displace the components of the biopsy assembly to obtain a sample once the biopsy assembly is actuated.

**[00137]** In some embodiments the biopsy assembly may be advanced within a sheath or other path with the components in the configuration shown in Figure 46A. In this configuration, the biopsy assembly may be advanced along a delivery lumen and/or be pushed into tissue until the biopsy assembly is adjacent tissue to be biopsied. The distal tip 762 of the stylet 760 may be configured to pierce or separate tissue as the assembly is advanced within tissue. While the assembly is advanced or operated, the slots (773a of Figure 45) and tabs (783a of Figure 45) may resist relative rotation of the outer tubular member 780 and cannula 770 with respect to each other.

**[00138]** As shown in Figure 46B, the cannula 770 may be advanced distally beyond the distal tip 762 of the stylet 760 when the assembly is actuated. The distal tip 772 of the cannula 770 may extend into tissue, severing a longitudinal length of

the tissue sample as the cannula 770 is advanced. As shown in Figure 46C, the outer tubular member 780 may then be advanced with respect to the cannula 770. As the outer tubular member 780 is advanced, the cutting member 785 of the outer tubular member 780 may pass through the opening 775 in the cannula 770 such that the cutting member 785 severs a distal end of the tissue sample. A tissue sample may then be disposed proximally of the cutting member 785 and distally of the distal tip 762 of the stylet 760. The entire assembly may then be retracted together, as shown in Figure 46D, to retrieve the tissue sample from the body. For example, the assembly may be pulled back along a delivery path in the configuration of Figure 46D with a sample disposed within the assembly, between the cutting member 785 and the distal tip 762 of the stylet 760.

**[00139]** Once removed from the patient, the cannula 770 may be advanced with respect to the outer tubular member 780 such that the cutting member 785 is no longer disposed within the cannula 770, as shown in Figure 46E. The stylet 760 may then be advanced with respect to the cannula 770 and outer tubular member 780 such that the stylet 760 forces the sample out of the cannula 770.

**[00140]** Without further elaboration, it is believed that one skilled in the art can use the preceding description to utilize the present disclosure to its fullest extent. The examples and embodiments disclosed herein are to be construed as merely illustrative and exemplary and not a limitation of the scope of the present disclosure in any way. It will be apparent to those having skill in the art, and having the benefit of this disclosure, that changes may be made to the details of the above-described embodiments without departing from the underlying principles of the disclosure herein.

Claims

1. A rotary indexing device, configured for use in connection with a medical instrument, the rotary indexing device comprising:
  - a base; and
  - a rotary member rotatably coupled to the base and configured to be coupled to the medical instrument, the rotary member and base engaged such that interaction between the base and the rotary member releasably couples rotation of the rotary member with respect to the base at one or more stops such that rotation of the rotary member is configured to rotate the medical instrument at positions corresponding to the one or more stops.
2. The rotary indexing device of claim 1, further comprising one or more detents configured to releasably couple rotation of the rotary member with respect to the base at the one or more stops.
3. The rotary indexing device of claim 2, wherein the detents comprise a protrusion disposed on the rotary member and one or more recesses disposed on the base.
4. The rotary indexing device of claim 3, further comprising a retainer coupled to the base and a biasing element disposed between the retainer and the rotary member, the biasing element configured to exert a biasing force on the rotary member such that the protrusion tends to remain in a recess of the one or more recesses when the protrusion is aligned with the recess.
5. The rotary indexing device of any of claims 3 or 4, wherein recesses are disposed in a complete circular path on the base.
6. The rotary indexing device of any of claims 3, 4, or 5, wherein the recesses are disposed such that the rotary member rotates between about 10 degrees and about 45 degrees between engagement with adjacent recesses, which corresponds to rotation of the medical instrument about 15 degrees between stops.
7. The rotary indexing device of any of claims 2-6, wherein the detents are configured to create an audible indicia of detent engagement or disengagement as the rotary member is rotated with respect to the base.
8. The rotary indexing device of any of claims 2-7, wherein the detents are configured to create a tactile indicia of detent engagement or disengagement as the rotary member is rotated with respect to the base.

9. The rotary indexing device of any of claims 2-8, wherein visual indicia comprising different colors correspond to different positions of detent engagement.
10. The rotary indexing device of any of claims 1-9, wherein the rotary member further comprises a ridge configured to engage a slot on the medical instrument.
11. A rotary indexing assembly, for positioning of a medical device, the rotary indexing assembly comprising:
  - a rotary indexing device comprising:
    - a base; and
    - a rotary member rotatably coupled to the base, the rotary member and base engaged such that interaction between the base and the rotary member releasably couples rotation of the rotary member with respect to the base at one or more stops; and
    - a medical device coupled to the rotary member such that rotation of the rotary member also rotates the medical device.
12. The rotary indexing assembly of claim 11, wherein the medical device comprises an elongate member having a curved distal tip and a pointer coupled adjacent a proximal end of the elongate member, the pointer disposed such that a portion of the pointer lies in substantially the same plane as the curved distal tip.
13. The rotary indexing assembly of claim 12, wherein visual indicia on the base are disposed such that the position of the pointer with respect to the indicia corresponds to the rotational position of the curved distal tip.
14. The rotary indexing assembly of any of claims 11, 12 or 13, wherein the medical device comprises a transvascular introducer sheath.
15. The rotary indexing assembly of claim 14, wherein the transvascular introducer sheath is configured to advance a liver biopsy device within a body lumen.
16. The rotary indexing assembly of any of claims 11, 12, or 13, wherein the medical device comprises a transvascular liver therapy device configured for use in a transjugular intrahepatic portosystemic shunt procedure.
17. The rotary indexing assembly of any of claims 1, 12, or 13, wherein the medical device comprises a drainage catheter.

18. A method of transvascular therapy, comprising:
  - introducing an elongate device into the vasculature of a patient; and
  - rotating the elongate device about a longitudinal axis between predetermined stop points.
19. The method of claim 18, wherein rotating the elongate device between predetermined stops comprises rotating the elongate device until a detent between the elongate device and a base member engages.
20. The method of any of claims 18 or 19, wherein rotating the elongate device between predetermined stops comprises rotating the elongate device until perceiving an audible indication of the position of the elongate device.
21. The method of any of claims 18, 19, or 20, wherein rotating the elongate device between predetermined stops comprises rotating the elongate device until perceiving a tactile indication of the position of the elongate device.
22. The method of any of claims 18-21, wherein the elongate device comprises an introducer sheath and the method further comprises inserting a biopsy device through the introducer sheath.
23. The method of claim 22, further comprising obtaining a first tissue sample and a second tissue sample with the biopsy device and wherein the elongate device is rotated between obtaining the first tissue sample and the second tissue sample.
24. The method of any of claims 18-23, further comprising determining the orientation of a distal tip of the elongate device by observing a pointer coupled adjacent a proximal end of the elongate device.
25. A transvascular liver access assembly, comprising:
  - an introducer sheath configured to traverse a portion of the vasculature between an insertion point and the liver, the introducer sheath having a curved distal tip and a pointer coupled adjacent a proximal end of the introducer sheath, the pointer disposed such that a portion of the pointer lies in substantially the same plane as the curved distal tip.
26. The transvascular liver access assembly of claim 25, further comprising a valve coupled adjacent a proximal end of the introducer sheath, the valve configured to be disposable in an open configuration and in a closed configuration.

27. The transvascular liver access assembly of claim 26, where the valve is configured to toggle between the open and closed configuration.
28. The transvascular liver access assembly of any of claims 26 or 27, wherein the pointer is disposed between the valve and the introducer sheath.
29. The transvascular liver access assembly of any of claims 26, 27 or 28, wherein the valve is configured to create an audible indicia when the valve is toggled between the open and closed configurations.
30. The transvascular liver access assembly of any of claims 26-29, wherein the valve is configured to create a tactile indicia when the valve is toggled between the open and closed configurations.
31. The transvascular liver access assembly of any of claims 26-30, further comprising a rotary indexing device operably coupled to the introducer sheath, the rotary indexing device comprising:
  - a base; and
  - a rotary member coupled to the introducer sheath and rotatably coupled to the base, the rotary member and base engaged such that interaction between the base and the rotary member releasably couples rotation of the rotary member with respect to the base at one or more stops.
32. A method of transvascular liver access, comprising:
  - inserting an introducer sheath into the vasculature of a patient;
  - toggling a valve coupled to the introducer sheath from a closed configuration to an open configuration after insertion of the introducer sheath;
  - inserting a liver therapy device into the introducer sheath while the valve is in the open configuration;
  - toggling the valve from an open to a closed configuration after insertion of the liver therapy device.
33. The method of transvascular liver access of claim 32, wherein the liver therapy device comprises a biopsy device, the method further comprising obtaining at least one tissue sample.
34. The method of claim 33, further comprising rotating the introducer sheath to obtain additional tissue samples.
35. The method of claim 34, wherein the valve is toggled to the closed configuration prior to rotating the introducer sheath.

36. The method of any of claims 33, 34, or 35, further comprising withdrawing the biopsy device from the introducer sheath.
37. The method of claim 36, where the valve is toggled from the closed to the open configuration prior to inserting the biopsy device and toggled from the closed to the open configuration prior to withdrawing the biopsy device.
38. The method of claim 37, further comprising determining if the valve has been toggled based on an audible indication of toggling.
39. The method of any of claims 37 or 38, further comprising determining if the valve has been toggled based on a tactile indication of toggling.
40. A transvascular liver therapy device comprising:
  - an elongate member comprising a proximal segment; a flexible segment distal of the proximal segment, the flexible segment being more flexible than the proximal segment; and an operative segment disposed distal of the flexible segment.
41. The transvascular liver therapy device of claim 40, wherein the flexible segment comprises a substantially circular cross-section, the proximal segment comprises a substantially circular cross-section, and wherein the flexible segment has a smaller outside diameter than the proximal segment.
42. The transvascular liver therapy device of claim 40, wherein the flexible segment comprises a hollow member having at least one spiral cut extending between an inside diameter and an outside diameter of the flexible segment.
43. The transvascular liver therapy device of claim 42, wherein the proximal segment comprises a hollow member integrally formed with the flexible segment.
44. The transvascular liver therapy device of any of claims 40-43, wherein the liver therapy device comprises a transjugular liver biopsy device.
45. The transvascular liver therapy device of any of claims 40, 41, or 44, wherein the elongate member comprises a stylet having a substantially circular cross-section, the operative segment comprising a partial core trough portion.
46. The transvascular liver therapy device of claim 45, further comprising a hollow cannula disposed around the stylet, the cannula comprising a proximal segment, a flexible segment distal of the proximal segment, and an operative segment disposed distal of the flexible segment.

47. The transvascular liver therapy device of claim 46, wherein the flexible segment of the elongate member and the flexible segment of the cannula are disposed adjacent each other.
48. The transvascular liver therapy device of any of claims 46 or 47, wherein the proximal and operative segments of the cannula comprise solid walls and the flexible segment of the cannula comprises a spiral cut in a wall of the flexible segment of the cannula.
49. The transvascular liver therapy device of claim 48, wherein the spiral cut is disposed about the flexible segment at between about four and about ten revolutions per inch.
50. The transvascular liver therapy device of claim 46, wherein the proximal and operative segments of the cannula comprise solid walls and the flexible segment of the cannula comprises more than one spiral cut in a wall of the flexible segment.
51. The transvascular liver therapy device of any of claims 48 or 49, wherein the spiral cut has a constant pitch along the entire flexible segment.
52. The transvascular liver therapy device of any of claims 48 or 49, wherein the spiral cut transitions from a shallow pitch at an end of the flexible segment and a steeper pitch at a midpoint of the flexible segment.
53. The transvascular liver therapy device of any of claims 48-49 or 51-52, wherein one or more struts are disposed along the path of the spiral cut.
54. The transvascular liver therapy device of claim 53, wherein at least one strut comprises a solid member.
55. The transvascular liver therapy device of any of claims 53 or 54, wherein at least one strut comprises a ball portion and a socket portion.
56. The transvascular liver therapy device of claim 44,  
wherein the elongate member comprises a cannula, and the biopsy device further comprises:
  - a stylet disposed within the cannula, the cannula configured to extend distally beyond the stylet to obtain a tissue sample; and
  - a cutting member coupled to an outer tubular member, the outer tubular member disposed around the cannula and the cutting member configured to sever a distal end of a tissue sample when the outer tubular member is displaced distally with respect to the cannula, the outer tubular

- member comprising a proximal segment; a flexible segment distal of the proximal segment, the flexible segment being more flexible than the proximal segment; and an operative segment disposed distal of the flexible segment.
57. The transvascular liver therapy device of claim 56, further comprising one or more engagement features configured to resist rotation of the outer tubular member with respect to the cannula and the cannula with respect to the outer tubular member.
58. The transvascular liver therapy device of claim 57, wherein the engagement features comprise a plurality of protrusions and slots.
59. A method of obtaining a tissue sample comprising:  
introducing a biopsy device into the vasculature of a patient, the biopsy device comprising an elongate member comprising a proximal segment; a flexible segment distal of the proximal segment, the flexible segment being more flexible than the proximal segment; and an operative segment disposed distal of the flexible segment;  
advancing the biopsy device to a tissue sample location, the biopsy device disposed such that the flexible segment facilitates advancement of the device along a curved path;  
obtaining a tissue sample by advancing the operative segment of the biopsy device into the tissue; and  
withdrawing the biopsy device from the vasculature.
60. The method of claim 59, wherein the flexible segment is disposed in a curved path when the tissue sample is obtained.
61. The method of any of claims 59 or 60, wherein introducing the biopsy device into the vasculature of a patient comprises inserting the biopsy device into an introducer sheath disposed within the vasculature of a patient.
62. The method of any of claims 59, 60 or 61, wherein the introducer sheath comprises a curved distal tip and wherein the flexible segment is at least partially disposed along a curved path defined by the curved distal tip when the tissue sample is obtained.
63. The method of any of claims 59-62,  
wherein the elongate member comprises a stylet having a partial core trough and wherein the biopsy device further comprises a hollow cannula

disposed around the stylet, a flexible segment of the cannula disposed adjacent the flexible segment of the stylet; and

wherein obtaining a tissue sample comprises advancing the stylet into the tissue such that tissue prolapses into the trough and advancing the cannula to sever the prolapsed tissue.

64. The method of claim 63, wherein the flexible segment of the stylet and the flexible segment of the cannula are disposed in curved paths when the tissue sample is obtained.

65. The method of any of claims 59-62,

wherein the elongate member comprises a cannula, and the biopsy device further comprises an outer tubular member disposed around the cannula and a cutting member coupled to the outer tubular member, a flexible segment of the outer tubular member disposed adjacent the flexible segment of the cannula; and

wherein obtaining a tissue sample comprises advancing the cannula into the tissue such that the cannula severs the outside diameter of a full core tissue sample and advancing the outer tubular member such that the cutting member severs a distal end of the tissue sample.

66. The method of claim 65, wherein the flexible segment of the cannula and the flexible segment of the outer tubular member are disposed in curved paths when the tissue sample is obtained.

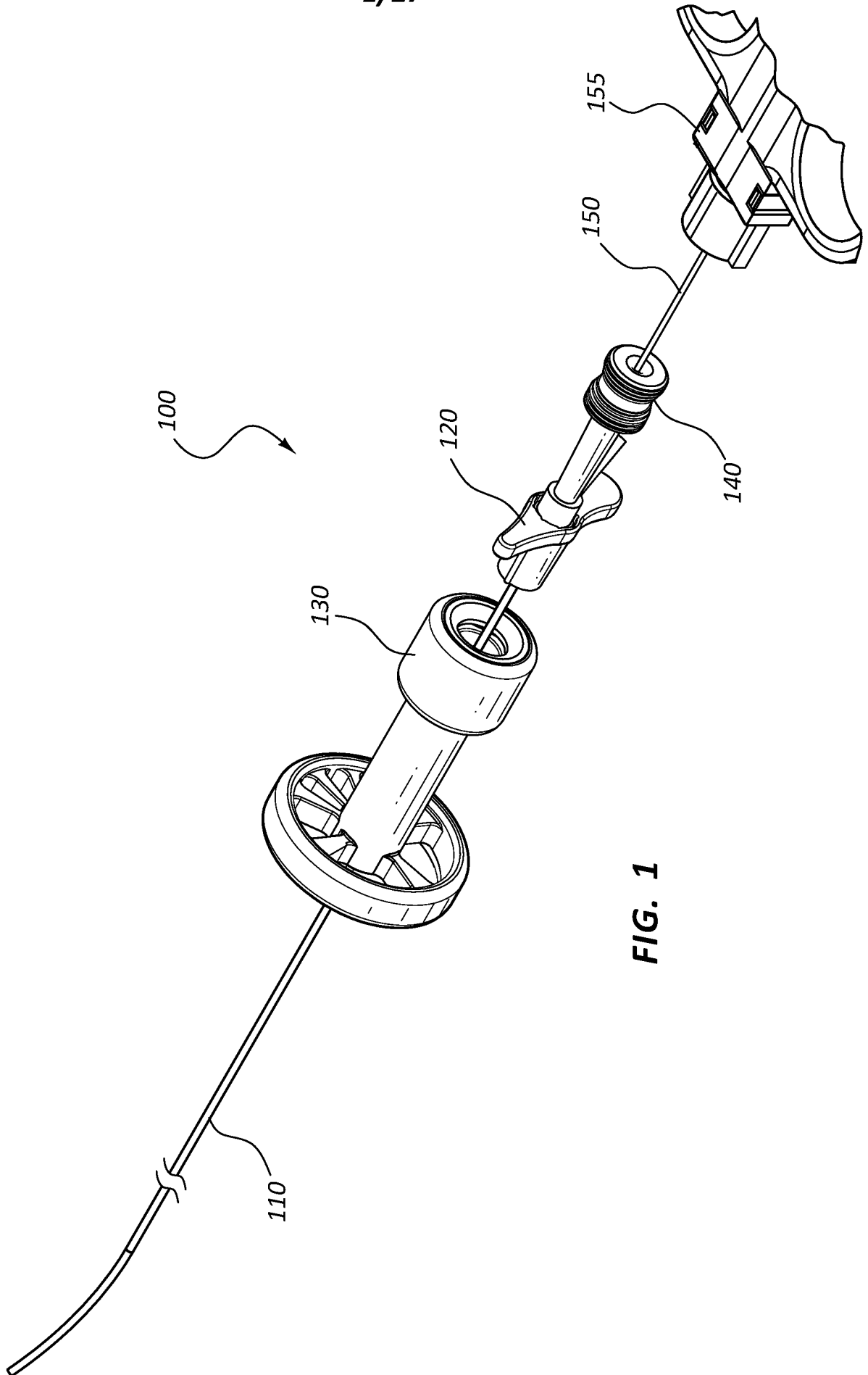


FIG. 1

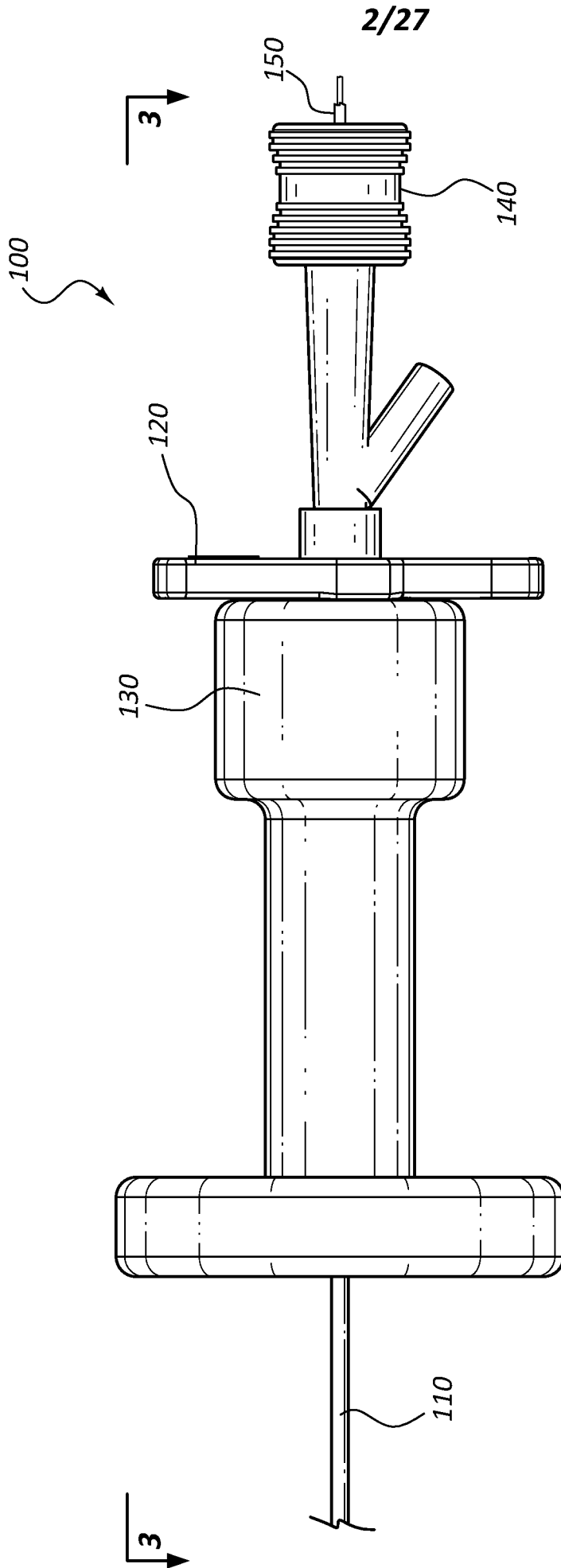


FIG. 2

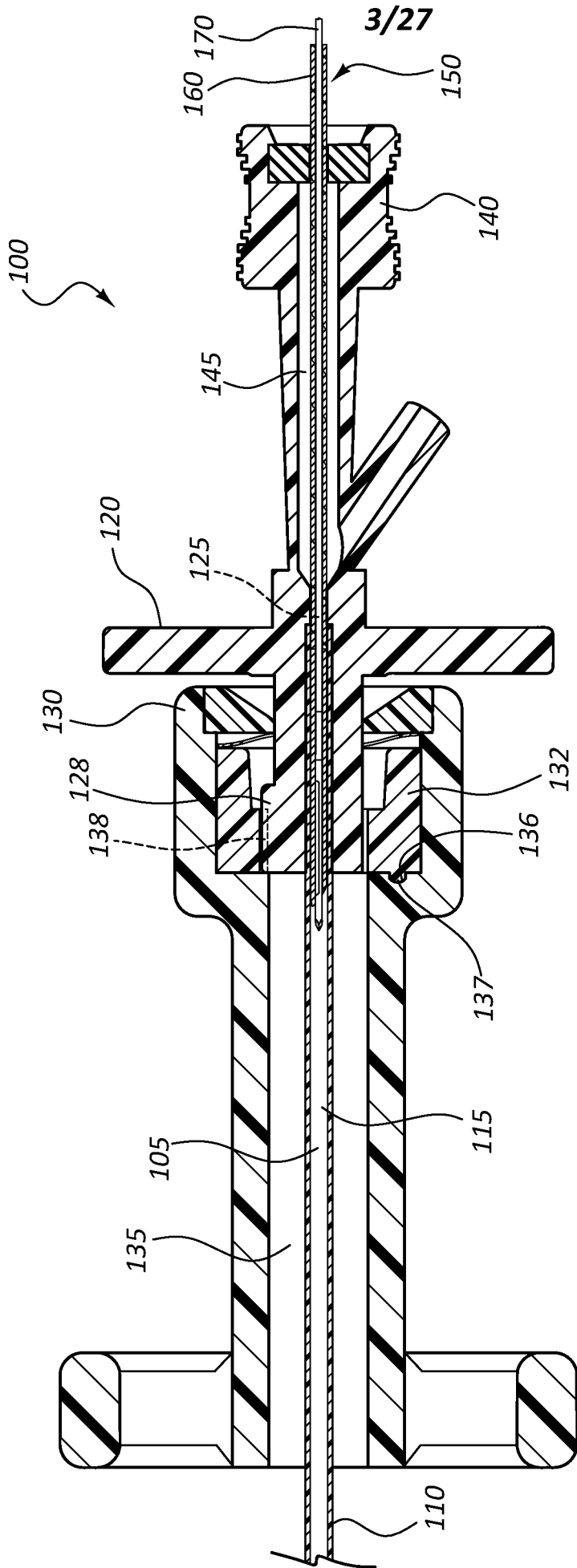


FIG. 3

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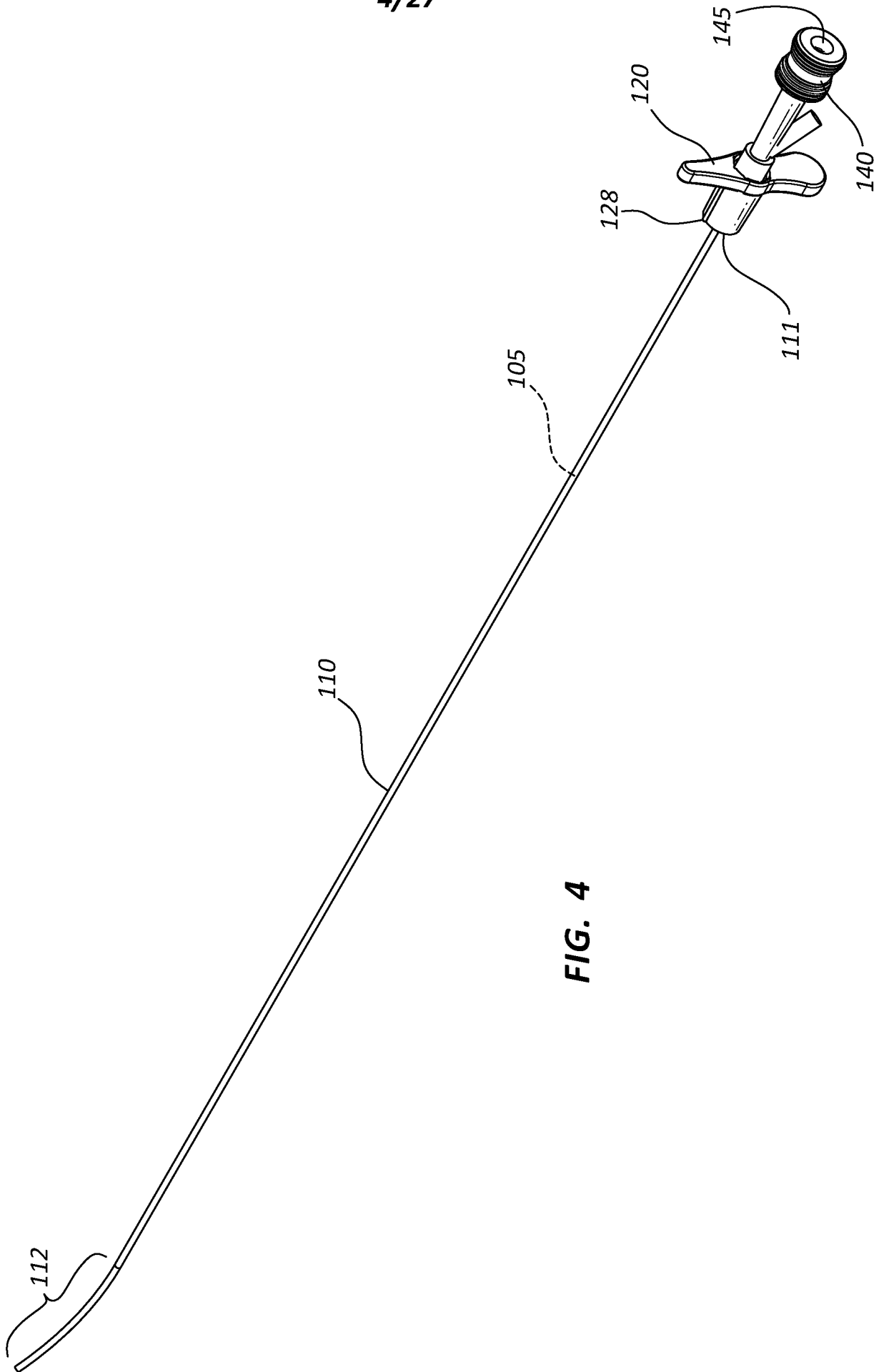


FIG. 4

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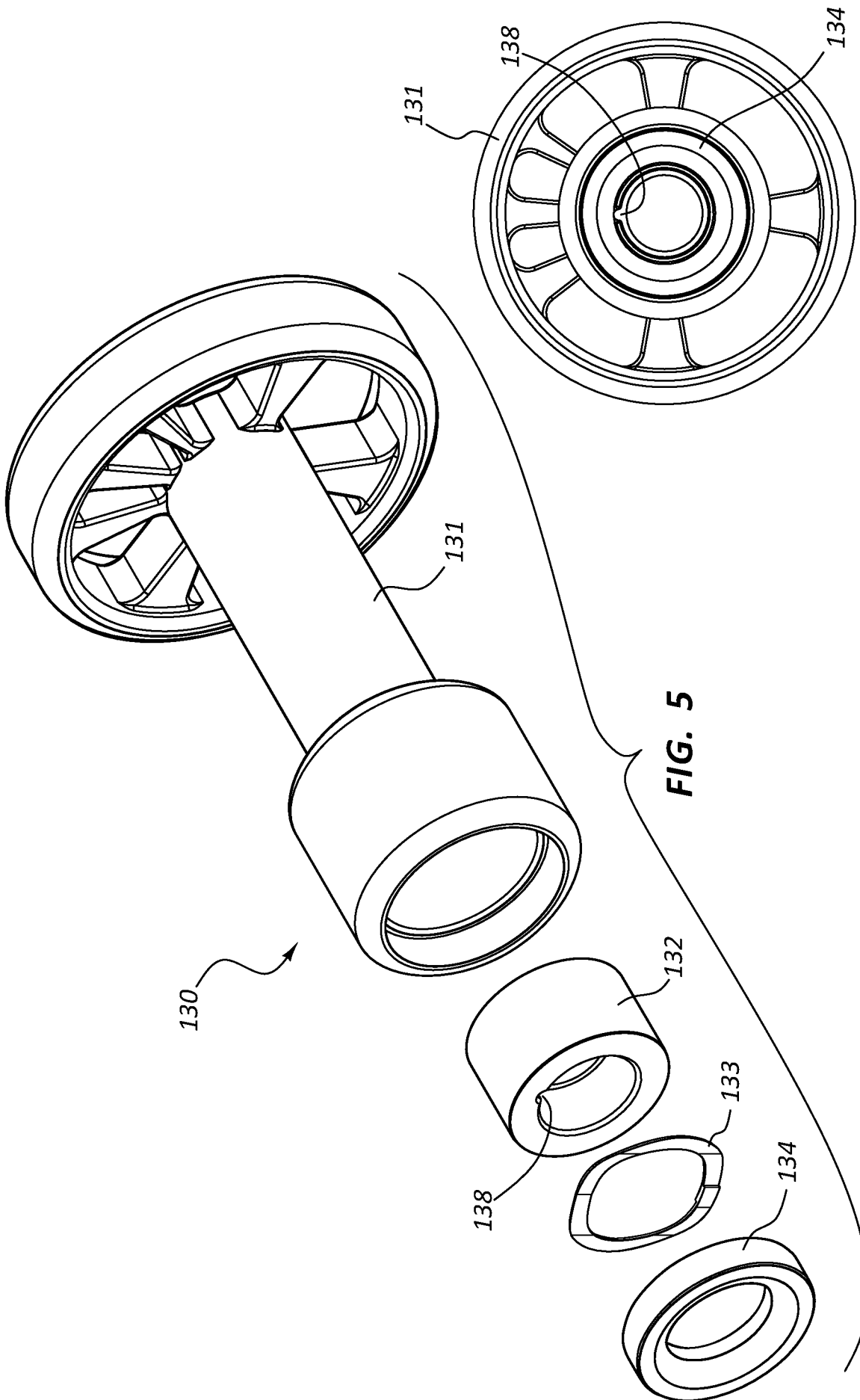


FIG. 5

FIG. 6

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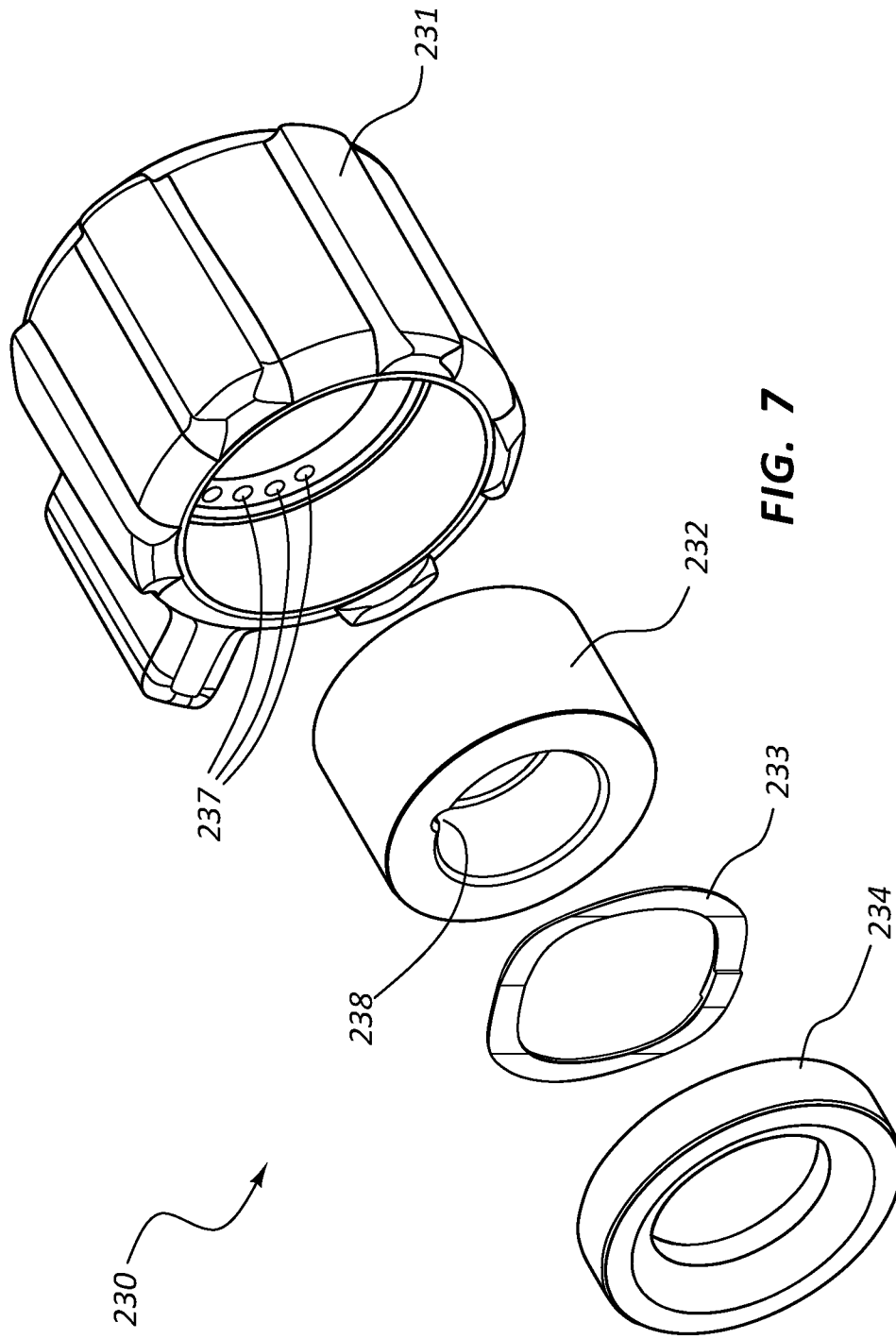


FIG. 7

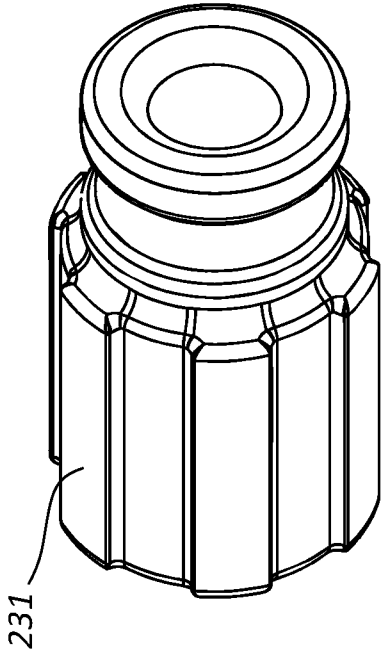


FIG. 9

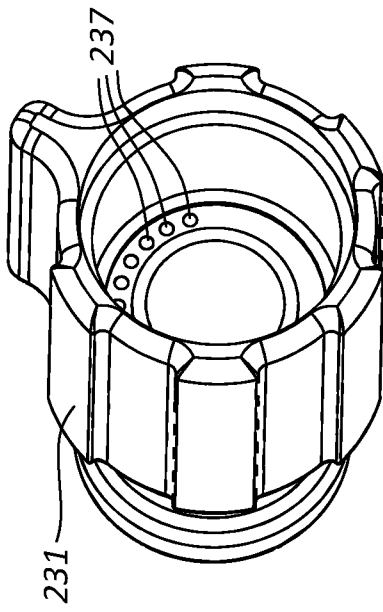


FIG. 8

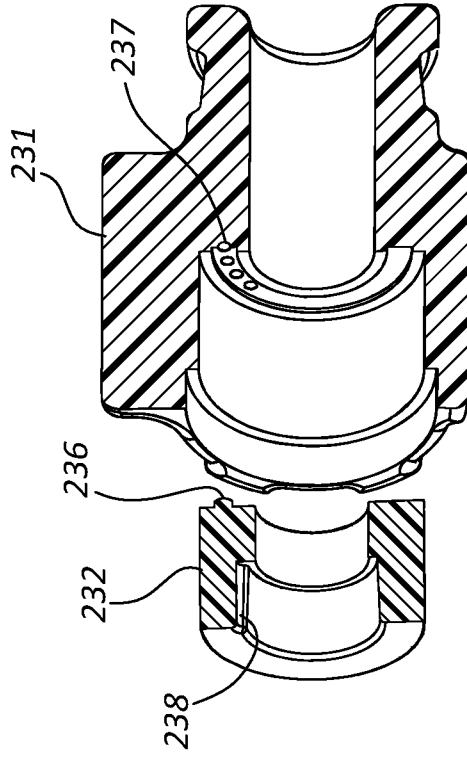


FIG. 12

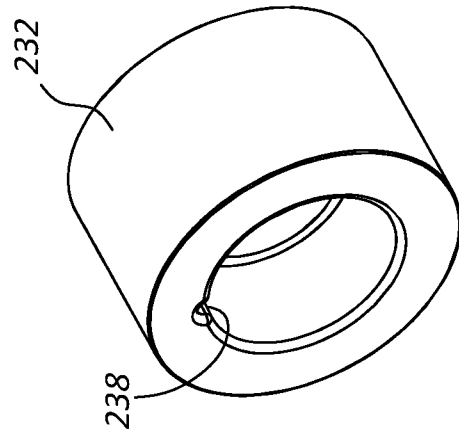


FIG. 11

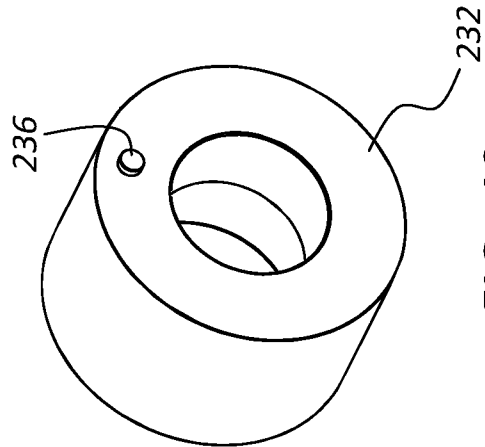


FIG. 10

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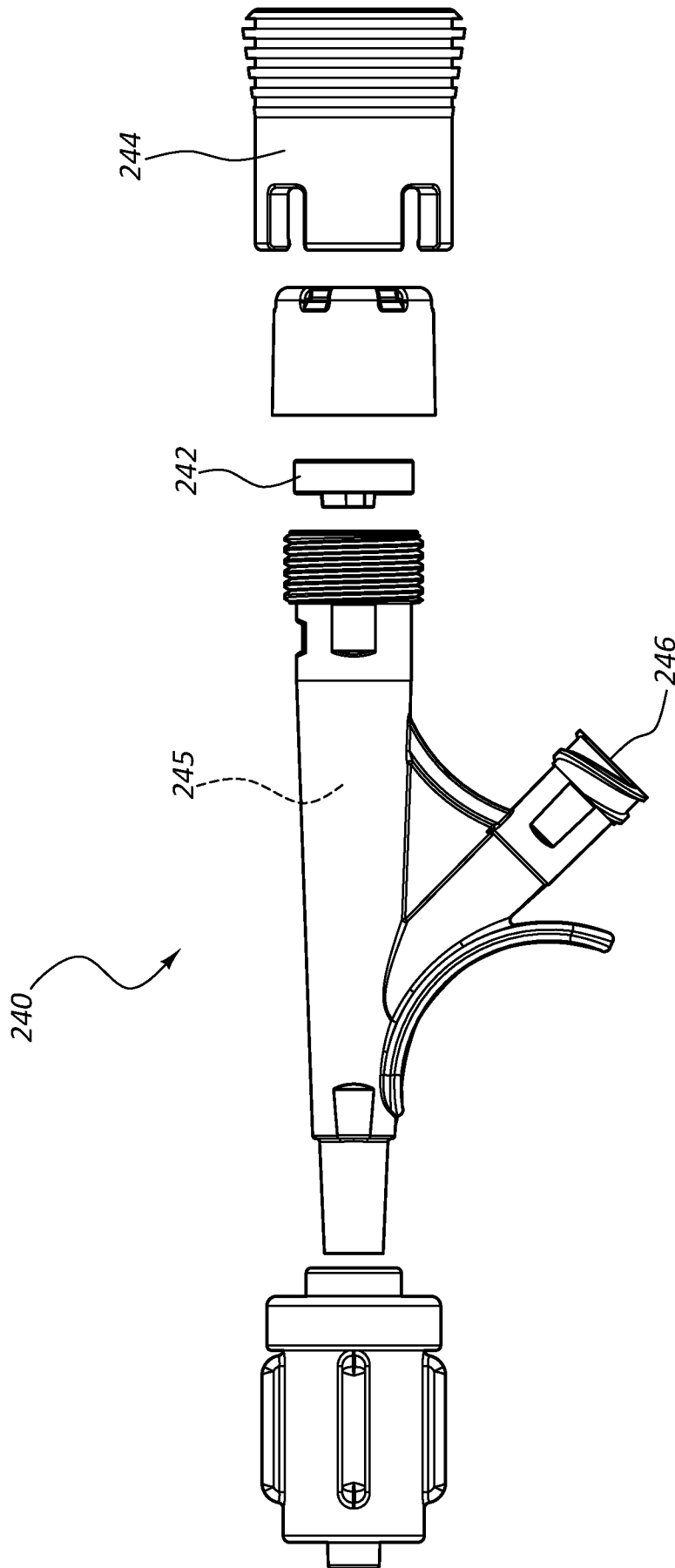


FIG. 13

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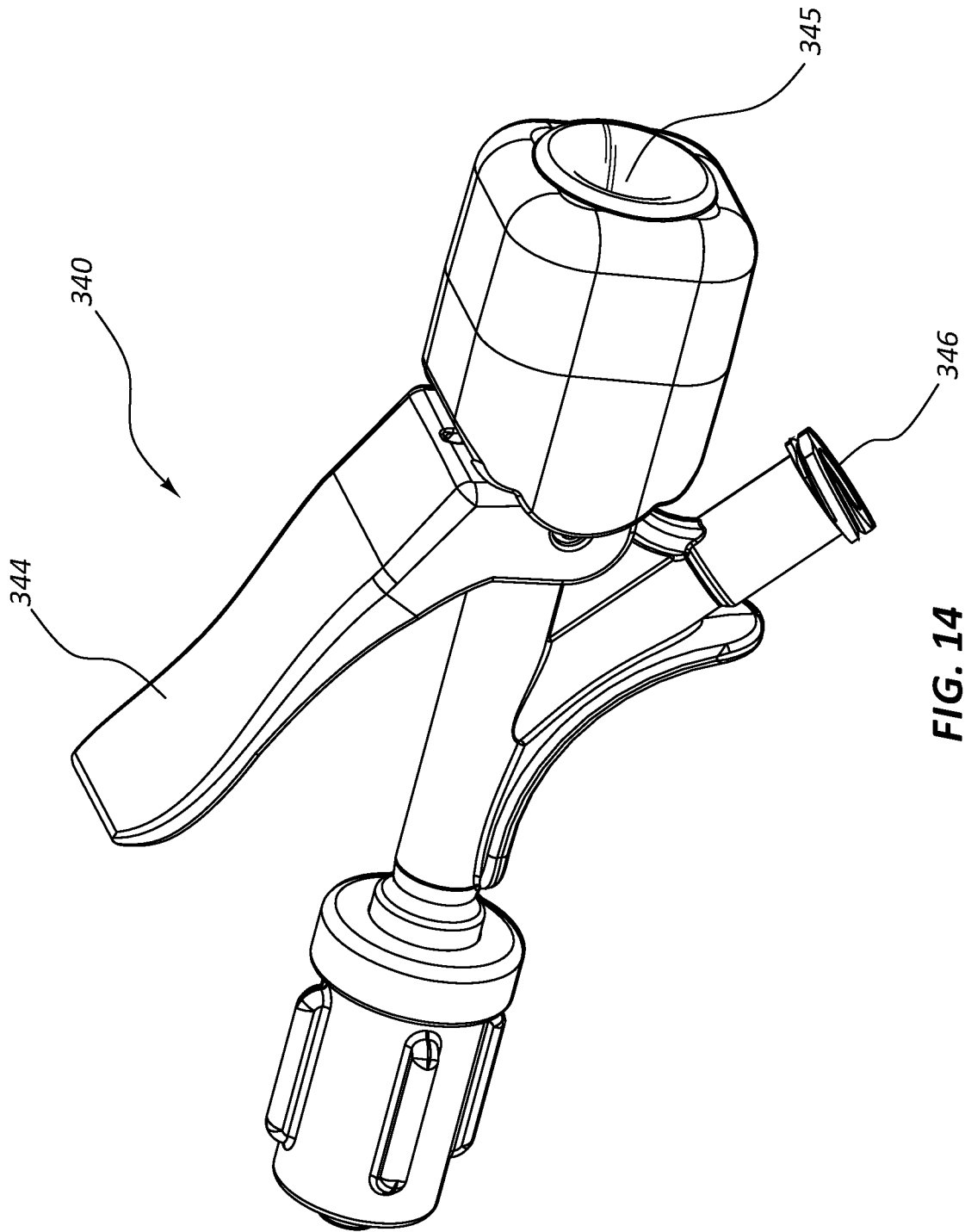


FIG. 14

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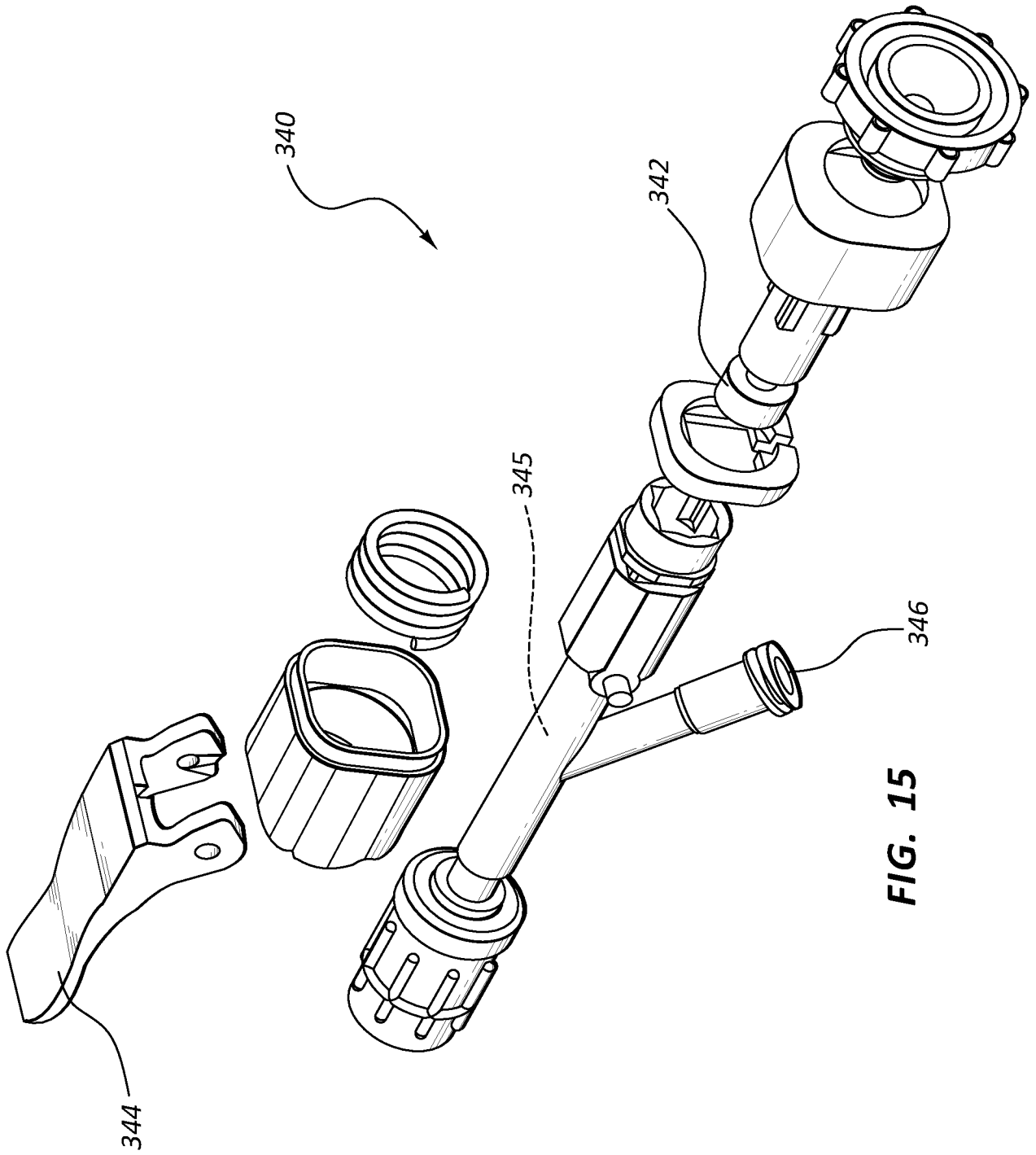


FIG. 15

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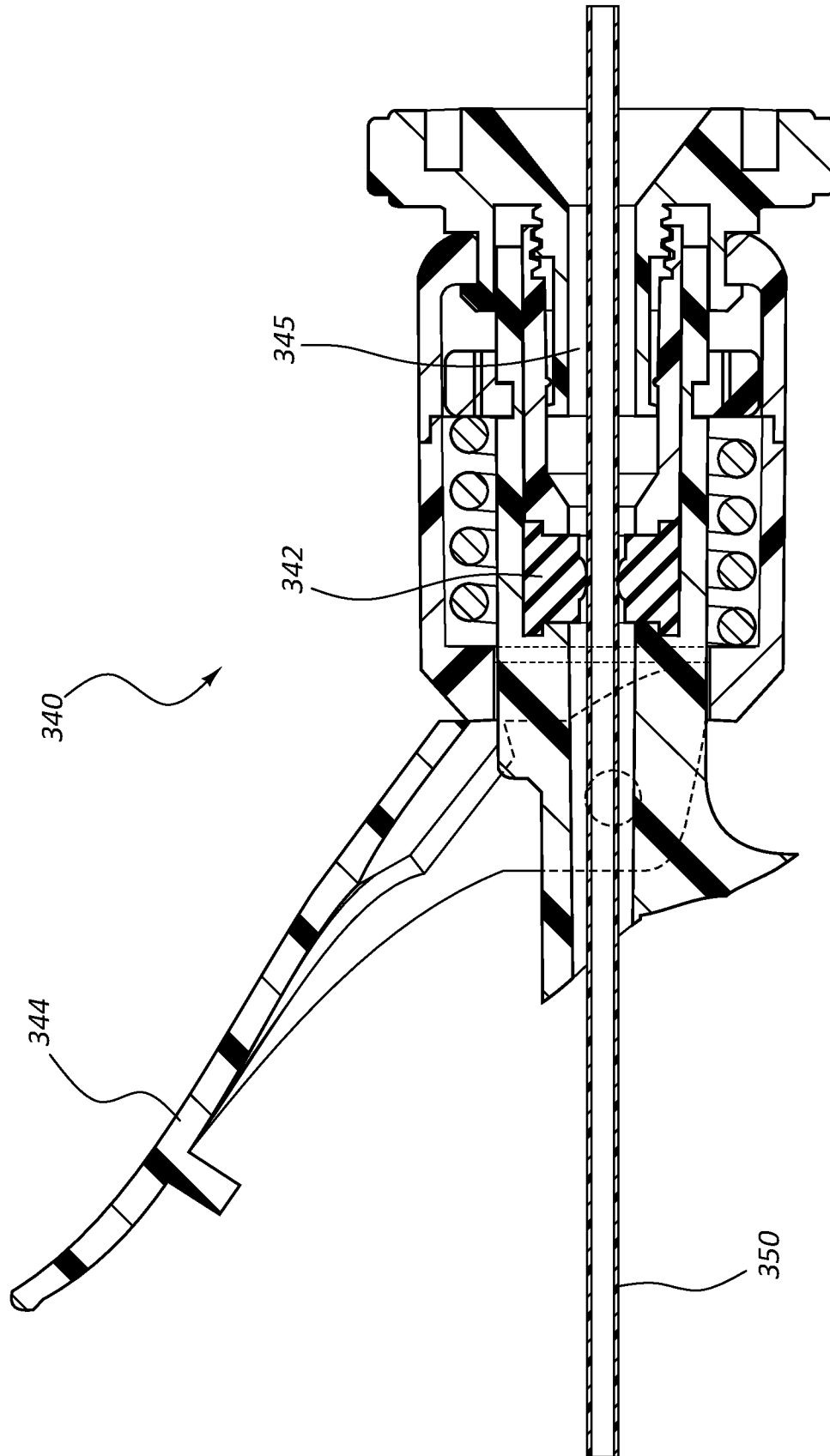


FIG. 16

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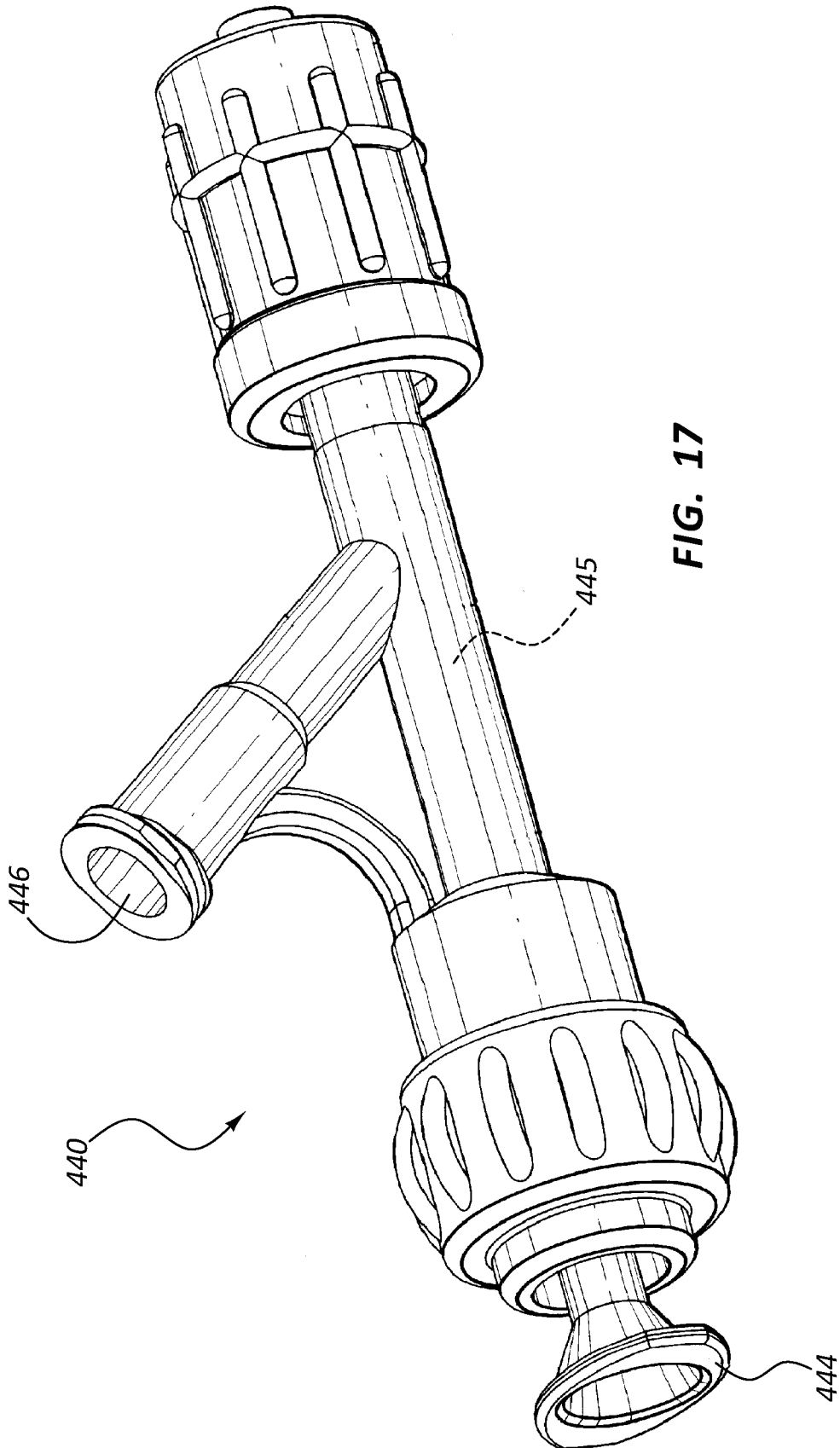


FIG. 17

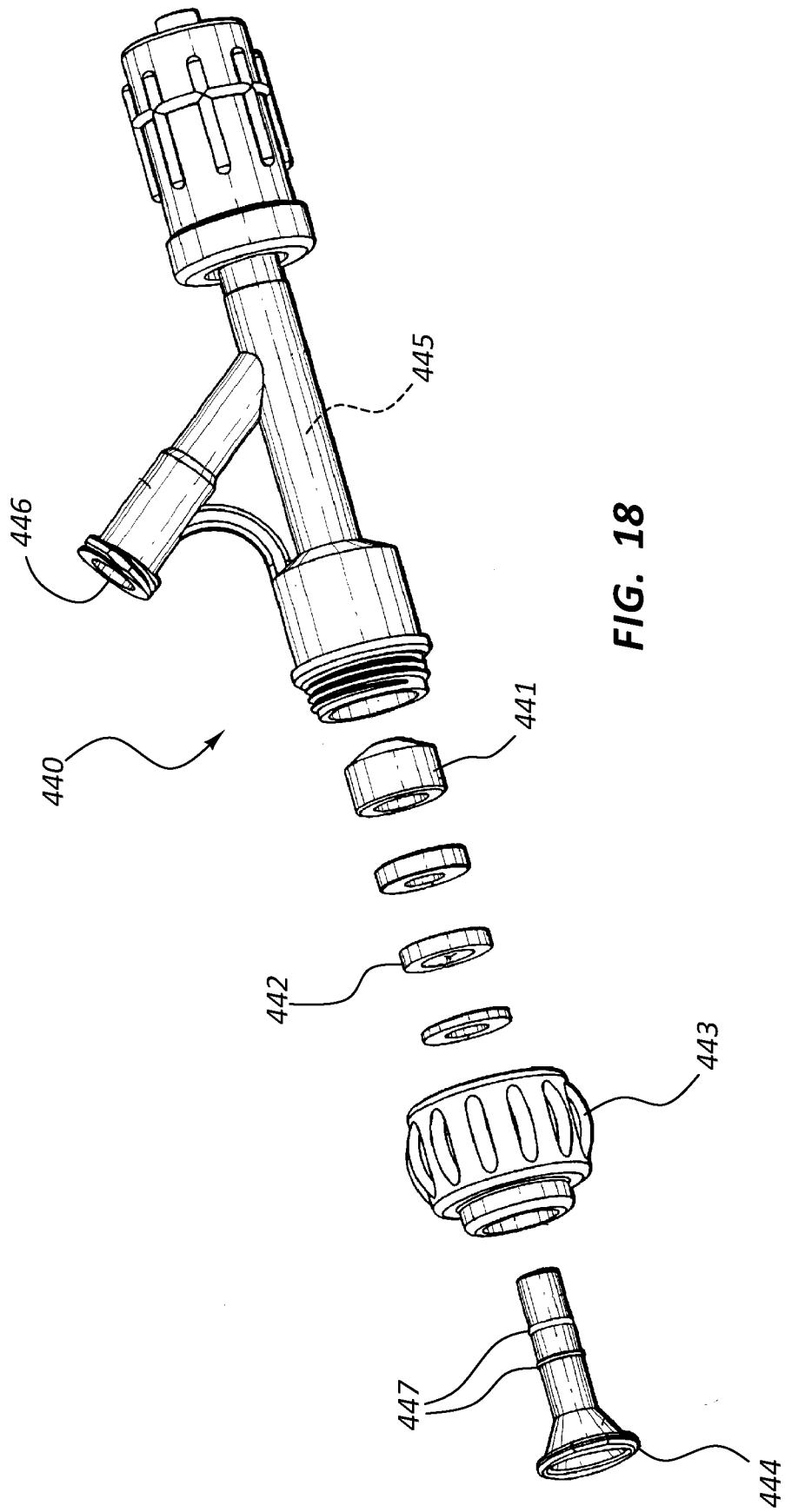


FIG. 18

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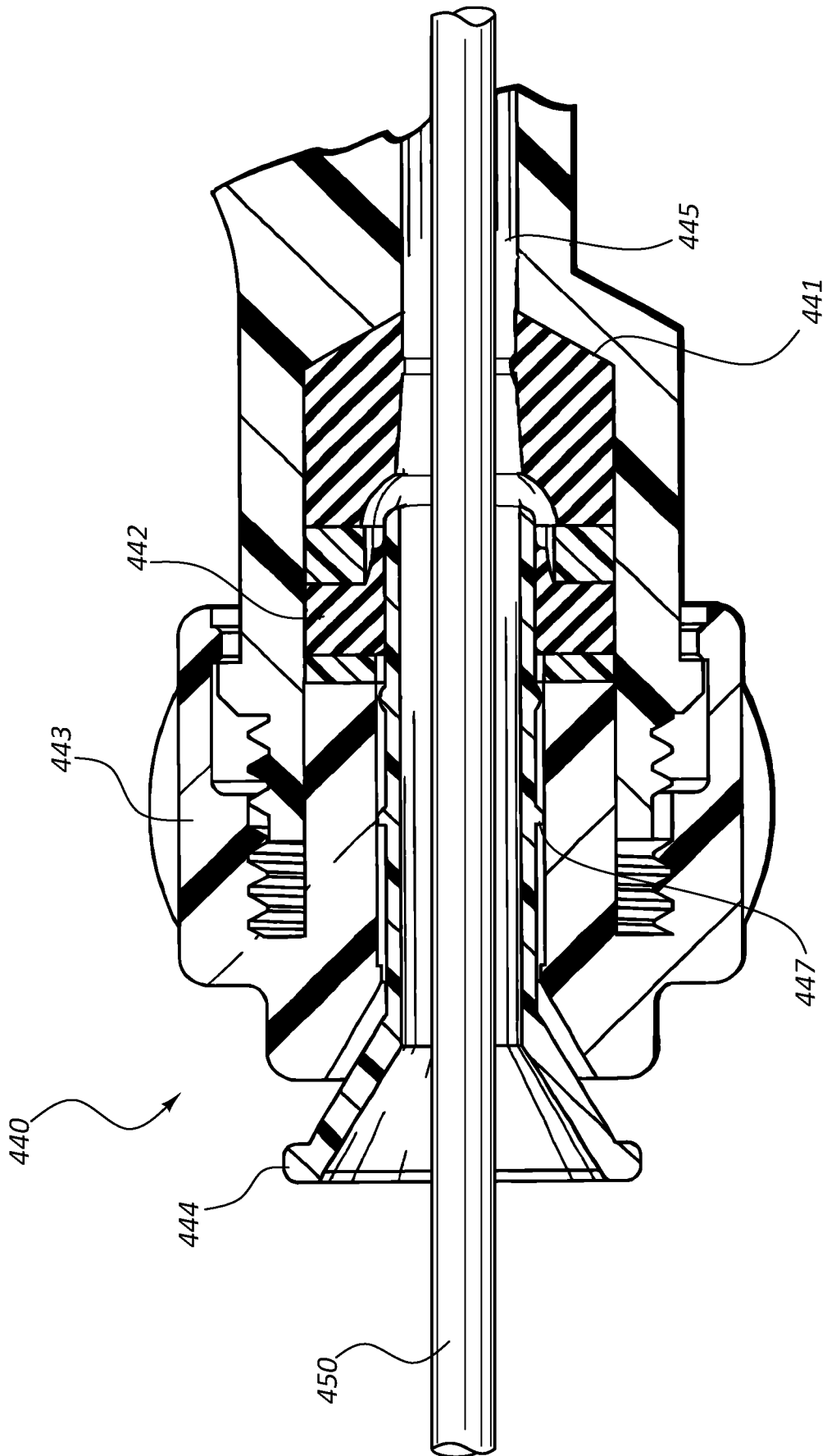


FIG. 19

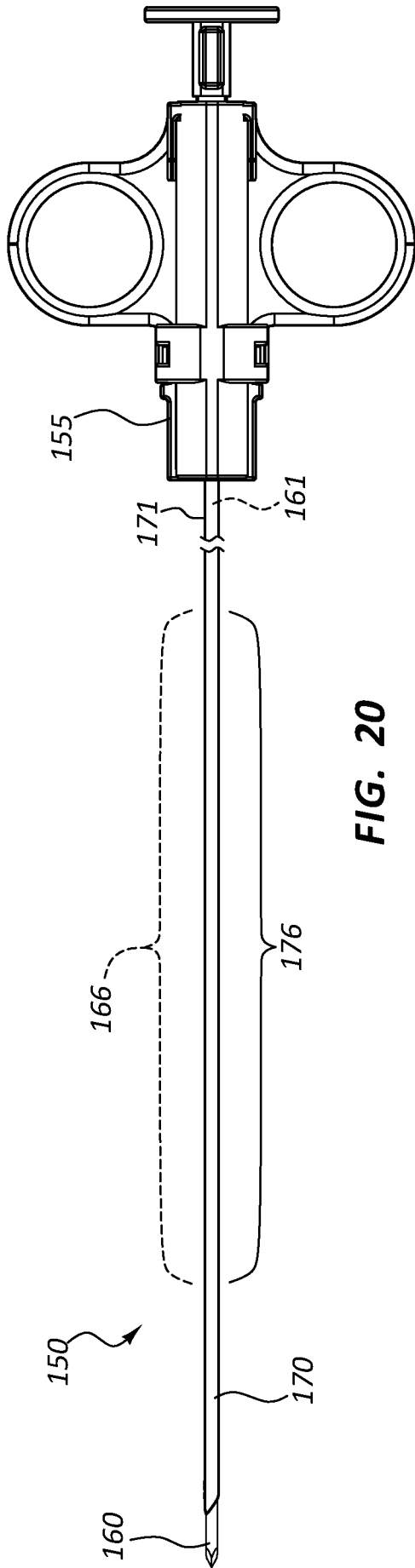


FIG. 20

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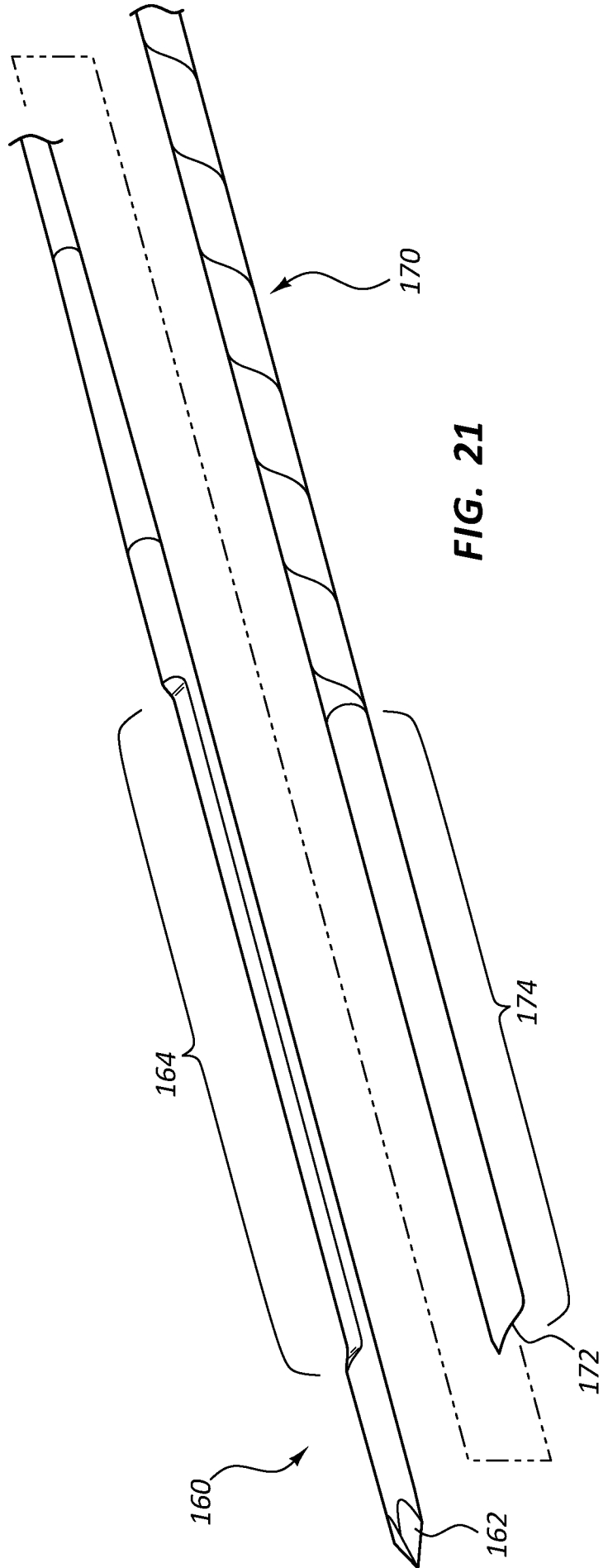


FIG. 21

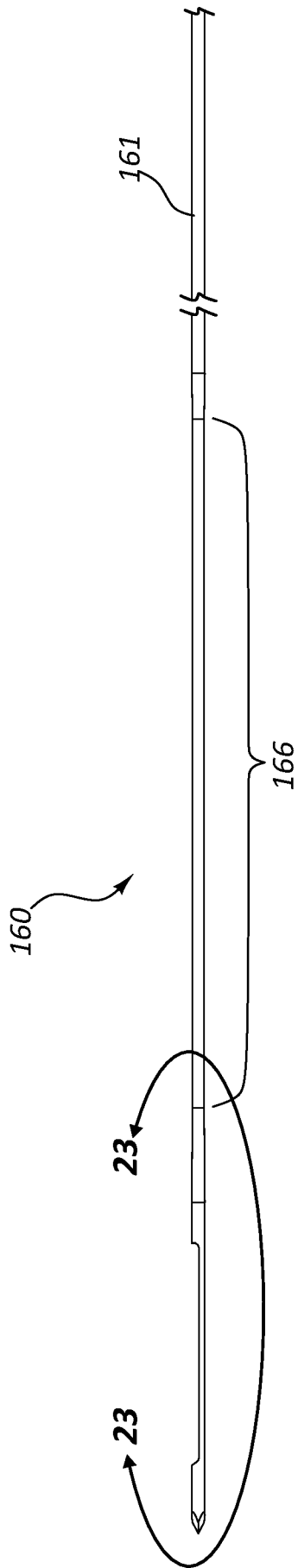


FIG. 22

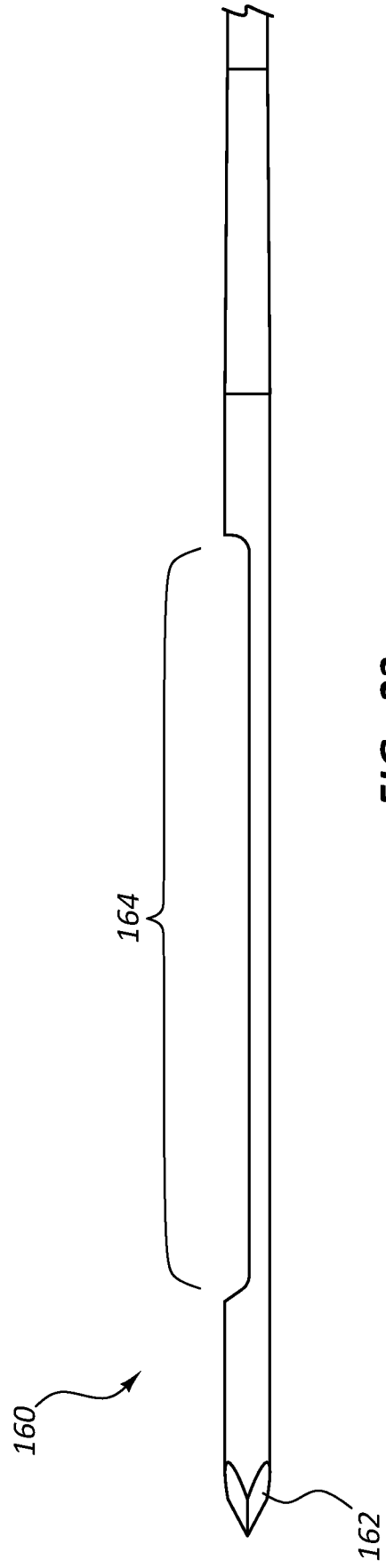


FIG. 23

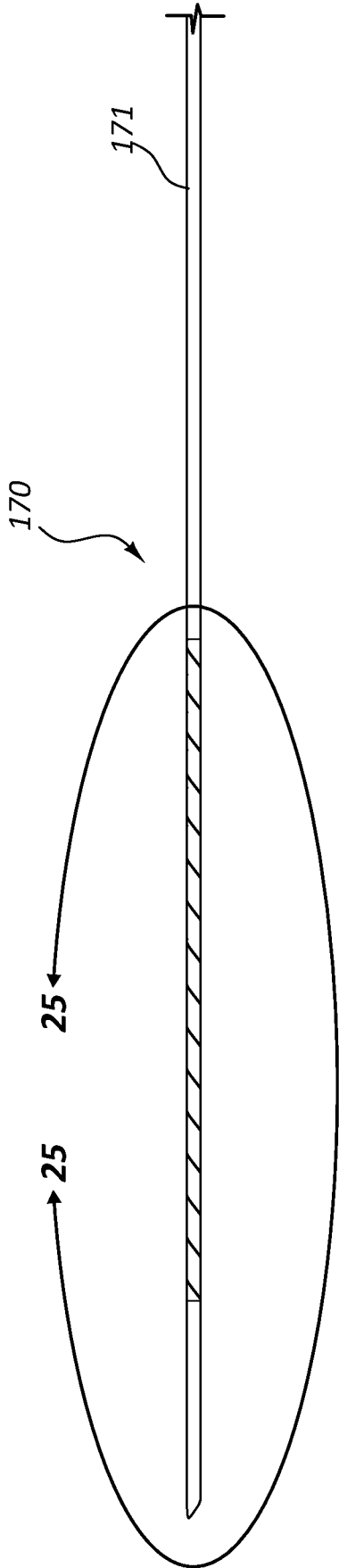


FIG. 24

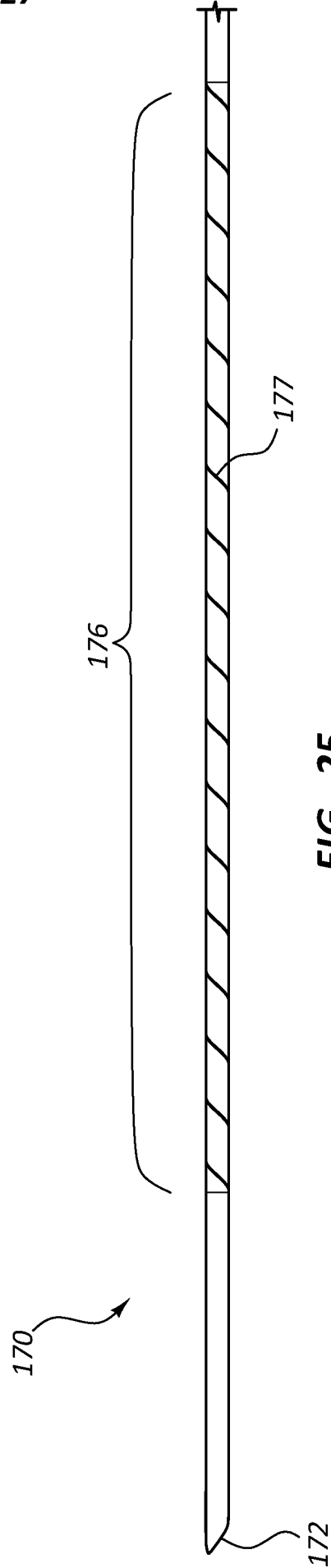
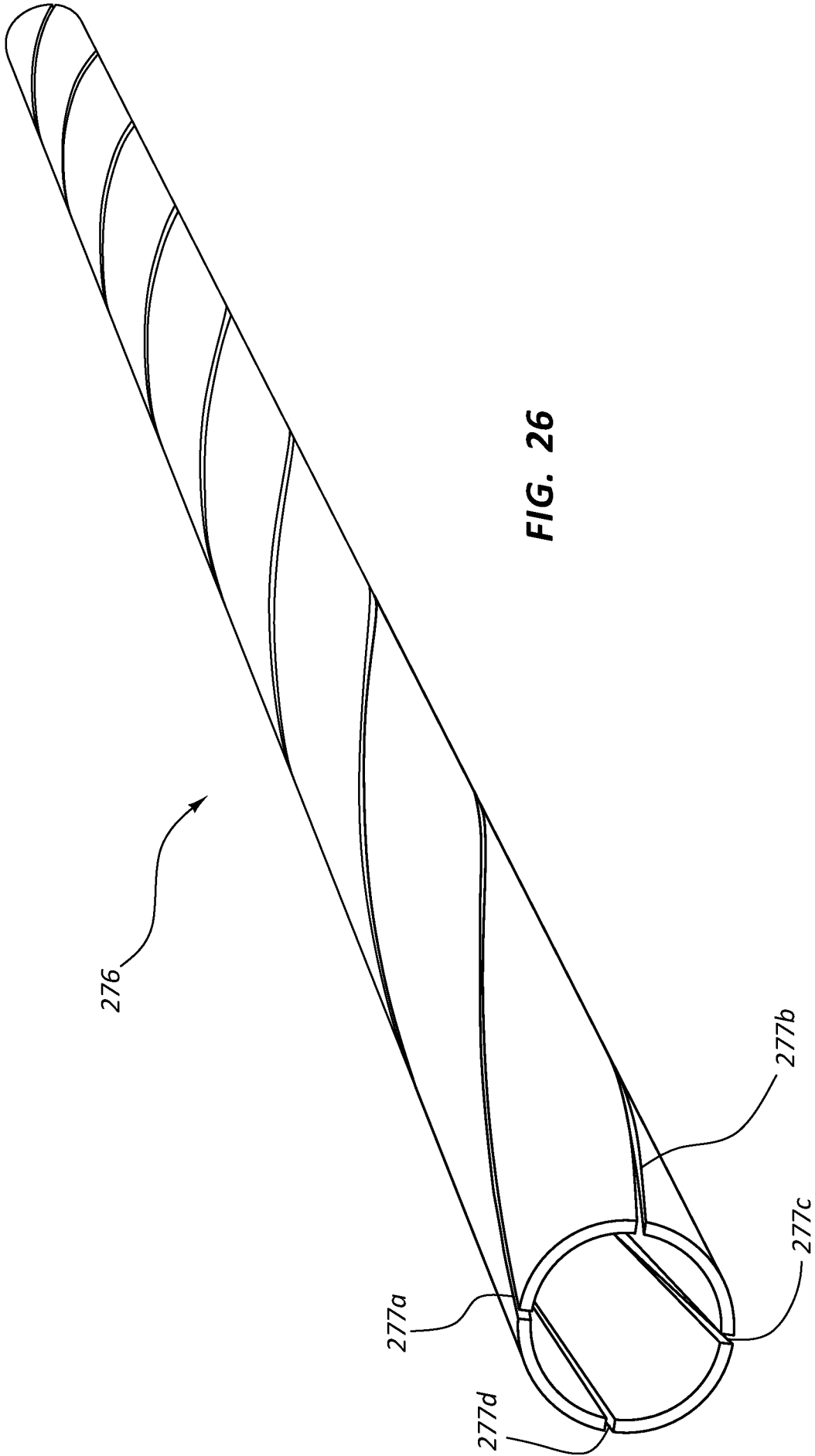
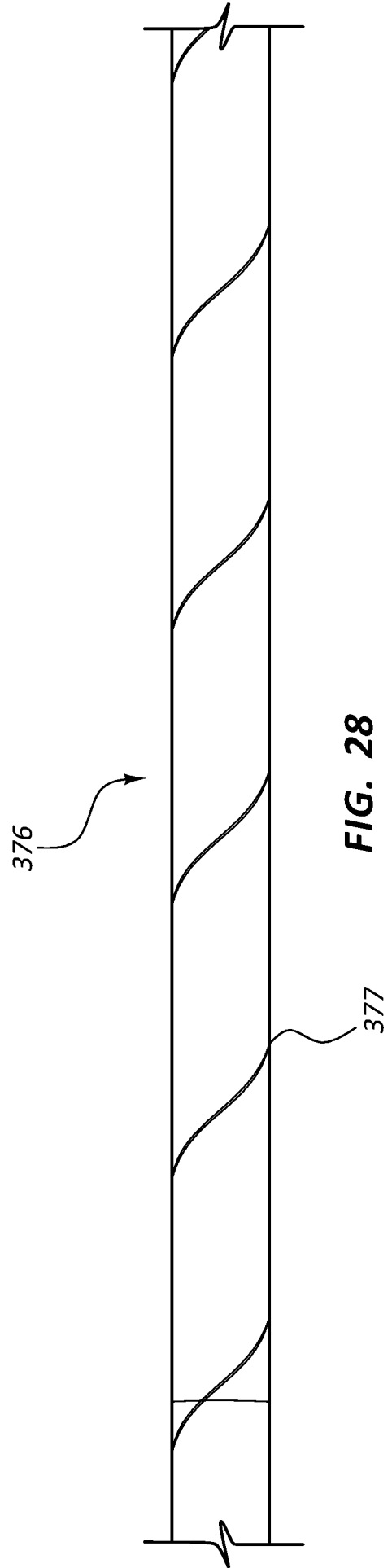
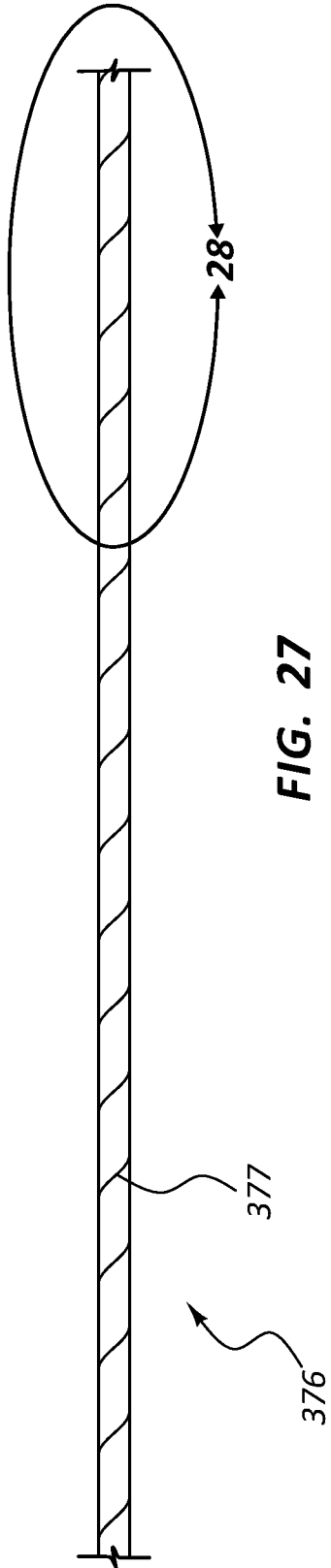


FIG. 25





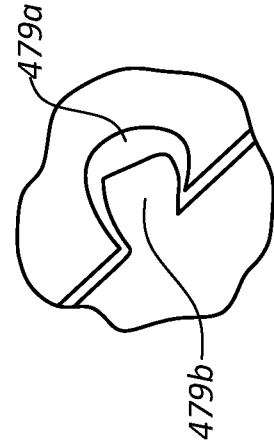
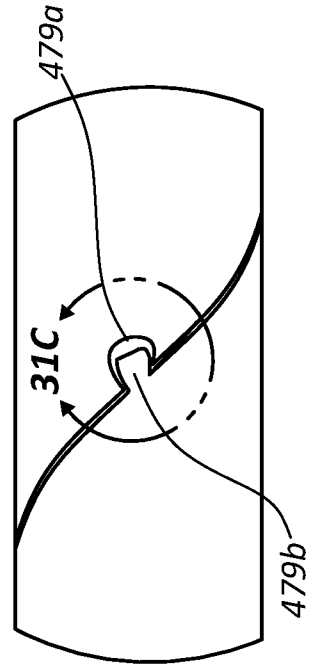
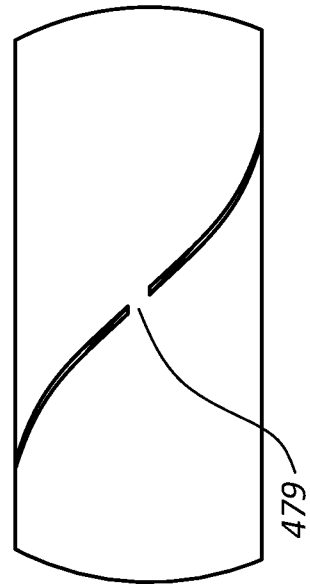
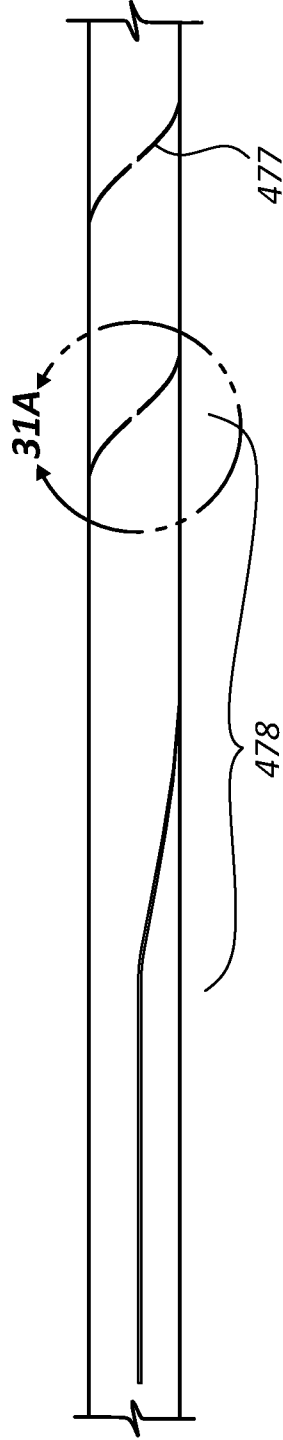
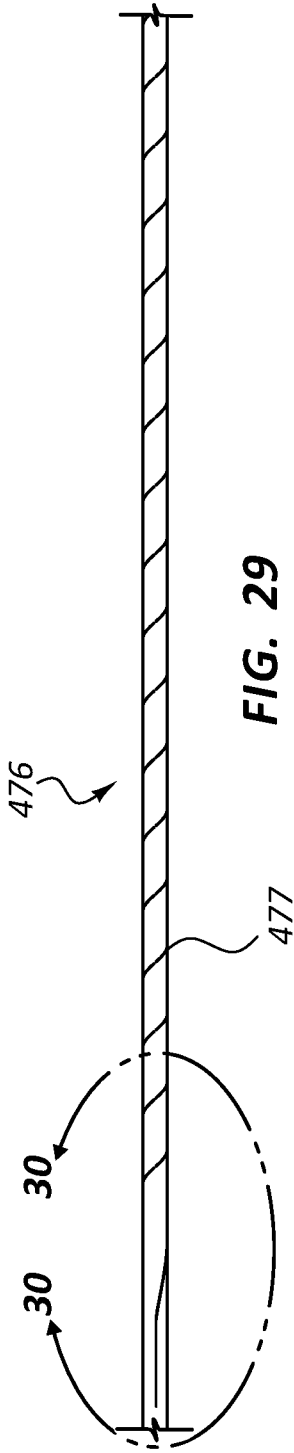




FIG. 32

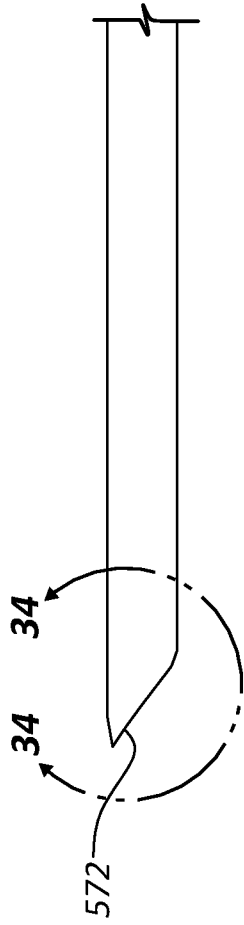


FIG. 33

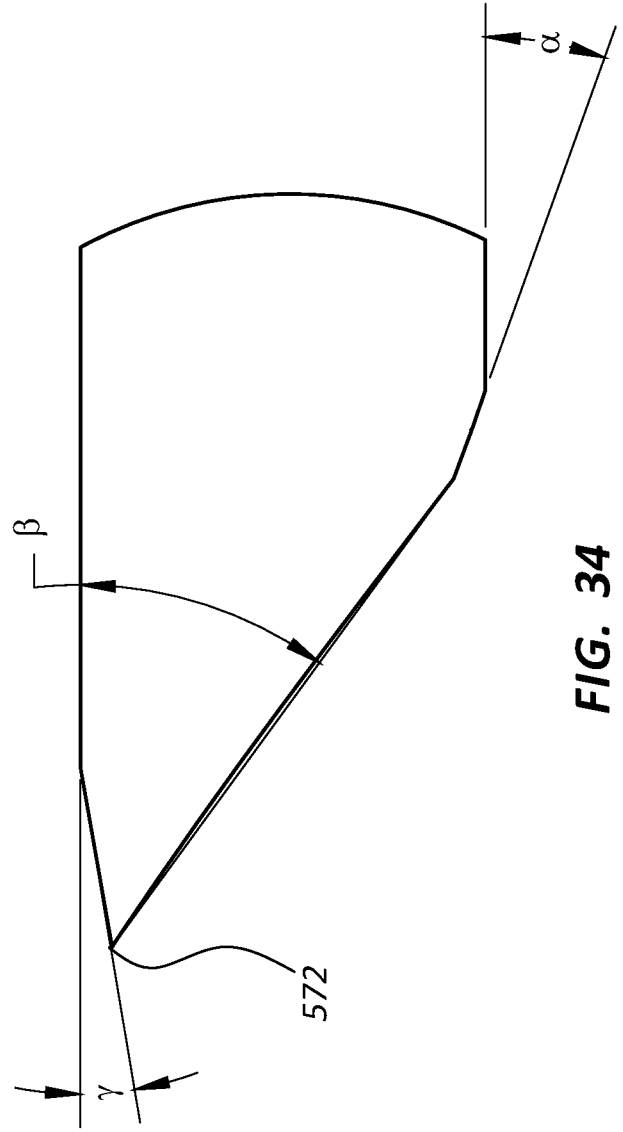
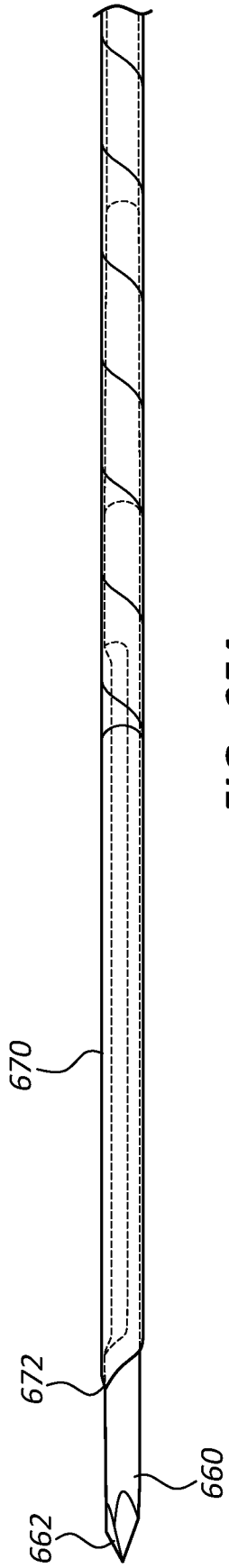
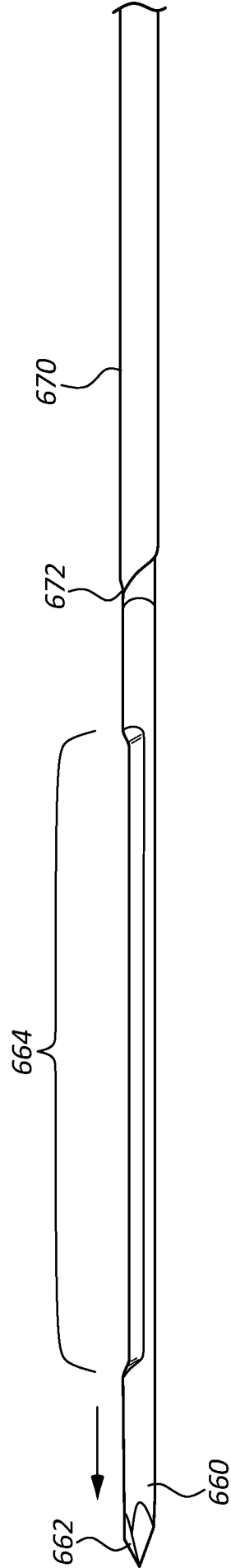


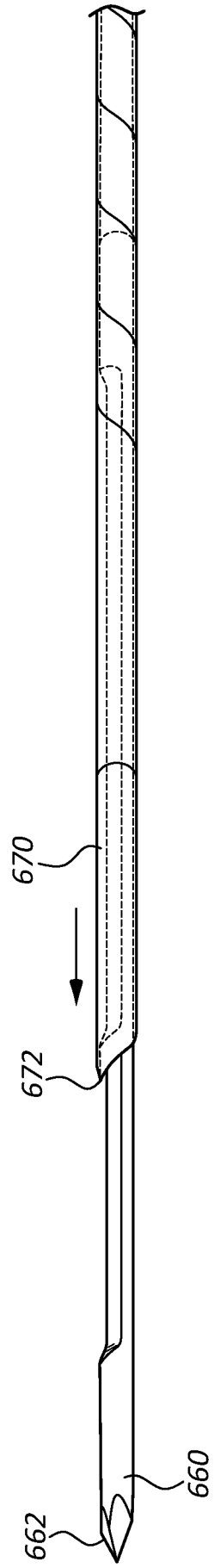
FIG. 34



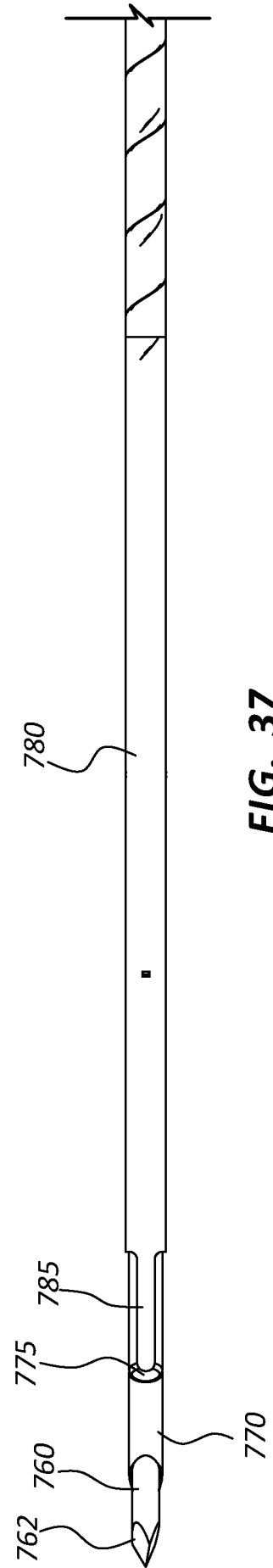
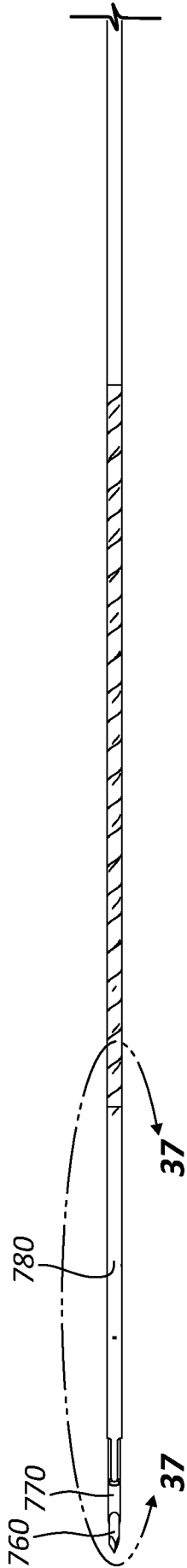
**FIG. 35A**

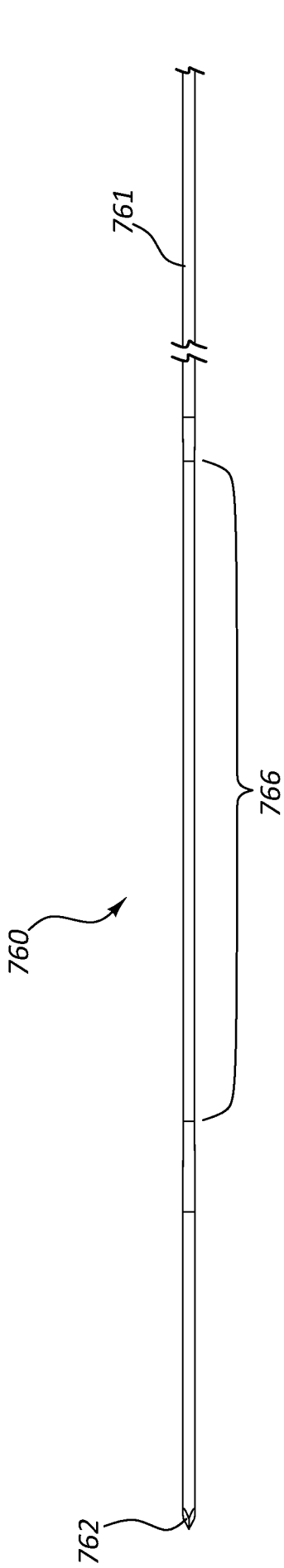


**FIG. 35B**

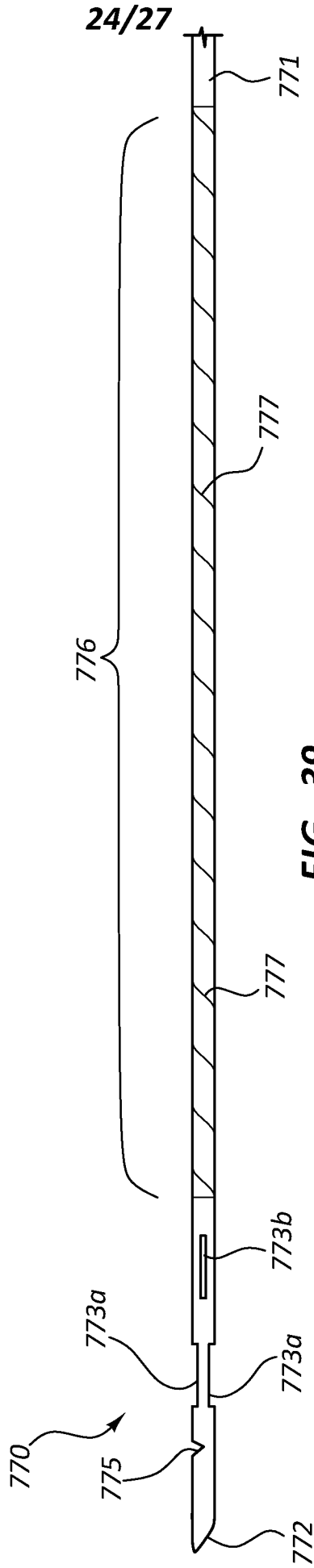


**FIG. 35C**

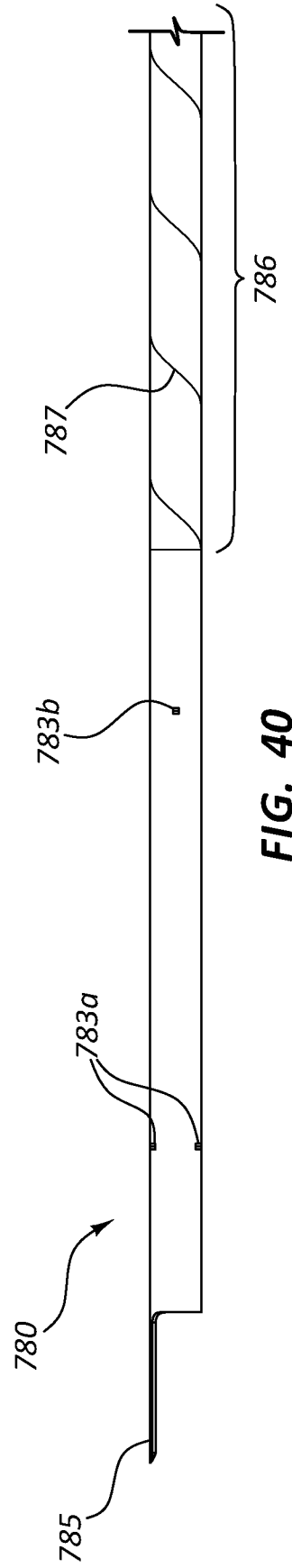




**FIG. 38**



**FIG. 39**



**FIG. 40**

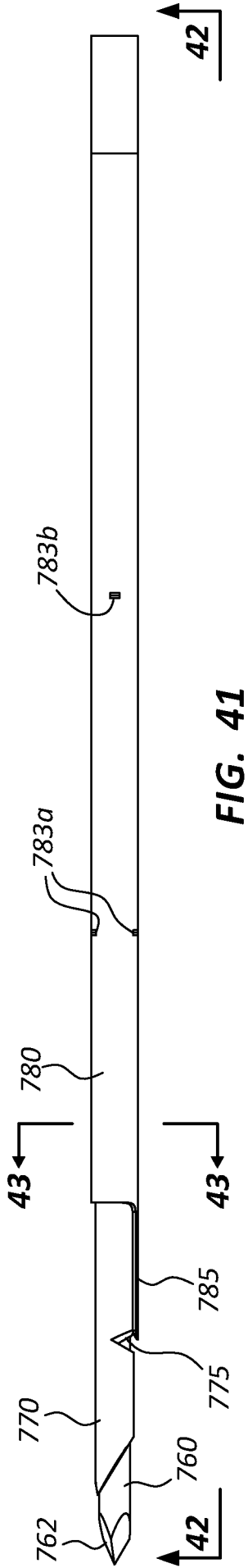


FIG. 41

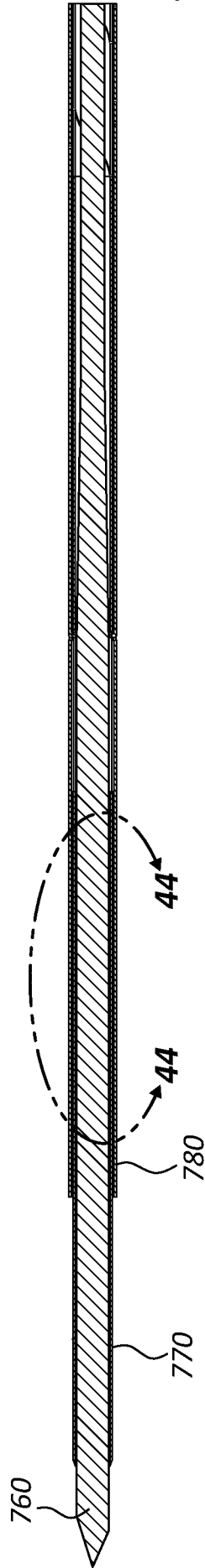


FIG. 42

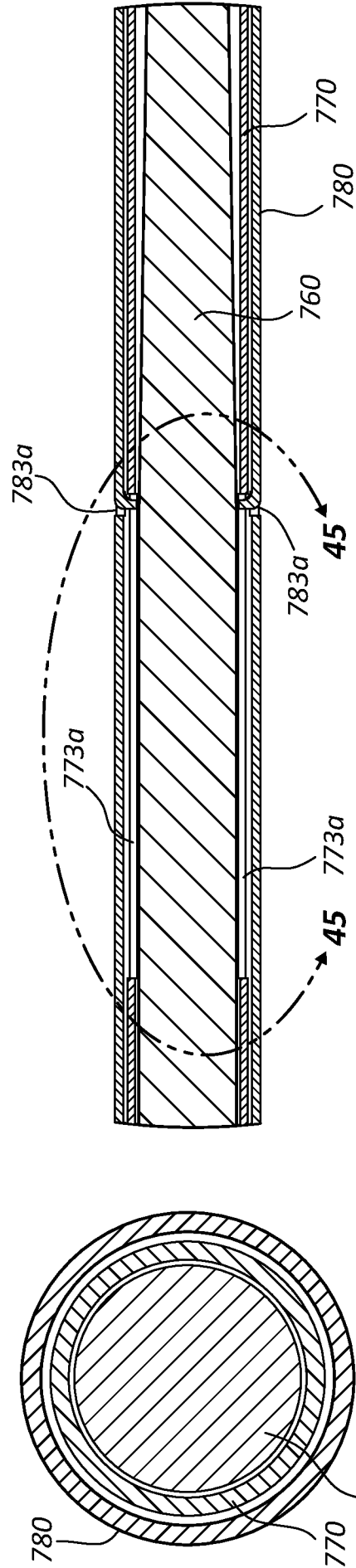


FIG. 43

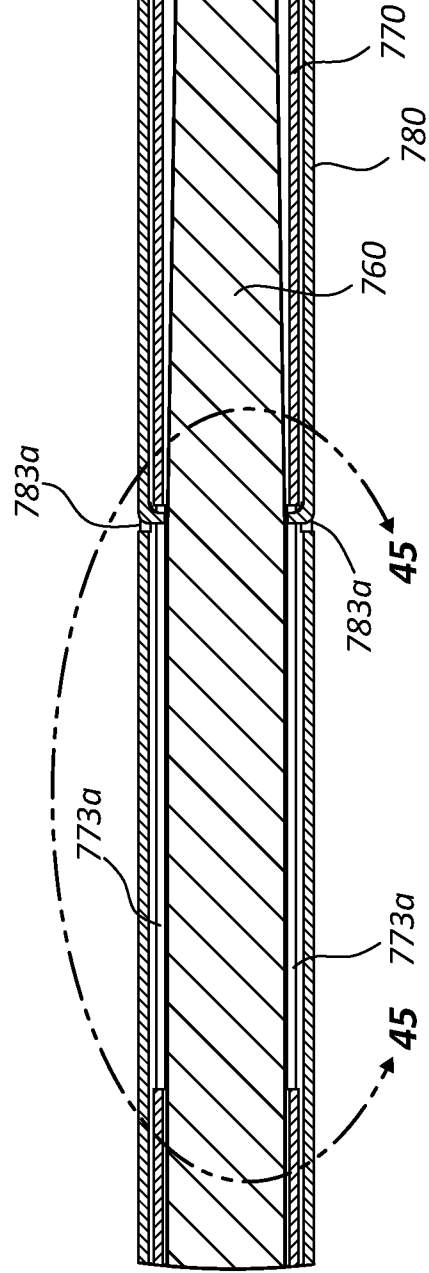


FIG. 44

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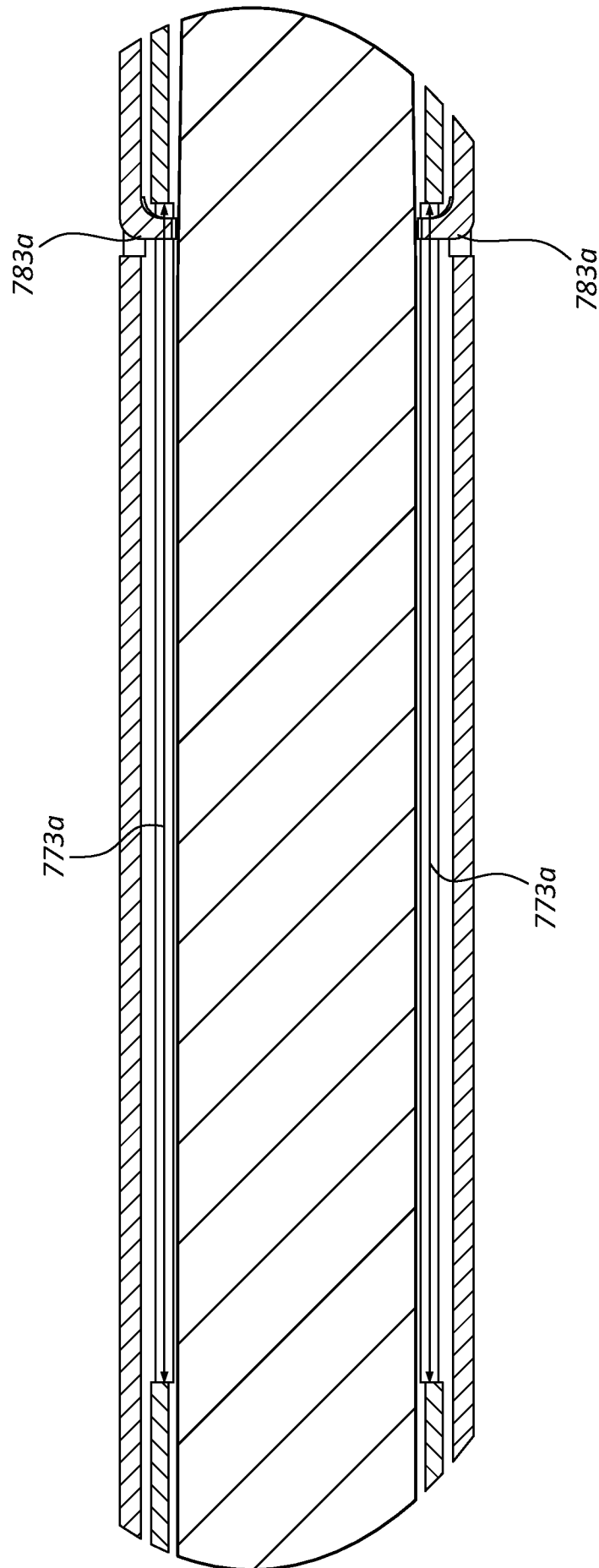


FIG. 45

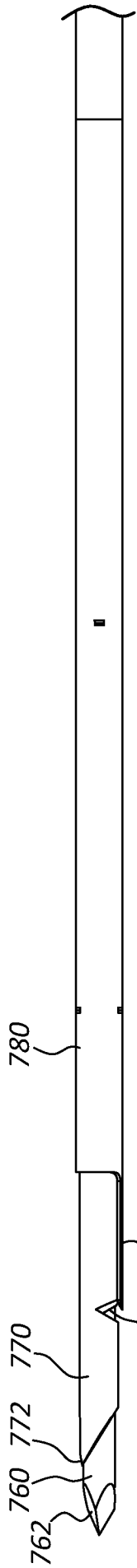


FIG. 46A

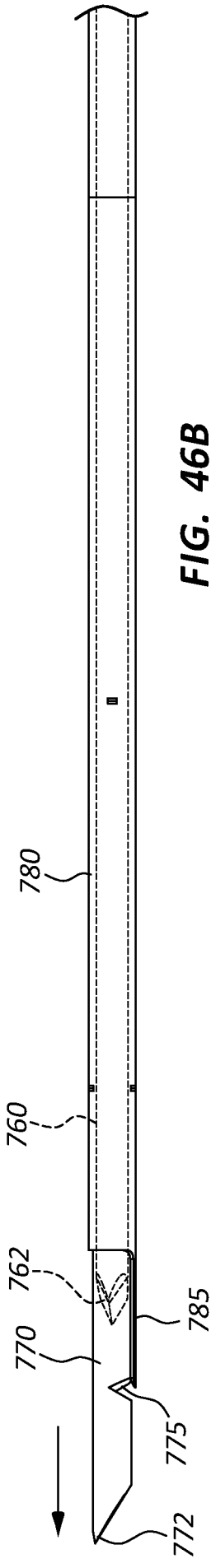


FIG. 46B

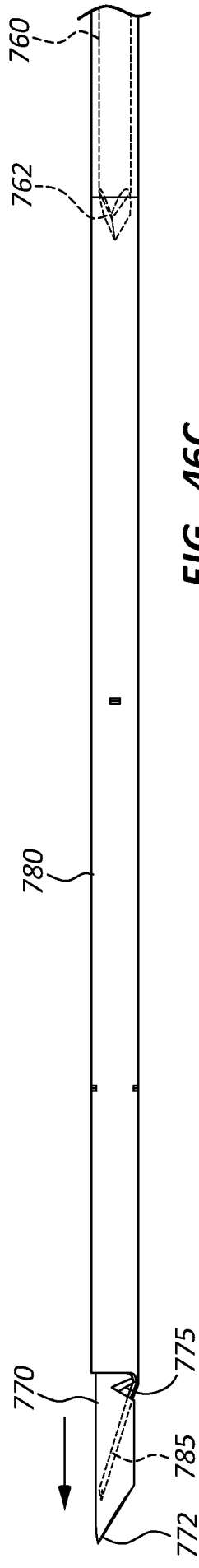


FIG. 46C

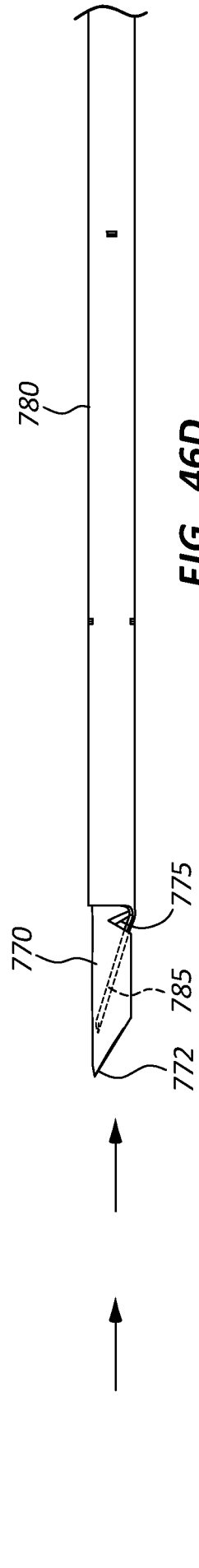


FIG. 46D

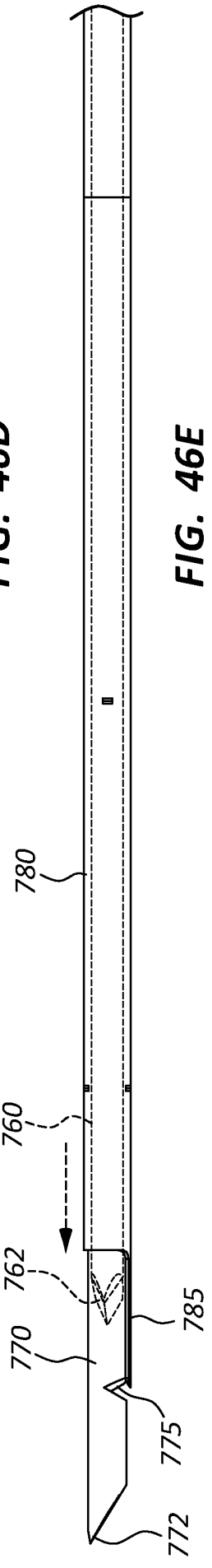


FIG. 46E

## INTERNATIONAL SEARCH REPORT

International application No.  
**PCT/US2013/076418****A. CLASSIFICATION OF SUBJECT MATTER****A61B 10/02(2006.01)i, A61B 17/34(2006.01)i, A61M 25/01(2006.01)i**

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

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

A61B 10/02; A61B 10/00; A61M 11/00; A61M 5/178; A61B 17/34; A61M 25/01

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

Korean utility models and applications for utility models  
Japanese utility models and applications for utility models

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

eKOMPASS(KIPO internal) &amp; keywords: biopsy, rotary, rotate, index, curved, flexible

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2012-0220894 A1 (MELSHEIMER, J. S.) 30 August 2012 See abstract; paragraphs [0024], [0028], [0030]; claims 1, 16; and figures 2-3, 9.	25,40-44,56
A		1-5,11-17,26-28 ,57-58
A	US 2009-0143698 A1 (JANSSENS, J. P.) 4 June 2009 See abstract; paragraphs [0013]-[0015], [0055], [0058]; claim 22; and figures 1, 3.	1-5,11-17,25-28 ,40-44,56-58
A	US 5800389 A (BURNEY, B. et al.) 1 September 1998 See abstract; column 4, lines 60-67; column 7, lines 19-23; claim 1; and figures 1-2.	1-5,11-17,25-28 ,40-44,56-58
A	US 2008-0300507 A1 (FIGUEREDO, S. et al.) 4 December 2008 See abstract; paragraph [0005]; claim 1; and figure 1.	1-5,11-17,25-28 ,40-44,56-58
A	US 2010-0130887 A1 (SELIS, J. E.) 27 May 2010 See abstract; paragraphs [0043]-[0044], [0065]; claim 17; and figures 10-15.	1-5,11-17,25-28 ,40-44,56-58
A	US 2004-0215103 A1 (MUELLER, JR., R. L. et al.) 28 October 2004 See abstract; paragraph [0028]; claims 1, 14; and figures 1-2.	1-5,11-17,25-28 ,40-44,56-58

 Further documents are listed in the continuation of Box C. See patent family annex.

\* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&amp;" document member of the same patent family

Date of the actual completion of the international search

02 April 2014 (02.04.2014)

Date of mailing of the international search report

**03 April 2014 (03.04.2014)**

Name and mailing address of the ISA/KR

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**INTERNATIONAL SEARCH REPORT**

International application No.

**PCT/US2013/076418****Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: 18-24,32-39,59-66  
because they relate to subject matter not required to be searched by this Authority, namely:  
Claims 18-24,32-39, and 59-66 pertain to methods for treatment of the human body and thus relate to a subject-matter which this International Searching Authority is not required, under Article 17(2)(a)(i) of the PCT and Rule 39.1(iv) of the Regulations under the PCT, to search.
2.  Claims Nos.: 23,46-47,49-50,54,64,66  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:  
Claims 23,46-47,49-50,54,64, and 66 are unclear because they refer to multiple dependent claims which do not comply with PCT Rule 6.4(a).
3.  Claims Nos.: 6-10,21-22,24,29-31,45,48,51-53,55,62-63,65  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2.  As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of any additional fees.
3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

- Remark on Protest**
- The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- No protest accompanied the payment of additional search fees.

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

**PCT/US2013/076418**

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2012-0220894 A1	30/08/2012	EP 2501297 A1 WO 2011-062736 A1	26/09/2012 26/05/2011
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