



(11)

EP 3 013 251 B1

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention
of the grant of the patent:
07.02.2018 Bulletin 2018/06

(51) Int Cl.:
A61B 17/02 ^(2006.01) **A61B 17/00** ^(2006.01)
A61B 17/34 ^(2006.01) **A61B 90/40** ^(2016.01)

(21) Application number: **14733617.6**

(86) International application number:
PCT/EP2014/063458

(22) Date of filing: **25.06.2014**

(87) International publication number:
WO 2014/207077 (31.12.2014 Gazette 2014/53)

(54) **AN INFLATABLE PNEUMOPERITONEUM DEVICE**

AUFBLASBARE PNEUMOPERITONEUMVORRICHTUNG

DISPOSITIF DE PNEUMOPÉRITOINE GONFLABLE

(84) Designated Contracting States:
**AL AT BE BG CH CY CZ DE DK EE ES FI FR GB
GR HR HU IE IS IT LI LT LU LV MC MK MT NL NO
PL PT RO RS SE SI SK SM TR**

(30) Priority: **26.06.2013 US 201361839461 P**
17.02.2014 US 201461940681 P
21.03.2014 US 201461968770 P
11.04.2014 US 201414251362

(43) Date of publication of application:
04.05.2016 Bulletin 2016/18

(73) Proprietor: **Atropos Limited**
Bray, County Wicklow (IE)

(72) Inventors:
• **BONADIO, Frank**
Bray
County Wicklow (IE)

• **HALPIN, Lucy Dolores**
Dublin 14 (IE)
• **WILLIAMS, Stephen**
Blackrock
County Dublin (IE)

(74) Representative: **O'Brien, John Augustine et al**
John A. O'Brien & Associates
Unit 4, The Courtyard Business Centre
Orchard Lane
Blackrock, County Dublin, A94 NV07 (IE)

(56) References cited:
WO-A1-95/09666 WO-A1-2010/099541
WO-A2-2011/090866 US-A- 5 618 296
US-A1- 2011 087 235 US-B2- 7 163 510

Note: Within nine months of the publication of the mention of the grant of the European patent in the European Patent Bulletin, any person may give notice to the European Patent Office of opposition to that patent, in accordance with the Implementing Regulations. Notice of opposition shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

Description

Introduction

[0001] This disclosure relates to a pneumoperitoneum device. Also described are methods involving the use of the device. WO 2010/099541 discloses tissue bags comprising a mouth portion having a first characteristic dimension and a body portion having a second characteristic dimension being greater than the first characteristic dimension. Such configurations can improve retention of tissue within tissue containment bags, thereby reducing the likelihood of undesirable contact between diseased and healthy tissue. WO2013/093030A (published June 27, 2013) describes an inflatable artificial pneumoperitoneum bag having an opening to receive tissue and a retainer for opening the bag. US5,618,296 describes a tissue morcellator system.

[0002] WO95/09666A describes an expandable and collapsible isolation bag.

[0003] WO2011/090866A describes a tissue retrieval device including an introducer and a tissue retrieval bag.

[0004] US2011/0087235A describes a tissue retrieval bag and an introducer for the bag.

Statements of Invention

[0005] The invention is defined in the appended claims. According to the disclosure there is provided an artificial pneumoperitoneum device (e.g., a bag) for receiving tissue, tissue isolation, and/or extraction in a laparoscopic procedure.

[0006] In one aspect the disclosure provides an apparatus for use in during laparoscopic surgery comprising an inflatable bag having a tissue-receiving opening at a proximal end thereof and a cuff at the proximal opening, the cuff having a closed configuration for delivery and retrieval of the bag and an open configuration to receive tissue, the cuff being biased into the open configuration.

[0007] In one aspect the disclosure provides an inflatable artificial pneumoperitoneum bag for receiving tissue during laparoscopic surgery, the bag having a tissue-receiving opening at a proximal end thereof and a cuff at the proximal opening, the cuff having a closed configuration for delivery and retrieval of the bag and an open configuration to receive tissue, the cuff being biased into the open configuration, the bag having a main body which extends from the cuff and the main body of the bag is more flexible than the cuff, and a biasing element to bias the cuff into the open configuration.

[0008] In one embodiment the bag has a main body which extends from the cuff and the main body of the bag is more flexible than the cuff. The cuff may be of a different material than that of the main body of the bag. The cuff may be of a stiffer material than that of the main body of the bag.

[0009] In one embodiment the main body of the bag includes a neck region extending from the cuff. The neck

region may be more flexible than the cuff.

[0010] In some embodiments the cuff extends for a distance of between 2 and 20 cm, between 2 and 10 cm, or about 5 cm.

[0011] In one embodiment the bag comprises a biasing element (e.g., a ring) that extends at least partially around the opening. The biasing element (e.g., ring) is preferably flexible to facilitate entry through an incision and/or an instrument access port. That is, the biasing element (e.g., ring) may bias the cuff into the open configuration.

[0012] In one case the biasing element (e.g., ring) comprises a loop (e.g., an O-ring) extending around the cuff. The loop may be of a shape memory material such as Nitinol. The loop may comprise a single loop element which is open or closed.

[0013] In one case the loop comprises a plurality of loop parts.

[0014] In some embodiments at least one of the loop parts is movable relative to another of the loop parts. At least one of the loop parts may be movable circumferentially relative to another ring element.

[0015] In one case the apparatus comprises a retainer for opening the bag.

[0016] The retainer may comprise at least one ring element which extends at least partially around the opening. The ring element may be flexible to facilitate entry through an incision and/or an instrument access port. In one case the ring element comprises an O-ring.

[0017] In one embodiment the retainer comprises ring parts.

[0018] There may be two separate ring parts.

[0019] In one embodiment the apparatus (e.g., bag) comprises a tether for each of the ring parts.

[0020] In some embodiments, the bag may comprise a tether extending proximally from the cuff. The cuff may comprise a tab for use in grasping the bag. In one case the bag comprises tabs arranged on opposite sides of the cuff.

[0021] The or each tab may comprise an indicator to indicate a preferred orientation of the bag.

[0022] In one embodiment the bag is configured to be foldable into a reduced size for insertion through an incision or an opening.

[0023] In one aspect the retainer has an insertion configuration and an expanded deployed configuration. The retainer may be biased into the deployed configuration.

[0024] The disclosure also provides an apparatus comprising an introducer sheath or pouch for containing the bag in an insertion configuration. The introducer sheath or pouch may be at least partially insertable through an opening and/or an incision and/or an access port.

[0025] In one case the apparatus comprises an activator for delivering the bag from the pouch, on insertion. The activator may comprise a tab. In one case the activator comprises a plunger.

[0026] In one embodiment the apparatus comprises a user tether attached to the bag.

[0027] In one case the bag comprises a neck region. The neck region may be adjacent to the retainer.

[0028] In one embodiment the bag itself comprises a port. The port may be an exit port and/or an entrance port. The bag may comprise a plurality of ports.

[0029] In some embodiments the port comprises a valve. The valve may comprise a choke valve or a cuff valve. In one case the valve comprises an elastomeric material such as a gel.

[0030] In some embodiments the apparatus comprises a proximal tether and a distal tether.

[0031] The distal tether may be movable relative to the proximal tether.

[0032] In one case the proximal tether comprises a loop through which the distal tether is movable.

[0033] There may be a lock to restrict movement of the distal tether. In one case the lock is provided by or on the proximal and/or the distal tether. The lock may comprise a projection on the distal tether which is engagable by the proximal tether.

[0034] In one embodiment the apparatus further comprises an access port to which the bag is mounted or mountable. The access port may comprise a retractor having a distal anchoring element for location within a wound interior, a proximal member for location externally of a wound opening and a retractor member extending proximally from the distal anchoring element to retract laterally the sides of an incision.

[0035] The bag may be mountable to the proximal member of the retractor.

[0036] In one embodiment the apparatus further comprises a cap for closing the proximal side of the retractor. The cap may comprise an access device for an instrument or a surgeon's hand/arm. The access device may be mountable to the proximal member of the retractor.

[0037] The disclosure also provides apparatus for use in laparoscopic surgery comprising a bag of the disclosure and a retractor. The apparatus may further comprise an access port.

[0038] The disclosure also provides a viscera retainer comprising an apparatus of the invention.

[0039] Also described is a method for performing a laparoscopic procedure comprising the steps of:-

inserting a bag according to the invention through an opening;
inflating the bag;
delivering tissue into the bag before or after inflating the bag; and
carrying out a procedure on the tissue located in the inflated bag.

[0040] In one embodiment the opening is an opening into a body cavity.

[0041] The opening may be provided, at least in part, by an incision.

[0042] In one embodiment the method comprises providing a trocar and inserting the bag through the trocar.

[0043] The method may comprise providing a retractor in the opening and inserting the bag through the retracted opening.

[0044] The tissue may be delivered into the bag before inflating the bag.

[0045] The method may comprise the step, either before or after delivery of the tissue into the bag, of mounting the bag to the retractor.

[0046] In one embodiment the method comprises passing an instrument into the inflated bag to carry out a procedure.

[0047] In one case the method comprises inserting a trocar into the bag.

[0048] The method may comprise the steps of providing an access port in the bag and passing an instrument and/or tissue through the access port.

[0049] In one embodiment the method comprises sealing the access port prior to and/or subsequent to passage of an instrument and/or tissue through the access port.

[0050] The device of the disclosure comprises at least one instrument seal to effect a seal around at least one instrument extended through the device, the instrument seal being configured to be arranged in sealing relationship to a body of a patient. The device preferably has a distal anchoring member for location within a wound interior. The device preferably also has a retractor member extending proximally from the distal anchoring member to retract laterally the sides of a wound opening. Preferably the device comprises a first instrument seal to effect a seal around a first instrument extended through the device, and a second instrument seal to effect a seal around a second instrument extended through the device. By providing the two seal arrangement, this ensures that insertion or manipulation or removal of the second instrument does not adversely effect the seal around the first instrument. The device may comprise a third instrument seal to effect a seal around a third instrument extended through the device. The first instrument seal may be spaced apart from the second instrument seal. The first instrument seal may be formed separately from the second instrument seal. The first instrument seal may have a larger radial dimension than the second instrument seal. The instrument seal may be a valve. Alternatively, the seal is of a gelatinous elastomeric material.

[0051] In one case the device comprises a proximal member for location externally of a wound opening. The retractor member may extend at least between the distal anchoring member and the proximal member. The retractor member may extend in two layers between the distal anchoring member and the proximal member. A first end portion of the retractor member may be fixed to the proximal member. The retractor member may be movable relative to the distal anchoring member. A second end portion of the retractor member may be movable relative to the proximal member. The retractor member may extend distally from the proximal member to the distal anchoring member, may be looped around the distal anchoring member, and may extend proximally from the

distal anchoring member to the proximal member. The proximal member may comprise an inner part and an outer part. The retractor member may extend between the inner part and the outer part.

[0052] In another embodiment the instrument seal is spaced proximally of the proximal member. The device may comprise at least one connector member to connect the proximal member to the at least one instrument seal. The connector member facilitates a degree of lateral movement of the instrument while maintaining the seal. The connector member may comprise a sleeve. The connector member may be of a laterally flexible material. The connector member may be of a longitudinally rigid material. The connector member may be of a rubber-like material. The connector member may be of a longitudinally flexible material.

[0053] In another case the instrument seal is mounted to the connector member. The instrument seal may be releasably mounted to the connector member. The instrument seal may comprise a mounting part to mount the instrument seal to the connector member. The mounting part may be of a rigid material. The instrument seal may comprise a sealing part to effect a seal around an instrument extended through the device, the sealing part being overmoulded over at least part of the mounting part.

[0054] A method for performing a laparoscopic procedure may include inserting a bag through a patient opening, delivering tissue into a bag opening of the bag, withdrawing the bag opening back through the patient opening, sealing the bag, inflating the bag to create an artificial pneumoperitoneum that extends the abdomen, conforms to the peritoneal cavity, and provides additional working and viewing space; and carrying out a procedure on the tissue located in the inflated bag.

[0055] A method for performing a laparoscopic procedure includes inserting a bag through a patient opening and into a peritoneal cavity, delivering tissue into the bag in an insufflated peritoneal cavity, sealing the bag, inflating the bag to apply a retracting force to the materials outside the bag thereby enlarging the peritoneal cavity, carrying out a procedure on the tissue located in the inflated bag; and allowing the peritoneal cavity to uninsufflate so that the inflated bag is located in an uninsufflated peritoneal cavity.

[0056] A method for performing a laparoscopic procedure includes inserting a bag through a patient opening and into a peritoneal cavity, delivering tissue into the bag, sealing the bag, inflating the bag to retract surrounding structures and organs and urge the bag against the abdominal wall, piercing the bag by one or more trocars at a location where the bag is urged against the abdominal wall, carrying out a procedure on the tissue located in the inflated bag, and communicating the peritoneal cavity with the atmosphere prior to carrying out a procedure on the tissue located in the inflated bag.

[0057] A method for performing a laparoscopic procedure includes insufflating the peritoneal cavity to provide a working and viewing space, excising tissue within the

working and viewing space, inserting a bag into the working and viewing space, delivering the excised tissue into the bag, sealing the bag, inflating the bag to replace the working and viewing space of the insufflated peritoneal cavity with a working and viewing space within the bag, and carrying out a procedure on the tissue located in the inflated bag.

[0058] The methods for performing a laparoscopic procedure may further include one or more of the following features: the opening may be a naked incision in the abdominal wall; the opening may be one of an opening through a retractor device coupled to an incision, or an opening through a trocar coupled to an incision; the inserting of a bag through an opening may include inserting the bag through a valve; the extending of the abdomen includes the bag contacting both the anterior abdominal wall and the abdominal viscera; the opening may be an opening through the retractor device, and the sealing of the bag may include sealing the bag to a proximal end of the retractor device with a cap; ; the delivering of tissue into the bag may be performed in an insufflated peritoneal cavity; the carrying out of a procedure in the inflated bag may be performed in an uninsufflated peritoneal cavity; the inflating of the bag may include inflating the bag to retract surrounding structures and organs and urge the bag against the abdominal wall; piercing the bag with one or more trocars at a location where the bag may be urged against the abdominal wall; the carrying out of a procedure may include sealably inserting a morcellator into the bag and morcellating the tissue; and retrieving the tissue by pulling the bag out through the opening.

[0059] In yet another embodiment, an inflatable artificial pneumoperitoneum bag includes a length and a width, an artificial pneumoperitoneum bag neck portion having a first end and a second end, the first end forming a bag opening, an artificial pneumoperitoneum bag body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion, and a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition.

[0060] In yet another embodiment, an inflatable artificial pneumoperitoneum bag, including a first planar sheet portion and a second planar sheet portion, the first and second planar sheet portions having joined edges to form the pneumoperitoneum bag having a length and a width, a neck portion having a first end and a second end, the first end forming a bag opening, a body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion, and a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition.

[0061] The inflatable artificial pneumoperitoneum bag may further include one or more of the following features: a length of the bag is between 300 and 600 mm when

the bag is uninflated, and a maximum width of the bag is between 200 and 500 mm when the bag is uninflated; the neck portion has a diameter of between 100 and 220 mm when in the open condition; the neck portion has a length of between 100 and 300 mm when the bag is uninflated; a width-extending cross-section of the body portion includes an oval shape when the bag is inflated; the bag includes sterilized polyester polyurethane planar sheet portions having joined edges; the bag is symmetric about two planes that are normal to one another; the neck portion includes edges that extend parallel to a lengthwise axis of the bag; and the ring is formed of a shape memory material and is received through loops in the neck portion.

[0062] In another embodiment, a method includes creating an artificial pneumoperitoneum in a patient using an artificial pneumoperitoneum bag, the artificial pneumoperitoneum bag including a length and a width, a neck portion having a first end and a second end, the first end forming a bag opening, a body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion, and a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition.

[0063] In another embodiment, a method includes creating an artificial pneumoperitoneum in a patient using an artificial pneumoperitoneum bag, the artificial pneumoperitoneum bag including a first planar sheet portion and a second planar sheet portion, the first and second planar sheet portions having joined edges to form the pneumoperitoneum bag having a length and a width, a neck portion having a first end and a second end, the first end forming a bag opening, a body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion, and a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition, and the creating of the artificial pneumoperitoneum includes positioning the joined edges in alignment with the lateral walls of the abdomen.

[0064] The methods for performing a laparoscopic procedure may further include one or more of the following features: the length of the bag is between 300 and 600 mm when the bag is uninflated, and a maximum width of the bag is between 200 and 500 mm when the bag is uninflated; the neck portion has a diameter of between 100 and 220 mm when in the open condition; the neck portion has a length of between 100 and 300 mm when the bag is uninflated; a width-extending cross-section of the body portion includes an oval shape when the bag is inflated; the bag is formed of sterilized polyester polyurethane planar sheet portions having joined edges; the bag is symmetric about two planes that are normal to one another; the neck portion includes edges that extend parallel to a lengthwise axis of the bag; the ring is formed of a shape memory material and is received through loops

in the neck portion.

[0065] According to the invention there is provided an inflatable artificial pneumoperitoneum bag, comprising:

- 5 a length and a width;
- an artificial pneumoperitoneum bag neck portion having a first end and a second end, the first end forming a bag opening;
- 10 an artificial pneumoperitoneum bag body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion; and
- 15 a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition.

The length of the bag is between 300 and 600 mm when the bag is uninflated, and a maximum width of the bag is between 200 and 500 mm when the bag is uninflated.

- 20 The neck portion has a diameter of between 100 and 220 mm when in the open condition.

[0066] The neck portion has a length of between 100 and 300 mm when the bag is uninflated.

- 25 **[0067]** In one embodiment a width-extending cross-section of the body portion includes an oval shape when the bag is inflated.

[0068] In one case the bag includes sterilized polyester polyurethane planar sheet portions having joined edges.

- 30 **[0069]** In one embodiment the bag is symmetric about two planes that are normal to one another. The neck portion includes edges that extend parallel to a lengthwise axis of the bag.

- 35 **[0070]** In one embodiment the ring is formed of a shape memory material and is received through loops in the neck portion.

[0071] The disclosure also provides an inflatable artificial pneumoperitoneum bag, comprising:

- 40 a first planar sheet portion and a second planar sheet portion, the first and second planar sheet portions having joined edges to form the pneumoperitoneum bag having
- a length and a width;
- 45 a neck portion having a first end and a second end, the first end forming a bag opening;
- a body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion; and
- 50 a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition.

- 55 **[0072]** In one embodiment the length of the bag is between 300 and 600 mm when the bag is uninflated, and a maximum width of the bag is between 200 and 500 mm when the bag is uninflated.

[0073] In one case the neck portion has a diameter of between 100 and 220 mm when in the open condition.

[0074] In one embodiment the neck portion has a length of between 100 and 300 mm when the bag is uninflated.

[0075] In one case a width-extending cross-section of the body portion includes an oval shape when the bag is inflated.

[0076] In one embodiment the bag includes sterilized polyester polyurethane.

[0077] In one case the bag is symmetric about two planes that are normal to one another.

[0078] In one embodiment the neck portion includes edges that extend parallel to a lengthwise axis of the bag.

[0079] In one case the ring is formed of a shape memory material and is received through loops in the neck portion.

[0080] Also described is a method, comprising:

creating an artificial pneumoperitoneum in a patient using an artificial pneumoperitoneum bag, the artificial pneumoperitoneum bag including

a length and a width,
a neck portion having a first end and a second end, the first end forming a bag opening;
a body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion; and
a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition.

[0081] In one embodiment the length of the bag is between 300 and 600 mm when the bag is uninflated, and a maximum width of the bag is between 200 and 500 mm when the bag is uninflated. In one case the neck portion may have a diameter of between 100 and 220 mm when in the open condition. The neck portion may have a length of between 100 and 300 mm when the bag is uninflated. In one case a width-extending cross-section of the body portion includes an oval shape when the bag is inflated. The bag may include sterilized polyester polyurethane planar sheet portions having joined edges. The bag may be symmetric about two planes that are normal to one another. The neck portion may include edges that extend parallel to a lengthwise axis of the bag. The ring may be formed of a shape memory material and is received through loops in the neck portion.

[0082] Also described is a method, comprising:

creating an artificial pneumoperitoneum in a patient using an artificial pneumoperitoneum bag, the artificial pneumoperitoneum bag including

a first planar sheet portion and a second planar sheet portion, the first and second planar sheet portions having joined edges to form the pneu-

moperitoneum bag having

a length and a width;

a neck portion having a first end and a second end, the first end forming a bag opening;

a body portion forming a closed cavity in fluid communication with the second end of the neck portion, the body portion having a length and width greater than a length and width of the neck portion; and

a ring located at the bag opening, the ring configured to bias the bag opening toward an open condition; and

the creating of the artificial pneumoperitoneum includes positioning the joined edges in alignment with the lateral walls of the abdomen.

[0083] In one embodiment the length of the bag is between 300 and 600 mm when the bag is uninflated, and a maximum width of the bag is between 200 and 500 mm when the bag is uninflated. The bag may be symmetric about two planes that are normal to one another.

[0084] Also described is a method for performing a laparoscopic procedure, comprising:

inserting a bag through a patient opening;
delivering tissue into a bag opening of the bag;
withdrawing the bag opening back through the patient opening;
sealing the bag;
inflating the bag to create an artificial pneumoperitoneum that extends the abdomen and provides additional working and viewing space; and
carrying out a procedure on the tissue located in the inflated bag.

[0085] In one embodiment the patient opening is a naked incision in the abdominal wall. The patient opening may be one of an opening through a retractor device coupled to an incision, or an opening through a trocar coupled to an incision. In one embodiment the inserting of a bag through a patient opening includes inserting the bag through a valve. In one embodiment the extending of the abdomen includes the bag contacting both the anterior abdominal wall and the abdominal viscera. In one case the patient opening is an opening through the retractor device, and the sealing of the bag includes sealing the bag to a proximal end of the retractor device with a cap. In one embodiment the delivering of tissue into the bag is performed in an insufflated peritoneal cavity. The carrying out of a procedure in the inflated bag may be performed in an uninsufflated peritoneal cavity. The inflating of the bag may create an artificial pneumoperitoneum that extends the abdomen includes urging the bag against the abdominal wall, and the method further includes piercing the bag with one or more trocars at a location where the bag is urged against the abdominal wall. In one embodiment the carrying out of a procedure

includes sealably inserting a morcellator into the bag and morcellating the tissue, and the method further includes retrieving the tissue by pulling the bag out through the patient opening.

[0086] Also described is a method for performing a laparoscopic procedure, comprising:

inserting a bag through a patient opening and into a peritoneal cavity;
 delivering tissue into the bag in an insufflated peritoneal cavity;
 sealing the bag;
 inflating the bag to apply a retracting force to the materials outside the bag thereby enlarging the peritoneal cavity;
 carrying out a procedure on the tissue located in the inflated bag; and
 allowing the peritoneal cavity to uninsufflate so that the inflated bag is located in an uninsufflated peritoneal cavity.

[0087] In one embodiment the application of the retracting force to the materials outside the bag includes the bag contacting both the anterior abdominal wall and the abdominal viscera. The patient opening may be one of an opening through a retractor device coupled to an incision, or an opening through a trocar coupled to an incision. The inserting of a bag through an opening may include inserting the bag through a valve. The patient opening may be an opening through the retractor device, and the sealing of the bag includes sealing the bag to a proximal end of the retractor device with a cap. The method may further include pulling an open end of the bag out through the patient opening prior to inflating the bag. The allowing of the peritoneal cavity to uninsufflate may be performed before the carrying out of a procedure on the tissue located in the inflated bag. The inflating of the bag may be performed in an uninsufflated peritoneal cavity. The inflating of the bag to apply a retracting force may include urging the bag against the abdominal wall, the method further including piercing the bag with one or more trocars at a location where the bag is urged against the abdominal wall. The carrying out of a procedure may include sealably inserting a morcellator into the bag and morcellating the tissue, and the method further includes retrieving the tissue by pulling the bag out through the opening.

[0088] Also described is a method for performing a laparoscopic procedure, comprising:

inserting a bag through a patient opening and into a peritoneal cavity;
 delivering tissue into the bag;
 sealing the bag;
 inflating the bag to retract surrounding structures and organs and urge the bag against the abdominal wall;
 piercing the bag by one or more trocars at a location where the bag is urged against the abdominal wall;

carrying out a procedure on the tissue located in the inflated bag; and
 communicating the peritoneal cavity with the atmosphere prior to carrying out a procedure on the tissue located in the inflated bag.

[0089] In one embodiment the patient opening is one of an opening through a retractor device coupled to an incision, or an opening through a trocar coupled to an incision. In one case the patient opening is an opening through the retractor device, and the sealing of the bag includes sealing the bag to a proximal end of the retractor device with a cap. The method may further include pulling an open end of the bag out through the patient opening prior to inflating the bag, and wherein the retraction of the surrounding structures and organs includes the bag contacting both the anterior abdominal wall and the abdominal viscera. The delivering of tissue into the bag may be performed in an uninsufflated cavity. The inflating of the bag may be performed in an uninsufflated peritoneal cavity.

[0090] Also described is a method for performing a laparoscopic procedure, comprising:

insufflating the peritoneal cavity to provide a working and viewing space;
 excising tissue within the working and viewing space;
 inserting a bag into the working and viewing space;
 delivering the excised tissue into the bag;
 sealing the bag;
 inflating the bag to replace the working and viewing space of the insufflated peritoneal cavity with a working and viewing space within the bag; and
 carrying out a procedure on the tissue located in the inflated bag.

[0091] In one embodiment the method further includes pulling an open end of the bag out through a patient opening prior to inflating the bag, and wherein the inflating of the bag includes the bag contacting both the anterior abdominal wall and the abdominal viscera. The method may further include uninsufflating the peritoneal cavity after delivering the excised tissue into the bag. The inflating of the bag may be performed in an uninsufflated peritoneal cavity.

Brief Description of the Drawings

[0092] The invention will be more clearly understood from the following description of some embodiments thereof, given by way of example only, with reference to the accompanying drawings, in which: -

Fig. 1 is an isometric view of a pneumoperitoneum device;

Fig. 2 is another isometric view of the device of Fig. 1;

Figs. 3 to 5 are views of another pneumoperitoneum device;

Figs. 6 to 18 are diagrams illustrating the use of the device of Figs. 1 to 5;

Figs. 19 to 20 are diagrams illustrating another use of the device of Figs. 1 to 5;

Figs. 23 to 24 are diagrams illustrating a further use of the device of Figs. 1 to 5;

Figs. 25 to 30 are diagrams illustrating various ways in which a device may be introduced;

Figs. 31 to 35 are diagrams illustrating the device, in use;

Fig. 36 is a diagram of another device;

Figs. 37 to 45 are diagrams illustrating the device of Fig. 36, in use;

Fig. 46 is a diagram of another device;

Figs. 47 to 54 are diagrams illustrating the device of Fig. 46, in use;

Fig. 55 is a diagram of a further device;

Figs. 56 to 63 are diagrams illustrating the device of Fig. 55, in use;

Fig. 64 is a diagram of another device;

Figs. 65 to 74 are diagrams illustrating the device of Fig. 64, in use;

Fig. 75 is a diagram of a device for use as a visceral retainer;

Figs. 76 to 78 are diagrams illustrating the device of Fig. 75, in use;

Fig. 79 is a diagram of the device of Figs. 75 to 78 with an associated grommet;

Figs. 80 and 81 are diagrams illustrating the device of Figs. 75 to 78 in use;

Figs. 82 to 87 are isometric views of alternative grommets;

Fig. 88 is an isometric view of another device with a multi-lumen access port removed;

Fig. 89 is an isometric, partially cut-away view of the device of Fig. 88 with an access port in position for

use;

Fig. 90 is an isometric view of a device;

Fig. 91 is another view of the device of Fig. 90 with an access port in position for use;

Figs. 92(a) and 92(b) are isometric views of single lumen access ports for use with the devices;

Fig. 93 is an isometric view of a pneumoperitoneum device with an exit port from the device;

Fig. 94 is an isometric view of another device similar to Fig. 93 with a number of exit ports;

Fig. 95 is an isometric view illustrating devices of the types of Figs. 93 and 94, in use;

Figs. 96 to 98 are views of various seals that may be used in association with the device;

Figs. 99 to 110 illustrate one method of use of devices;

Figs. 111 to 113 are views illustrating a locking detail of the device of Figs. 99 to 110;

Figs. 114 to 117 are views of a device in use in the colon;

Fig. 118 is an isometric view of another device;

Fig. 119 is an isometric view of a further device;

Fig. 120 is an isometric view of a pneumoperitoneum device according to the disclosure loaded in an introducer;

Fig. 121 is an isometric view of a bag according to the invention;

Fig. 122 is a cross sectional view on the line A-A in Fig. 121;

Figs. 123 to 126 are isometric views illustrating the insertion of the bag through an opening;

Figs. 127 and 128 are isometric views illustrating the delivery of tissue into the bag;

Figs. 129 and 130 are isometric views illustrating the closing of the bag;

Figs. 131 to 133 are isometric views illustrating the presentation of the bag to the tissue opening, subsequent inflation of the bag, and procedures such as morcellation being carried out on tissue in the

inflated bag;

Figs. 134 to 138 are diagrams illustrating various alternative arrangements of trocar(s) and instrument(s) used in performing procedures on tissue in the inflated bag; 5

Figs. 139 to 143 are various views of a biasing loop comprising loop sections which are movable relative to one another; 10

Figs. 144 to 149 are various views of another biasing loop;

Figs. 150 to 153 are various views of a further biasing loop; 15

Figs. 154 to 156 are various views of another biasing loop; 20

Figs. 157 to 160 are diagrams illustrating another bag device of the disclosure;

Figs. 161 to 164 illustrate another bag device of the disclosure in use; 25

Fig. 165 is an isometric view illustrating closing of a bag device;

Figs. 166 to 168 are isometric views of further bag devices according to the disclosure with a retaining ring in place in a cuff of the bags; 30

Figs. 169(a)-169(c) and 170 illustrate the mounting of a ring to the cuff of the bag of Fig. 168; 35

Fig. 171 shows a protector cap for a bag device;

Figs. 172 to 176 illustrate an introducer for use in deploying and manipulating a bag device; 40

Figs. 177 to 180 are views of further introducers with various pressure dissipating features;

Figs. 181 to 184 illustrate another bag device having a generally ovoid shape; 45

Figs. 185 and 186 illustrate a retaining ring of the bag;

Figs. 187(a) to 187(f) are views illustrating the loading of a bag device into an introducer; 50

Fig. 188 is an illustration of a bag device with a reinforced section; 55

Fig. 189 shows an introducer with a valve;

Figs. 190(a) to 190(c) illustrate the use of a bag de-

vice in trans-vaginal procedures;

Fig. 191 is a view of a part of a bag with a threaded bung or trocar in place;

Figs. 192(a) and 192(b) illustrate the closing of a chimney of a bag, prior to deflation;

Figs. 193(a) to 193(c) illustrate the use of the bag devices of the invention with various access devices; and

Fig. 194 is an isometric view of an artificial pneumoperitoneum device according to the invention;

Fig. 195 is a view of an introducer sheath for the device of Fig. 194;

Fig. 196 is a view of a plunger for use with the introducer sheath;

Figs. 197 and 198 show the introducer sheath and plunger, in use;

Fig. 199 is a view of a bag part of the device of Fig. 194;

Fig. 200 is a cross sectional view of a detail of the bag device of Fig. 199;

Figs. 201a and 201b are views of an open end of the bag device;

Fig. 202 illustrates the threading of a wire through a collar of the bag device;

Fig. 203 is a cross sectional view of a detail of Fig. 202;

Fig. 204 illustrates the joint between ends of the wire;

Fig. 205 shows the attachment of a tether to the joint of Fig. 204;

Figs. 206 to 209 illustrate the folding of the device for loading into an introducer;

Fig. 210 illustrates the introducer and plunger after delivery of the device;

Figs. 211 to 218 illustrate the artificial pneumoperitoneum device at different stages of use;

Figs. 219 and 200 illustrate a seal between the introducer and the plunger;

Fig. 221 and 222(a) to 222(c) illustrate the mounting of a solid ring to the collar of a device of the invention;

Fig. 223 illustrates another method of attachment of a ring to a device of the invention;

Figs. 224 to 253 illustrate the artificial pneumoperitoneum device of the invention at various stages during use;

Figs. 254 to 256 illustrate the device of the invention used trans-vaginally;

Fig. 257 illustrates the closing of the device within the pneumoperitoneum;

Fig. 258 illustrates the use of the device in a procedure which involves the uses of an additional port;

Figs. 259 and 260 illustrate an artificial pneumoperitoneum device of the invention being used in different ways;

Figs. 261 to 266 illustrate further artificial pneumoperitoneum devices of the invention;

Fig. 267 illustrates the use of the device with a threaded trocar;

Figs. 268 to 276 illustrate various bungs for use with the trocar;

Figs. 277 to 280 illustrate an inflatable bung and its use; and

Figs. 281 to 285 illustrate a trocar cover.

Detailed Description

[0093] The disclosure provides an artificial pneumoperitoneum device for tissue isolation and/or extraction in a laparoscopic procedure

[0094] The device is used to safely reduce and remove resected tissue from within the abdomen via small laparoscopic incisions. The bag creates an artificial pneumoperitoneum containing the specimen and eliminating the dissemination of tissue and cellular fluids within the peritoneal cavity. The device facilitates effective and safe isolation of tissue/organs within an artificial pneumoperitoneum for improved surgical procedures and subsequent safe tissue extraction.

[0095] A tissue bag is inserted within the peritoneal cavity through an incision in the abdominal wall or vagina.

[0096] In one case the bag with one or more openings is placed within the abdomen. Excised tissue is placed within the opening of a deflated bag. One or more openings of the bag are withdrawn outside the abdomen and the bag is inflated. Instruments including laparoscopic visualization are placed within the inflated bag that remains within the peritoneal cavity. Visualisation tools may also be provided external of the bag. The tissue retained

within the bag is morcellated/crushed/reduced and removed. The bag is deflated and removed with residual tissue/blood/fluids inside. A major advantage is that the tissue to be removed is retained in the bag which prevents potentially harmful material such as cancerous cells from being released in the body cavity.

[0097] Referring to the drawings, and initially to Figs. 1 and 2 thereof, there is illustrated an apparatus for use in laparoscopic surgery comprising a bag 1 having an opening 2 to receive tissue and a ring element 3 extending around the opening 2. The bag is inflatable.

[0098] Referring to Figs. 3 to 5, there is illustrated another bag device according to the invention which is similar to the bag device of Figs. 1 and 2 and like parts are assigned the same reference numerals. In this case the bag 1 has a necked region 5 to reduce the amount of material near the ring 3. This facilitates attachment of the bag 1 to an external element.

[0099] The bag device 1 is suitable for use during laparoscopic surgery to facilitate procedures on tissue in an insufflated cavity while maintaining pneumoperitoneum.

[0100] The bag device 1 may be mounted to a retractor. One such retractor comprises a distal anchoring ring 10, a retractor member such as a sleeve 11, and a proximal ring assembly 12.

[0101] One such retractor is described in US 2005-0090717 A. The distal anchoring ring 10 is located within a wound interior, in use. In this case the distal anchoring ring 10 is provided in the form of an O-ring. The proximal ring assembly 12 is located externally of a wound opening, in use. The retractor member 11 may be employed to retract laterally the sides of a wound opening. In this case, the retractor member is provided in the form of a sleeve.

[0102] The proximal end of the retractor 11 is closable by a cap which in this case comprises an instrument access device 30 which may have a number of instrument ports 31 to effect a seal around an instrument extended through the device 30. The instrument access device 30 may be releasably mountable to the proximal ring assembly 12. At least some of the instrument ports may include a stalk 32 which is laterally flexible and longitudinally rigid.

[0103] Fig. 6 illustrates an instrument 51 being introduced under vision provided by a camera 52 through an instrument access port 31.

[0104] Fig. 7 shows an organ or tissue such as an uterus 35 which has been severed from its retaining structures.

[0105] Fig. 8 illustrates the bag device 1 being inserted into the abdominal cavity at the beginning of a procedure or as and when required. The bag 1 is inserted in a small flattened state for ease of insertion through a small opening such as an incision. As shown in Fig. 8, the bag device 1 is inserted through an opening that is open to the atmosphere, and thus the peritoneal cavity is uninsufflated in this embodiment. The bag may also be introduced through a valve without the need to remove the access

cap 30. One such arrangement is illustrated in Fig. 10, and this allows the bag device to be inserted into an insufflated peritoneal cavity.

[0106] When the bag 1 is inserted in the insufflated or uninsufflated peritoneal cavity, it is opened up (Fig. 9). An organ is then readily manipulated for insertion into the bag 1 as illustrated in Fig. 11. The rigidity of the O-ring 3 keeps the bag open to facilitate insertion of an organ.

[0107] Fig. 12 shows the organ located in the bag 1 and the O-ring 3 being grasped to facilitate manipulation of the bag towards the opening. As shown in Fig. 12, removal of access cap 30 provides an uninsufflated peritoneal cavity prior to inflating the bag device 1. The uninsufflated nature of the peritoneal cavity helps the bag device 1 expand as shown in Fig. 16, and apply a retracting force to the materials outside the bag thereby creating additional space (Fig. 76). It is understood that the bag device 1 could be inflated while the peritoneal cavity is insufflated if there is one or more openings to the peritoneal cavity that allow the insufflation gas to escape to the cavity. One such opening could be provided through an open trocar device or trocar incision opening that was used to carry out the tissue excising explained above. The O-ring 3 is pulled out through the opening (Fig. 13) and the bag 1 is mounted to the proximal ring assembly 12 and the cap 30 is mounted to the proximal assembly 12 (Fig. 14). Fig. 15 illustrates the device in place with an organ enclosed within the bag 1.

[0108] The bag 1 is then inflated through an insufflation port 40. The inflation of the bag has the additional benefit of applying a retracting force to the materials outside the bag 1 thereby creating additional space (Fig. 16). Thus, the artificial pneumoperitoneum created in the uninsufflated peritoneal cavity, and within the inflated bag device 1, serves to extend the abdomen and provide additional working and viewing space within the bag device 1.

[0109] Fig. 17 shows an organ being worked on in the inflated bag 1. In this case the organ is morcellated. The material is all retained safely within the bag and is not released into the cavity which could cause major difficulties.

[0110] When the organ has been morcellated, the bag is readily removed through the original opening. All waste, blood, tissue and the like are safely removed and sealed within the bag 1.

[0111] Fig. 19 shows the bag device being inserted through a standard naked incision. Once the specimen has been inserted into the bag 1 (Fig. 20) the ring 3 is pulled back out through the incision (Fig. 21) and a trocar 60 is inserted to create a gas seal (Fig. 22). It may also be possible to insert the bag device 1 directly through a trocar.

[0112] In all cases there may be one or more access trocars used in addition to the primary port. Thus, the invention includes procedures which involve two or more incision laparoscopy.

[0113] For example, Figs. 23 and 24 show one ar-

angement in which an additional trocar 70 is inserted. In some cases, the additional trocar 70 may be extended through the bag whilst maintaining a seal.

[0114] A bag 1 is illustrated which has some depth which is preferred. However, a flat material can be used to form a holder in situ and the edges of the material pulled out through an incision and sealed outside, for example by an access device 30.

[0115] Also described is a method of inserting a large bag into the abdominal cavity to allow the insertion of a specimen into the bag. The bag is then sealed and inflated and procedure carried out within the bag.

[0116] Figs. 25 to 30 show various ways a bag 79 may be introduced into the abdomen.

[0117] In Fig. 25 the device may be inserted directly through an incision 80.

[0118] Referring to Fig. 26, the device may be inserted through a trocar 81.

[0119] In Fig. 27 a device may be inserted through a base retractor 82.

[0120] Referring to Fig. 28 a device may be inserted through a low profile port 83.

[0121] As shown in Fig. 29 the device may be inserted through a Multi-port device 84. The multiport device may, for example, be of the type described in US 8,187,178 or US 20110071389A.

[0122] Referring to Fig. 30 the device may be inserted through the base 85 of a multi-port device.

[0123] As illustrated in Fig. 31, once the bag 79 has been inserted the specimen is placed inside.

[0124] Fig. 32 the lip of the bag 79 is pulled out through the opening.

[0125] Fig. 33 the bag 79 is sealed by re-inserting the trocar 60, replacing the cap or inserting a morcellator 78.

If necessary an extra seal may be applied to the neck of the bag 79.

[0126] As shown in, Fig. 34, once the bag 79 is inflated additional trocars may be inserted into the abdomen as normal and pierced through the bag 79. Fig. 34 also shows a morcellator 78 inserted through a trocar 60.

[0127] Fig. 35 shows the morcellator 78 being inserted without the need for a trocar. A sealing ring 77 may be applied around the shaft of the morcellator 78 if necessary to hold back gas.

[0128] Referring to Figs. 36 to 45 a method of inserting a large bag 100 into an abdominal cavity which may automatically open to allow the insertion of a specimen is illustrated. The bag 100 is foldable and has a top opening 105 which may be biased into the open configuration by retaining elements which in this case comprise semi-circular ring parts 101, 102 which have attached tether elements 104, 103 respectively. A pouch 110 is used to house the bag 100 in a folded/retracted configuration. The pouch 110 has a grasping tab 111 and a pull string 115.

[0129] Fig. 36 illustrates the main components of the automatically opening bag device.

[0130] Fig. 37 illustrates a folded bag 100 inside the

pouch 110. In Fig. 38 the pouch 110 is inserted into the abdominal cavity with the aid of the grasping tab 111. When the pouch is inside, the distal pull tether 104 is pulled forward and the bag 100 is released. A rear pull string 115 is pulled in the opposite direction to aid release.

[0131] Referring to Fig. 40, it will be noted that as the distal end of the bag 100 is pulled forward the rear of the bag 100 is pulled in the opposite direction as it is attached to the pouch 110 with the connecting tether 103. This action opens the mouth of the bag 100 sufficient to ease the inserting of specimens.

[0132] Fig. 41 shows specimens being placed on top of the bag opening 105.

[0133] Referring to Fig. 42, by pulling the distal pull tether 104 back and over the specimen, the bag 100 begins to unroll and the specimen travels deeper inside the bag 100. Referring to Fig. 43, as the front and back retaining elements 101, 102 of the bag opening are pulled outwards, the specimen travels further into the bag 100.

[0134] Fig. 44 shows the rim of the bag being opened up and the incision being cleared of excess bag material.

[0135] Referring to Fig. 45, the opening is re-sealed by attaching a cap, by inserting a trocar, or by inserting a morcellator through the opening.

[0136] Referring to Figs. 46 to 54 there is illustrated another device according to the invention. The device is similar to that of Fig. 36 to 45 and like parts are assigned the same reference numerals. In this case a bag 120 is housed within a cartridge 121 for delivery and automatically opens when it exits the cartridge 121 on insertion into the abdominal cavity. In this case the ring part 102 remains attached to the cartridge 121. A tether 125 extends between the distal end of the cartridge 121 and the ring element 102. The ring element 101 has a tether element 126 which is grasped by an instrument 127 to pull the bag 120 from the cartridge 121.

[0137] Figs. 46 to 54 show the bag 120 housed in the cartridge 121 which can be inserted into a valve on an access port/trocar 130. The cartridge 121 remains in place during the procedure.

[0138] Referring to Fig. 46, the loaded cartridge 121 is placed through a valve on the port 130. Fig. 47 shows the distal pull tether 126 that is positioned so that it is easily grasped with an instrument 127. In Fig. 48 an instrument 127 is inserted and the pull tether 126 is grasped.

[0139] Referring to Fig. 49, as the instrument/grasper 127 is pushed forward the bag 120 is released from the cartridge 121. As shown in Fig. 50, once the bag is in far enough, the tether 125 which connects the back side of the bag 120 to the cartridge 121 begins to open the bag 120 up.

[0140] Referring to Fig. 51, when the mouth of the bag 120 is sufficiently open a specimen may be placed inside. When the distal pull tether 126 is pulled back as illustrated in Fig. 52 this forces the bag 120 to unroll and the specimen to travel deeper into the bag 120.

[0141] Referring to Fig. 53, the cap/trocar 130 is then

removed and the rim of the bag 120 is pulled out through the incision and mounted to the retractor 135. Fig. 54 shows the cap, trocar, or morcellator reconnected. The bag 120 is then inflated.

[0142] Referring to Figs 55 to 63, there is illustrated a removable cartridge 140 with a manually opened bag 141 for insertion through a single port 142. These drawings illustrate a method of inserting a large bag 141 which will be manually opened by the user when inserted into the abdominal cavity. Figs. 55 to 63 show a bag 141 housed in a cartridge 140 which plugs into a valve 142 on an access port/trocar 130. The bag 141 is ejected from cartridge 140 using a plunger 145 and the cartridge 140 is removed. As shown, for example, in Fig. 54, inflating the bag device 1 creates the artificial pneumoperitoneum and urges the bag against the abdominal wall. This arrangement allows one or more additional trocars 70 to extend through the abdominal wall and directly and sealingly pierce the inflated bag device 1 where the bag device is urged against the abdominal wall. See, for example, Figs. 24, 34, and 35. Further facilitating this sealing of the additional trocars 70 to the inflated bag device 1 is the fact that the inflated bag device 1 is applying a retracting force to the materials outside the bag device and is thus secured in place against the abdominal wall.

[0143] Referring to Fig. 55, the bag 141 is loaded into a cartridge 140 which is then inserted through a valve 142 on the port/trocar 130. When the cartridge 140 is in place of the plunger 145 is inserted through the proximal end of the cartridge 140 as illustrated in Fig 56. Pushing the plunger 145 down as illustrated in Fig 57 forces the bag 141 to eject into the abdominal cavity.

[0144] Referring to Fig. 58, when the bag 141 has been ejected, the cartridge 140 may be removed, leaving an activation tether 147 in place. An instrument 148 is inserted as illustrated in Fig 59 and the instrument 148 is used to grasp the distal pull tether 149 which is attached to the front band or ring part/element 101 on the bag 141.

[0145] Referring to Fig. 60, the specimen is then lifted into the open mouth of the bag 141. The surgeon can control the mouth of the bag 141 using the activation tether 147. When both the front and the back ring elements 101, 102 of the bag 141 are grasped as illustrated in Fig. 61, the bag 141 can be pulled towards the incision, forcing the specimen to travel deeper into the bag 141.

[0146] Fig. 62 shows the valve/trocar 130 being removed and the rim of the bag being pulled out through the incision. In Fig. 63, the rim of the bag 141 is opened up, and the valve/trocar 130 are replaced to seal the bag 141. The bag 141 is then inflated and the procedure carried out within.

[0147] Referring to Figs. 64 to 74 there is illustrated a removable cartridge 150 with a manually opened bag 151 (laparoscopic). These drawings show a method of inserting a large bag 151 which can be manually opened by the user when inserted into the abdominal cavity.

[0148] Referring to Fig. 64, the loaded cartridge 150 is inserted through a pre-made incision. When the cartridge

150 is in place a plunger 152 is inserted as illustrated in Fig. 65. The plunger 152 is pushed all the way down and the bag 151 is ejected as shown in Figs. 66 and 67.

[0149] Referring to Fig. 68 an instrument 155 is inserted through a trocar/access port 156 and is used to grasp a distal pull tether 157 which is attached to the front band on the bag. Once the distal tether 157 is held, pulling on the activation tether 158 opens the mouth of the bag 151 and forces the excess material to unroll. The specimen may now be lifted into the open mouth of the bag as shown in Fig. 69. The surgeon can control the mouth of the bag 151 to some degree with an activation tether 158.

[0150] Referring to Fig. 70, with the back end of the bag 151 grasped, the bag 151 can be pulled towards the incision, forcing the specimen to travel deeper into the bag 151. The valve/trocar is removed and the rim of the bag is pulled out through the incision as illustrated in Fig. 71. The rim of the bag 151 is opened up, and the morcellator is inserted to seal the bag as shown in Fig. 72. The bag is then inflated and the procedure carried out within.

[0151] As illustrated in Figs. 73 and 74, when the bag 151 is inflated trocars 159 can be pierced through to allow access for additional instruments 160.

[0152] Referring to Figs. 75 to 87 there is illustrated the use of a bag 170 as described above as a visceral retainer. The bag 170 is first inserted and positioned where required (Fig. 75). As the bag 170 is inflated, surrounding structures and organs (abdominal viscera) are retracted as shown in Fig. 76.

[0153] As shown in Figs. 77 and 78, it may be of benefit to fill, or partially fill the bag 170 with a liquid. These benefits may include: 1) The specimen floats to the top of the bag and therefore the risk of bag damage at the base may be reduced. 2) Liquid may reduce smoke build up in the bag. 3) Blood will be diluted and may therefore allow for enhanced visibility.

[0154] Referring to Fig. 79, when the bag 170 is in place and the neck has been pulled through the incision there is often a lot of excess material in the incision. A grommet 175 may be inserted through the bag/incision to keep excess material away from the incision as illustrated in Fig. 80. This will help prevent damage to the bag 170 and aid visibility and gas flow. With the grommet 175 in place instruments can be inserted with ease as shown in Fig. 81.

[0155] The grommet 175 may be used with multiport or single port access devices (Fig. 82).

[0156] In some cases the grommet 175 have an insufflation/desufflation line 176 built in (Fig. 83).

[0157] As illustrated in Fig. 84 the grommet may include a series of slits 177 which allow it to conform to various incision dimensions.

[0158] The grommet may include a valve system 178 as illustrated in Fig. 85.

[0159] An instrument locking mechanism 179 may also be included (Fig. 86).

[0160] In some cases, as illustrated in Fig. 87, the

grommet may have a series of lumens 180 to aid with ventilation/insufflation.

[0161] Referring to Figs. 88 and 89 there is illustrated a bag device 200 according to the disclosure. In this case, the bag 200 is shown in the inflated configuration within a body cavity such as the abdomen. A tissue sample 201 is contained within the bag. An incision is made in the abdomen 202 and the incision is retracted using a retractor 203 as described above. In this case the retractor 203 has an outer proximal ring 204 and a multilumen access port 205 is releasably mounted to the ring 204. The bag 200 extends through the retracted incision and terminates in a retainer ring 206.

[0162] Figs. 90 and 91 illustrate a bag device similar to that shown in Figs. 88 and 89 but in this case a single instrument lumen access port 211 is mountable to a proximal part of the retractor assembly. The access port 211 may have a cannula section that extends through the retractor or may be an access port 212 with a short proximal leg.

[0163] The bag device may itself have an access port to facilitate passage of instruments into and out of the bag and/or to facilitate passage of a tissue sample into the bag.

[0164] Referring to Fig. 93 the bag has a single access port 220. However, there may be a plurality of such access ports as illustrated in Fig. 94. Passage of a tissue sample 225 through an access port 220 is illustrated in Fig. 95. The access port 220 may be provided with any suitable valve such as a choke valve, for example, for example a drawstring 226 as illustrated in Fig. 96, a cuff valve 227 as illustrated in Fig. 97, or an elastomeric valve 228 as illustrated in Fig. 98. The valve 228 may be of any suitable plastics, rubber or gel material.

[0165] Referring to Figs. 99 to 110 there is illustrated various steps in methods involving the use of the bag devices of the disclosure. In the example illustrated the device is of the type described above. The methods involve the use of a bag device 250, a retractor 251, an external access port system 252 and is used to access tissue 253 such as a specimen or an organ through an opening 254 in the body, in this particular case in the abdomen 255. The bag device has a delivery configuration in which it is housed in a retracted condition in a cartridge 260. A plunger 261 is used to deliver the retracted bag device out of the cartridge 260. The bag device 250 has an opening which is biased into an open configuration by a retainer ring 265. The ring 265 may be of a shape memory material as described above. A proximal tether which in this case is in the form of a ring or loop 267 is provided on one side of the ring 265 and a distal tether 268 extends from the side of the ring 265 generally opposite to the proximal tether 267.

[0166] In Fig. 99 the bag device is placed in the delivery configuration in the cartridge or pouch 260. In this case the access port device 252 is in situ on top of the retractor 251 and the cartridge 260 is inserted through one lumen of the access port (Fig. 100). The plunger 261 is used to

push the bag device 250 out of the cartridge 260 (Fig. 101). In this particular case the bag device is not tethered to the user, on delivery. On delivery into the body cavity, such as the abdomen, the retractor ring is free to move to its normally expanded configuration in which it opens up the bag opening (Fig. 102). The bag is folded in the delivery configuration. Using various instruments 268 a clinician manipulates a tissue specimen, organ or the like and then delivers it into the bag 250 through the open mouth of the bag (Figs. 103, 104). Figs. 105 and 106 illustrate one particular way in which the bag containing the tissue is retrieved. A grasper type instrument 273 is led through the proximal tether loop 267 and is used to grasp the distal tether 268 (Fig. 105). The distal tether 268 is pulled through the proximal tether loop 267 which ensures that the clinician has control over the bag as it is moved up towards the body opening (Fig. 106). As the retaining ring 265 engages with the retractor 251 it retracts allowing it to be pulled up through the body opening (Fig. 107). The access port 252 is removed and the retaining ring 265 is again free to expand (Fig. 108).

[0167] The access port 252 is re-attached and the bag is inflated to increase the operative field. The tissue sample can then readily be worked on (Figs. 109, 110) without the risk of any potentially harmful material being released into the body cavity.

[0168] In some cases there may be a lock feature which prevents movement of one tether relative to the other in some directions. One such lock feature is illustrated in Figs. 111 to 113. The distal tether has a one-way step feature 280 which permits the distal tether to pass through the proximal loop tether but once it has passed through this reverse movement is prevented as illustrated in Fig. 113. This ensures even greater control on the movement of the retaining ring 265 to aid closing of the bag as the ring 265 is being withdrawn.

[0169] As discussed above, the devices of the invention may be used in any suitable body cavities. One such use is in the colon and one embodiment for this use is illustrated in Figs. 114 to 117. The device may be inserted as described above. Once in place and inflated a clinician can inspect the wall of the colon for any unusual features such as a growth. One such growth 280 is illustrated in Fig. 115. In this case, when a growth 280 is identified some or all of the growth 280 may be accessed by cutting a hole in the wall of the bag which remains in place by virtue of its engagement with the rest of the colon. Using various instruments, at least a portion of the growth 280 can be excised and removed through the bag. As in the other embodiments described a major advantage is that the tissue to be removed is retained in the bag which prevents potentially harmful material such as cancerous cells from being released in the body cavity.

[0170] Referring to Fig. 118 there is illustrated another bag device 400 of the disclosure. The bag device has a neck or collar region 401 between a retaining ring 402 and the main body of the bag. Because the retaining ring 402 is of smaller diameter than that of the bag it is more

easily inserted through an access port. As shown in Fig. 118, the bag device 400 includes joined planar sheet portions forming the neck region 401 and a body portion 403. Also as shown in Fig. 118, the body portion 403 forms a closed cavity in fluid communication with the neck region 401, and the body portion 403 has a length and width greater than a length and width of the neck region 401. As also shown in Fig. 118, the retaining ring 402 is configured to bias the bag opening toward an open condition.

[0171] Fig. 119 illustrates another bag device 410 and shows how the main body of the bag may be folded in the retracted delivery configuration.

[0172] Referring to Figs. 121 and 122, there is illustrated a bag 500 for use in laparoscopic surgery having an opening 502 to receive tissue and a cuff or collar 503 extending around the opening 502. The bag 500 may be inflatable.

[0173] The bag 500 comprises a main body 505 which extends from the cuff 503. The main body 505 is more flexible than the cuff 503 to assist in retaining the bag 500 open in the deployed configuration. The cuff 503 may be of a different material than that of the bag 500 or may comprise the same or a similar material which is thicker than that of the main body 505 of the bag 500. In one case the cuff 503 is of a plastics material and the main body 505 of the bag 500 is of a plastics material which is more flexible than that of the cuff 503. However, the cuff 503 is sufficiently flexible to allow closing of the opening 502. A joint 507 between the cuff 503 and the main body 505 of the bag may be reinforced in any suitable manner such as by using a double layer of the cuff material as illustrated in Fig. 22.

[0174] The cuff 503 has an axial extent which is important in maintaining the opening 502 in a fully open configuration. This has the major advantage that tissue can be more readily manoeuvred into the opening 502 by the surgeon performing a laparoscopic procedure through a small opening. When the bag 500 is in the open configuration the surgeon is able to concentrate on manipulation of the tissue/material to be inserted into the bag 500 without the added task of complex manipulation of the bag 500 at the same time as the material is being manipulated. Thus, the bag 500 greatly facilitates the laparoscopic surgical procedure.

[0175] The cuff 503 extends axially for a length which is sufficient to ensure that the opening 502 remains open to provide an axially extending delivery mouth into the main body 505 of the bag 500. For example the cuff 503 may extend for an axial length of from 2 to 20 cm, 2 to 10 cm, or about 5 cm. Alternative size dimensions for the cuff or collar 503, as well as sizes for other features of the bag, will be discussed in more detail below.

[0176] In the disclosure the cuff 503 is biased into the open configuration. In this case the bag 500 comprises a biasing element to bias the cuff 503 into the open configuration.

[0177] The biasing element may comprise a loop 510

extending around the cuff 503. The loop may be of a shape memory material such as Nitinol. The loop 510 extends round the cuff 503 in any suitable manner. It may, for example, be threaded through the cuff as illustrated or may extend through a table or track provided in or on the cuff 503.

[0178] The cuff 503 has tabs 511, 512 which may be used for more readily grasping the bag during a laparoscopic procedure.

[0179] A tether 515 extends from the cuff 503 and may be used to activate the opening and/or closing of the bag 500.

[0180] Referring to Fig. 120, the bag 500 is folded and housed in an introducer sheath or pouch 520 ready for deployment. Any suitable insertion tool may be used to deliver the bag 500 through an opening. The bag 500 may be deployed in any suitable manner such as by using a plunger 516 which a user activates to deliver the bag from the introducer 520.

[0181] The deployment of the bag 500 through an opening such as a retracted incision 525 is illustrated in Figs 123 to 126. As soon as the bag starts to exit the pouch the cuff 503 opens up and when fully deployed defines an axially extending open mouth. Figs. 127 and 128 illustrate tissue 530 being manipulated into the open mouth of the bag 500 using an instrument such as a grasper 531. When the tissue 530 is in the bag 500 the tether 515 is used to pull the cuff 503 which initially closes the opening and is then pulled out through the tissue opening 525. When the proximal end of the bag 500 has passed through the tissue opening the bag 500 may be inflated and a procedure carried out on the tissue isolated within the bag 500. This may be carried out in several different ways, depending on the procedure being performed. For example, the bag entrance may be closed with an access port 540 as illustrated in Fig. 133 which facilitates access of any suitable instruments to perform procedures such as morcellation of the tissue within the bag 500. Alternatively access is gained to the captured tissue without a requirement for an access port - see Figs. 134, 135 and 138. Alternatively a trocar 545 may be provided through which an instrument is passed (Fig. 136). There may be additional trocars 550 used (see Fig. 137, for example).

[0182] The collar/cuff 503 may be collapsible to allow tissue to be rolled into the bag opening 502 rather than being lifted into the bag opening 502.

[0183] The bag 500 may be mounted to a retractor. One such retractor comprises a distal anchoring ring, a retractor member such as a sleeve, and a proximal ring assembly. One such retractor is described in US 2005-0090717 A, the entire contents of which are incorporated herein by reference. The distal anchoring ring is located within a wound interior, in use. In this case the distal anchoring ring is provided in the form of an O-ring. The proximal ring assembly is located externally of a wound opening, in use. The retractor member may be employed to retract laterally the sides of a wound open-

ing. In one case the retractor member is provided in the form of a sleeve.

[0184] The proximal end of the retractor is closable by a cap which may comprise an instrument access device 540 which may have a number of instrument ports to effect a seal around an instrument extended through the device. The instrument access device may be releasably mountable to a proximal ring assembly of a retractor. At least some of the instrument ports may include a stalk which is laterally flexible and longitudinally rigid. One such instrument access port is described in US8,187,178 or US2011-0071389A.

[0185] Fig. 127 shows an organ or tissue such as an uterus which has been severed/amputated from its retaining structures.

[0186] Figs. 123 to 126 illustrate the bag device being inserted into the abdominal cavity at the beginning of a procedure or as and when required. The bag is inserted in a small flattened state for ease of insertion through a small opening such as an incision. The bag may also be introduced through a valve without the need to remove an access cap.

[0187] When the bag is inserted it is opened up. An organ 530 is then readily manipulated for insertion into the bag as illustrated in Fig. 128. The relative rigidity of the cuff 503 keeps the bag open to facilitate insertion of an organ.

[0188] Fig. 129 shows the organ located in the bag and the cuff 503 being grasped to facilitate manipulation of the bag towards the opening. The cuff 503 is pulled out through the opening. The bag may be mounted to a proximal ring assembly of a retractor and a cap may be mounted to the proximal assembly. Fig. 131 illustrates the device in place with an organ enclosed within the bag.

[0189] The bag is then inflated (Fig. 132) through an insufflation port. The inflation of the bag has the additional benefit of applying a retracting force to the materials outside the bag thereby creating additional space.

[0190] Fig. 133 shows an organ being worked on in the inflated bag. The organ may be morcellated. The material is all retained safely within the bag and is not released into the cavity which could cause major difficulties. The bag is retained externally, for example by clamping/connecting to a retractor.

[0191] When the organ has been morcellated the bag is readily removed through the original opening. All waste, blood, tissue and the like are safely removed and sealed within the bag.

[0192] The bag device may be inserted through a standard naked incision. Once the specimen has been inserted into the bag an opening such as a cuff 503 is pulled back out through the incision and a trocar may be inserted to create a gas seal. The bag device may also be inserted directly through a trocar as illustrated in Fig. 136.

[0193] In some cases there may be one or more access trocars used in addition to the primary port. Thus, the invention includes procedures which involve two or more

incision laparoscopy.

[0194] For example, Fig. 137 shows one arrangement in which an additional trocar is inserted. In some cases, the additional trocar may be extended through the bag whilst maintaining a seal.

[0195] The disclosure provides a method of inserting a large bag into the abdominal cavity to allow the insertion of a specimen into the bag. The bag is then sealed and inflated and procedure carried out within the bag.

[0196] As discussed above, the devices of the disclosure may be used in any suitable body cavities. One such use is in the colon. The device may be inserted as described above. Once in place and inflated a clinician can inspect the wall of the colon for any unusual features such as a growth.

[0197] In this case, when a growth is identified some or all of the growth may be accessed by cutting a hole in the wall of the bag which remains in place by virtue of its engagement with the rest of the colon. Using various instruments, at least a portion of the growth can be excised and removed through the bag. As in the other embodiments described a major advantage is that the tissue to be removed is retained in the bag which prevents potentially harmful material such as cancerous cells from being released in the body cavity.

[0198] Referring to Figs. 139 to 155 various embodiments of biasing loop elements which may be used in the bag device of the invention to assist in maintaining the bag open are illustrated. In these cases the loop comprises a number of loop parts which are movable relative to one another for loading deployment, and/or retrieval.

[0199] Referring to Figs. 139 to 143 in this case there are two joined loop parts 551a, 551b which are movable circumferentially through hoops 552. In some cases a single piece loop is sufficiently flexible to allow manipulation.

[0200] Referring to Figs. 144 and 149 in this case a loop 560 is configured in the manner of a noose with a leg 561 extending from the loop which may be pulled to reduce the diameter of the loop. There is a closed loop 562 on one end and a kink 563 on the other end which links into the loop 562 to facilitate reducing the diameter of the loop 560.

[0201] Referring to Figs. 150 to 153 in this case there may be closed loops 570, 571 at each end of the retaining loop 572. These may be used as tether attachment points.

[0202] Referring to Figs. 154 to 156 in this case the ends of a retaining loop 580 have bends 581, 582 and extend to define arms 583, 584 which facilitate manipulation of the loop for retrieval.

[0203] The device of the invention may include features to isolate the contents of the bag from the wound opening and hence protect from escape of material as could occur with seeding of cancer cells and the like.

[0204] Referring to Figs. 157 to 160 in this case the bag comprises a small opening 600 which remains closed under insufflation pressure. The opening 600 is

opened on insertion of an implement such as a trocar 601 through the opening 600. The bag has excess material in the region of the opening 600 which defines a sleeve or chimney 603 which seals along part of the length of the trocar shaft when the shaft is in place in the opening 600. The opening 600 is located at the top of the bag, in use i.e. in the region that touches the abdominal wall. A trocar 601 may be inserted in a region of the abdominal wall remote from key organs and manipulated under full vision to locate the opening. The opening stays closed whilst there is internal insufflation pressure and the sleeve or chimney 603 collapses around the trocar to create a seal.

[0205] Referring to Figs. 161 to 163, in this case any excess bag material 604 may be grasped using a grasper 605 inserted through a trocar 606. The end of the excess material may then be cut (Fig. 162) creating a chimney 607 extending from the bag. The chimney 607 is then sealed - for example using a threaded trocar 608 which is twisted inside the chimney 607 to create a seal. The chimney 607 prevents contact between the wound and debris created during morcellation. In this case the surgeon is provided with an additional port into the bag for use during the procedure. One such procedure may involve insertion of a laparoscope or a grasper, for example through the secondary port.

[0206] Referring to Fig. 164, in another embodiment a bung 610 may be placed inside the trocar 606, sealing the bag to the trocar. An implement such as a laparoscope 611 may be inserted through the bung 610.

[0207] Referring now to Fig. 165, in this case the bag has a cuff 620 and a tether 621 loop is threaded through the cuff. The tether 621 has a strap 622 which may be used to pull on the tether from a location external of the abdomen. The cuff 620 also has grasping tabs 623, 624. In this case a grasper 625 may be used to grip the tab 623 as illustrated and then push the cuff towards the opposite side of the bag, towards the tab 624, thus closing the bag. The strap 622 enables the bag to be held firmly whilst being closed within the pneumoperitoneum.

[0208] In some cases the bag is retained in the normally open configuration by a retainer ring of a shape memory material such as Nitinol. It has been found that a single ring formed by attaching, for example by welding, together the ends of a ring-forming member is most efficient in achieving rapid deployment. Referring to Figs. 185 and 186 the retaining ring of the bag may be manufactured in any suitable manner. In one case, for example, a shape memory material 675 is formed into a loop, shape set and welded. A tether strap 676 may also be attached in any suitable manner.

[0209] In some cases the cuff of the bag may have features to facilitate mounting of the pre-formed ring to the bag.

[0210] In one embodiment illustrated in Fig. 166 the bag has a cuff 630 through which a loop 631 of a shape memory material such as Nitinol is led. Internal capturing regions 632 for the retaining loop 631 are created by

cutting panels in the material of the bag and folding down over the retaining loop 631 and sealing using a suitable tool such as a circular sealing tool. A tether strap 632 is mounted to the loop 631 and extends through an access port 633 to maintain control over the bag.

[0211] Referring to Fig. 167, in this case pressed out attachments 635 are folded over a closed retaining loop 636 of a shape memory material such as Nitinol and sealed, for example by heating. A front grasping tab 637 includes reinforced attachment loops 638.

[0212] Figs. 168 to 170 illustrate another method of attaching a retaining loop 640 to a cuff 641 of a bag according to the invention. In this case the material of the collar or cuff 641 is punched out with a cut detail 642 extending from a hole 643 (through which the retaining ring 640 is extended) to the edge of the cuff. The cuff is formed into a collar 641 as illustrated in Figs. 169(a) to 169(c). The retaining ring 640 is inserted through the cut details 642 and into the receiving holes 643. Using a circular sealing tool, the tabs are folded down to create an alternating overlap. The cuff is then sealed, for example by heatsealing to close the holes 643. In this way a closed loop 640 of a shape memory material such as Nitinol is attached to the bag to maximise the opening and facilitate insertion of tissue laparoscopically when the bag is within the abdomen.

[0213] In some embodiments the abdominal bag is protected from damage during use of a morcellator. For example, as illustrated in Fig. 171 an insert/cannula 650 may be inserted to isolate a morcellator from the material of the bag. The insert 650 may for example be rigid, flexible or of a segmented rigid material. An access port or cap 651 may be releasably mountable to the insert 650.

[0214] The device used to introduce the bag into an insufflated abdomen may be adapted to control the opening of the bag. For example, as illustrated in Figs. 172 to 176 a tip 655 of an introducer sheath 656 may have mirrored slots 657 which allow a retaining ring of the bag to start to expand before the full bag is inserted. Such an introducer will aid manoeuvring of the bag into a desired position before the bag becomes detached from the introducer. In addition, the deployed bag can be partially withdrawn into the introducer to enable further rigid movement of the bag to optimise positioning.

[0215] To limit blood or other material being pushed into the introducer by the positive insufflation pressure within the abdomen the tip of the introducer may have pressure dissipating features. For example, as illustrated in Figs. 177 to 180 the introducer tip may have various slots 660, holes 661 and/or tapering features 662.

[0216] In another embodiment as illustrated in Fig. 189 to restrict blood or other material from entering the introducer, a shaft 695 of the introducer may have a valve 696 at the proximal end. The valve 696 may be openable on insertion of a pusher 697 used to deploy the bag device from the introducer.

[0217] It will be appreciated that the bag of the invention may comprise different materials to fulfil different re-

quirements for sections of the bag. For example, as illustrated in Figs. 181 and 182 a main body 665 of the bag may be made from a relatively thin flexible material and a collar 666 of a thicker, more rigid material. For example, the main body 665 may include a sterilized polyester polyurethane, such as SS-1495-95 (95 shore A polyester polyurethane blown film with high slip surface and good clarity), and the collar or neck portion 666 may include a sterilized polyether polyurethane, such as ST-625-85 (85 Shore A polyether polyurethane formulated for medium/high slip with good sealability).

[0218] The bag may also be of any desired shape/profile. For example, the bag may be of oval or elliptical profile as illustrated in Figs. 181 to 184. This shape assists in pushing organs such as a section of bowel away from the abdominal wall which prevents the bowel from migrating over the top of the bag. As shown in Fig. 184, the creation of the artificial pneumoperitoneum includes positioning the joined edges 671 of the bag in alignment with the lateral walls of the abdomen 672 such that one of the planar sheet portions of the bag faces, and is in contact with, the anterior abdominal wall, and the other sheet portion of the bag faces and contacts the viscera. Tissue 670 is illustrated in Figs. 183 and 184 within the bag.

[0219] The bag may be sized to be larger than the peritoneal cavity of the patient so as to provide apposition against the abdominal wall after inflation. The oversized nature of the bag, and the loading of the bag into the introducer as described below in connection with Figs. 187(a) to 187(f), allows any additional bag material after inflation to remain rolled at the bottom of the bag. This helps to minimize pleats or folds of the inflated bag from extending into the artificial pneumoperitoneum. Consistent with this, and as shown in Fig. 181, the bag may be sized to include a total bag length 661 of between 300 and 600 mm, or between 400 and 550 mm, or approximately 500 mm, all in an uninflated condition. The bag may also include a maximum width 677, for example in the body portion 665, of between 200 and 500 mm, or between 300 and 400 mm, or approximately 350 mm, all in the uninflated condition. The collar or neck 666 may include a constant width 663 of between 150 and 350 mm, or between 180 and 320 mm, or approximately 250 mm, all in a bag uninflated condition and when the neck 666 is flattened as shown in Fig. 181. The collar or neck 666 may also include a length 664 of between 100 and 300 mm, or between 100 and 200 mm, or approximately 150 mm, in a bag uninflated condition. The main body 665 may include a main body length 667 (not including the neck) of between 200 and 500 mm, or between 300 and 400 mm, or approximately 350 mm. Also, main body 665 may be configured to have a maximum width to maximum length ratio of approximately 1:1.

[0220] Also, as noted above, the bag may be retained in the normally open configuration by a retainer ring of a shape memory material such as Nitinol. Referring to Figs. 166-170, the retaining ring of the bag may be formed into

a loop, shape set and welded. The diameter of the bag opening when the retaining ring is in an open configuration may be between 100 and 220 mm, or between 120 and 200 mm, or approximately 160 mm.

[0221] Referring again to Figs. 181 and 182 the bag may be formed from shaped flat sheet laid on top of one another and attached at the edges 671, for example by welding. When inflated (Fig. 182), the bag expands into an ovoid type shape with rounder corners which can more closely align with the contours of the abdomen. Also as shown in Figs. 118, 181, and 182, the bag may be shaped to be symmetric about two different planes (668, 669) that are normal to one another. In addition, as shown in Fig. 118 and 181, neck or collar portion (401, 666) may be shaped to extend parallel to a longitudinal axis 271 of the bag, and thus edges 674 of the neck or collar portion (401, 666) may extend parallel to the longitudinal axis 271 of the bag.

[0222] The bag of the invention in some cases may be manipulated to optimise ease of loading into an introducer and/or to optimise deployment. For example, as illustrated in Figs. 187(a) to 187(f) the bag may be wider at certain sections 680. These extra width sections may be folded inwardly from the side to form pleats 681. The bag may then be rolled from the bottom up towards the neck 682 of the bag. The neck 682 of the bag may then be scrunched down and loaded into an introducer 683.

[0223] Referring to Fig. 188 in some cases sections of the bag may be of a different material and or material thickness in order to provide desired properties. For example, a bottom section 690 of the bag may be of a stronger / more rigid material to resist the action of a morcellator. The section 690 may be of a rip-stop nylon material, by way of example.

[0224] Figs. 190(a) to 190(c) illustrate the use of the bag device of the disclosure in trans-vaginal procedures, for example hysterectomies. In some cases (Figs. 190(a) and 190(b)) the bag device may be inserted and used directly without an access port. In other cases, the device may be used as described in other embodiments with an access port 730 (Fig. 190(c)).

[0225] Referring to Fig. 191, in some cases the twisting action of a threaded trocar or bung 700 may assist in pulling the bag up through the wound.

[0226] In some embodiments, in order to seal the hole into the chimney before the bag is deflated a closure device or valve may be deployed. For example, as illustrated in Figs. 192(a) and 192(b) a closure device 710 may be inserted into the chimneys in a collapsed configuration and deployed at the distal end of the chimney. The closure device may be retained in place using any suitable retaining means such as an adhesive pad(s) 711.

[0227] The abdominal bag device of the disclosure may be used with any suitable access port system 730 as illustrated, for example, in Figs. 193(a) to 193(c). In some cases the access port is mounted within the neck of the bag.

[0228] Referring to Figs. 194 to 285 there are illustrated

further views of various artificial pneumoperitoneum devices and systems of the invention. These devices and systems have several features in common with the devices described above and the relevant disclosures should be read in conjunction with the following.

[0229] Fig. 194 to 198 illustrate an artificial pneumoperitoneum device 800, an introducer sheath 801 for the device 800 and a plunger 802 for delivery of the device 800 from the sheath 801. The sheath 801 has an open proximal end 803 with a tapered profile and an indicator arrow to indicate to the user the direction of travel of the plunger, and a distal end 804. The plunger 802 has a handle 805 at the proximal end and an enlarged sealing head 806 at the distal end. There are also seals 807 spaced-apart along the length of the plunger 802. Referring also to Fig. 210 the plunger 802 is sized to overshoot the distal end of the introducer 801 to ensure that the device 800 is always fully ejected from the introducer sheath 801. When the distal seal 806 has been ejected from the introducer sheath 801 the plunger 802 will sufficiently fill the introducer tube 801 to prevent significant loss of pneumo whilst being sufficiently loose to ensure that a tether 809 part of the device can be pulled freely.

[0230] As described in detail above and referring also to Figs. 199 and 200 the shape and size of the device has several unique benefits. Figs. 199 and 200 show a bag part of the device in a flattened configuration (i.e. without an opening ring structure). The bag has a main body region 810, a collar region 811 and a neck region 812 between the collar 811 and the main body 810. A length L of the main body is the same as the width W of the main body 810 whilst the neck region 812 is narrowed. The neck region 812 in use is used to line the incision or opening through which it is introduced. The collar 811 is stiffer than the neck and main body and is typically welded to the neck 81 region.

[0231] A biasing element in the form of a Nitinol loop 815 extends through and in this case is threaded through the collar 811. Pull tabs 816, 817 are mounted to and extend proximally from the collar 811. As described above, the collar is of stiffer material than the main body of the bag and in addition to assisting in maintaining an opening it also facilitates secure mounting of various elements such as the biasing loop 815 and the tabs 86, 817.

[0232] Referring to Fig. 201, in use a seal 820 is pressed out of closed cell foam with the slit in the middle. The tether is passed through this slit and glued into place. The seal 820 seals the introducer preventing blood being sucked up onto the introducer and possibly out of the handle. The proximal tab provides reinforcement for the two holes that the Nitinol wire passes.

[0233] Referring to Figs. 202 to 205, the procedure for fitting the Nitinol ring 815 is as follows:

- The collar 811 is formed before the Nitinol wire 815 is added.
- The Nitinol wire 815 is threaded through holes in the collar, alternating inside and outside in a pattern as

illustrated in Fig. 202.

- The Nitinol wire 815 is on the inside of the collar at the distal side of the collar 811.
- The Nitinol wire 815 is on the outside of the collar at the proximal side, where the tether 809 is attached.
- The distal tab 817 captures the Nitinol 815 wire on the inside of the collar 811.
- Due to the requirement to thread the Nitinol through the holes, a cold crimping process is used.
- Both ends of the Nitinol wire 815 are placed inside a tube 825 (which may be of Nitinol) which is then cold pressed to hold the wire 815.
- The tether 807 is formed over the crimp to ensure the crimp comes out first and to cover any sharp edges.
- The crimp region is the first part of the wire to be pulled outside of the body by the tether 809.

[0234] Figs. 206 to 209 illustrate the folding/rolling of the device 800 to ready it for loading. The procedure is as follows:

- Fold in the excess width to make same width as flattened opening.
- Roll from the bottom as far as the shoulder / neck.
- Scrunch down the neck. This ensures that the tissue specimen has sufficient space to allow the device to close without the tissue getting caught in the opening into the device.
- The device is then loaded into the introducer 801.

[0235] Referring to Figs. 211 and 212 it will be apparent that the shape and the width of the bag device 800 retracts the viscera 840 whilst ideally preventing the viscera from creeping over the device to become trapped between the device and the abdominal wall.

[0236] Referring to Figs. 213 to 215 it will be apparent that once inflated the bag device has no pleats, folds, curtains or excess material.

[0237] Referring to Fig 216 it will be noted that the bag device 800 is larger than most abdominal cavities therefore ensuring all available space within the abdominal cavity is taken by the bag device. The device, its loading and how it inflates minimises internal folds in material and that any excess material is kept on the outside, therefore not providing any accidental grabbing hazards on the inside. As the bag device 800 inflates, it only unrolls as much material as necessary to fill the available space, leaving all excess material neatly rolled on the outside.

[0238] Referring in particular to Figs. 217 and 218 it will be apparent that the viscera 840 is retracted away from a morcellator, making it difficult for the morcellator to accidentally touch off the body of the bag device 800.

[0239] It will be apparent from Figs. 219 and 220 that a zero seal and lip seal introducer 845 may be moulded

into the introducer handle which is pierced by the plunger 802. In his case the tether 809 extends out from the front of the introducer 801 and up along the outside.

[0240] Figs. 221 and 222 illustrate an alternative manufacture described above in which for assembly with a solid Nitinol ring; the cuff 811 is attached to a pre-formed Nitinol ring 850.

[0241] Fig. 223 illustrates an alternative manufacturing procedure for assembly with a solid Nitinol ring 850. This uses loops 851 instead of threading the Nitinol through holes.

[0242] Referring to Fig. 224 there is illustrated a bag device 800 being introduced through an access port 860. A scope 861 is also inserted.

[0243] Fig. 225 and 226 illustrates introduction of the device 800 in a multiport configuration in which a scope 861 is inserted through a separate port or trocar.

[0244] Referring to Figs. 227 and 228 it will be apparent that the bag device 800 opens automatically as it is ejected from the introducer 801.

[0245] Figs. 228 to 230 illustrate that as the bag device 800 is ejected from the introducer 801, the distal tab 817 pops up with an arrow printed on both sides to aid the surgeon with orientation - i.e. prevents the bag device being accidentally deployed upside down.

[0246] Figs. 231 and 232 illustrate tissue 870 being manipulated into the opening of the device, in some cases using a flipping or a rolling action.

[0247] Fig. 233 shows a surgeon pulling the tether 809 until the opening ring is partially pulled through a large instrument valve 875.

[0248] Fig. 234 and 235 illustrates that the device closes automatically as it is pulled up against the wound retractor 880.

[0249] Figs. 236 and 237 illustrate removal of the port and the surgeon continuing to pull the tether 209 until the device collar is exposed which is indicated by a line.

[0250] Referring to Figs. 238 and 239 the surgeon then pulls the tabs 816, 817 apart to open the device.

[0251] Figs. 240 and 241 are internal views of the bag device 200 before inflation. All gas is expelled from the abdomen before the access port 860 is reattached.

[0252] The reattachment of the port 860 is illustrated in Fig. 242.

[0253] Referring to Figs. 243 and 244 an insufflation supply line 885 is connected to the access port 860 and the bag device 800 is inflated.

[0254] Referring to Fig. 245 it will be noted that the viscera 840 is retracted and the bag device 800 lines the abdominal wall when it is inflated.

[0255] Referring to Figs. 246 and 247 it will be noted that the bag device 800 prevents bowel 840 from creeping up over the device.

[0256] Referring to Figs. 248 and 249 it will be apparent that the bag device 800 retracts bowel 840 to a safe distance avoiding accidental connection with the morcellator or its tenaculum under vision significantly.

[0257] Figs. 250 and 251 illustrate that under vision,

the surgeon morcellates the tissue until no large chunks remain.

[0258] The bag device of the invention provides an artificial pneumoperitoneum in which procedures can be carried out on tissue isolated in the bag.

[0259] Referring to Figs. 252 and 253 illustrate removal of the access port and, once deflated, the bag device 800 can be pulled out, bringing with it all the debris from the morcellation.

[0260] The device of the disclosure can also be used in trans-vaginal procedures in which the device is introduced into the vagina as illustrated in Figs. 254 to 256.

[0261] In some cases the bag device 800 may be closed within the insufflated abdomen as illustrated in Fig. 257.

[0262] Referring to Fig. 258 in some cases an additional port 890 may be used to pass through the bag device 800 to provide access for an instrument or a laparoscope.

[0263] Referring to Fig. 259 illustrates that the key device 800 can be deployed before the tissue has been resected. The device can be placed under the uterus 895 during dissection to catch any droplets of blood or tissue.

[0264] Fig. 260 illustrated that whilst partially deployed, the device can scoop up tissue. This is facilitated by the relatively stiff collar 811.

[0265] Referring to Figs. 261 and 262 in this case the device is made with excess material in the top area which can be grabbed with a grasper to be pulled through a trocar.

[0266] Referring to Figs. 263 to 266 illustrates a supplementary or second skin 900 on the device. Through a trocar incision 902, a grab tab 901 on the excess material 900 is pulled up through the wound. A hole is cut into the excess material 900 and the trocar is reinserted inside the extra material - this material then protects the wound from any insemination. The trocar then punctures into the main chamber of the device. This ensures that the insertion of the trocar does not cause material to egress through the hole created by the trocar.

[0267] Fig. 267 illustrates the puncturing of the device with a threaded trocar 910 twisting the trocar will pull excess material in the device up through the wound, therefore protecting it from dissemination.

[0268] Figs. 268 to 270 illustrate various bungs 940, 941, 942 which may be of closed cell foam in several shapes - this is pushed down through the trocar once the surgery is complete and before the insufflation is removed.

[0269] Figs. 271 to 273 illustrate that a bung 945 may have a rigid core to aide pushing the bung through the trocar.

[0270] Figs. 274 to 276 a bung 945 may be placed inside the inflated device. A grasper introduced via a second trocar 950 grabs the bung and pulls it up to the trocar. The trocar is pulled back and the bung is pulled into the puncture hold to seal it.

[0271] Referring to Figs. 277 to 280 illustrate an inflat-

able bung 960. Deflated - the bung 960 is passed through the trocar. Once in place, the bung 960 is inflated using a large syringe. The bung may contain a one way valve.

[0272] Figs. 281 to 285 illustrate a cover 970 for a trocar 971 that can be inflated to block the hole in the device once the trocar 971 has been removed. Fig. 281 show the cover 970 uninflated. Fig. 282 shows the cover 970 inflated.

[0273] Various features of the invention are described and illustrated. It will be appreciated that at least some of the features described in relation to one embodiment may be used not only in the embodiment specifically described but also in other appropriate embodiments. As used herein, the term "approximately" or "about" is understood to mean close in value or amount, but not precise. To the extent that such a definition for "approximate" or "about" is not considered sufficiently definite, approximate is understood to mean plus or minus 5% of the relevant parameter.

[0274] The invention is not limited to the embodiments hereinbefore described, with reference to the accompanying drawings, which may be varied in construction and detail.

Claims

1. An inflatable artificial pneumoperitoneum bag, comprising:

a length (661) and a width (677);
 an artificial pneumoperitoneum bag neck portion (666) having a first end and a second end, the first end forming a bag opening;
 an artificial pneumoperitoneum bag body portion (665) forming a closed cavity in fluid communication with the second end of the neck portion (666), the body portion (665) having a length (667) and width (677) greater than a length (664) and width (663) of the neck portion; and
 a ring (675) located at the bag opening, the ring configured to bias the bag opening toward an open condition,
 wherein the neck portion includes edges (674) that extend parallel to a lengthwise axis of the bag,
 wherein the length (661) of the bag is between 300 and 600 mm when the bag is uninflated, a maximum width (677) of the bag is between 200 and 500 mm when the bag is uninflated, and the length of the neck portion (666) is between 100 and 300 mm when the bag is uninflated, and
 wherein the neck portion has a diameter of between 100 and 220 mm when in the open condition.

2. An inflatable artificial pneumoperitoneum bag as claimed in claim 1 wherein a width-extending cross-

section of the body portion (665) includes an oval shape when the bag is inflated.

3. An inflatable artificial pneumoperitoneum bag as claimed in claim 1 or 2 wherein the bag is symmetric about two planes (668, 669) that are normal to one another.
4. An inflatable artificial pneumoperitoneum bag as claimed in any of claims 1 to 3 wherein the ring (875) is formed of a shape memory material and is received through loops in the neck portion.
5. An inflatable artificial pneumoperitoneum bag as claimed in any of claims 1 to 4 comprising:

a first planar sheet portion and a second planar sheet portion, the first and second planar sheet portions having joined edges to form the pneumoperitoneum bag.
6. An inflatable artificial pneumoperitoneum bag as claimed in any of claims 1 to 5 comprising a cuff (811) at the proximal opening, the cuff (811) having a closed configuration for delivery and retrieval of the bag and an open configuration to receive tissue, the cuff (811) being biased into the open configuration by the ring (815) which extends around the cuff (811).
7. A bag as claimed in claim 6 wherein the cuff (811) is of a stiffer material than that of the body portion (810) of the bag.
8. A bag as claimed in any of claims 6 or 7 wherein the main body of the bag includes a neck region (812) extending from the cuff (811) and wherein the neck region (812) is more flexible than the cuff (811).
9. A bag as claimed in any of claims 6 to 8 wherein the cuff (811) extends for a distance of between 2 and 20 cm, for a distance of between 2 and 10 cm, or wherein the cuff extends for a distance of about 5 cm.
10. A bag as claimed in any of claims 1 to 9 wherein the ring (811) is of a shape memory material such as Nitinol.
11. A bag as claimed in any of claims 6 to 10 comprising a tether (809) extending proximally from the cuff (811), the tether may be mounted to the ring (815).
12. A bag as claimed in any of claims 6 to 11 wherein the cuff (811) comprises a tab (816, 817) for use in grasping the bag.
13. A bag as claimed in claim 12 comprising tabs (816, 817) arranged on opposite sides of the cuff (811).

14. A bag as claimed in claim 12 or 13 wherein the or each tab (816, 817) comprises an indicator to indicate a preferred orientation of the bag.
15. A bag as claimed in any of claims 1 to 14 wherein the bag is configured to be foldable into a reduced size for insertion through an incision or an opening.
16. A bag as claimed in any of claims 1 to 15 wherein the bag comprises a port.
17. A bag as claimed in claim 16 wherein the port is an exit port and/or wherein the port is an entrance port.
18. A bag as claimed in claim 16 or 17 wherein the port comprises a valve, the valve may comprise a choke valve or a cuff valve.
19. A system comprising a bag as claimed in any of claims 1 to 18 and an introducer (801) for containing the bag in an insertion configuration.
20. A system as claimed in claim 19 wherein the introducer (801) is configured to be at least partially insertable through an incision and/or an access port.
21. A system as claimed in claim 19 or 20 comprising an activator (802) for delivering the bag from the introducer (801), on insertion.
22. A system as claimed in claim 21 wherein the activator comprises a tab, or wherein the activator comprises a plunger.
23. Apparatus for use in laparoscopic surgery comprising a bag as claimed in any of claims 1 to 18 and a retractor (880).
24. Apparatus as claimed in claim 23 further comprising an access port (860).

Patentansprüche

1. Aufblasbarer Pseudopneumoperitoneumbeutel, der Folgendes umfasst:

eine Länge (661) und eine Breite (677),
einen Pseudopneumoperitoneumbeutel-Halsabschnitt (666) mit einem ersten und einem zweiten Ende, wobei das erste Ende eine Beutelöffnung bildet,
einen Pseudopneumoperitoneum-Beutelhauptabschnitt (665), der einen geschlossenen Hohlraum bildet, welcher mit dem zweiten Ende des Halsabschnitts (666) in Fluidverbindung steht, wobei der Hauptabschnitt (665) eine Länge (667) und eine Breite (677) aufweist, die größer

- sind als eine Länge (664) und eine Breite (663) des Halsabschnitts, und einen Ring (675), der sich an der Beutelöffnung befindet und so konfiguriert ist, dass er die Beutelöffnung in einen offenen Zustand vorspannt, wobei der Halsabschnitt Ränder (674) aufweist, die parallel zu einer Längsachse des Beutels verlaufen, wobei die Länge (661) des Beutels zwischen 300 und 600 mm beträgt, wenn er nicht aufgeblasen ist, eine maximale Breite (677) des Beutels zwischen 200 und 500 mm beträgt, wenn er nicht aufgeblasen ist, und die Länge des Halsabschnitts (666) zwischen 100 und 300 mm beträgt, wenn der Beutel nicht aufgeblasen ist, und wobei der Halsabschnitt einen Durchmesser zwischen 100 und 220 mm aufweist, wenn er sich im offenen Zustand befindet.
2. Aufblasbarer Pseudopneumoperitoneumbeutel nach Anspruch 1, wobei ein in Breitenrichtung verlaufender Querschnitt des Hauptabschnitts (665) eine ovale Form aufweist, wenn der Beutel aufgeblasen ist.
 3. Aufblasbarer Pseudopneumoperitoneumbeutel nach Anspruch 1 oder 2, wobei der Beutel um zwei Ebenen (668, 669) symmetrisch ist, die senkrecht zueinander sind.
 4. Aufblasbarer Pseudopneumoperitoneumbeutel nach einem der Ansprüche 1 bis 3, wobei der Ring (875) aus einem Formgedächtnismaterial gebildet ist und von Schlaufen im Halsabschnitt aufgenommen wird.
 5. Aufblasbarer Pseudopneumoperitoneumbeutel nach einem der Ansprüche 1 bis 4 mit einem ersten und einem zweiten flachen Folienabschnitt, die zum Bilden des Pneumoperitoneumbeutels miteinander verbundene Ränder aufweisen.
 6. Aufblasbarer Pseudopneumoperitoneumbeutel nach einem der Ansprüche 1 bis 5 mit einer Manschette (811) an der proximalen Öffnung, wobei die Manschette (811) eine geschlossene Konfiguration für das Einsetzen und das Herausnehmen des Beutels und eine offene Konfiguration für das Aufnehmen von Gewebe aufweist, wobei die Manschette (811) von dem um die Manschette (811) herum verlaufenden Ring (815) in die offene Konfiguration vorgespannt wird.
 7. Beutel nach Anspruch 6, wobei die Manschette (811) aus einem steiferen Material besteht als der Hauptabschnitt (810) des Beutels.
 8. Beutel nach einem der Ansprüche 6 oder 7, wobei der Hauptteil des Beutels einen Halsbereich (812) aufweist, der von der Manschette (811) ausgeht, und der Halsbereich (812) flexibler ist als die Manschette (811).
 9. Beutel nach einem der Ansprüche 6 bis 8, wobei sich die Manschette (811) über eine Distanz zwischen 2 und 20 cm, eine Distanz zwischen 2 und 10 cm oder sich über eine Distanz von etwa 5 cm erstreckt.
 10. Beutel nach einem der Ansprüche 1 bis 9, wobei der Ring (811) aus einem Formgedächtnismaterial wie Nitinol besteht.
 11. Beutel nach einem der Ansprüche 6 bis 10 mit einem Halteband (809), das proximal zur Manschette (811) verläuft und an dem Ring (815) angebracht sein kann.
 12. Beutel nach einem der Ansprüche 6 bis 11, wobei die Manschette (811) eine Lasche (816, 817) zur Verwendung beim Ergreifen des Beutels umfasst.
 13. Beutel nach Anspruch 12 mit Laschen (816, 817), die an gegenüberliegenden Seiten der Manschette (811) angeordnet sind.
 14. Beutel nach Anspruch 12 oder 13, wobei die oder jede Lasche (816, 817) eine Angabe zum Angeben einer bevorzugten Ausrichtung des Beutels umfasst.
 15. Beutel nach einem der Ansprüche 1 bis 14, wobei der Beutel so konfiguriert ist, dass er sich zum Einführen durch einen Einschnitt oder eine Öffnung auf eine verringerte Größe zusammenfalten lässt.
 16. Beutel nach einem der Ansprüche 1 bis 15, wobei der Beutel einen Port umfasst.
 17. Beutel nach Anspruch 16, wobei es sich bei dem Port um einen Ausgangsport und/oder um einen Eingangsport handelt.
 18. Beutel nach Anspruch 16 oder 17, wobei der Port ein Ventil umfasst, das ein Drossel- oder ein Manschettentventil umfassen kann.
 19. System mit einem Beutel nach einem der Ansprüche 1 bis 18 und einer Einführvorrichtung (801) zum Aufnehmen des Beutels in einer Einführkonfiguration.
 20. System nach Anspruch 19, wobei die Einführvorrichtung (801) so konfiguriert ist,

dass sie sich zumindest teilweise durch einen Einschnitt und/oder einen Zugangsport einführen lässt.

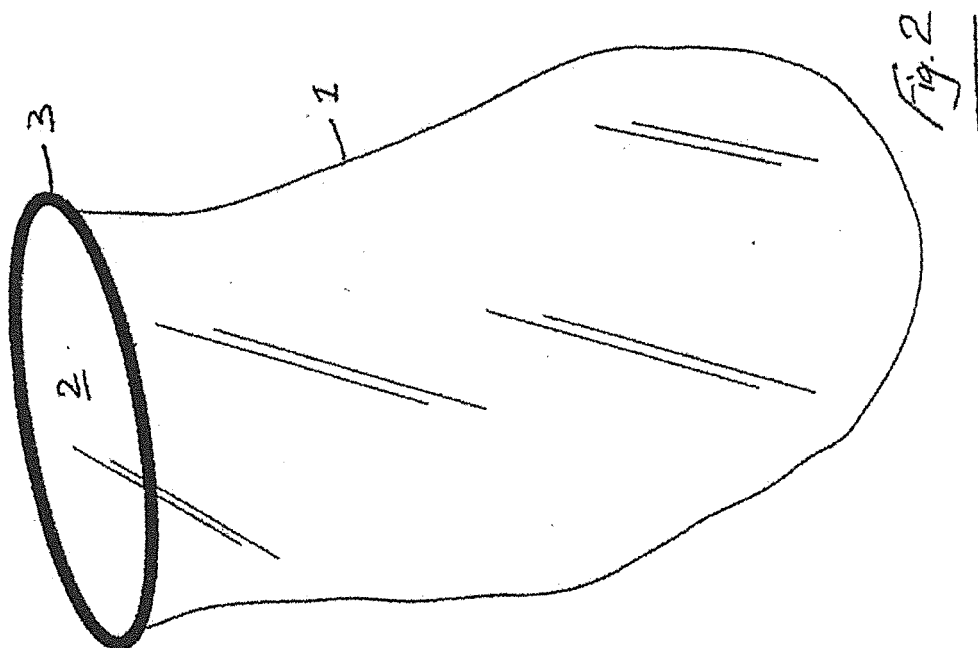
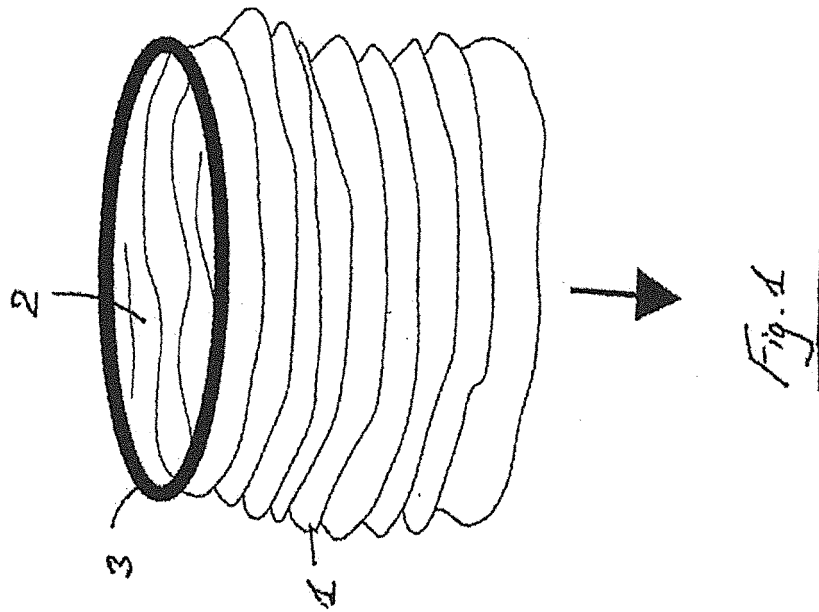
21. System nach Anspruch 19 oder 20 mit einer Aktivierungsvorrichtung (802) zum Einbringen des Beutels aus der Einführvorrichtung (801) nach dem Einführen. 5
22. System nach Anspruch 21, wobei die Aktivierungsvorrichtung eine Lasche oder eine Druckstange umfasst. 10
23. Vorrichtung zur Verwendung bei der laparoskopischen Chirurgie mit einem Beutel nach einem der Ansprüche 1 bis 18 und einer Einziehvorrichtung (880). 15
24. Vorrichtung nach Anspruch 23, die ferner einen Zugangsport (860) umfasst. 20

Revendications

1. Sac de pneumopéritoine artificiel gonflable, comprenant : 25
 - une longueur (661) et une largeur (677) ;
 - une partie de col de sac de pneumopéritoine artificiel (666) ayant une première extrémité et une deuxième extrémité, la première extrémité formant une ouverture de sac ; 30
 - une partie de corps de sac de pneumopéritoine artificiel (665) formant une cavité fermée en communication fluide avec la deuxième extrémité de la partie de col (666), la partie de corps (665) ayant une longueur (667) et une largeur (677) supérieures à une longueur (664) et une largeur (663) de la partie de col ; et 35
 - un anneau (675) situé au niveau de l'ouverture du sac, l'anneau configuré pour solliciter l'ouverture du sac vers une condition ouverte, dans lequel la partie de col inclut des bords (674) qui s'étendent parallèles à un axe longitudinal du sac, 40
 - dans lequel la longueur (661) du sac est située entre 300 et 600 mm quand le sac n'est pas gonflé, une largeur maximale (677) du sac est située entre 200 et 500 mm quand le sac n'est pas gonflé, et la longueur de la partie de col (666) est située entre 100 et 300 mm quand le sac n'est pas gonflé, et 45
 - dans lequel la partie de col a un diamètre situé entre 100 et 220 mm dans la condition ouverte. 50
2. Sac de pneumopéritoine artificiel gonflable selon la revendication 1 dans lequel une section transversale s'étendant sur la largeur de la partie de corps (665) inclut une forme ovale quand le sac est gonflé. 55
3. Sac de pneumopéritoine artificiel gonflable selon la revendication 1 ou 2 dans lequel le sac est symétrique par rapport à deux plans (668, 669) qui sont normaux l'un par rapport à l'autre.
4. Sac de pneumopéritoine artificiel gonflable selon l'une quelconque des revendications 1 à 3 dans lequel l'anneau (875) est formé dans un matériau à mémoire de forme et est reçu à travers des boucles dans la partie de col.
5. Sac de pneumopéritoine artificiel gonflable selon l'une quelconque des revendications 1 à 4 comprenant :
 - une première partie de feuille plane et une deuxième partie de feuille plane, les première et deuxième parties de feuille plane ayant des bords joints pour former le sac de pneumopéritoine.
6. Sac de pneumopéritoine artificiel gonflable selon l'une quelconque des revendications 1 à 5 comprenant un manchon (811) au niveau de l'ouverture proximale, le manchon (811) ayant une configuration fermée pour la livraison et le retrait du sac et une configuration ouverte pour recevoir le tissu, le manchon (811) étant sollicité jusque dans la configuration ouverte par l'anneau (815) qui s'étend autour du manchon (811).
7. Sac selon la revendication 6 dans lequel le manchon (811) est formé dans un matériau plus raide que celui de la partie de corps (810) du sac.
8. Sac selon l'une quelconque des revendications 6 ou 7 dans lequel le corps principal du sac inclut une région de col (812) s'étendant à partir du manchon (811) et dans lequel la région de col (812) est plus souple que le manchon (811).
9. Sac selon l'une quelconque des revendications 6 à 8 dans lequel le manchon (811) s'étend sur une distance située entre 2 et 20 cm, sur une distance située entre 2 et 10 cm, ou dans lequel le manchon s'étend sur une distance d'environ 5 cm.
10. Sac selon l'une quelconque des revendications 1 à 9 dans lequel l'anneau (811) est formé dans un matériau à mémoire de forme tel que le Nitinol.
11. Sac selon l'une quelconque des revendications 6 à 10 comprenant une attache (809) s'étendant de manière proximale à partir du manchon (811), l'attache pouvant être montée sur l'anneau (815).
12. Sac selon l'une quelconque des revendications 6 à 11 dans lequel le manchon (811) comprend une pat-

te (816, 817) pour utilisation lors de la saisie du sac.

13. Sac selon la revendication 12 comprenant des pattes (816, 817) agencées sur des côtés opposés du manchon (811). 5
14. Sac selon la revendication 12 ou 13 dans lequel la ou chaque patte (816, 817) comprend un indicateur pour indiquer une orientation préférée du sac. 10
15. Sac selon l'une quelconque des revendications 1 à 14 dans lequel le sac est configuré pour être repliable en une taille réduite pour être inséré à travers une incision ou une ouverture. 15
16. Sac selon l'une quelconque des revendications 1 à 15 dans lequel le sac comprend un orifice. 20
17. Sac selon la revendication 16 dans lequel l'orifice est un orifice de sortie et/ou dans lequel l'orifice est un orifice d'entrée. 25
18. Sac selon la revendication 16 ou 17 dans lequel l'orifice comprend une soupape, la soupape pouvant comprendre une soupape d'étranglement ou une soupape de manchon. 30
19. Système comprenant un sac selon l'une quelconque des revendications 1 à 18 et un introducteur (801) pour contenir le sac dans une configuration d'insertion. 35
20. Système selon la revendication 19 dans lequel l'introducteur (801) est configuré pour être capable d'être inséré au moins partiellement à travers une incision et/ou un orifice d'accès. 40
21. Système selon la revendication 19 ou 20 comprenant un activateur (802) pour délivrer le sac à partir de l'introducteur (801), à l'insertion. 45
22. Système selon la revendication 21 dans lequel l'activateur comprend une patte, ou dans lequel l'activateur comprend un plongeur. 50
23. Appareil pour utilisation dans la chirurgie laparoscopique comprenant un sac selon l'une quelconque des revendications 1 à 18 et un rétracteur (880). 55
24. Appareil selon la revendication 23 comprenant en outre un orifice d'accès (860).



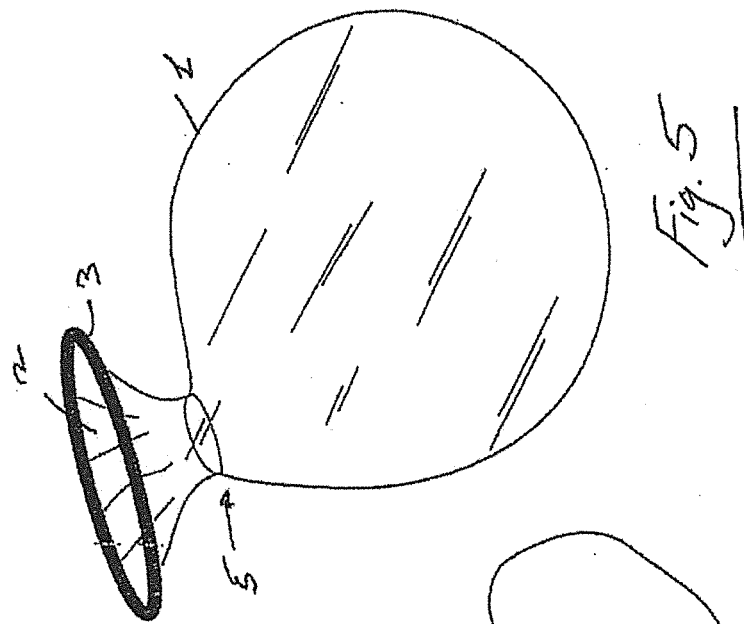


Fig. 5



Fig. 4

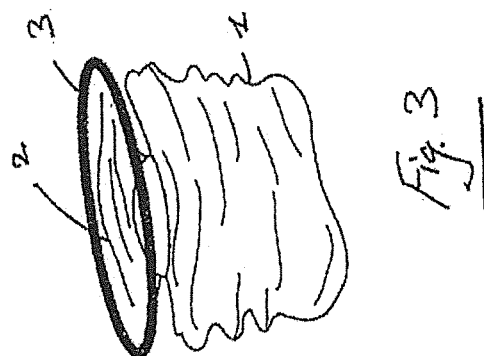


Fig. 3

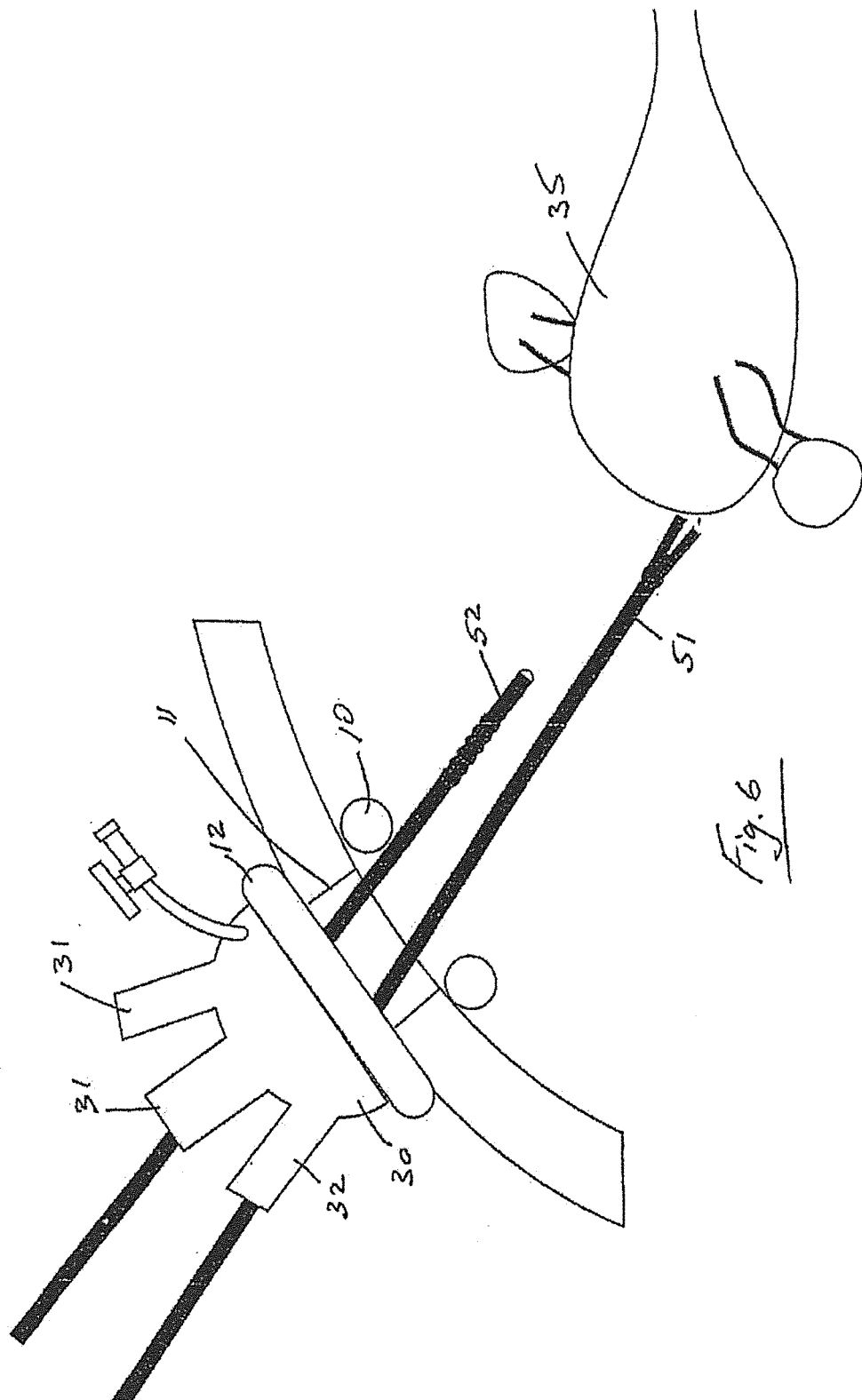
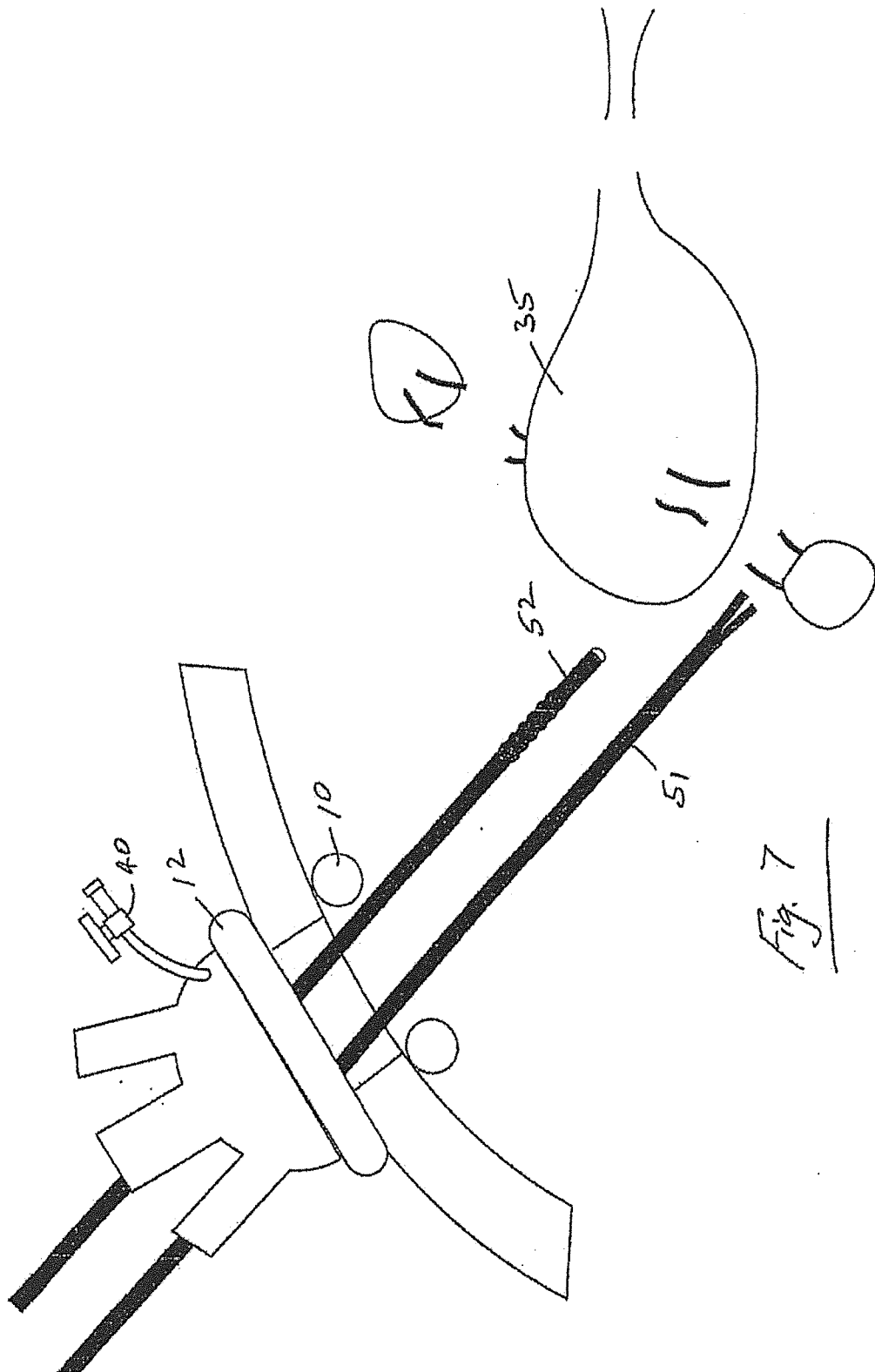


Fig. 6



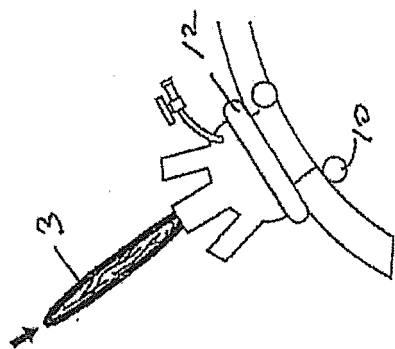


Fig. 10

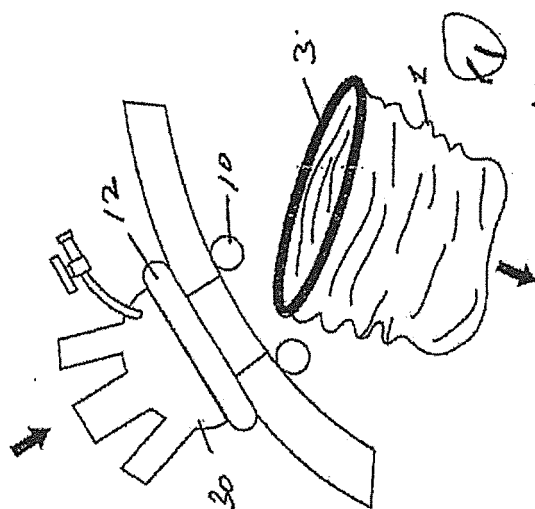


Fig. 9

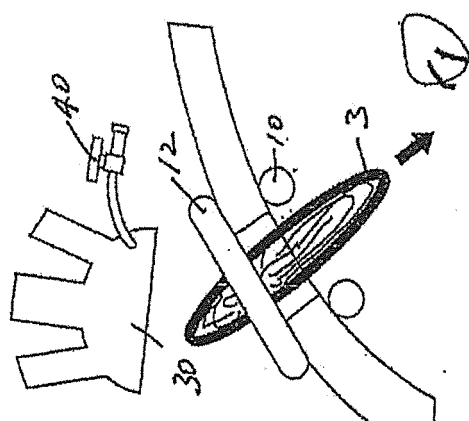
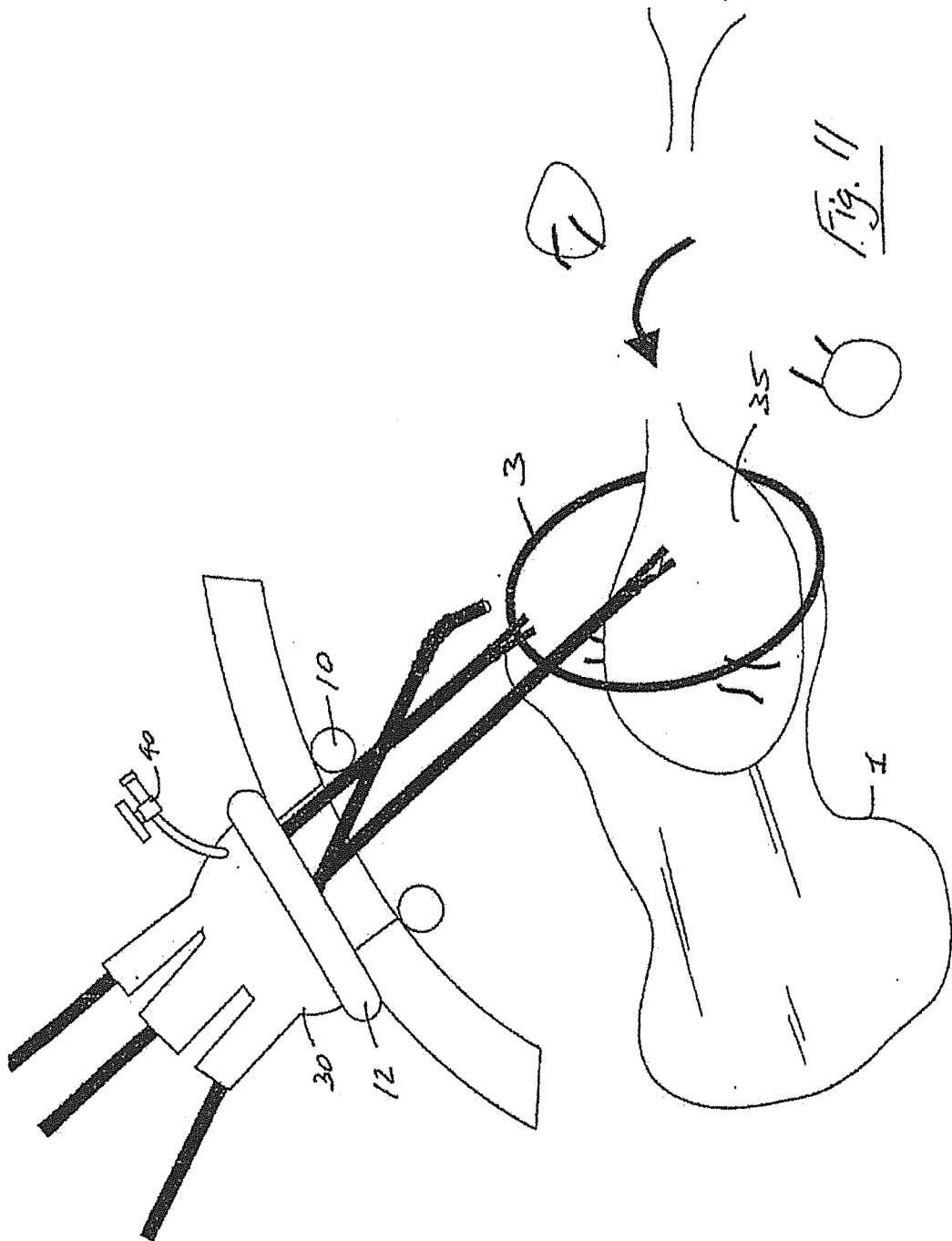
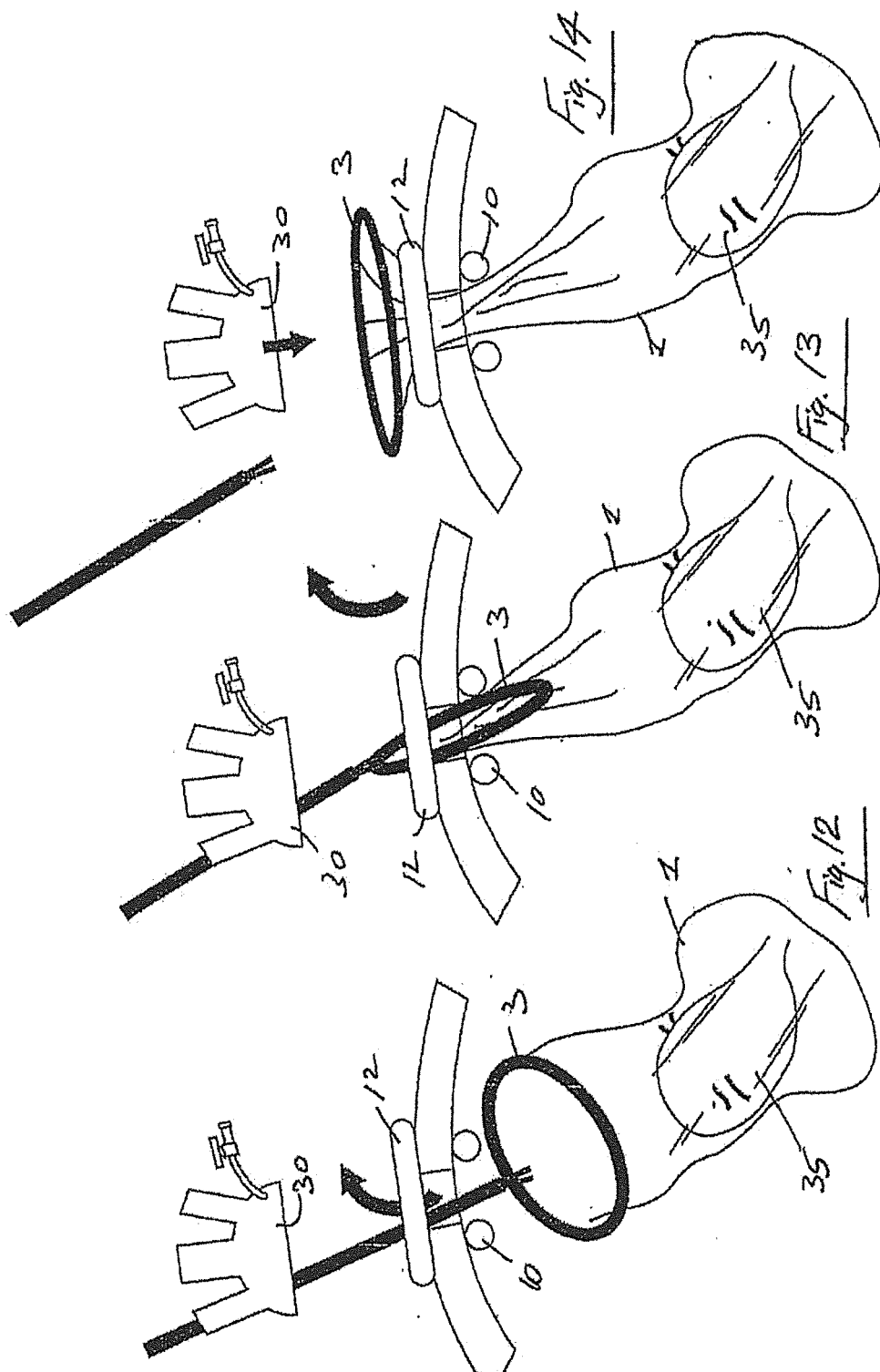
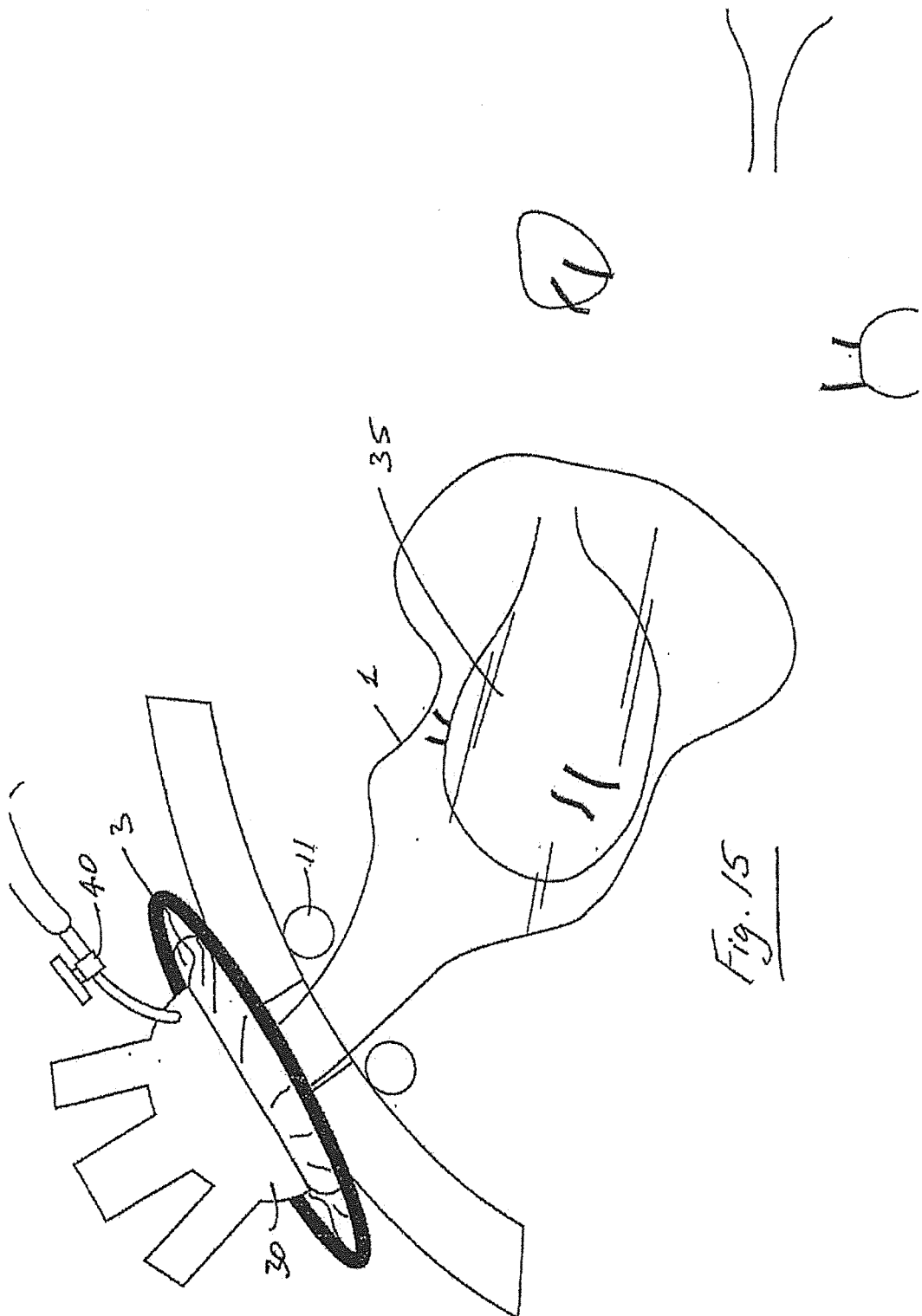


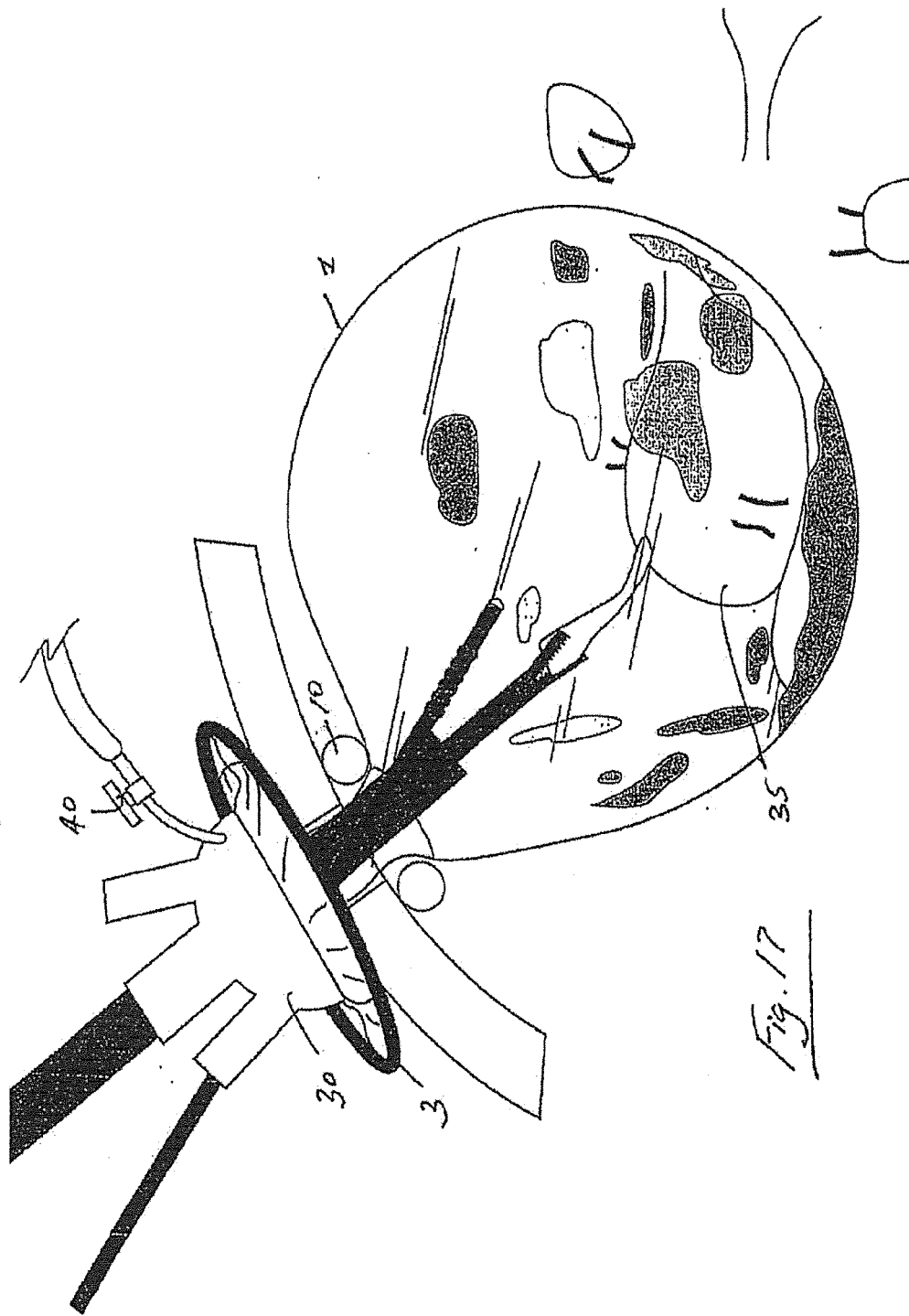
Fig. 8

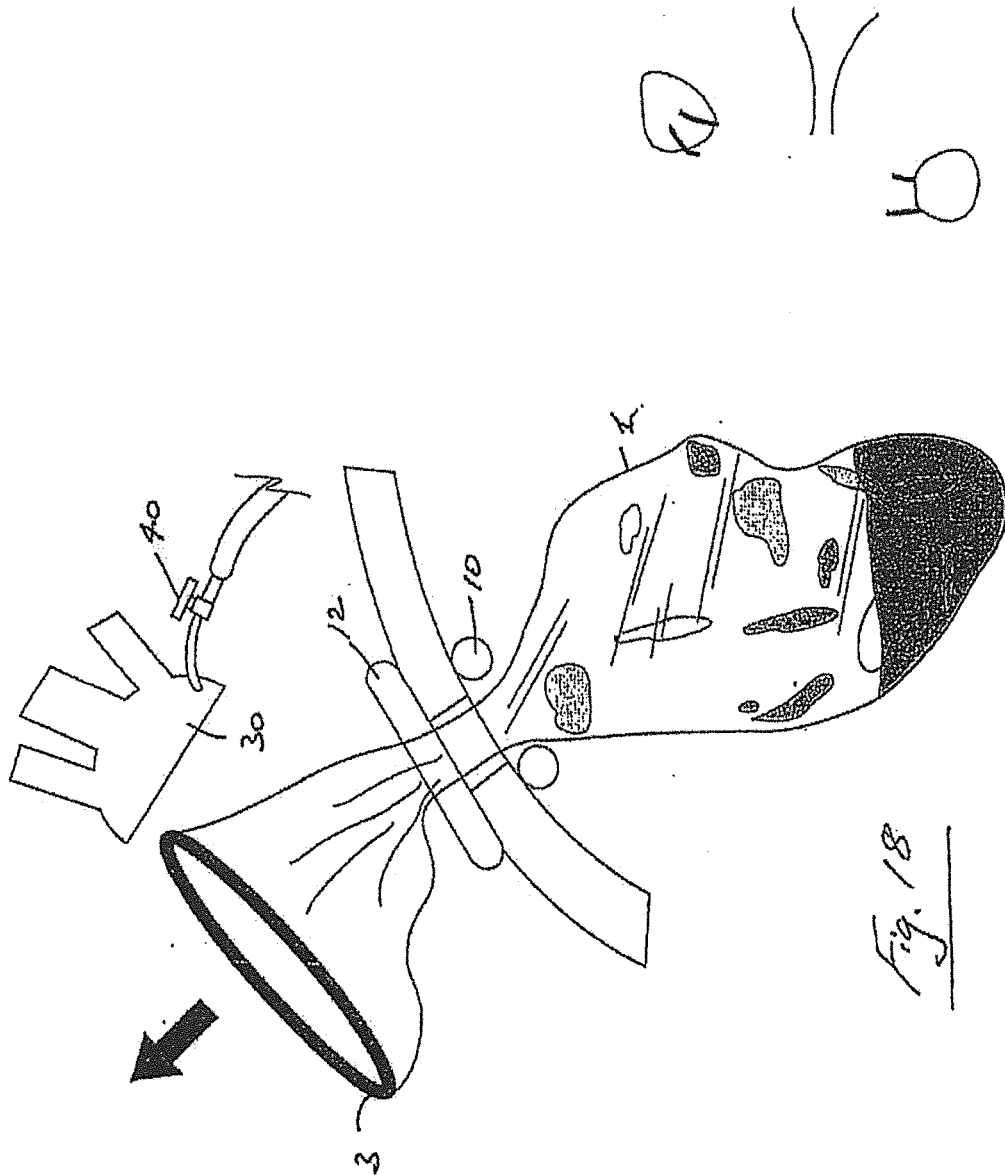


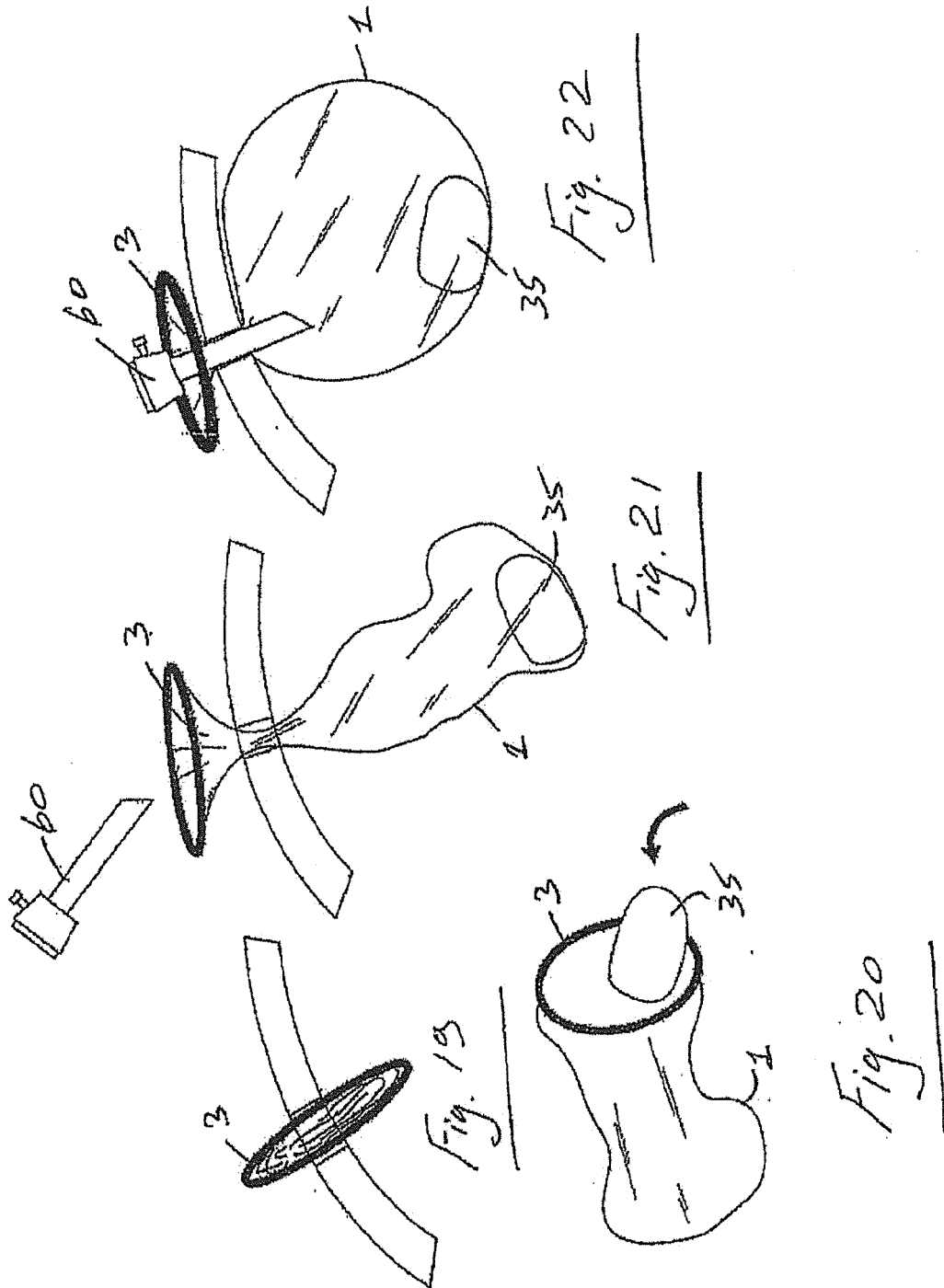


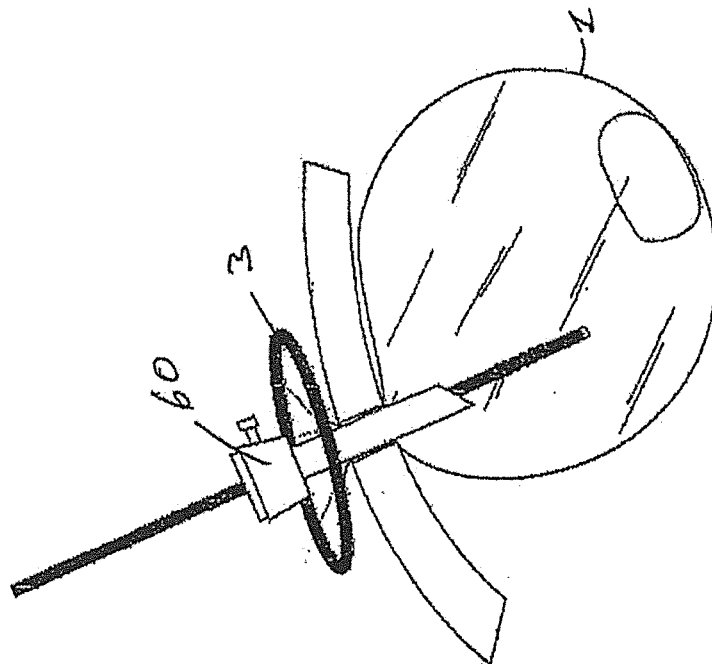
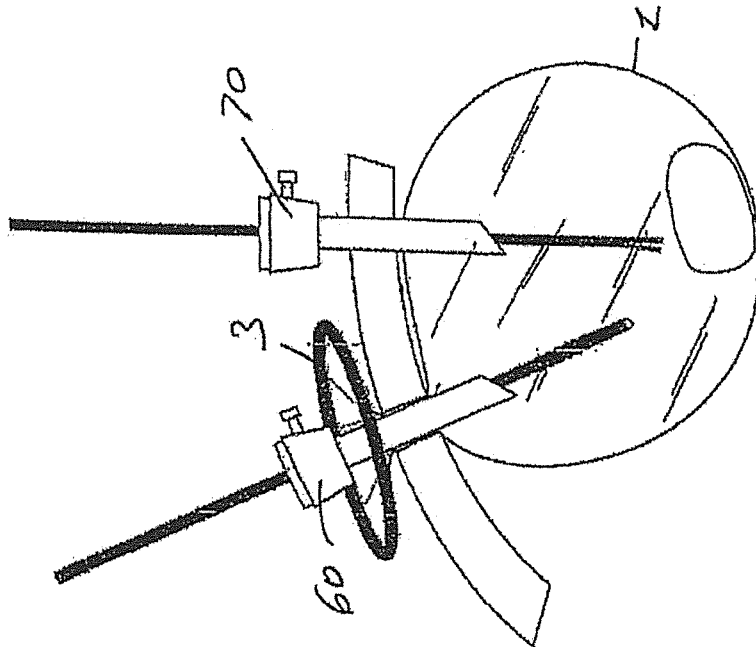












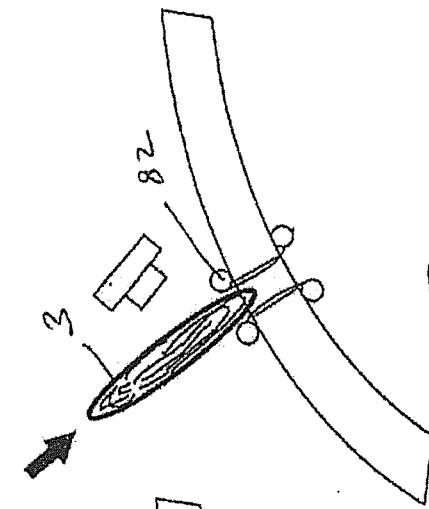


Fig. 25

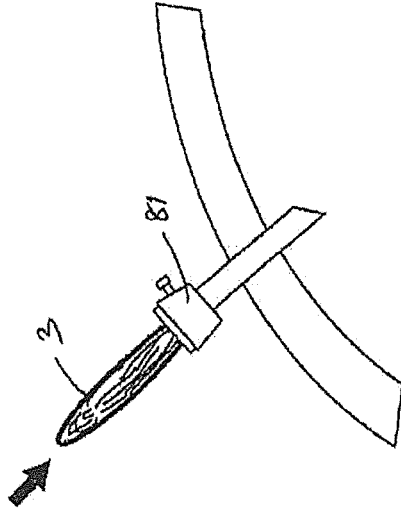


Fig. 26

Fig. 27

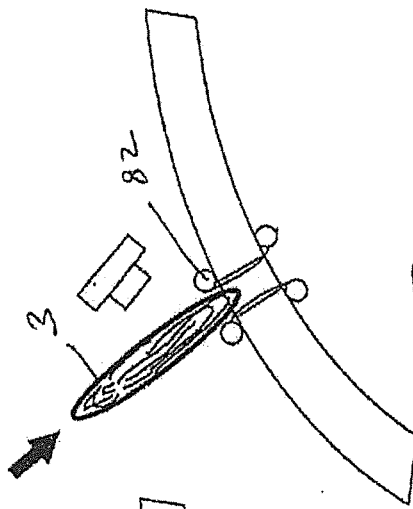


Fig. 28

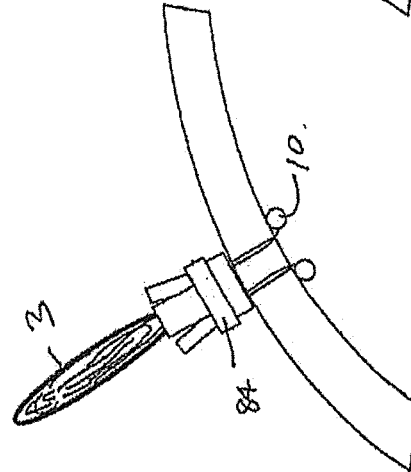


Fig. 29

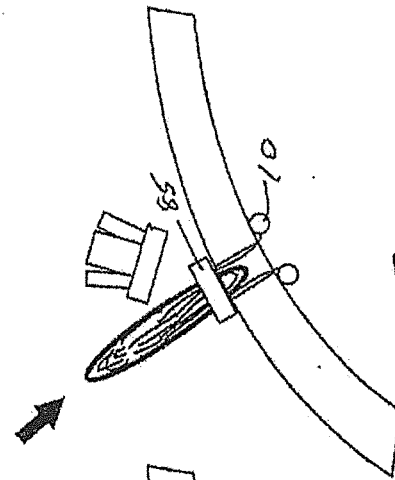


Fig. 30

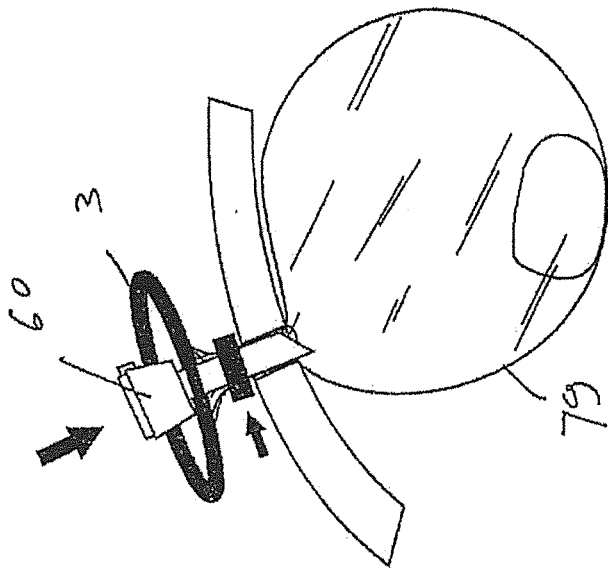


Fig. 33

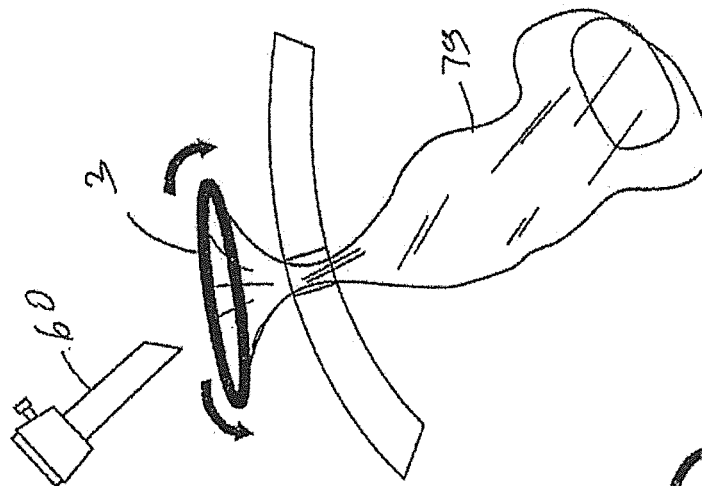


Fig. 32

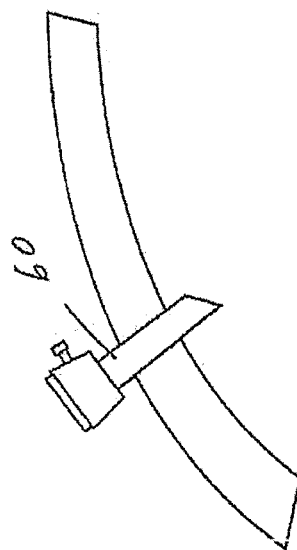
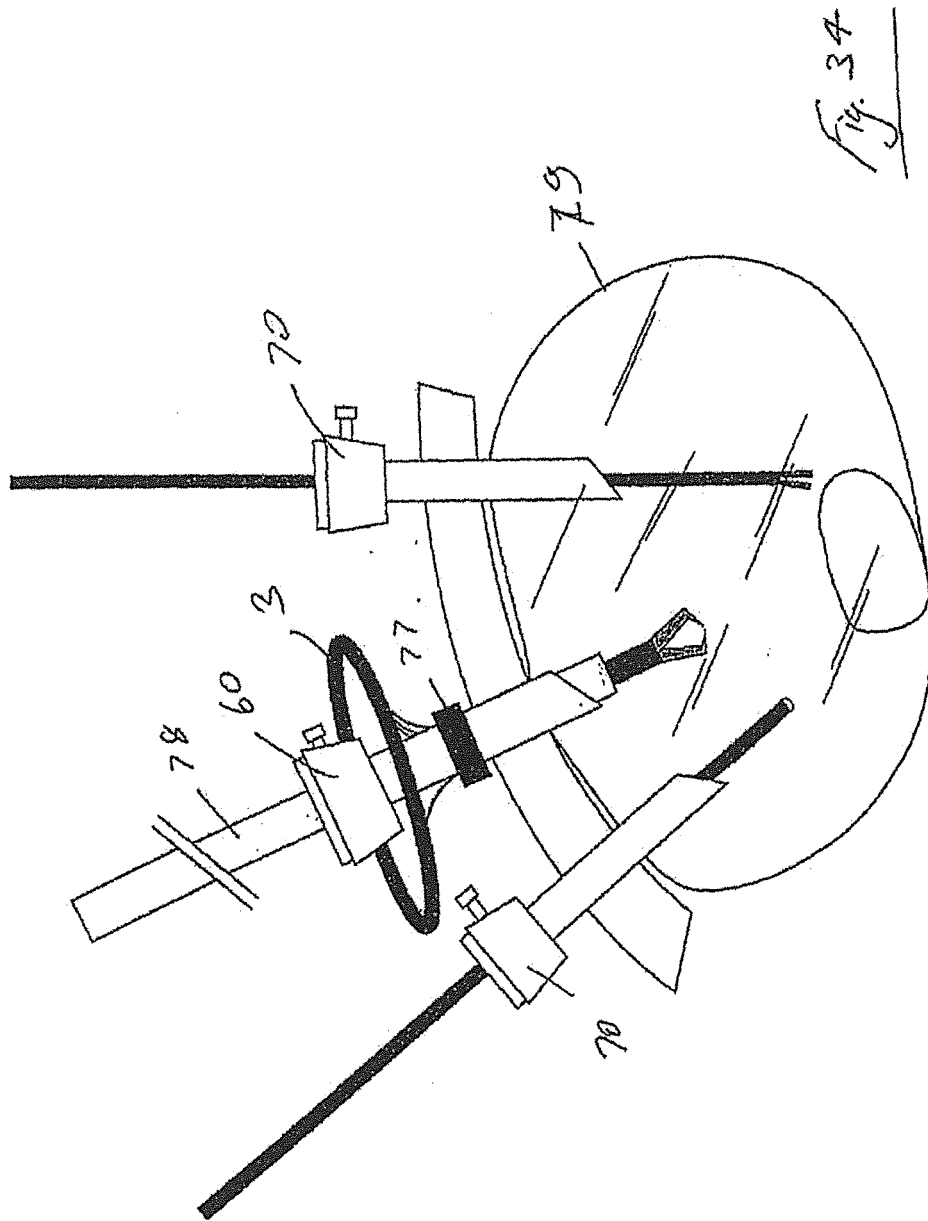
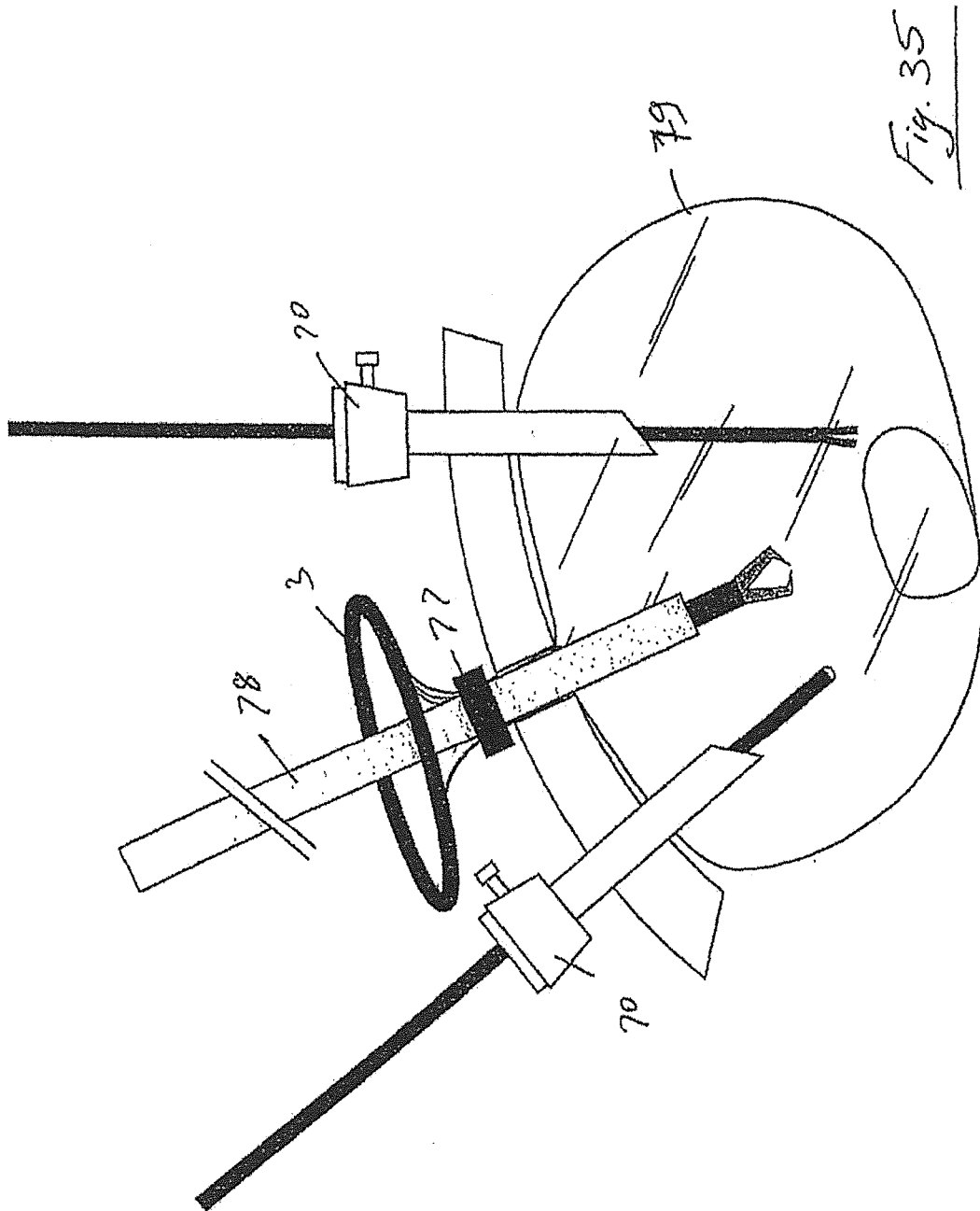


Fig. 31





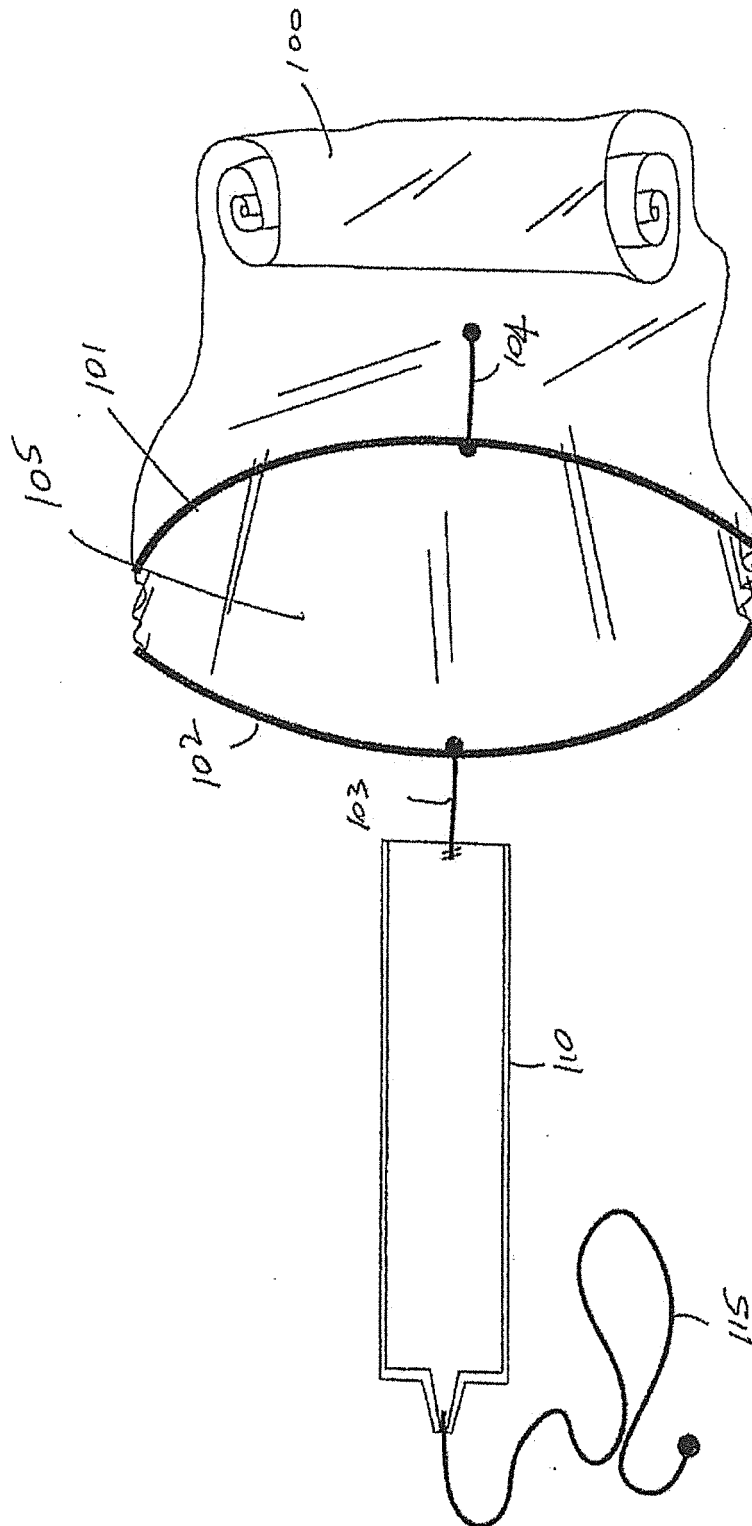


Fig. 36

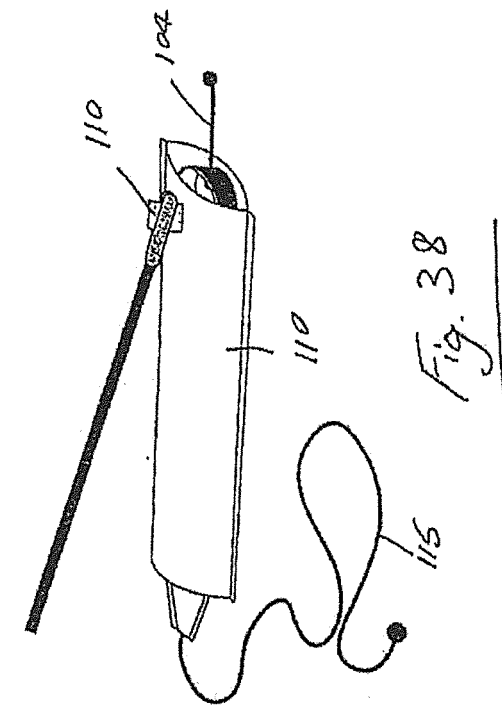


Fig. 37

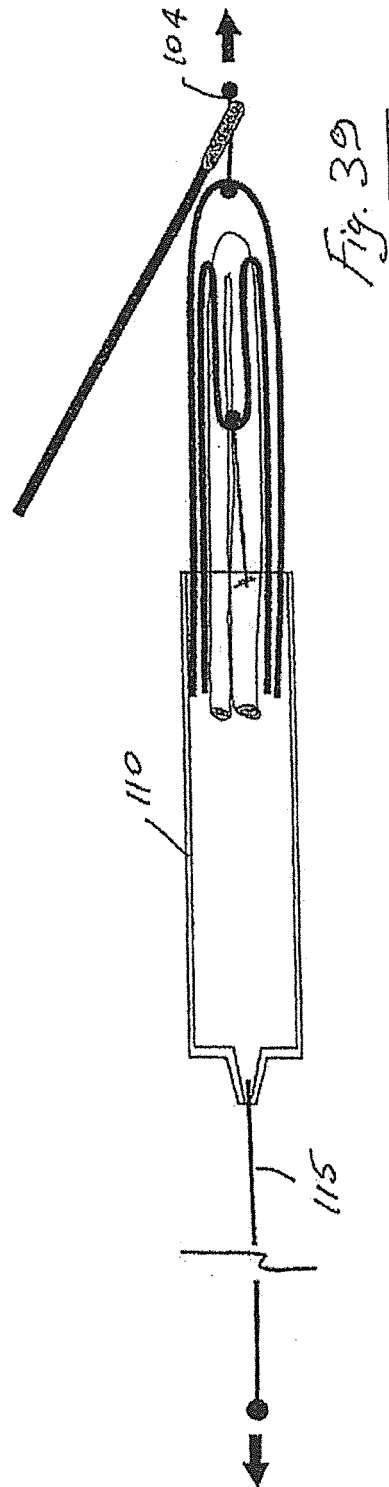
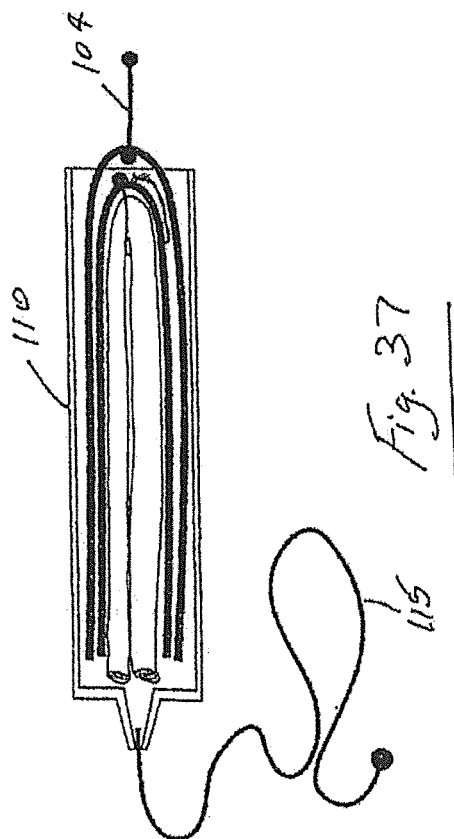
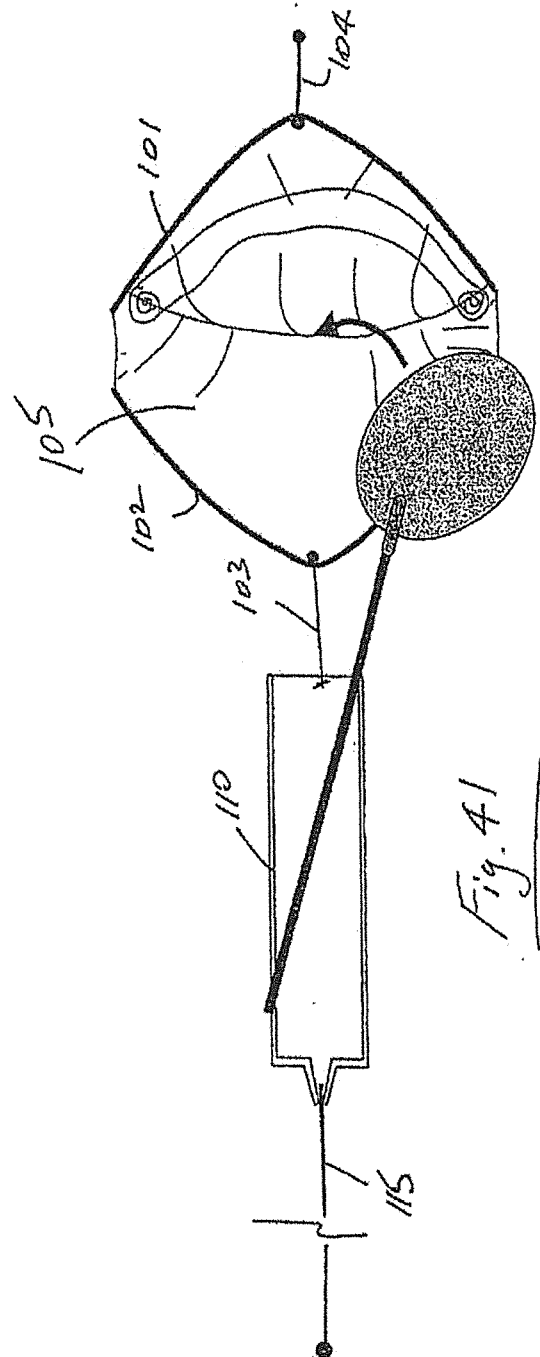
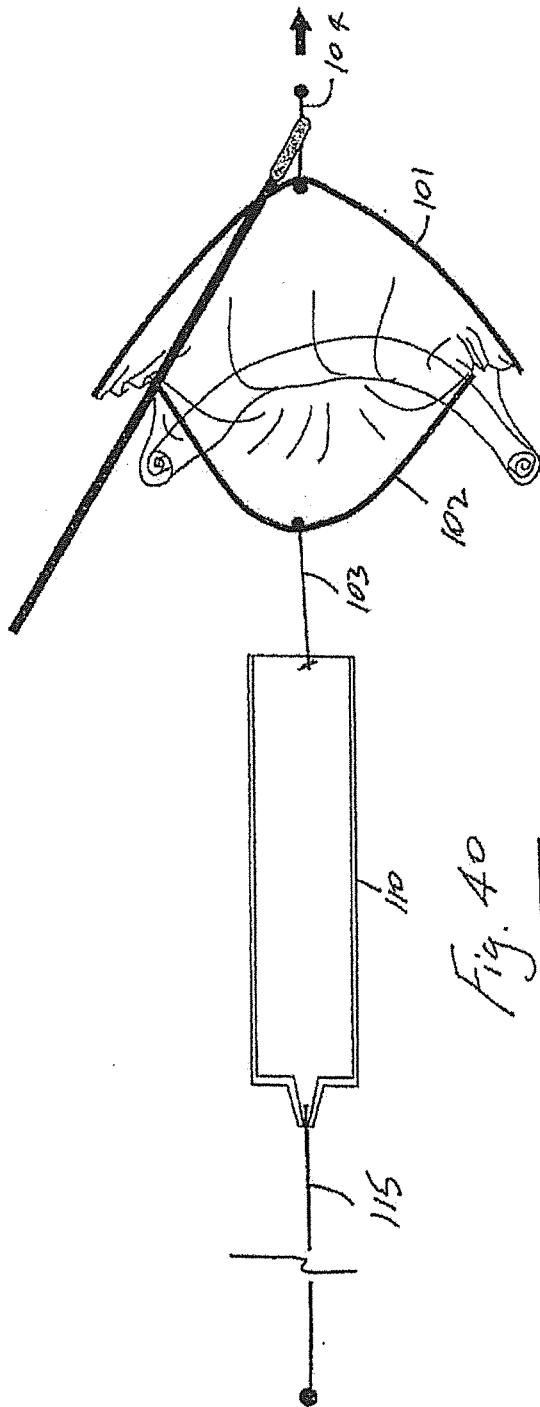
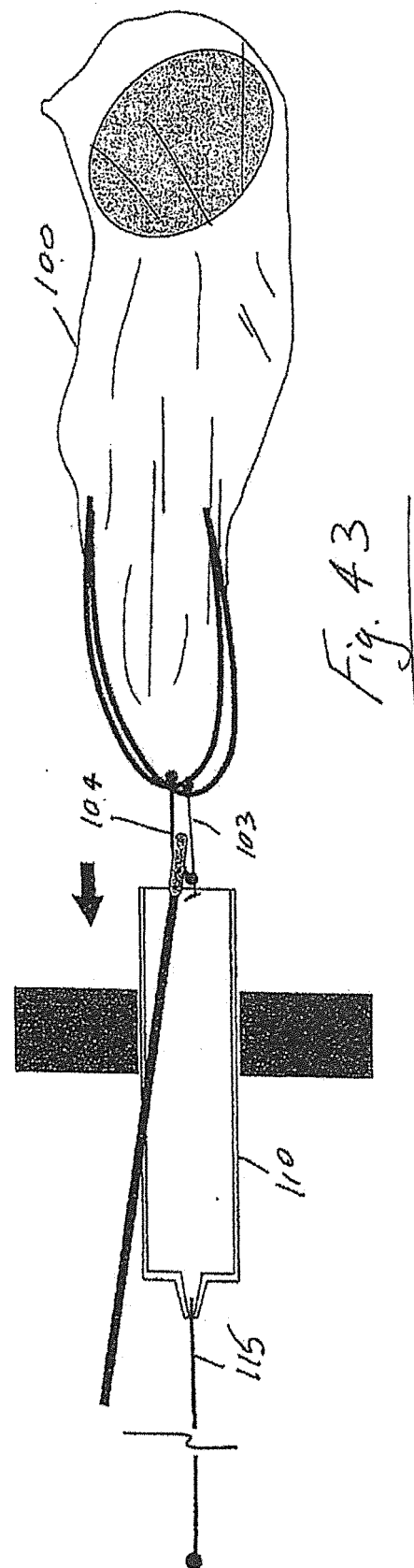
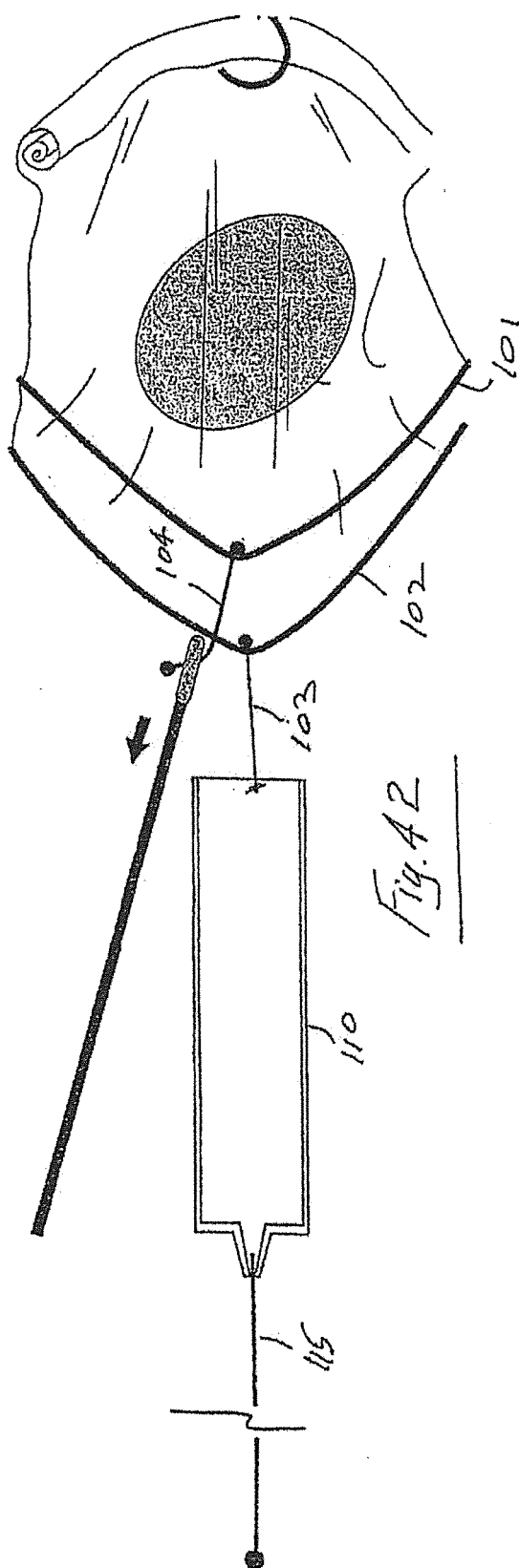
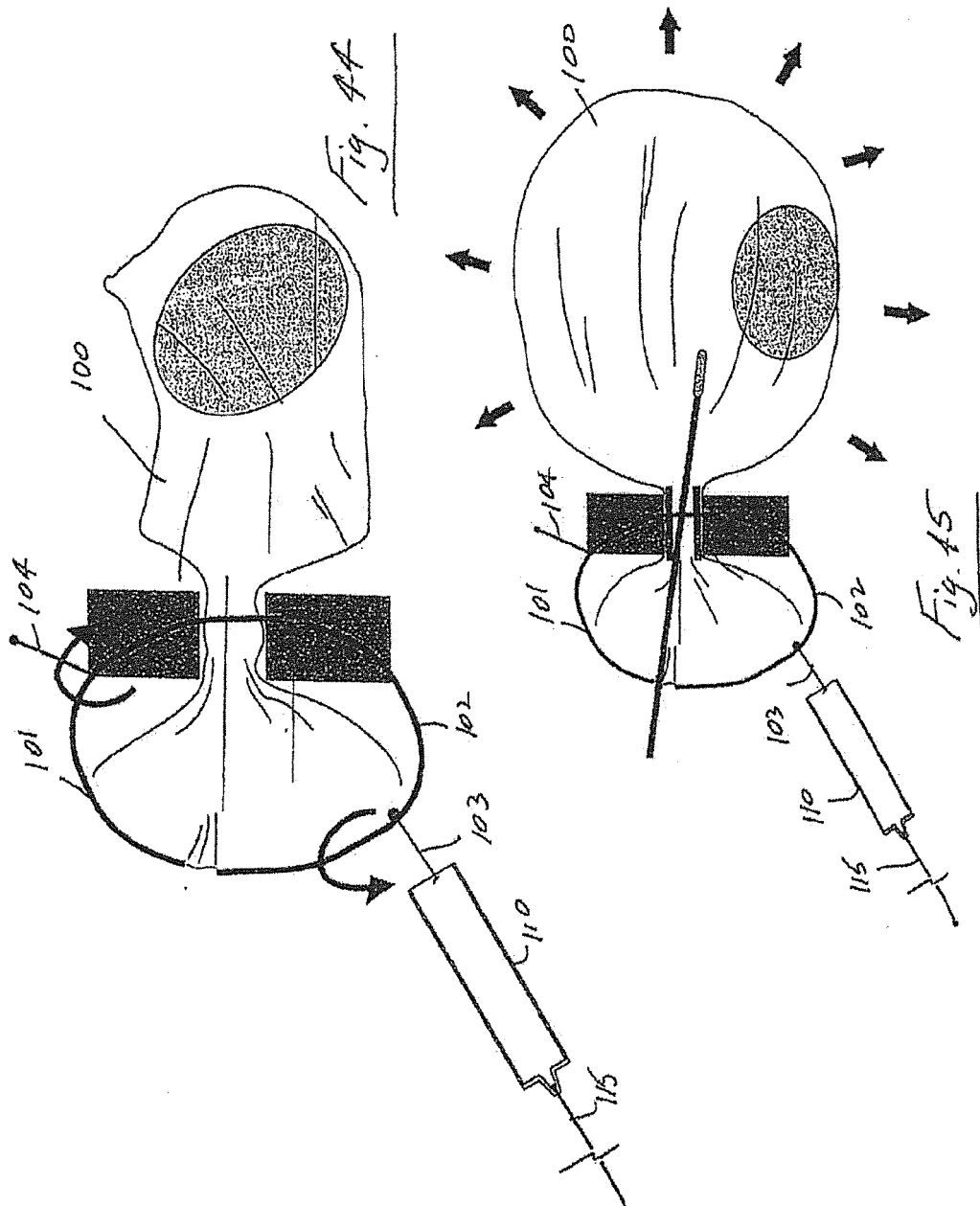


Fig. 38









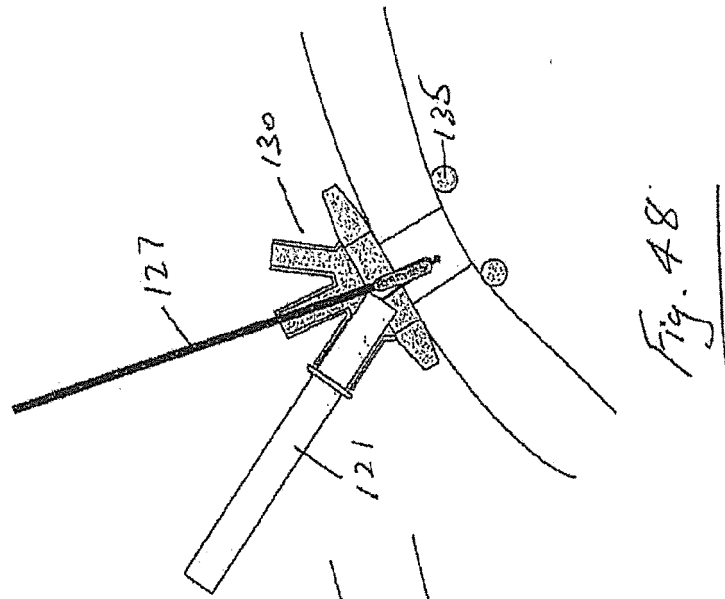


Fig. 46

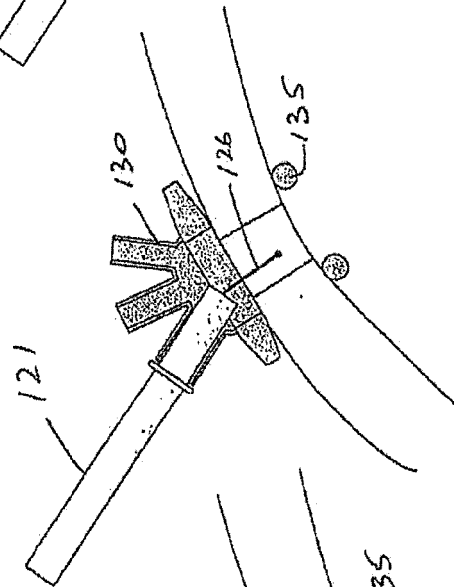


Fig. 47

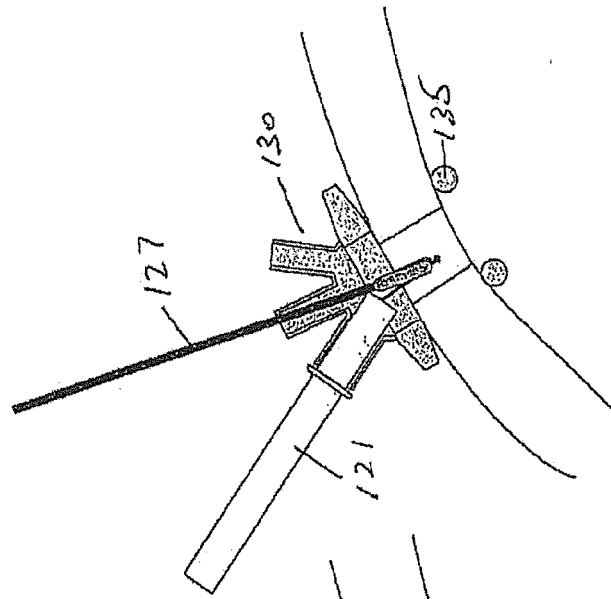
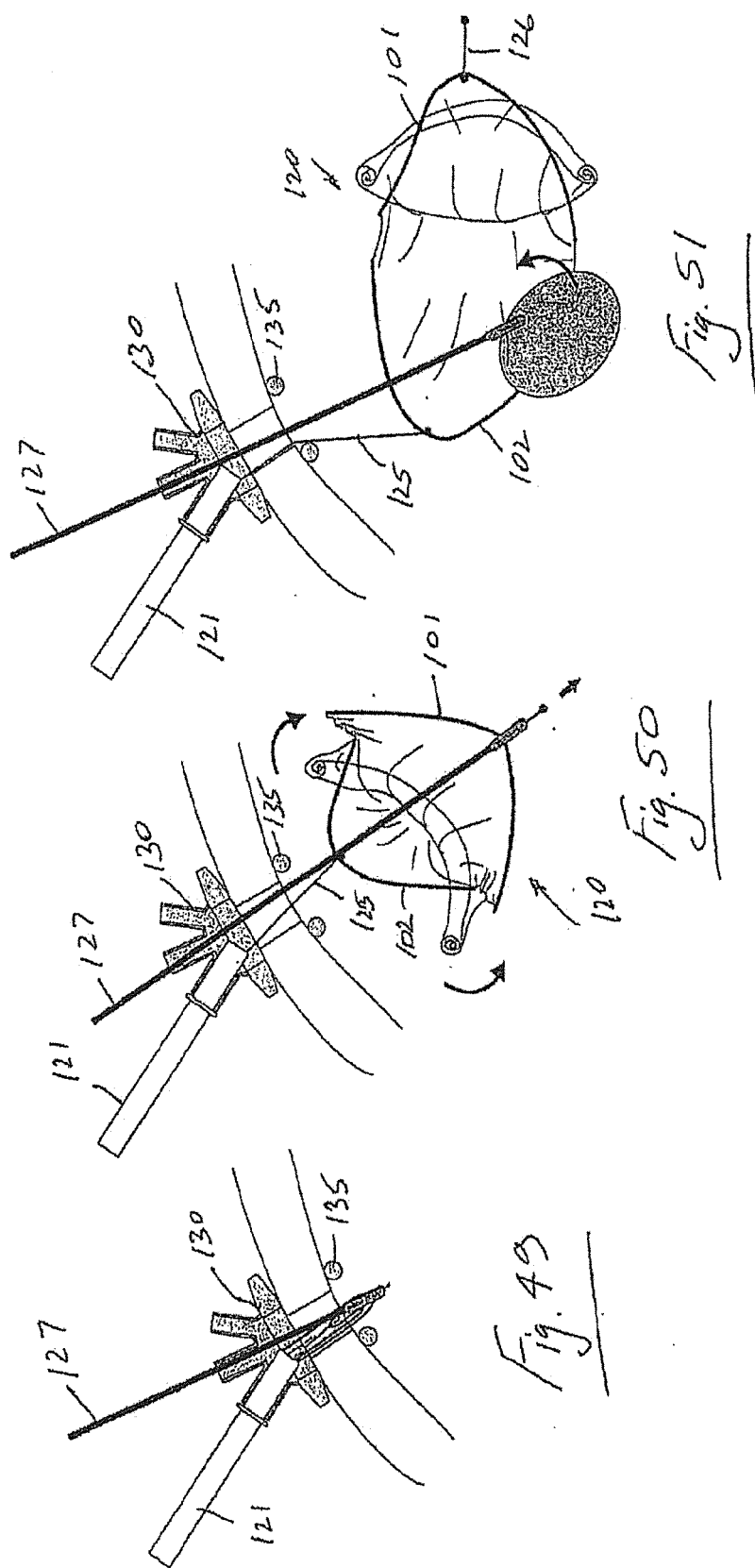
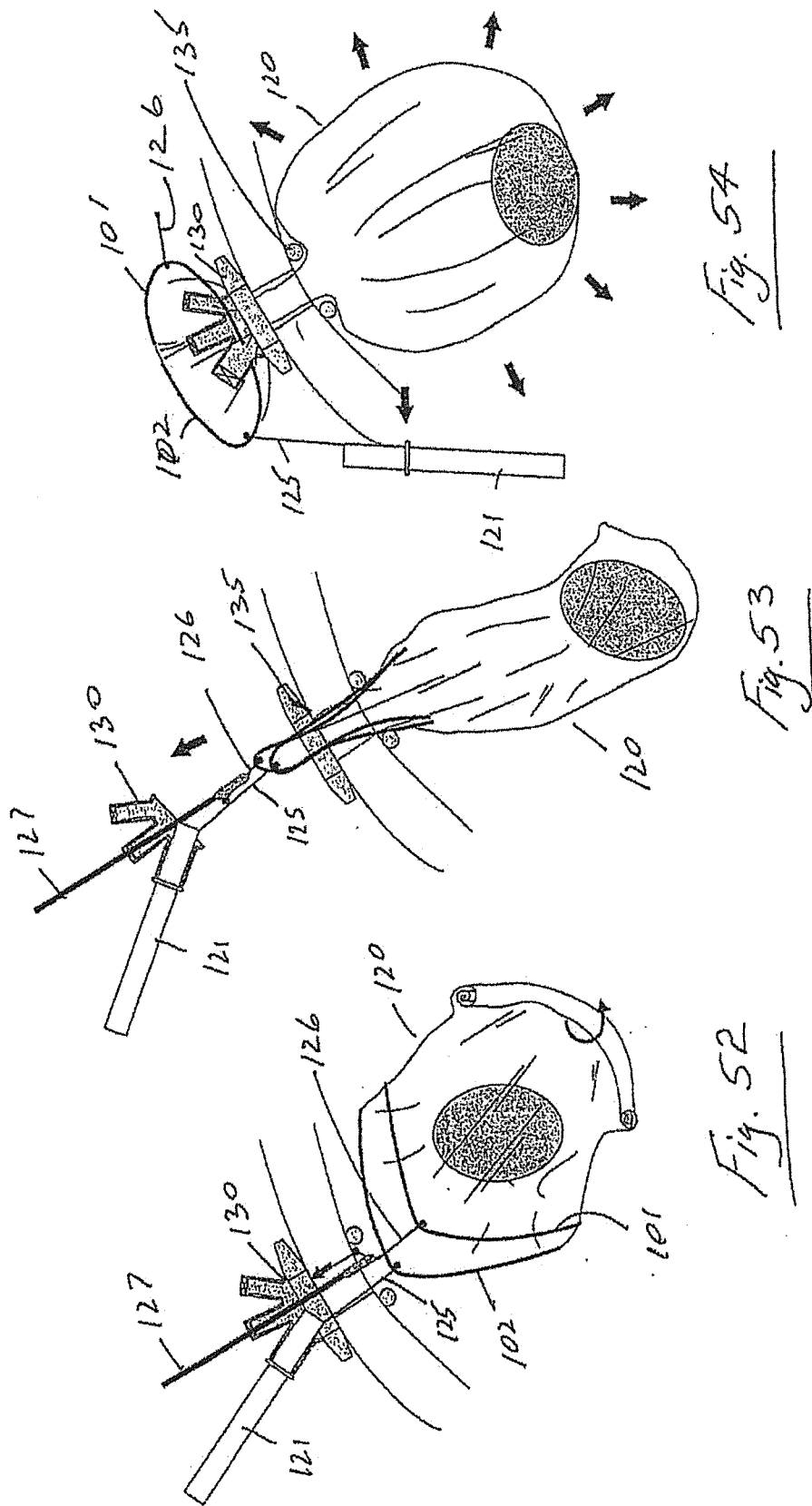
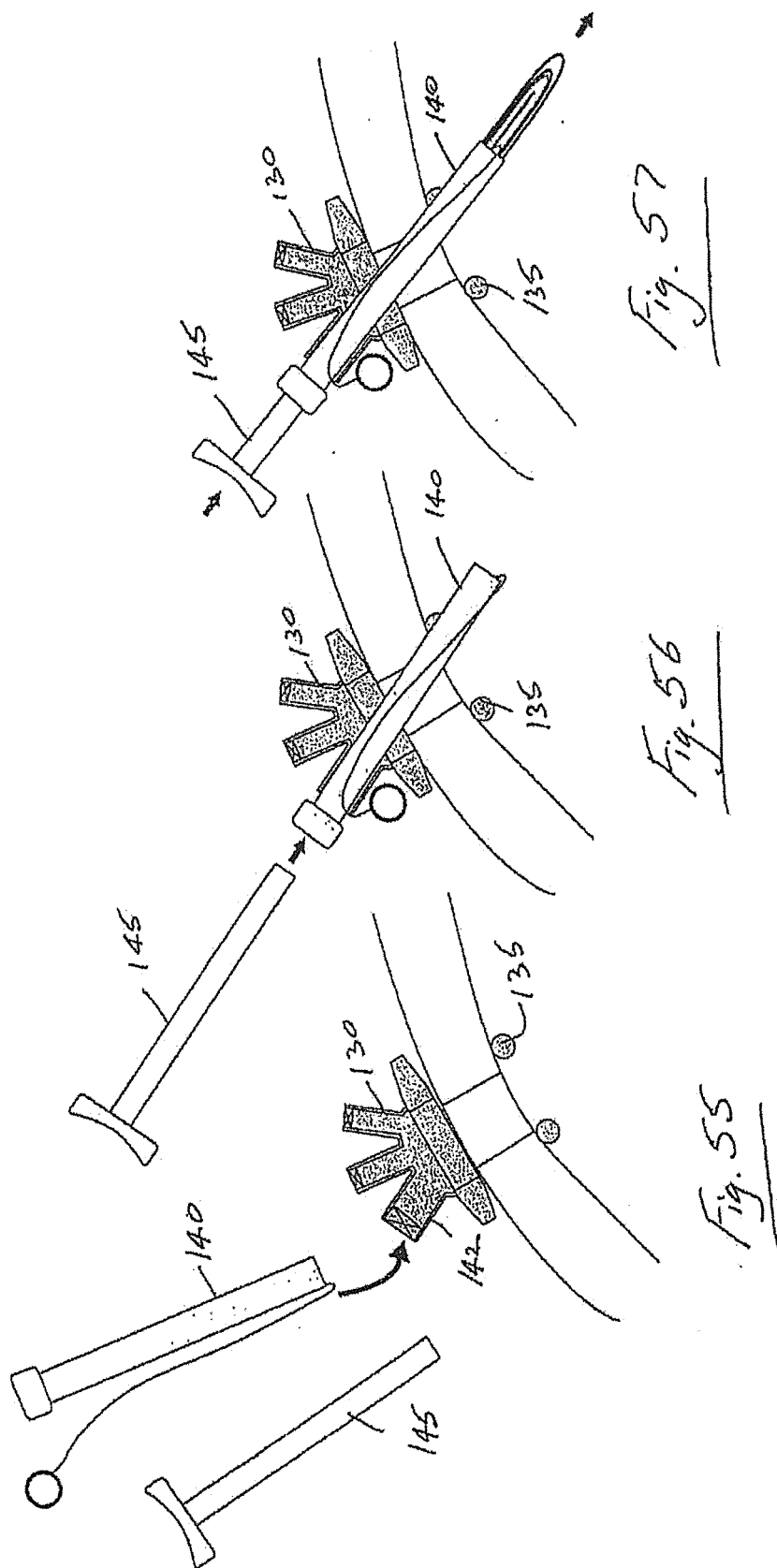


Fig. 48







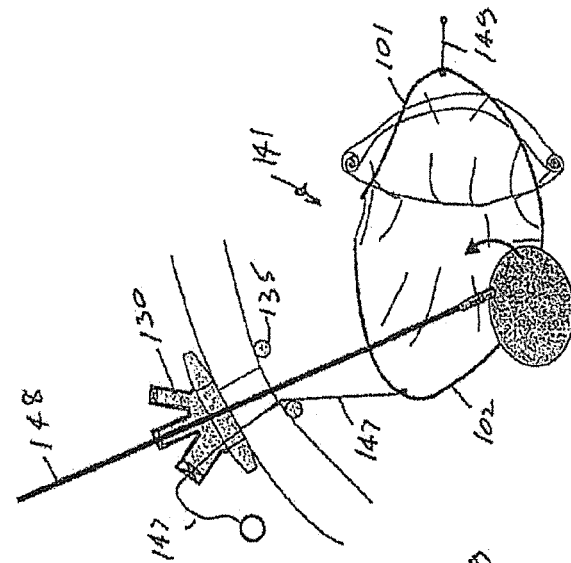


Fig. 58

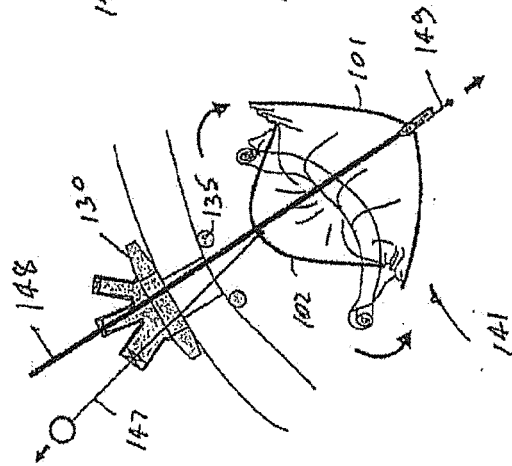


Fig. 59

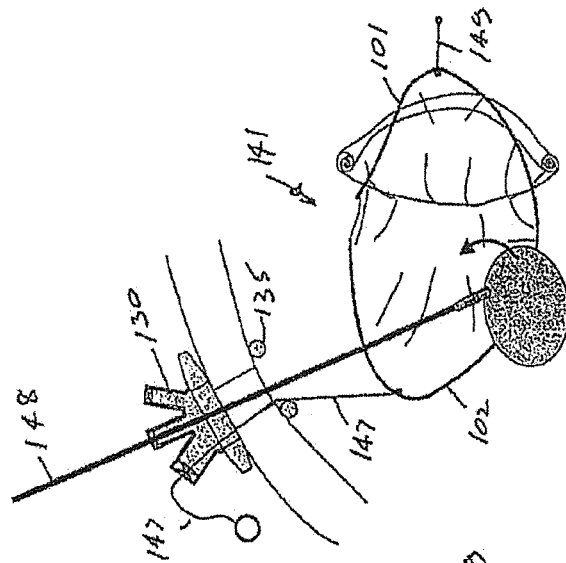


Fig. 60

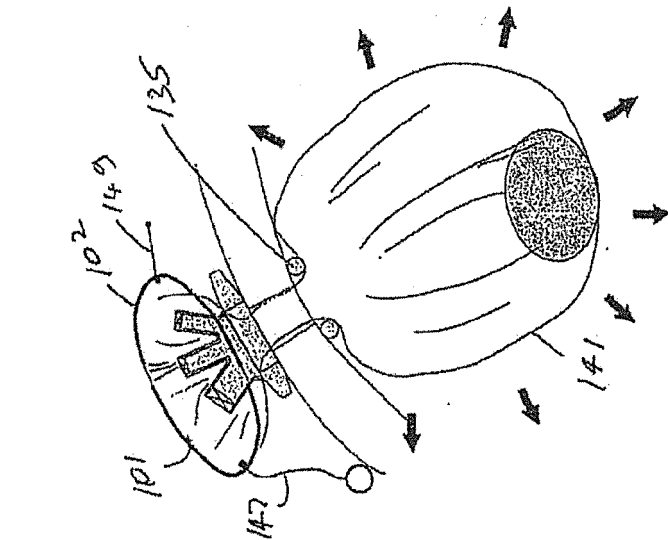


Fig. 61

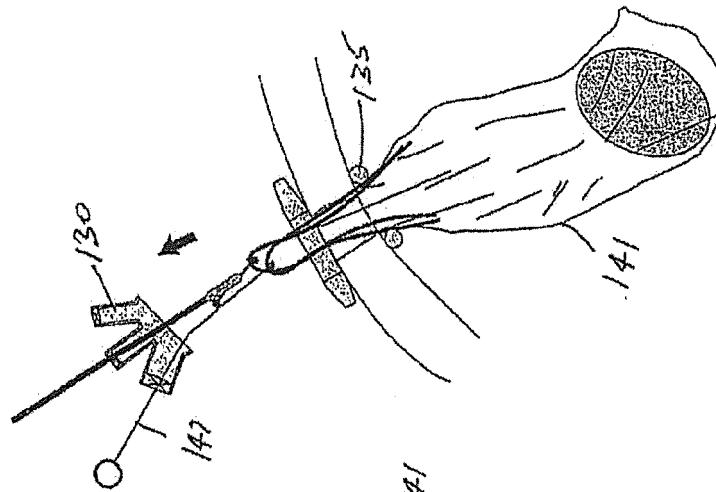


Fig. 62

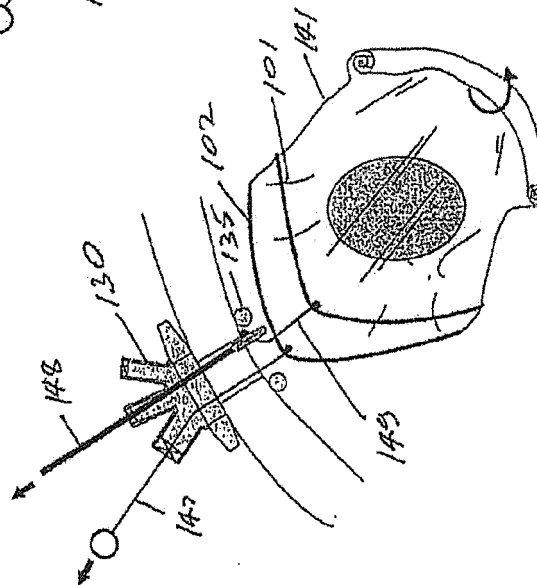
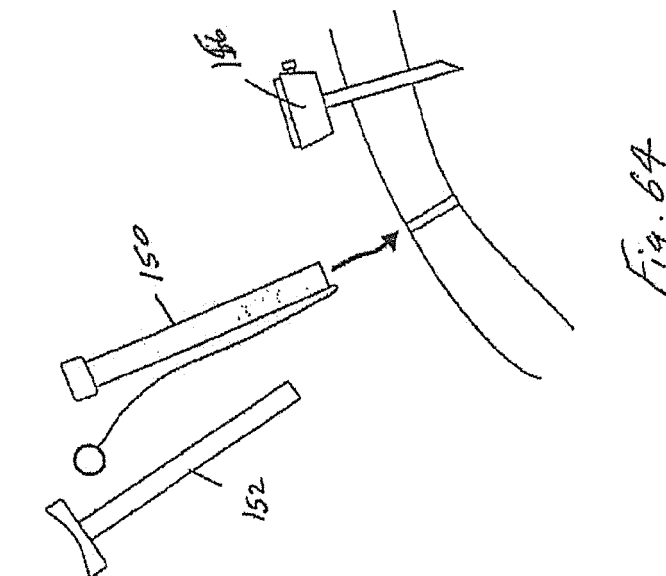
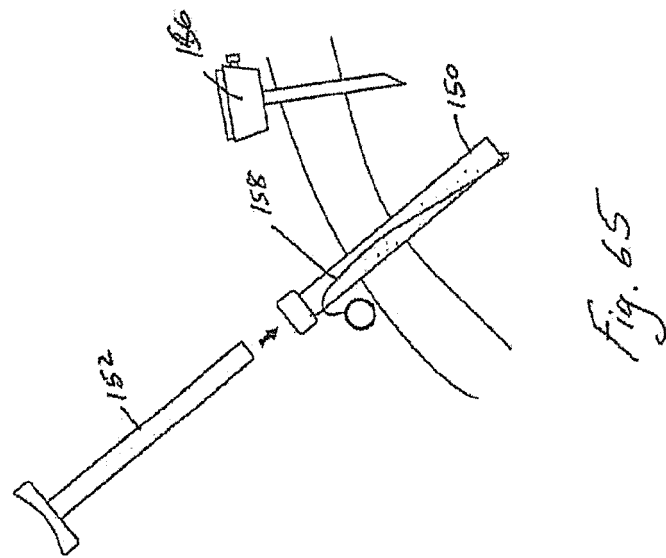
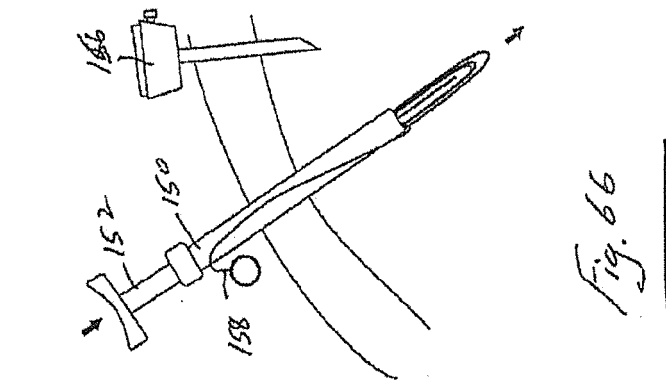


Fig. 63



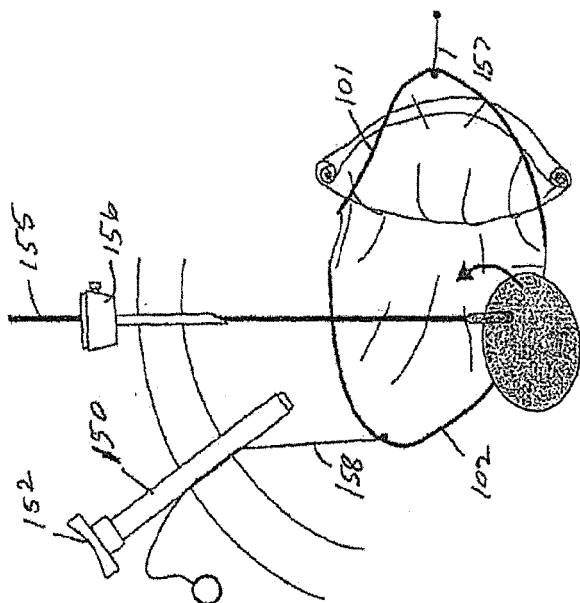


Fig. 67

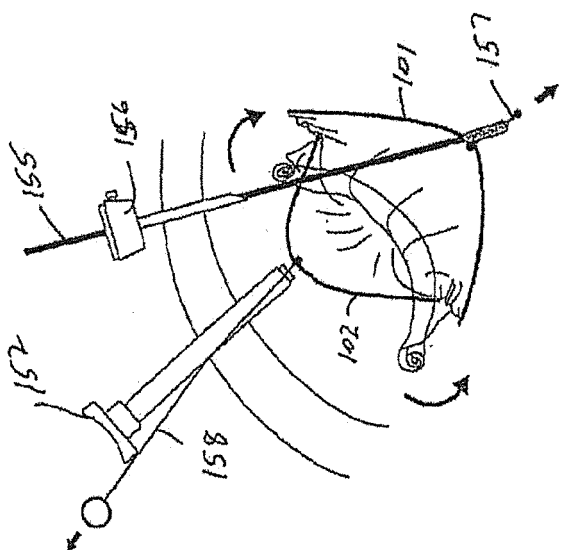


Fig. 68

Fig. 69

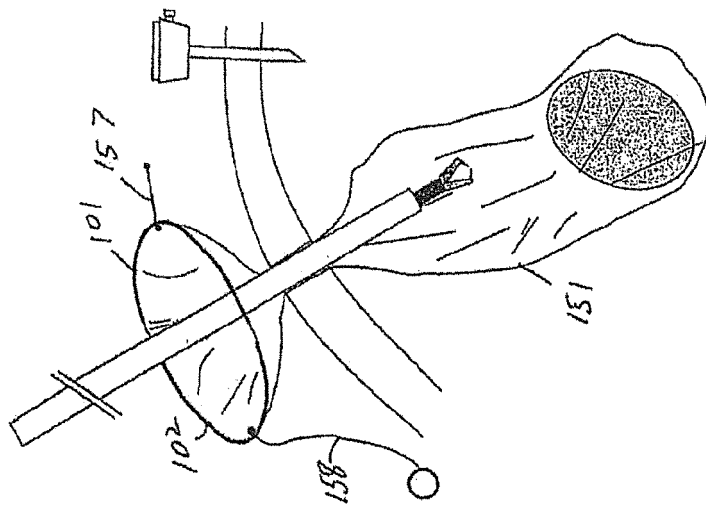


Fig. 72

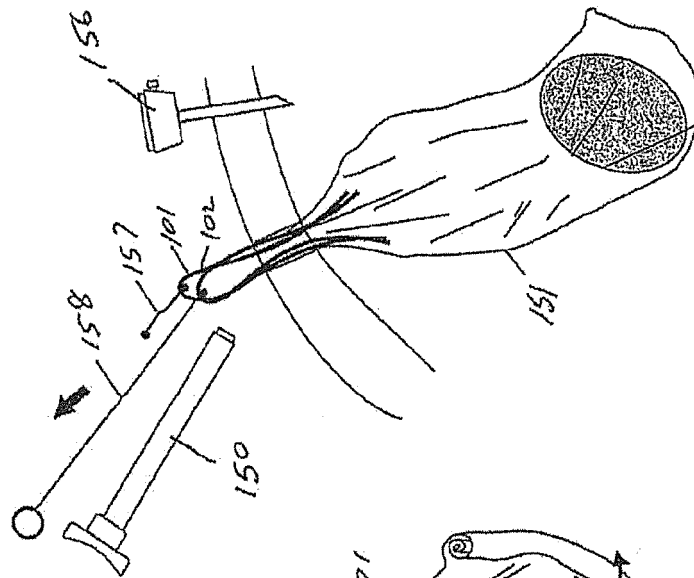


Fig. 71

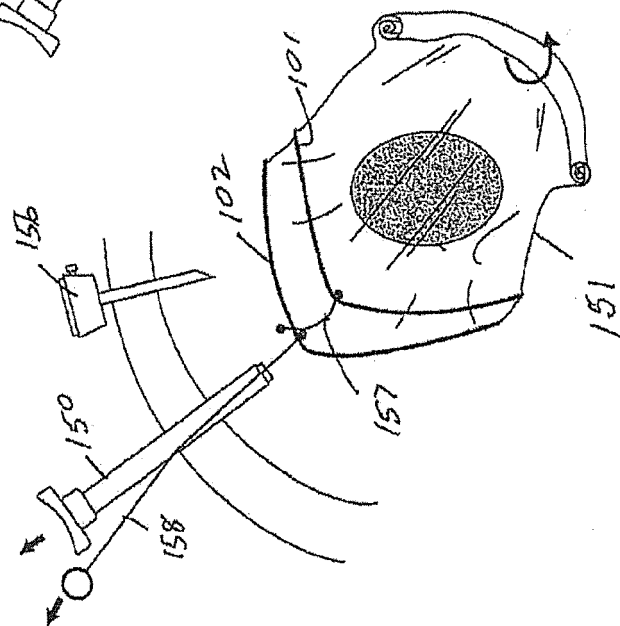


Fig. 70

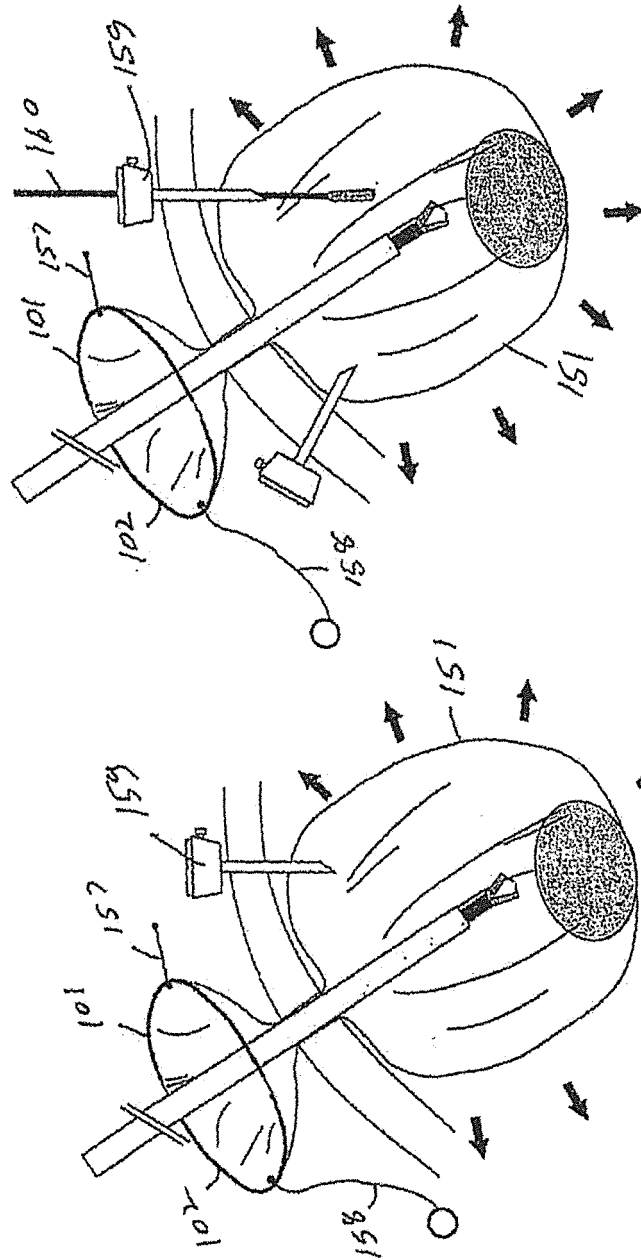


Fig. 74

Fig. 73

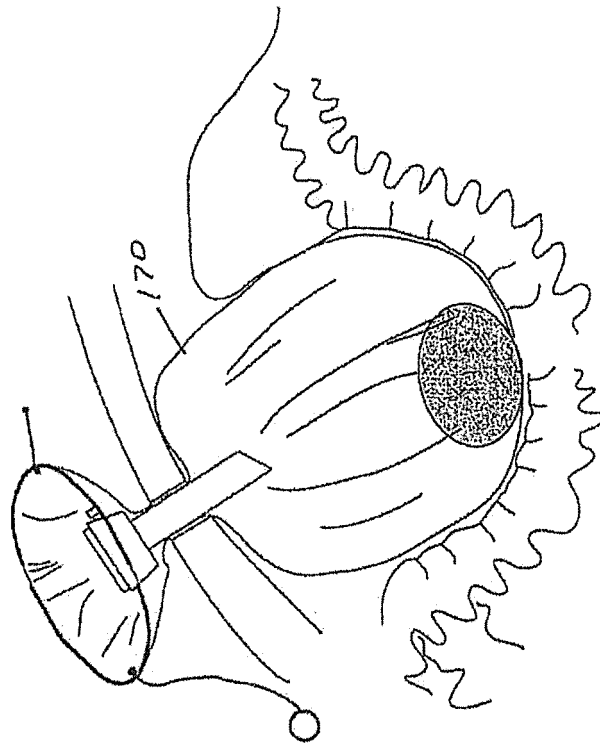


Fig. 76

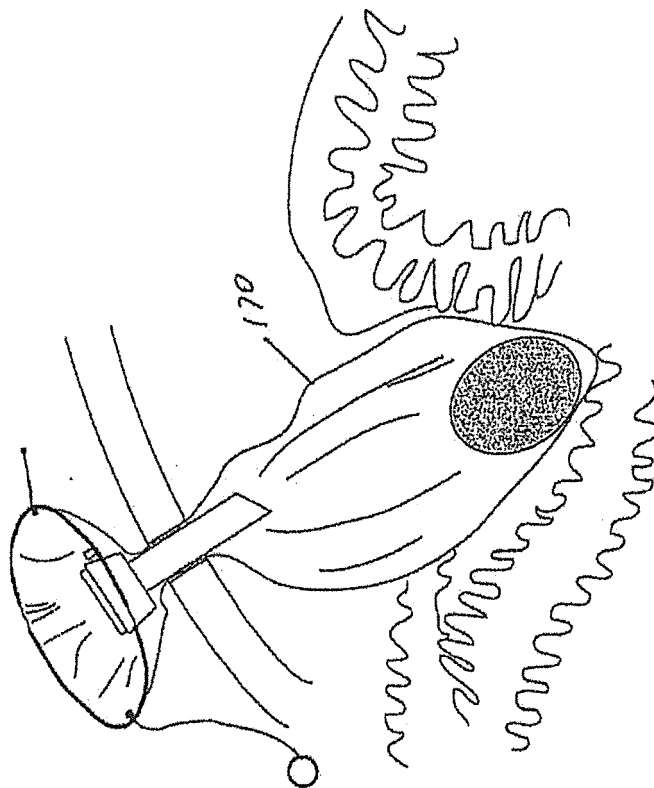


Fig. 75

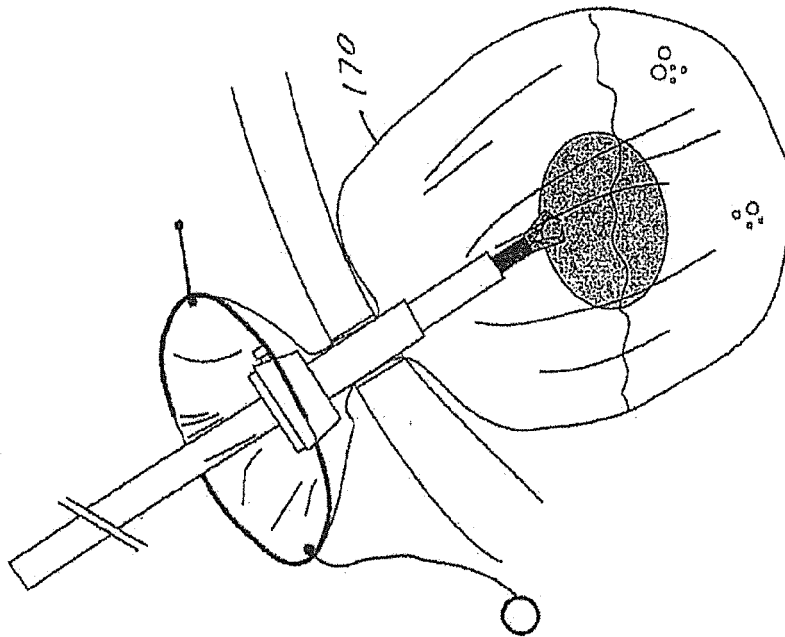


Fig. 78

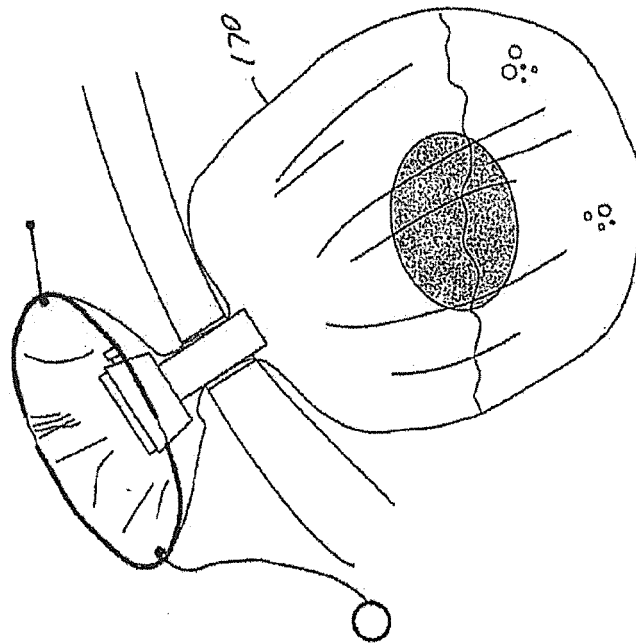


Fig. 77

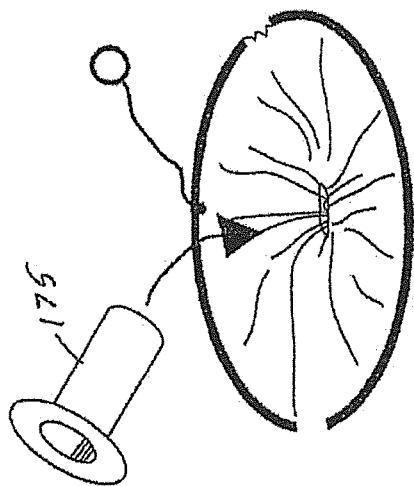


Fig. 79

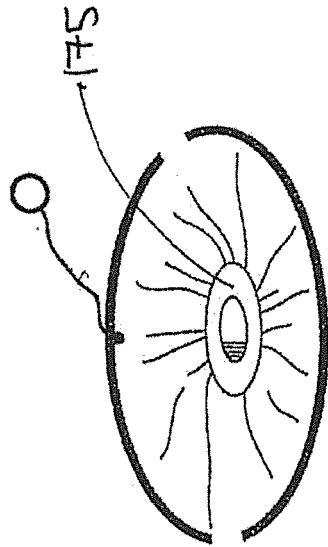


Fig. 80

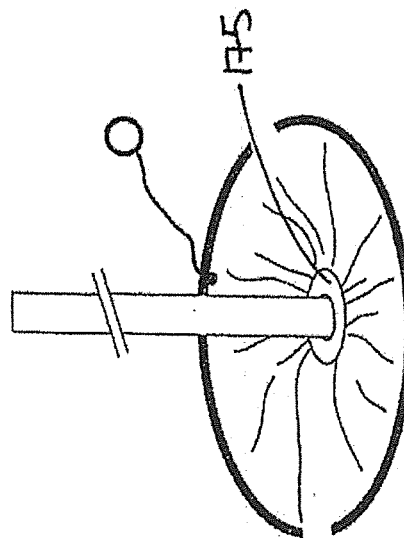


Fig. 81

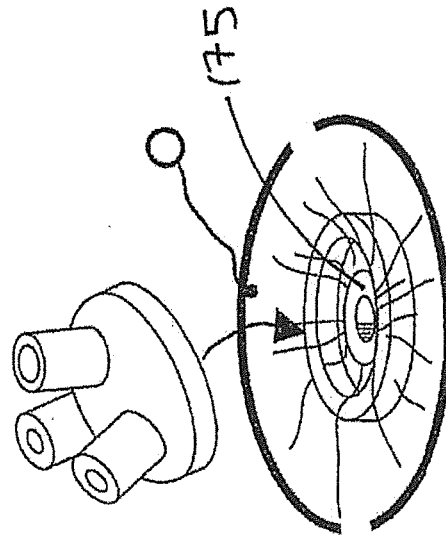


Fig. 82

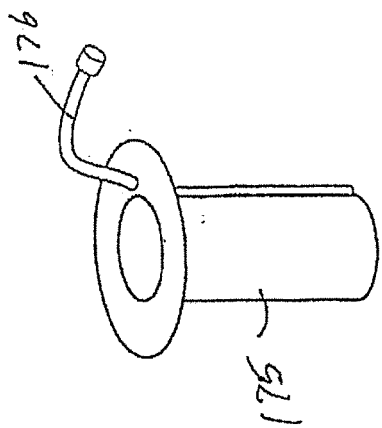


Fig. 83

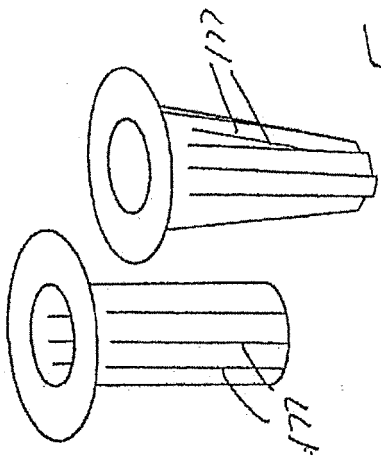


Fig. 84

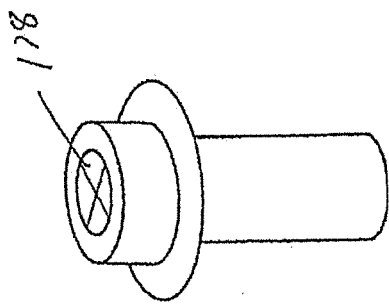


Fig. 85

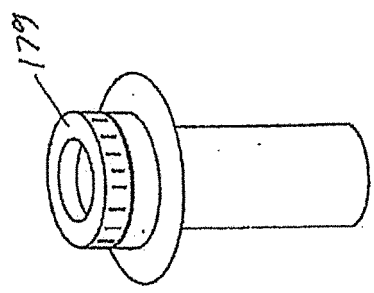


Fig. 86

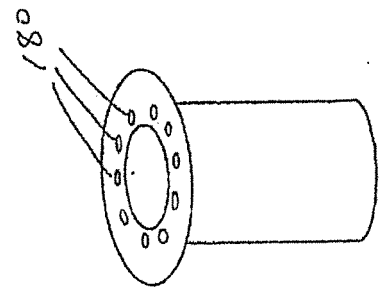
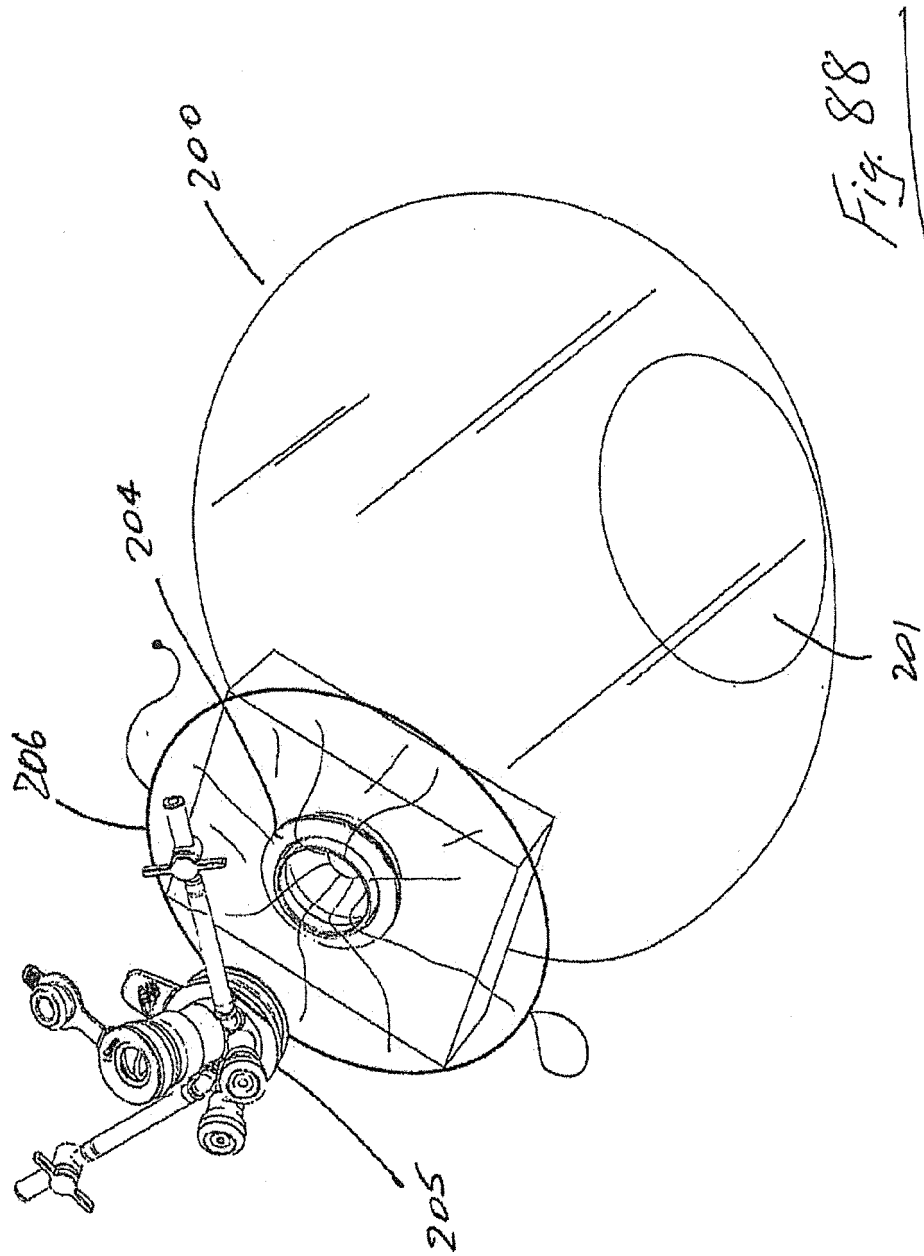
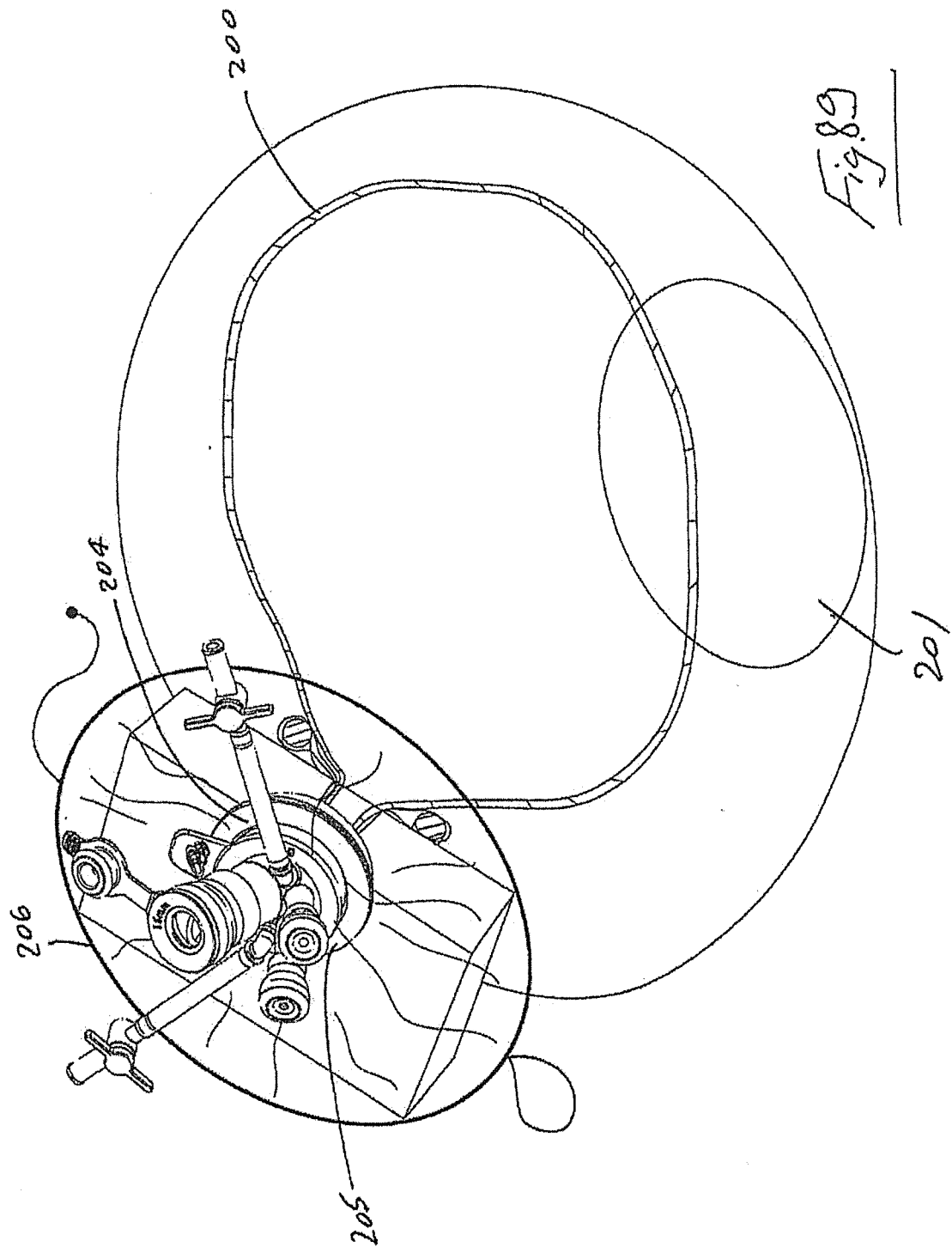
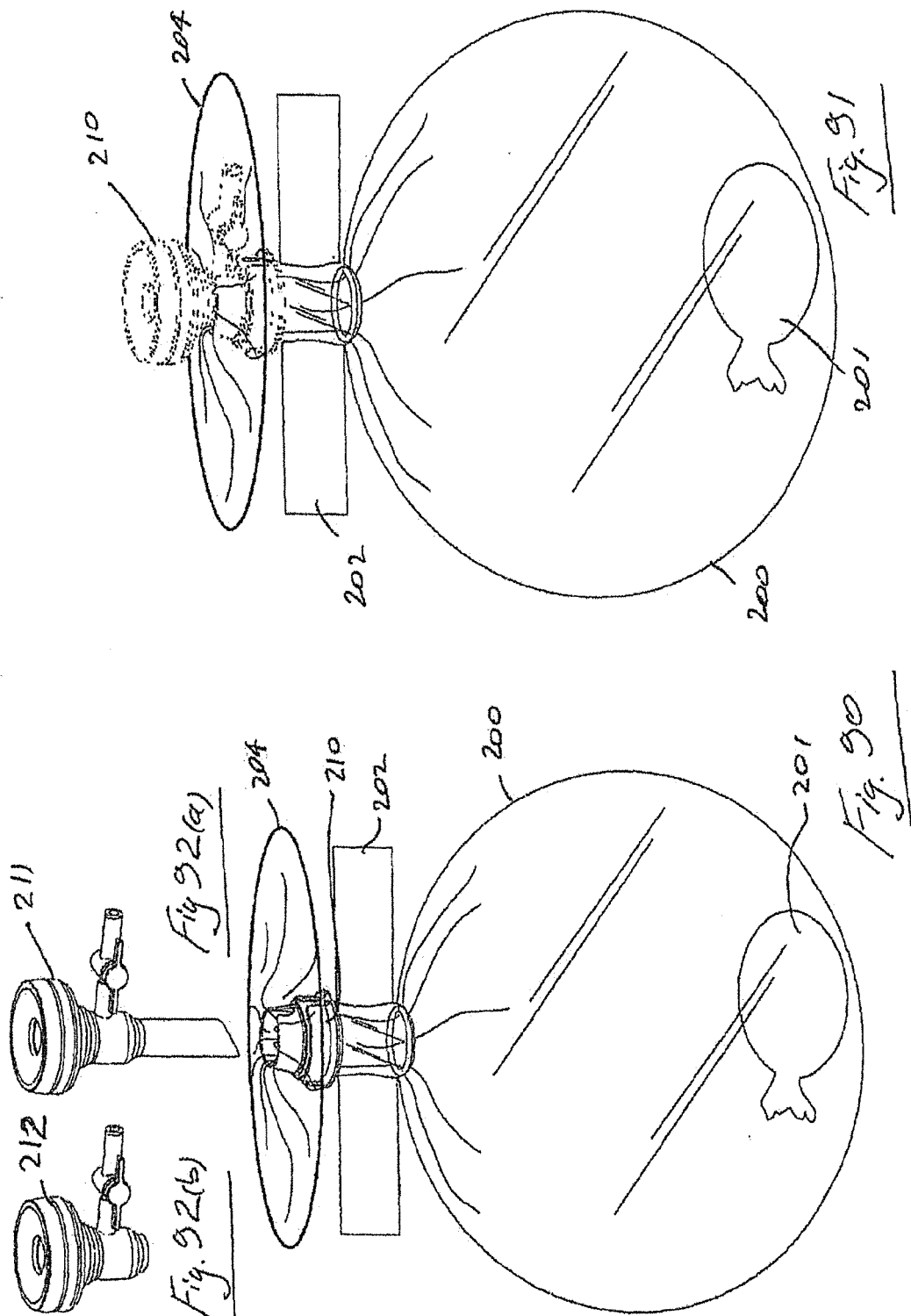
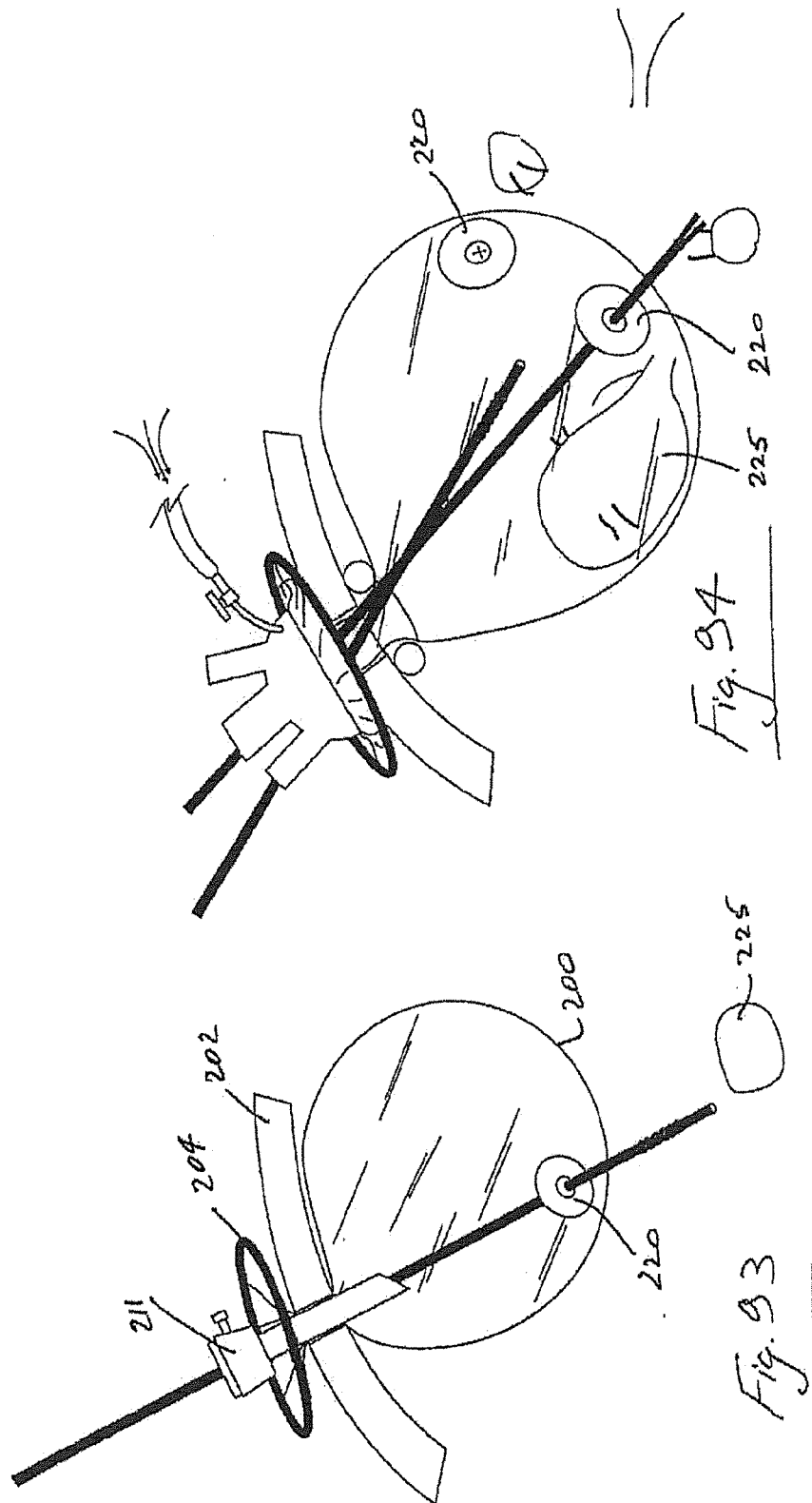


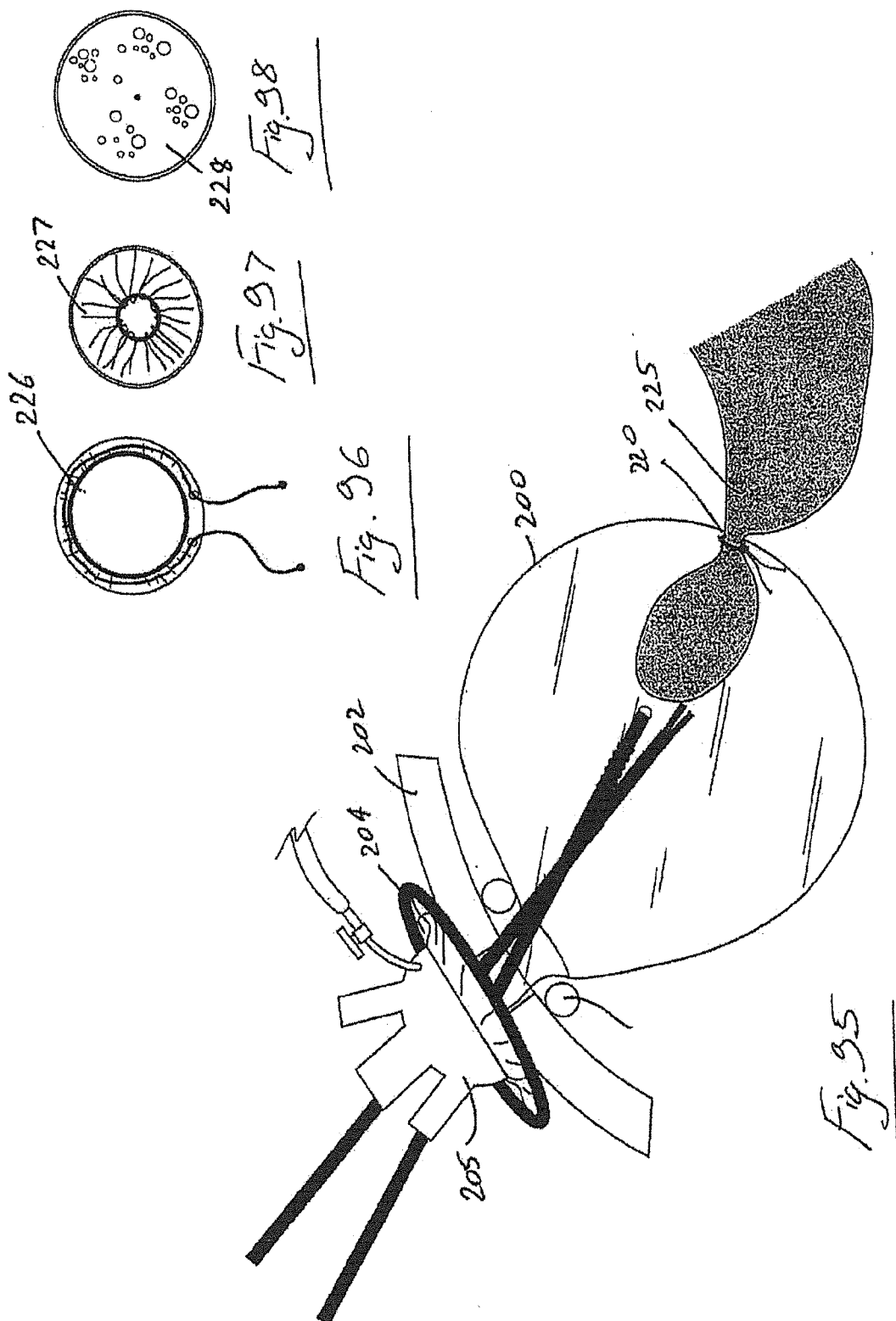
Fig. 87

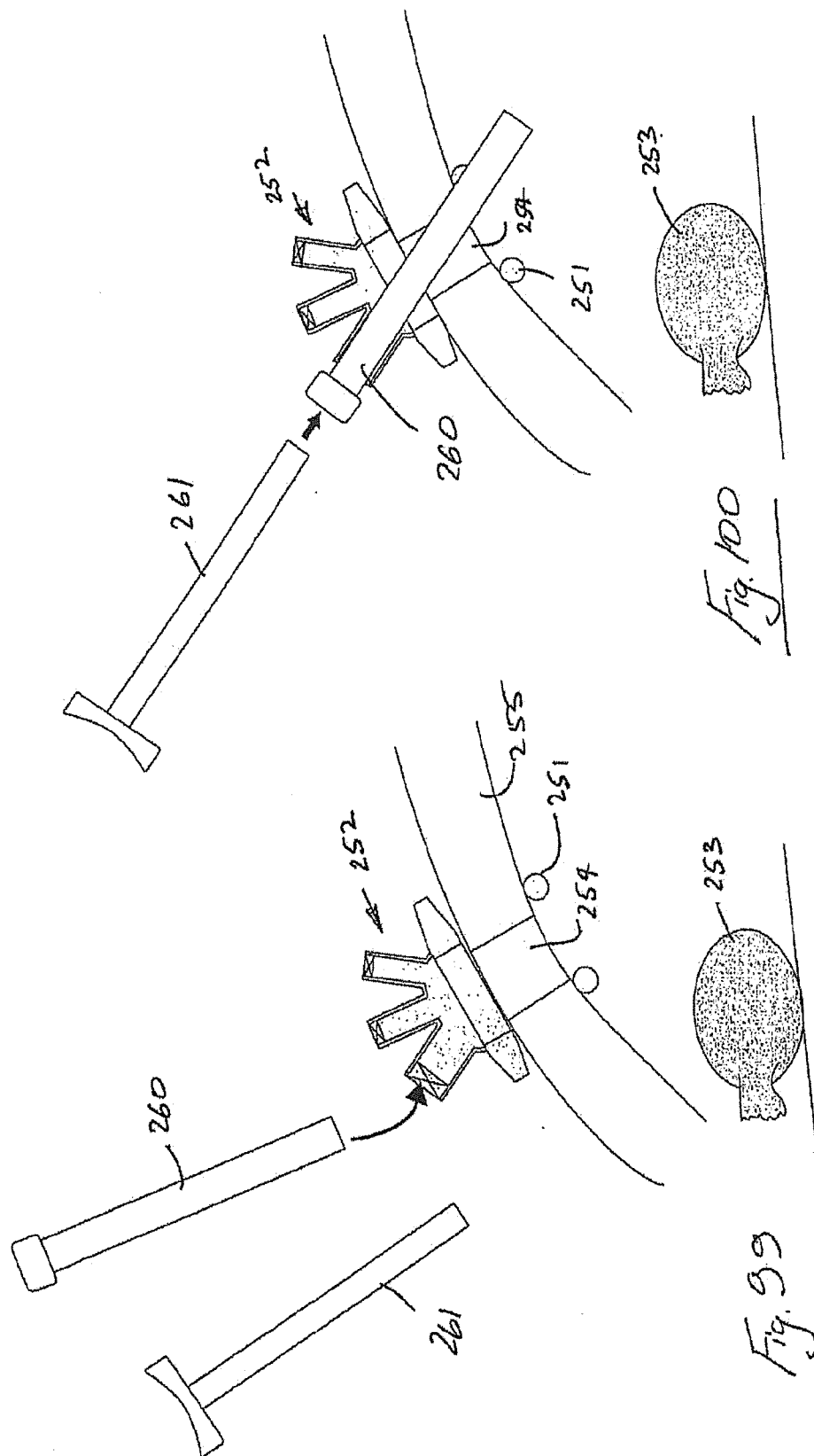


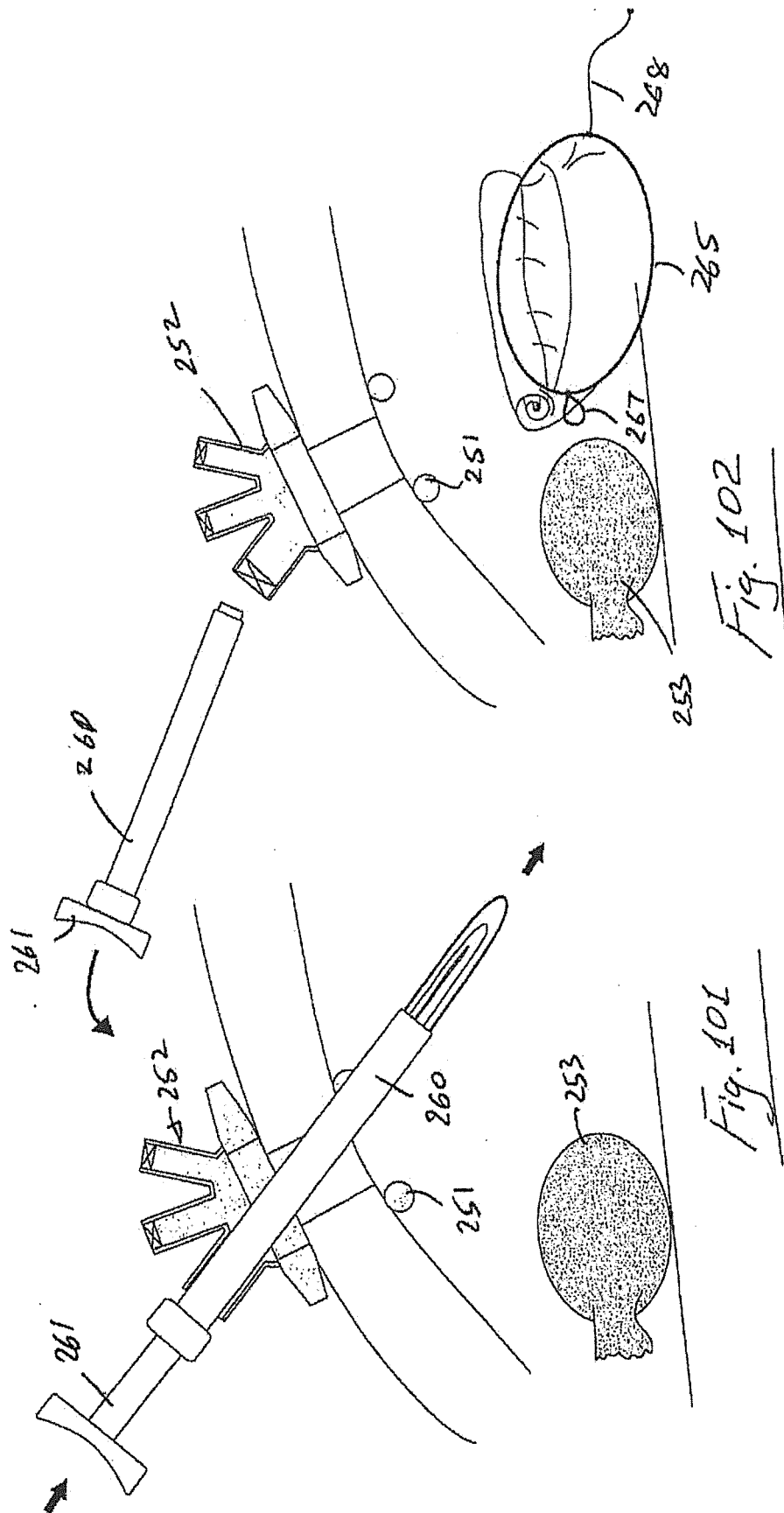


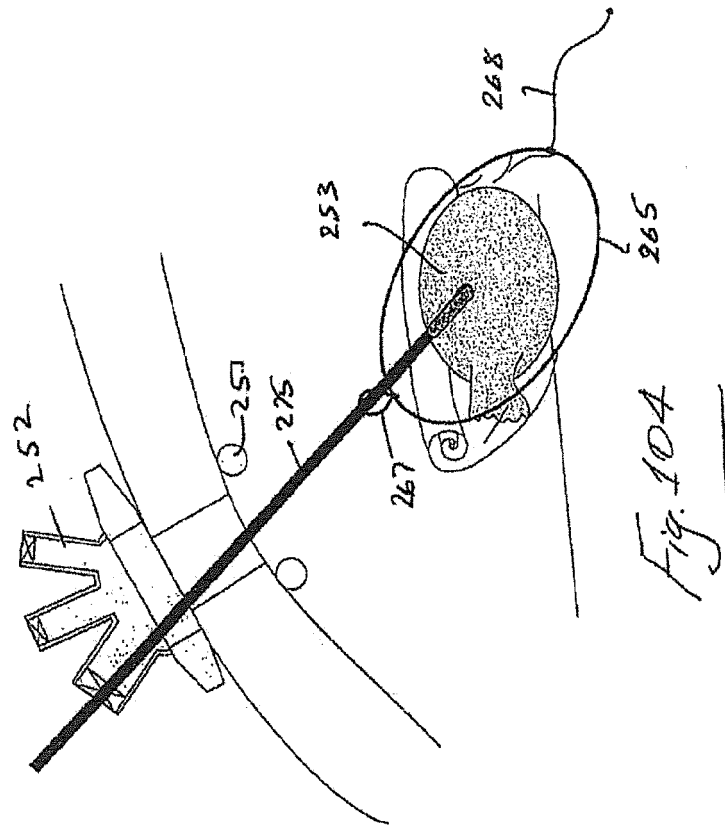
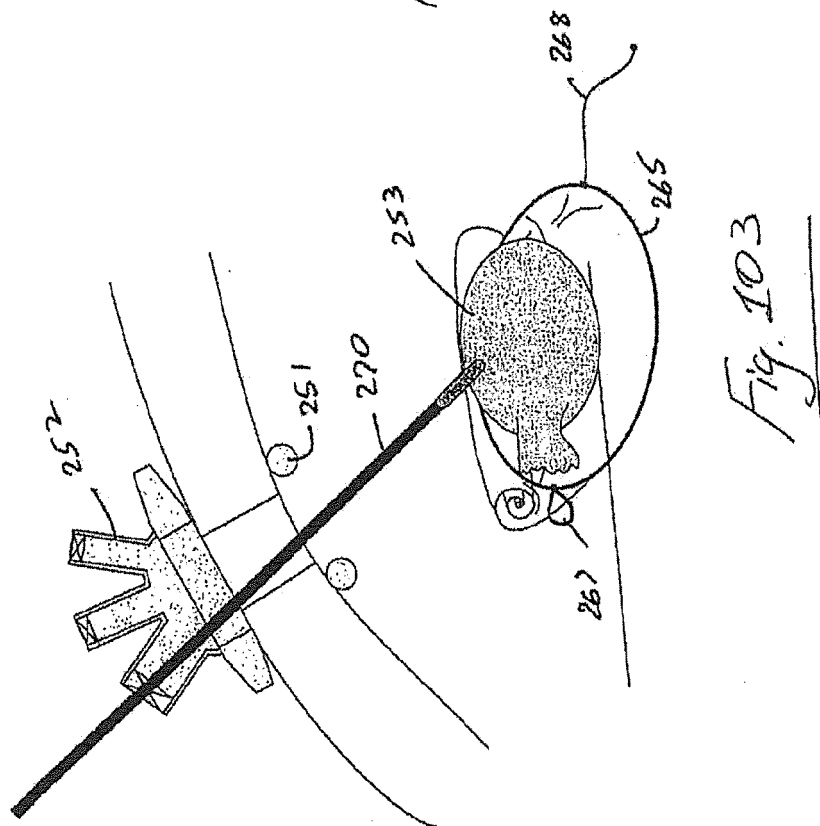












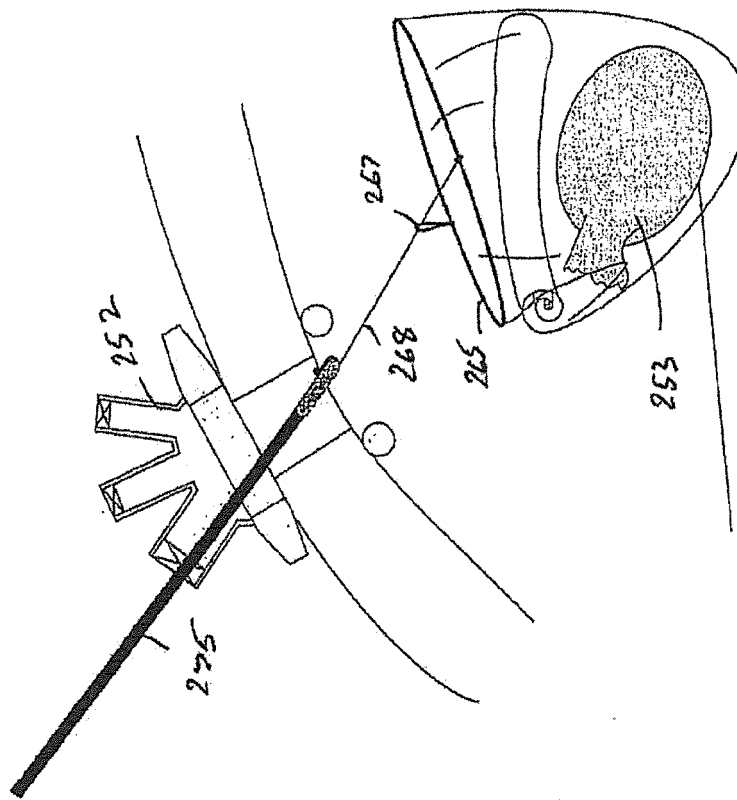


Fig. 105

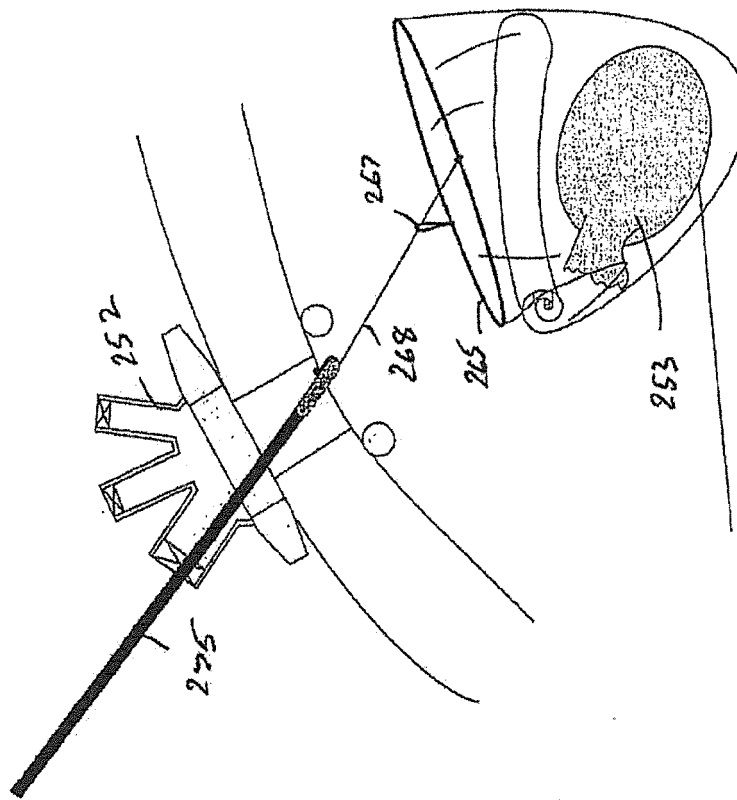
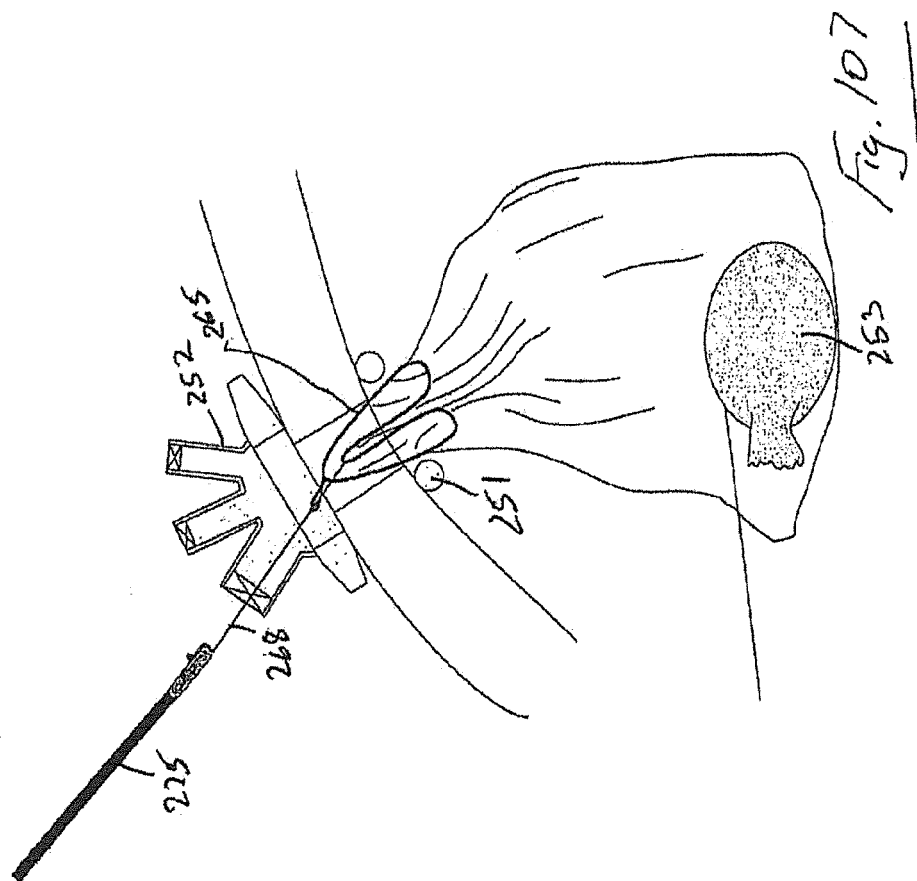
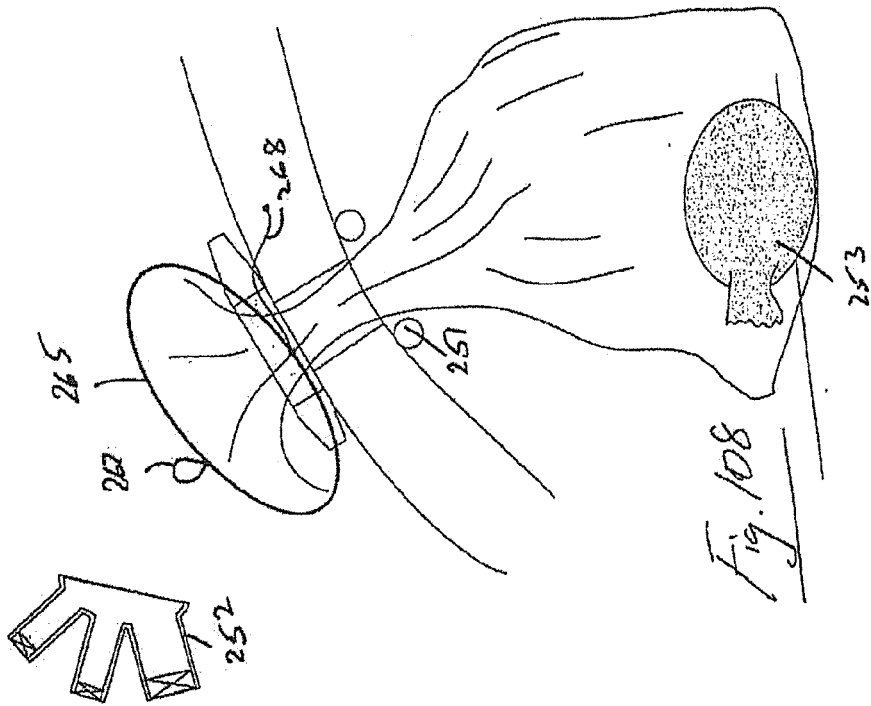
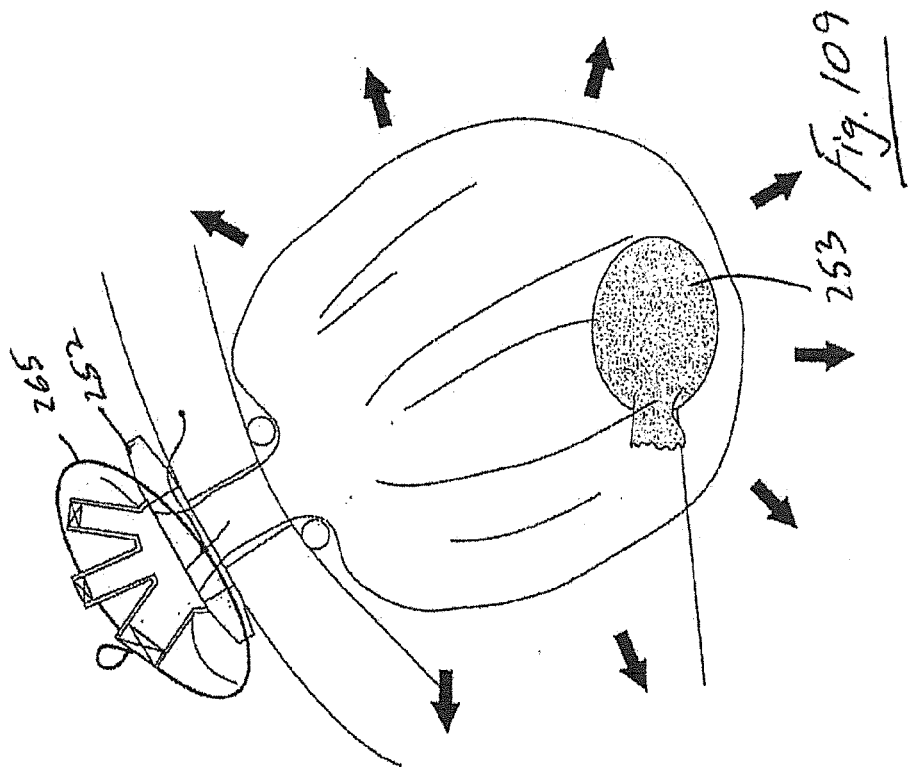
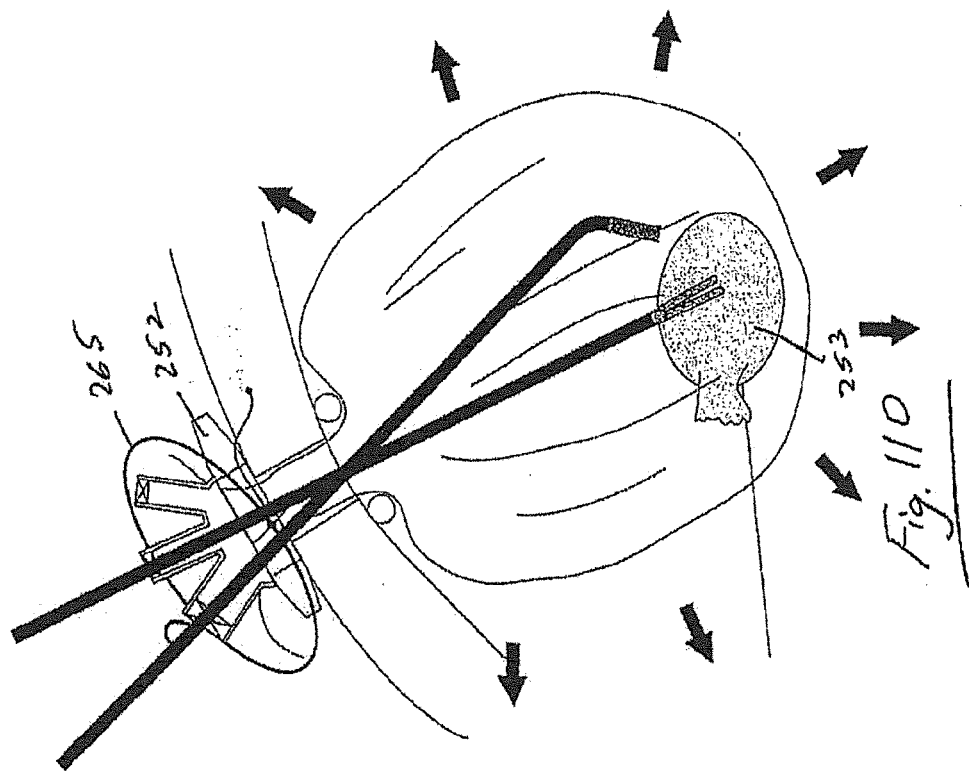


Fig. 106





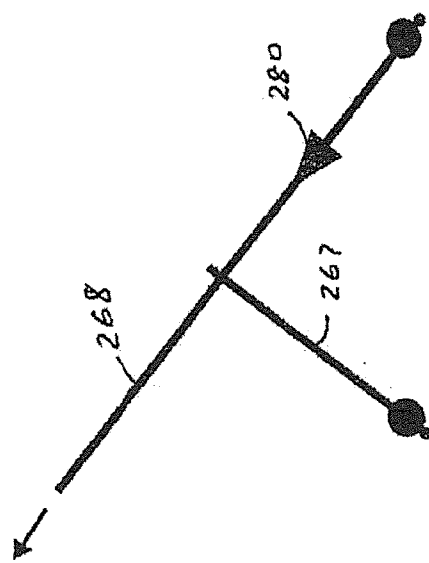


Fig. 112

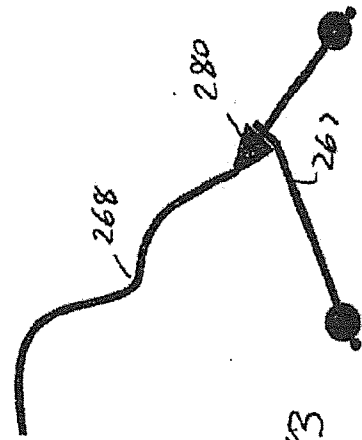


Fig. 113

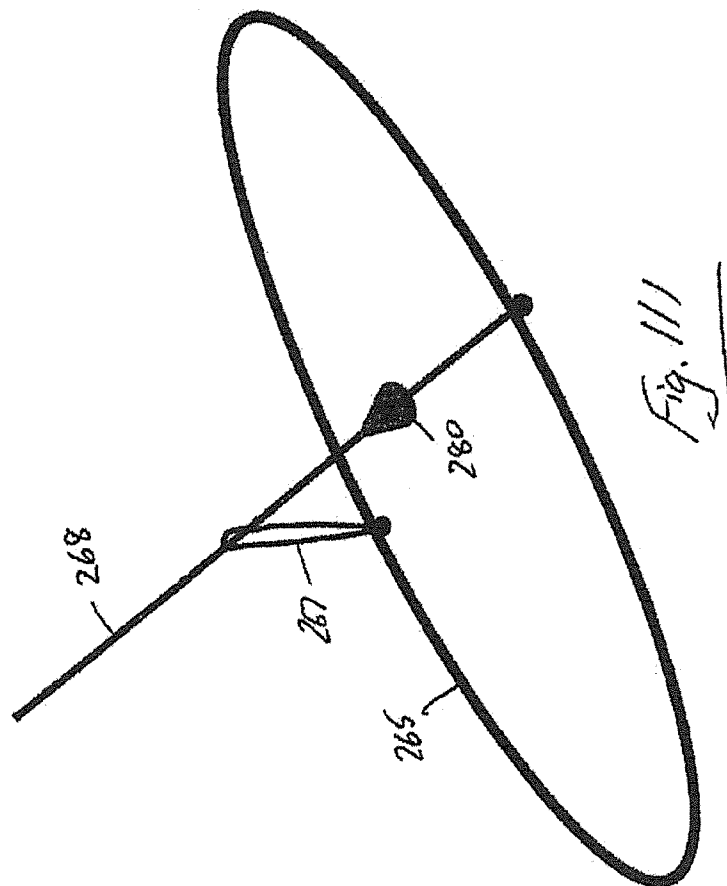
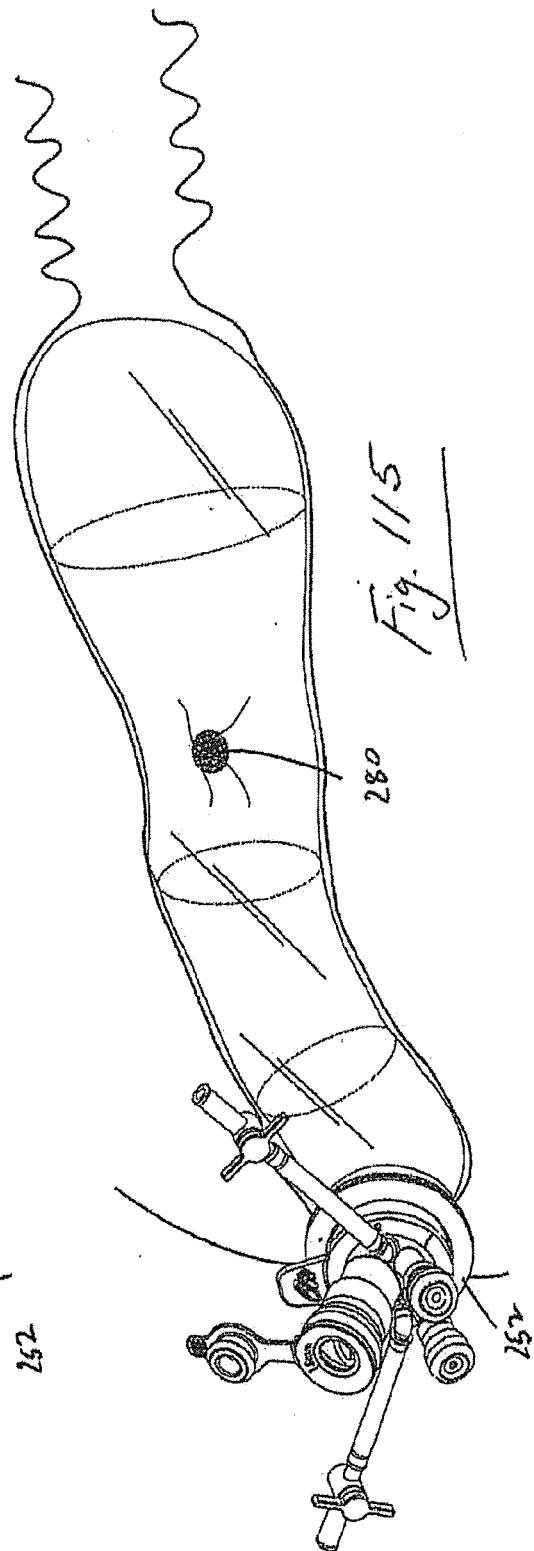
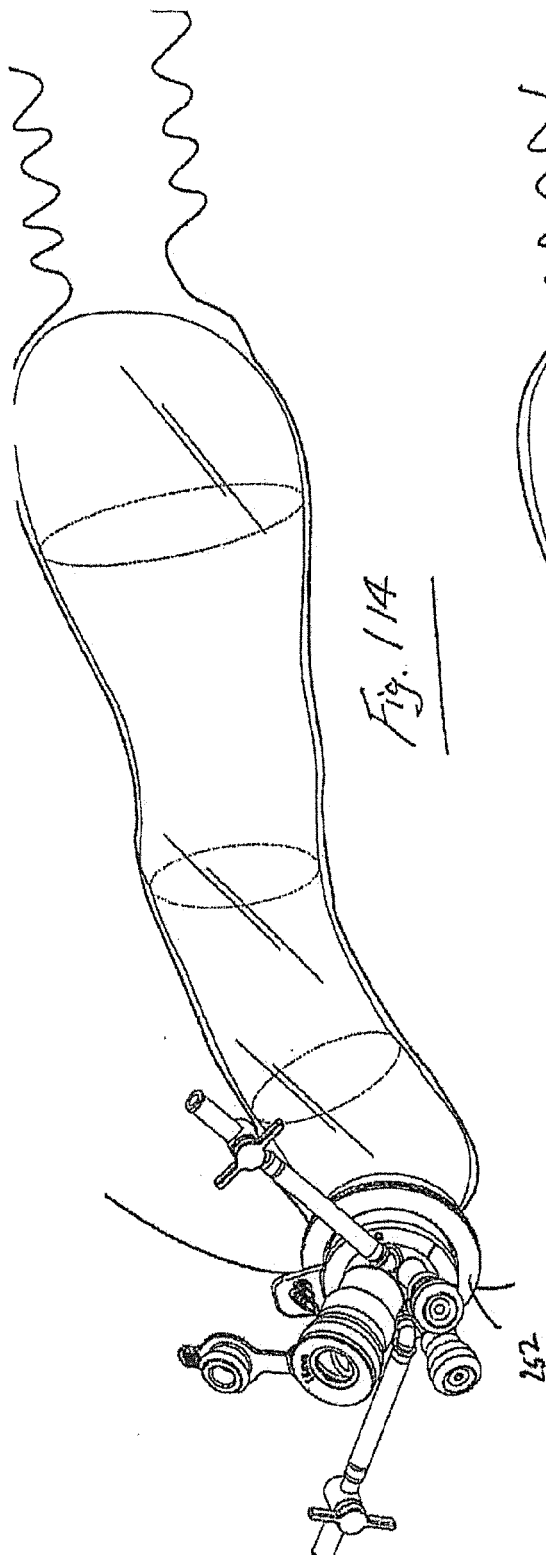
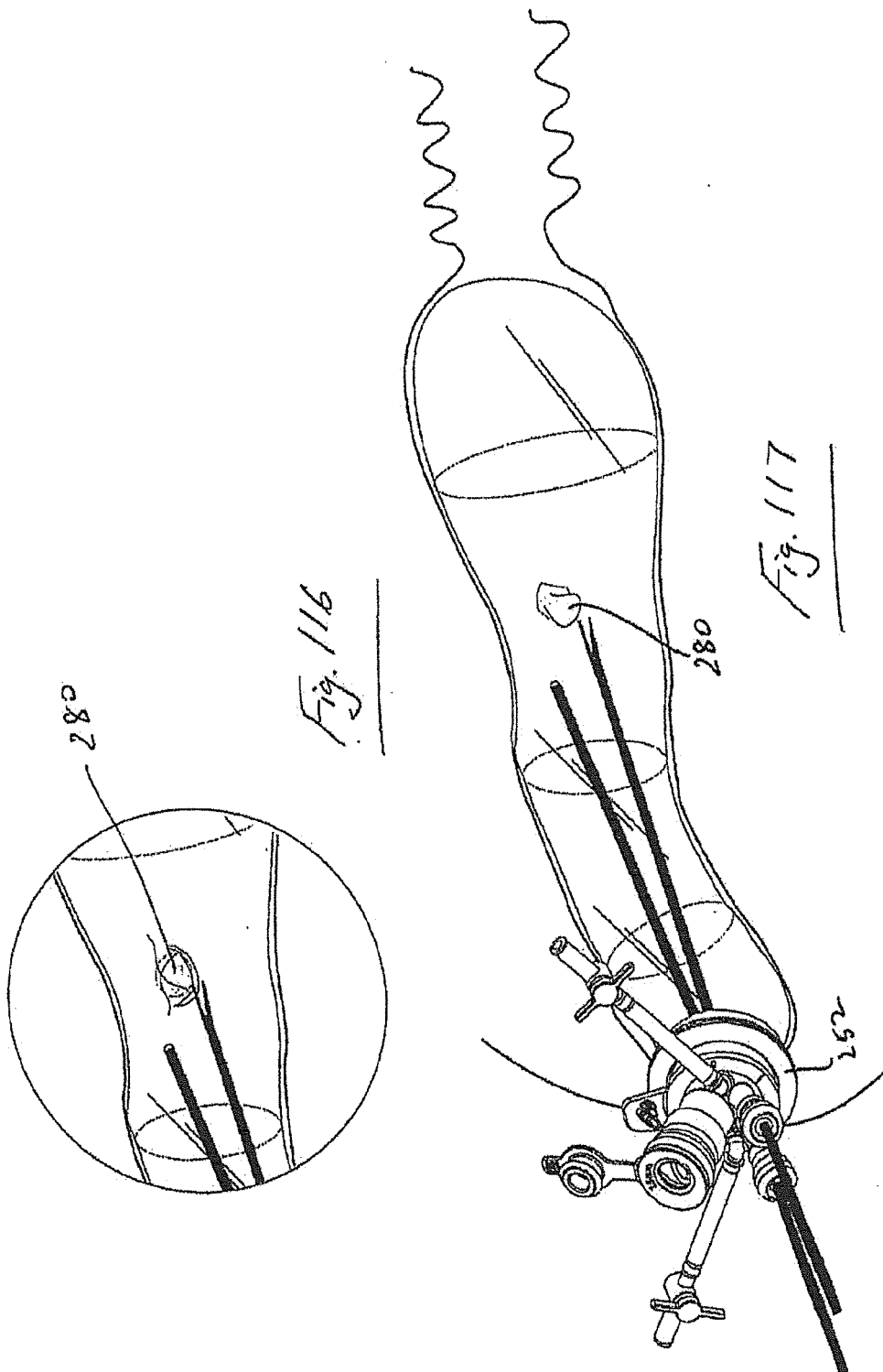
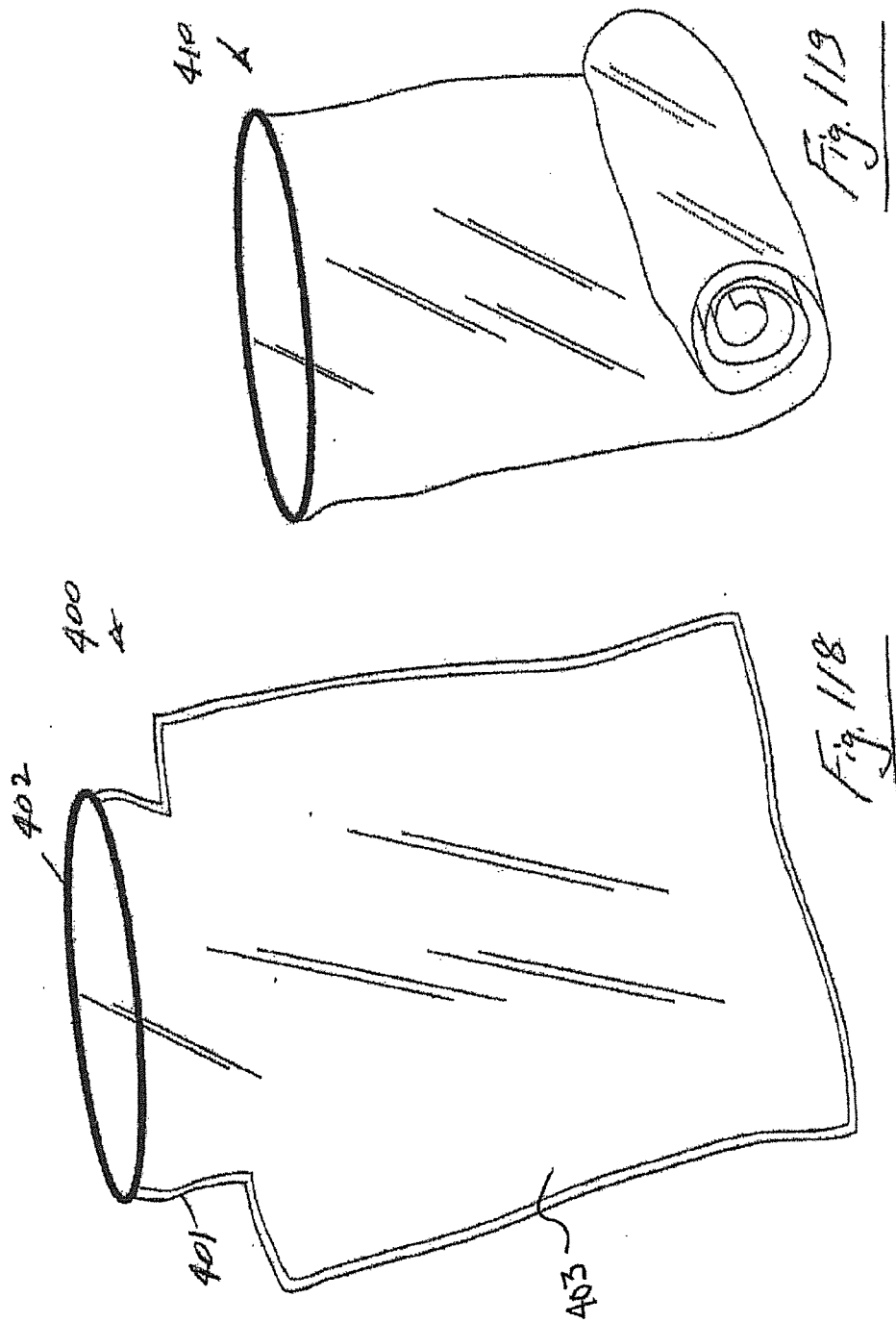
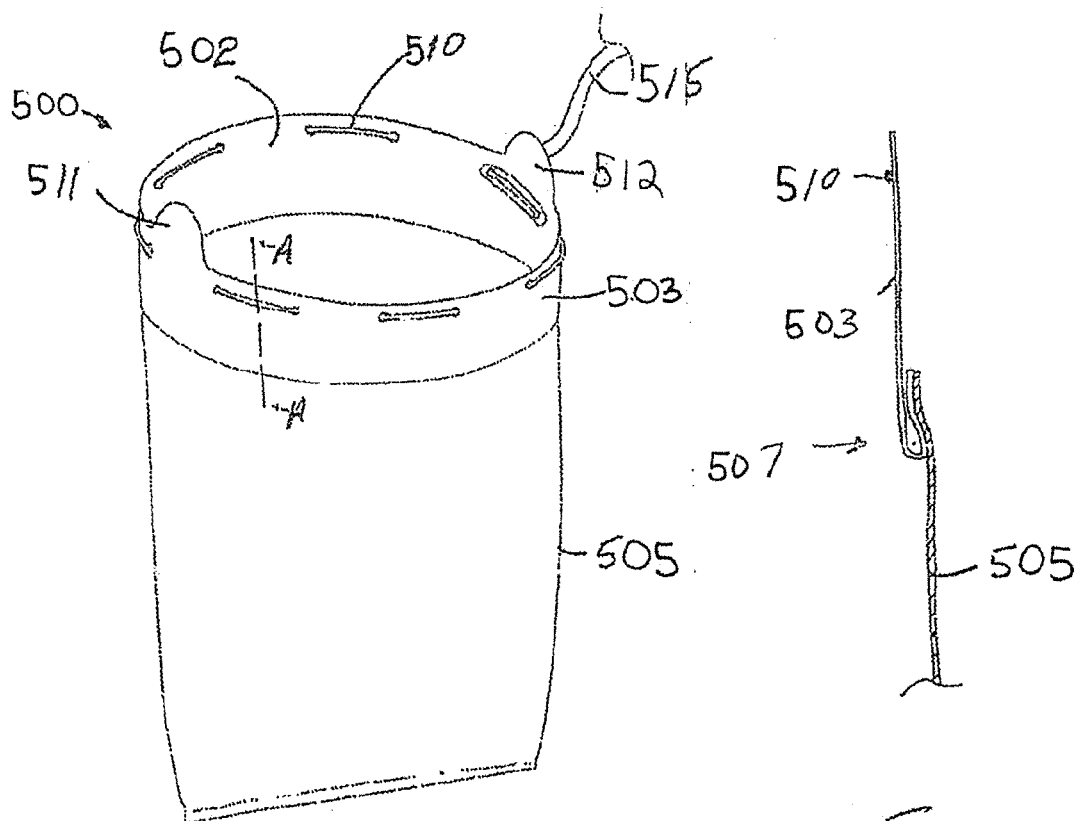
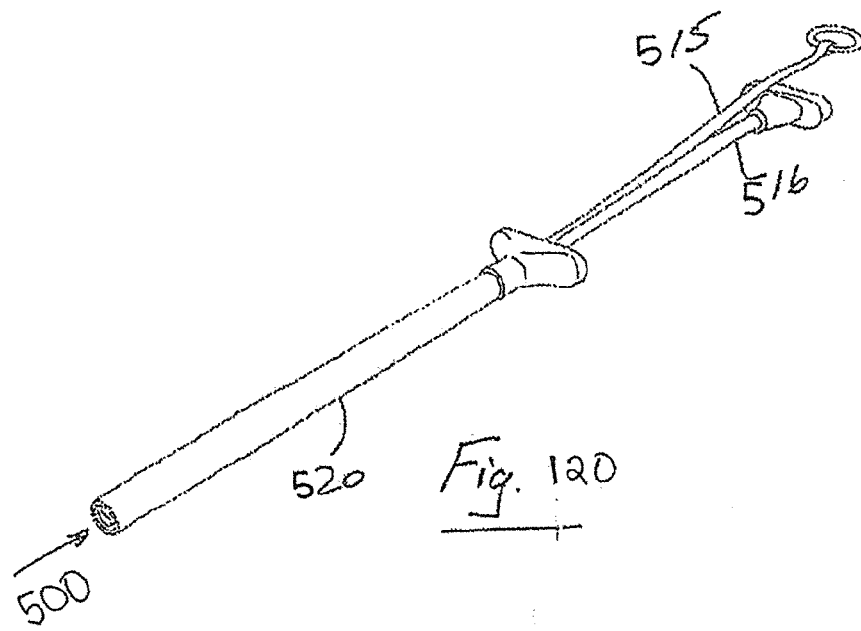


Fig. 111









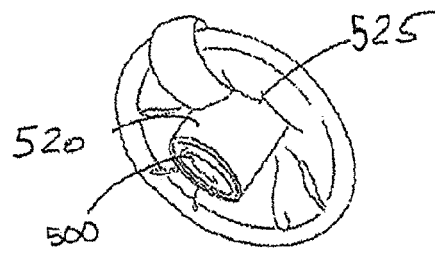


Fig. 123

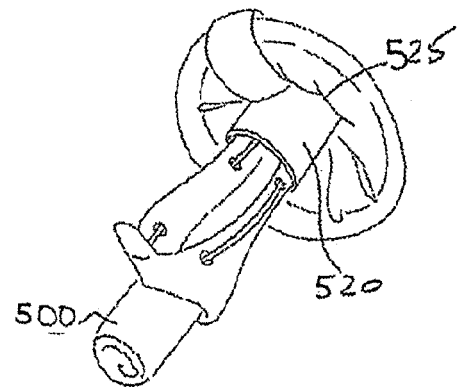


Fig. 124

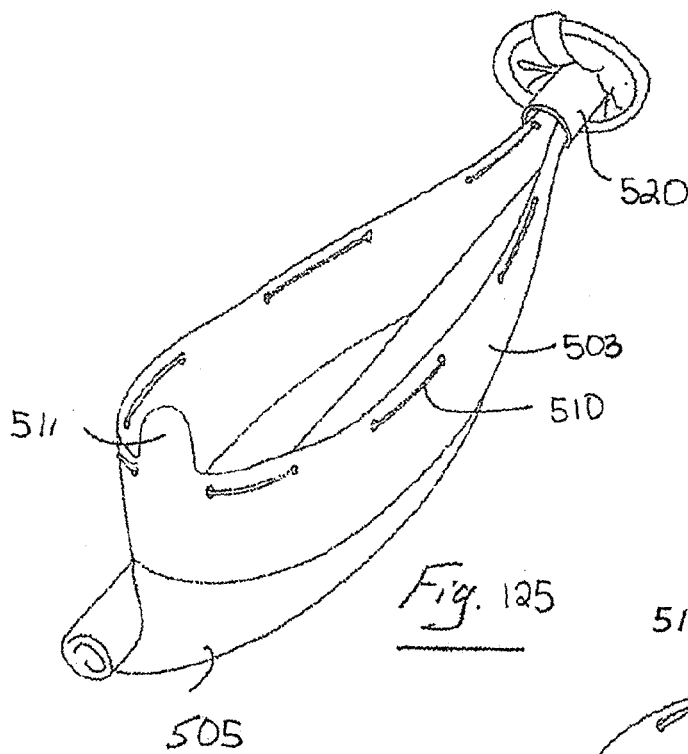


Fig. 125

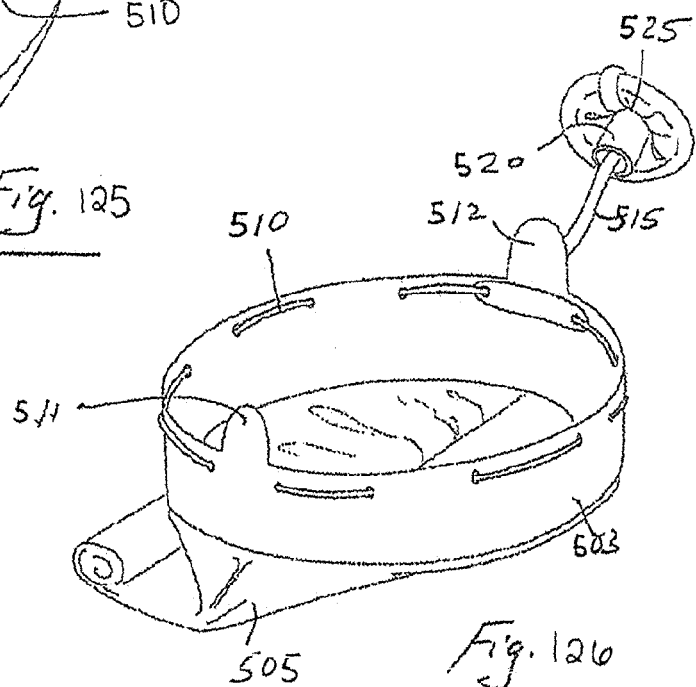
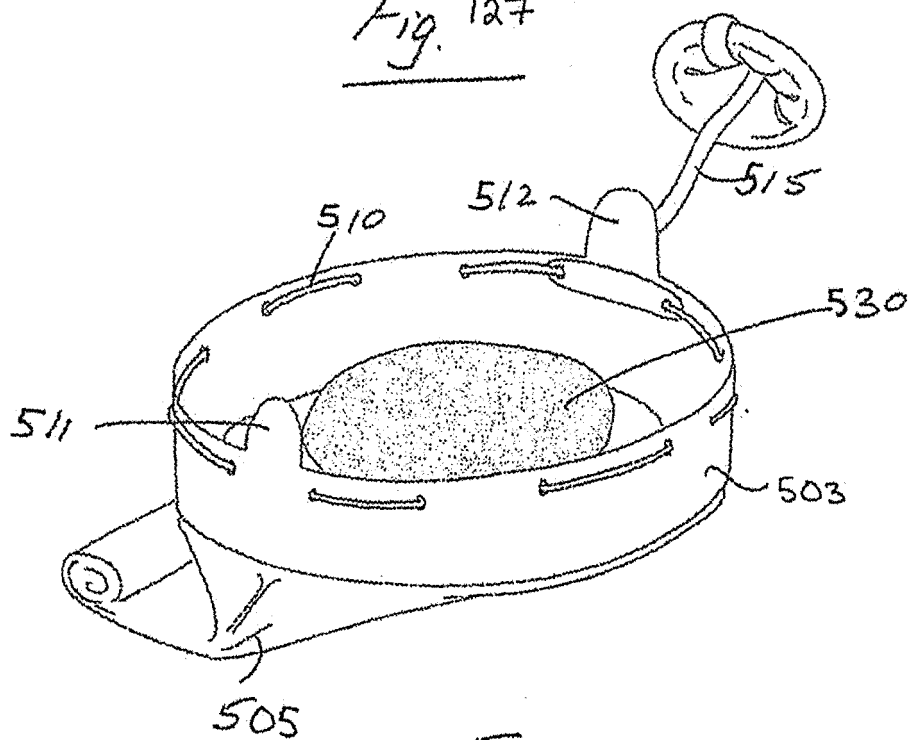
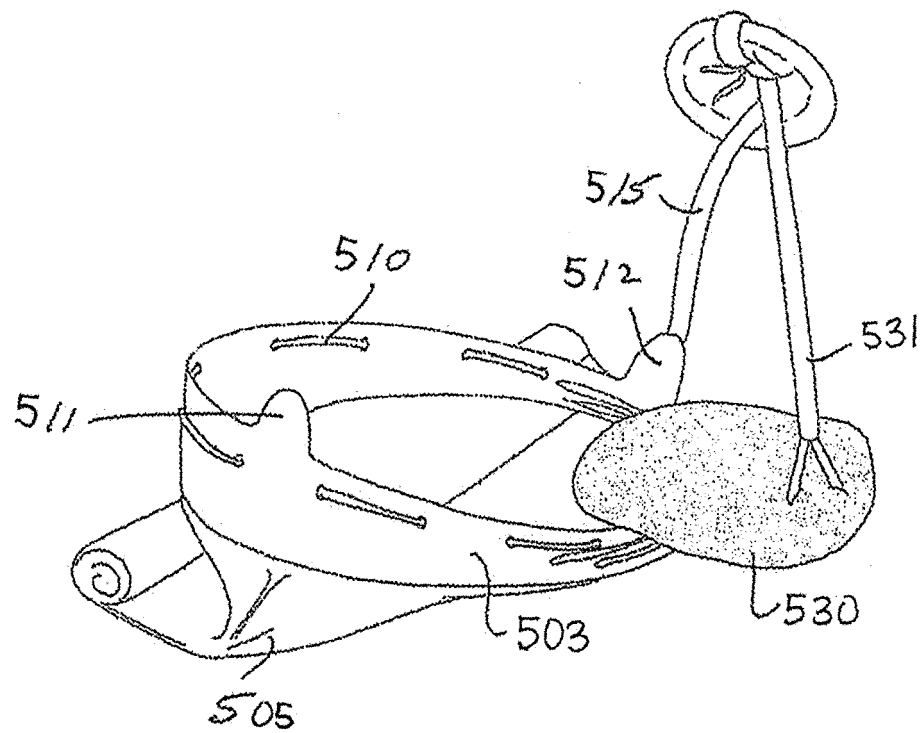


Fig. 126



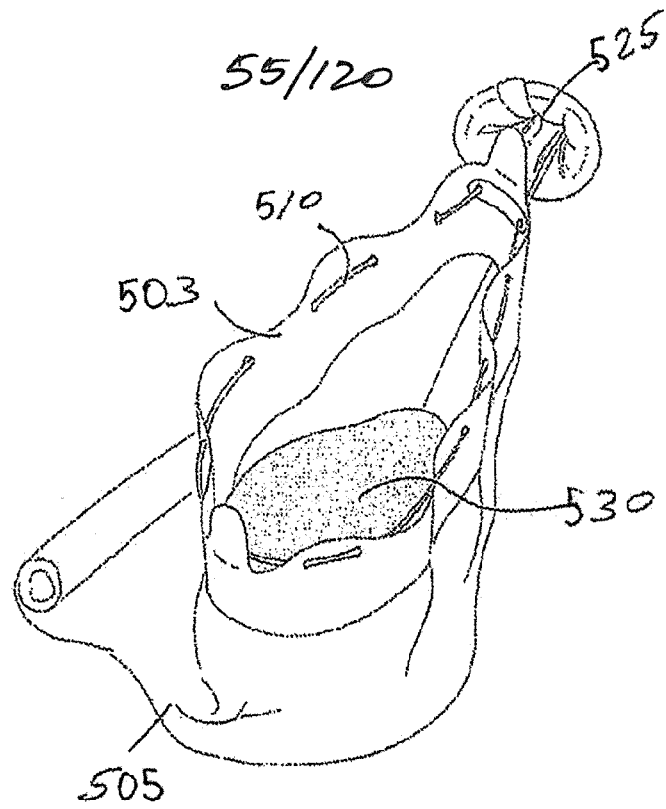


Fig. 129

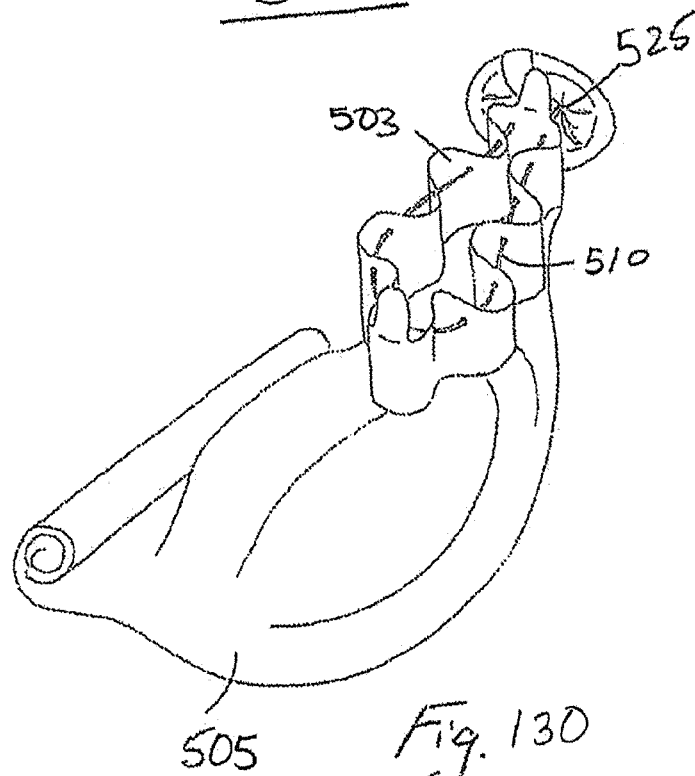


Fig. 130

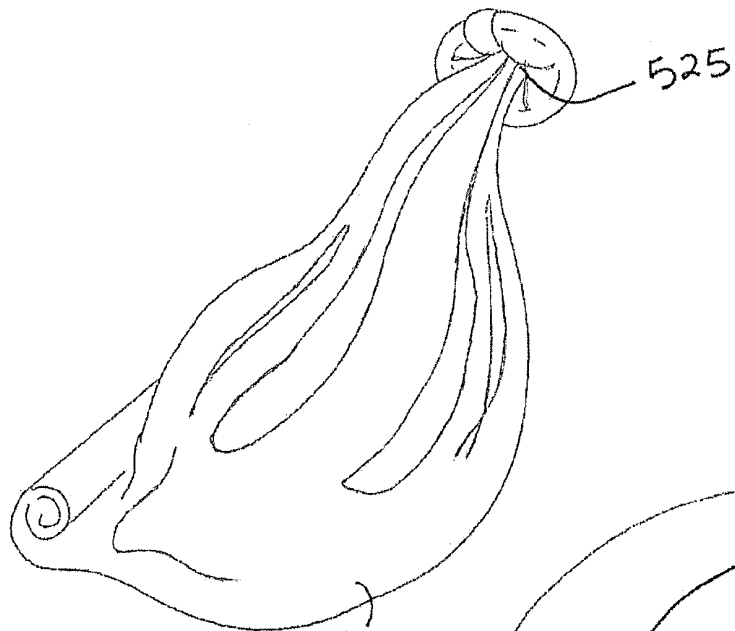


Fig. 131

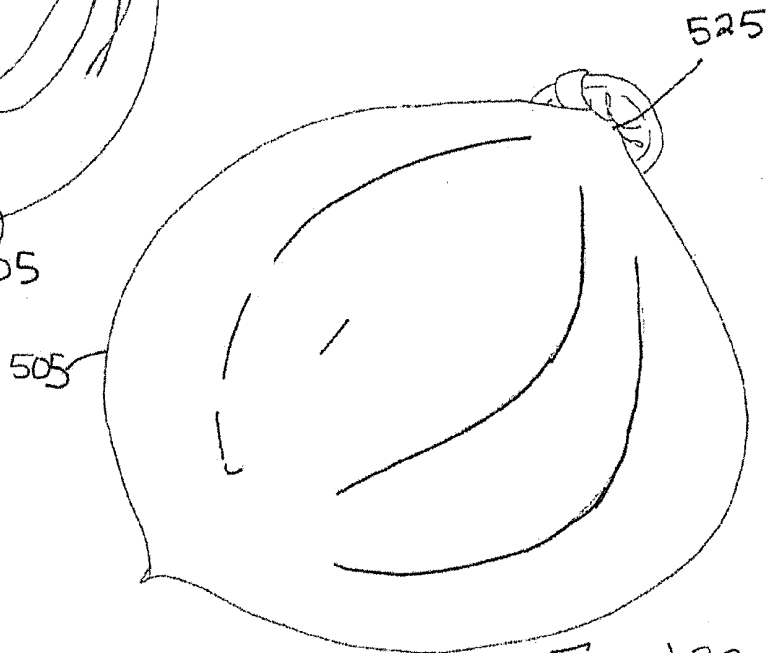


Fig. 132

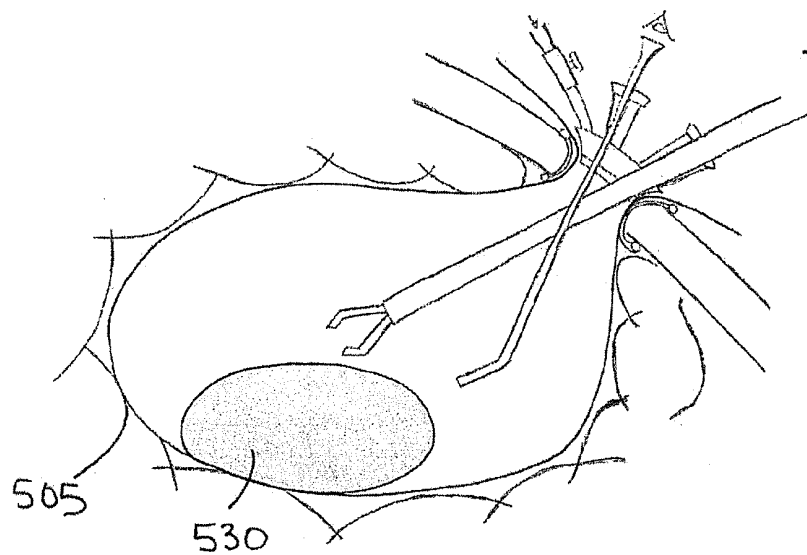


Fig. 133

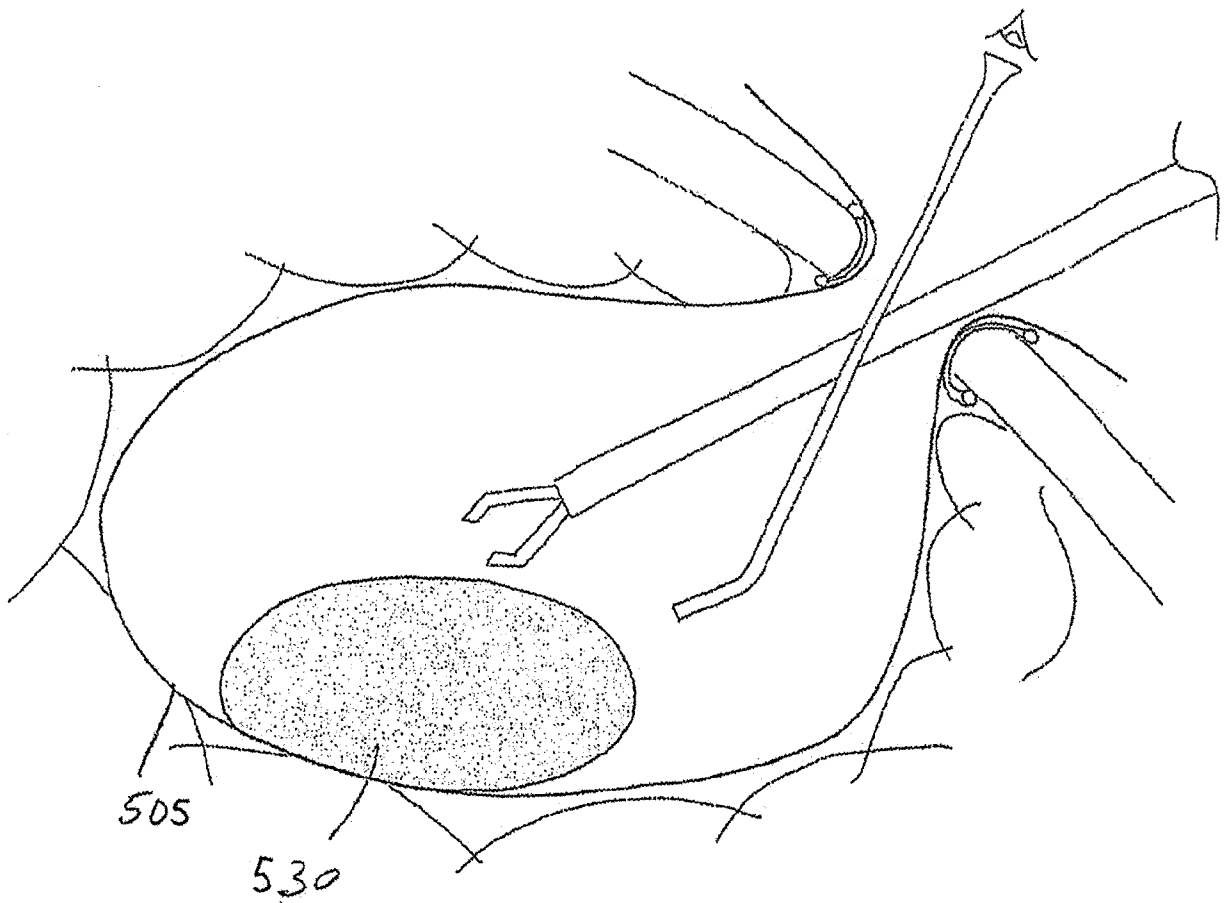


Fig. 134

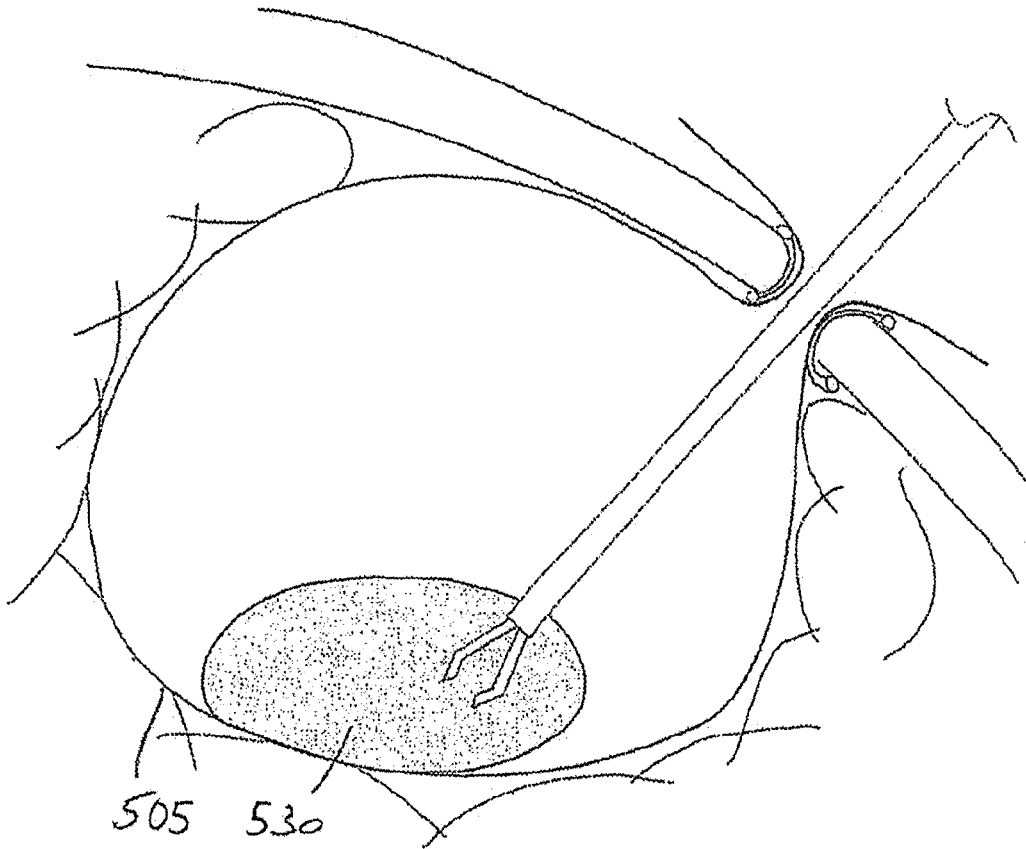


Fig. 135

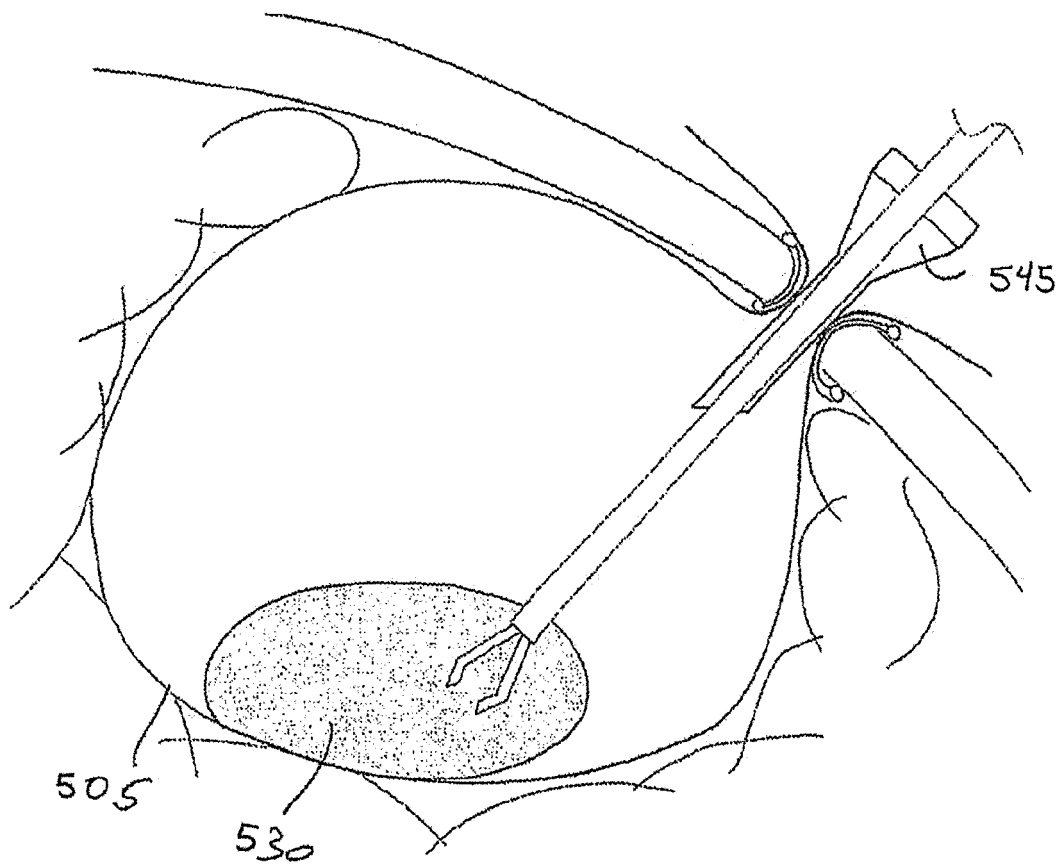


Fig. 136

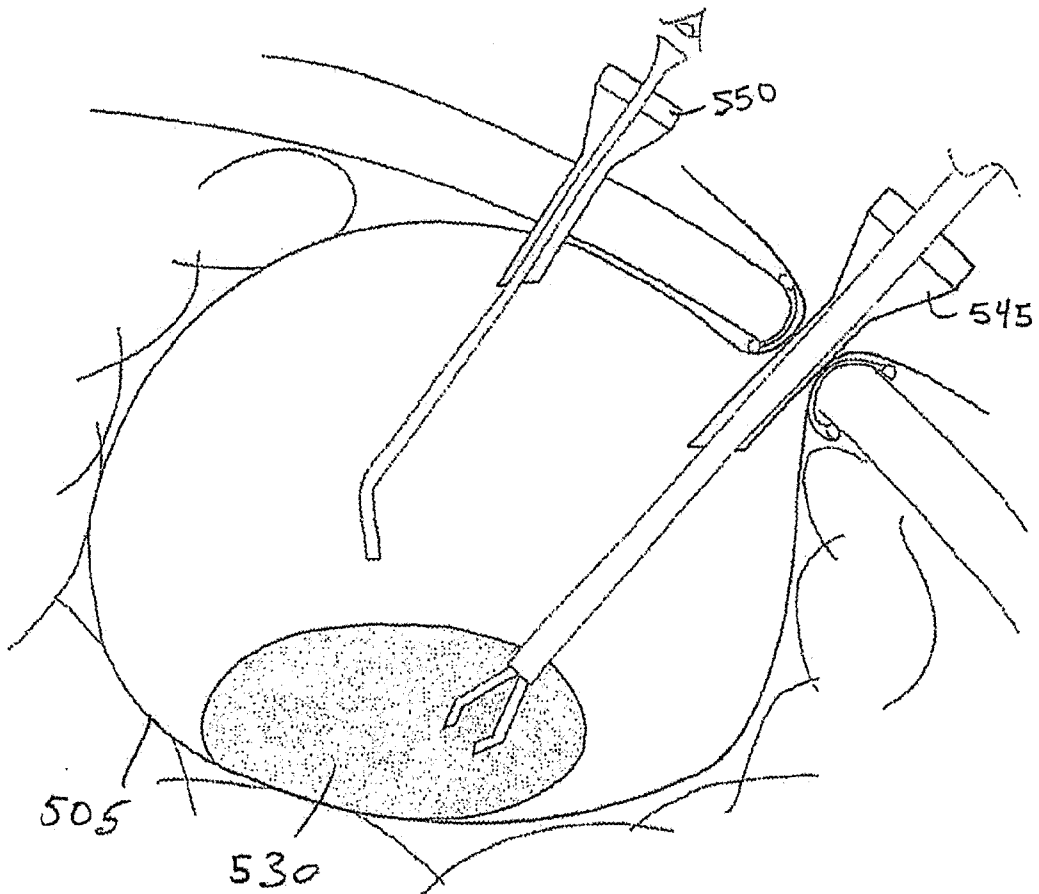


Fig. 137

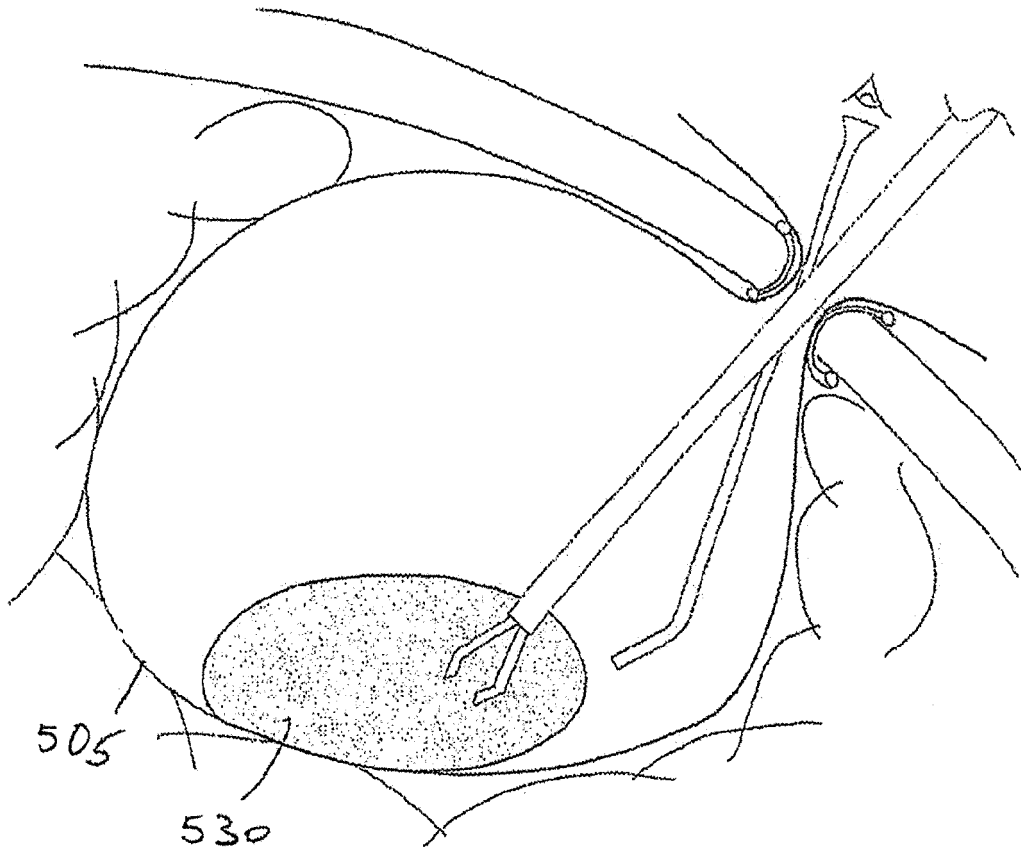
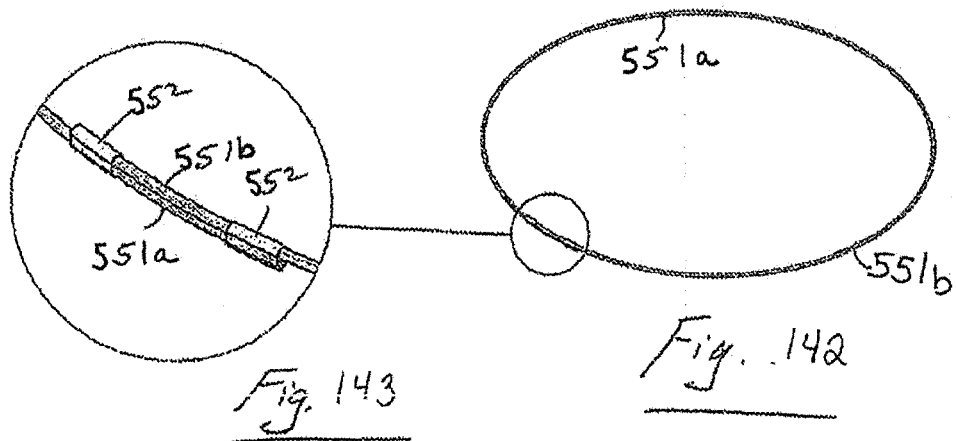
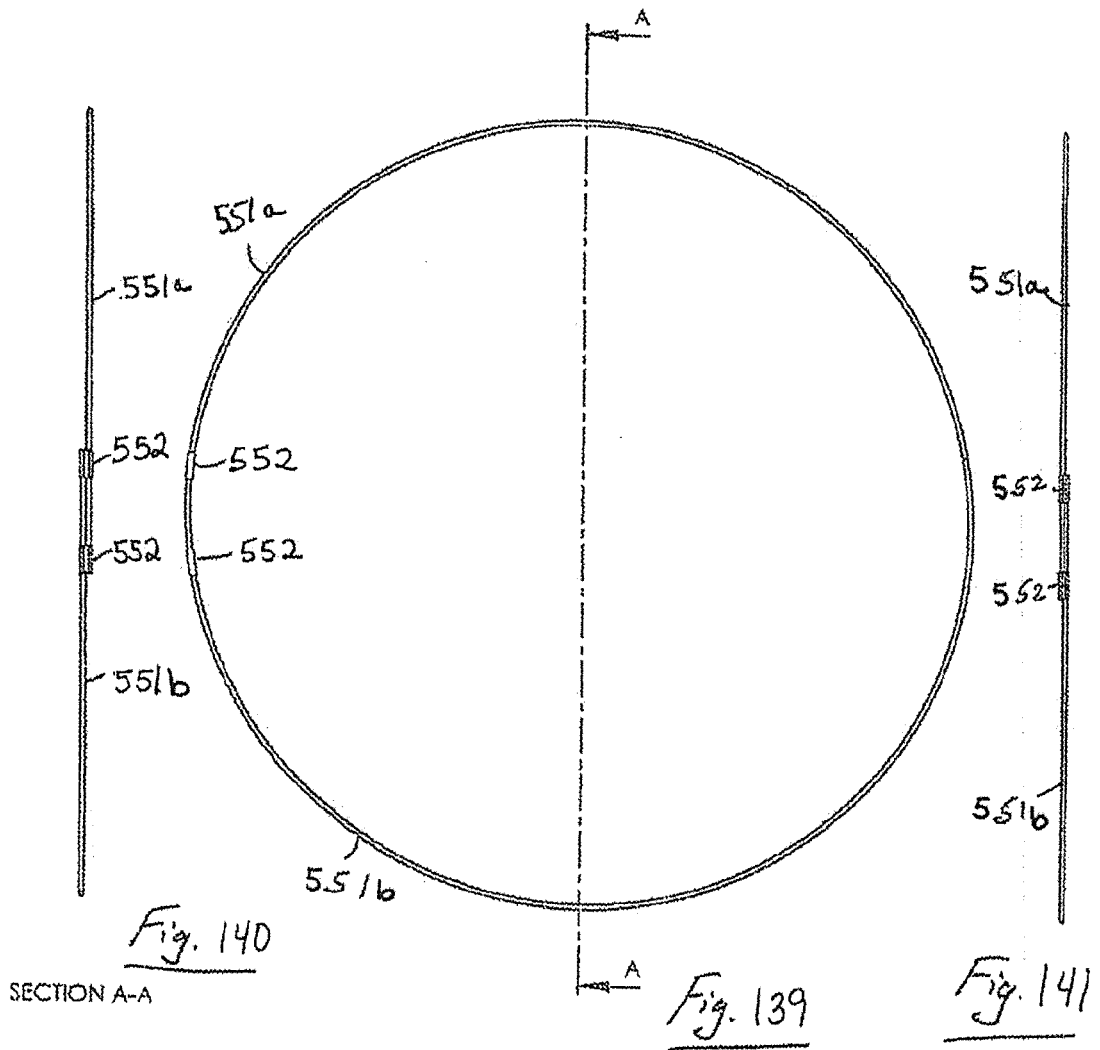
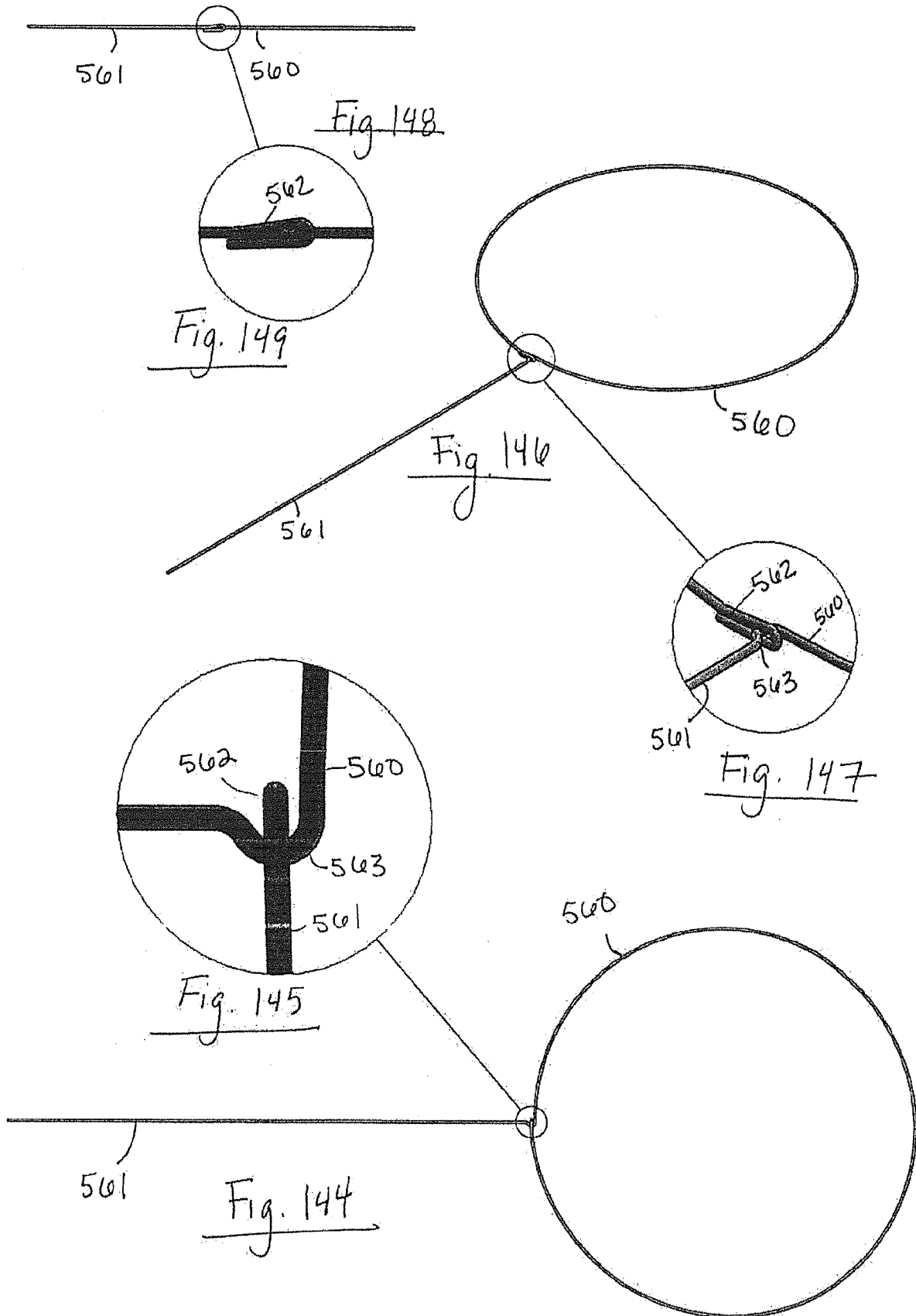


Fig. 138





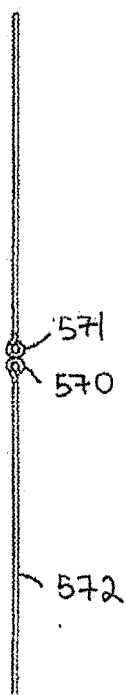


Fig. 151

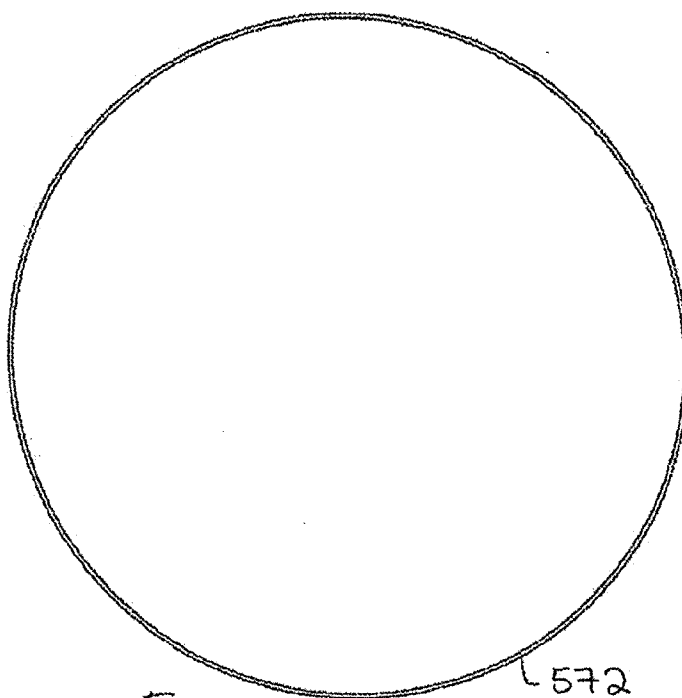


Fig. 150

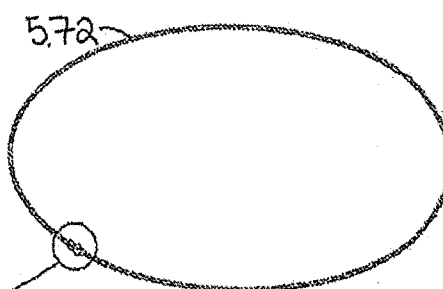


Fig. 152

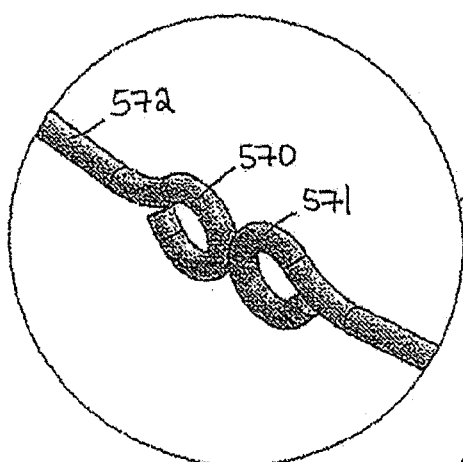


Fig. 153

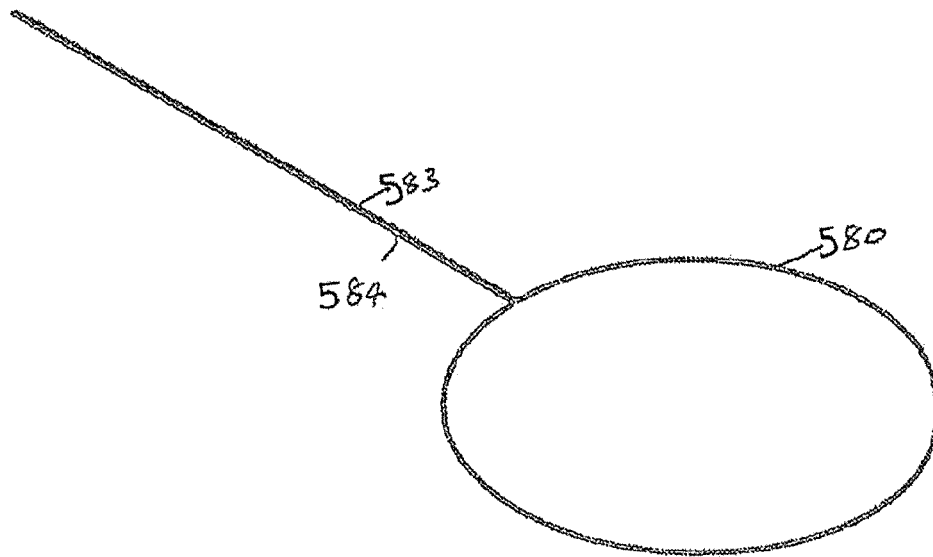


Fig. 154

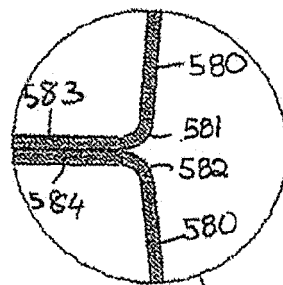


Fig. 156

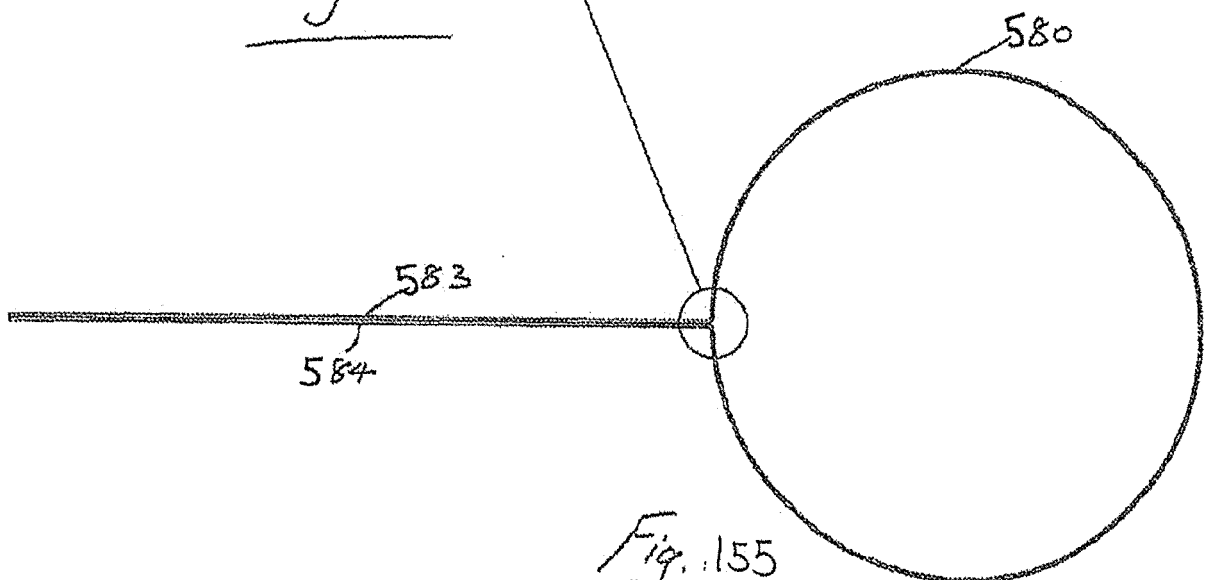
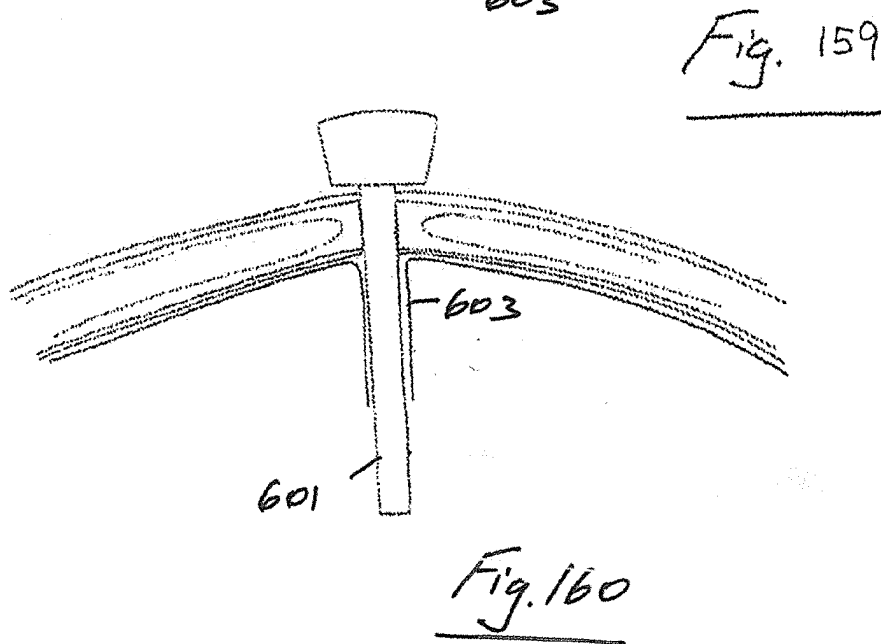
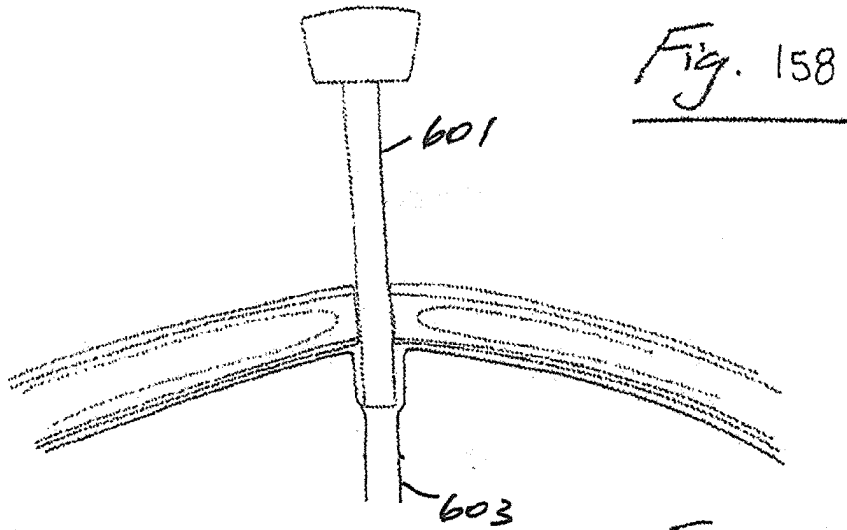
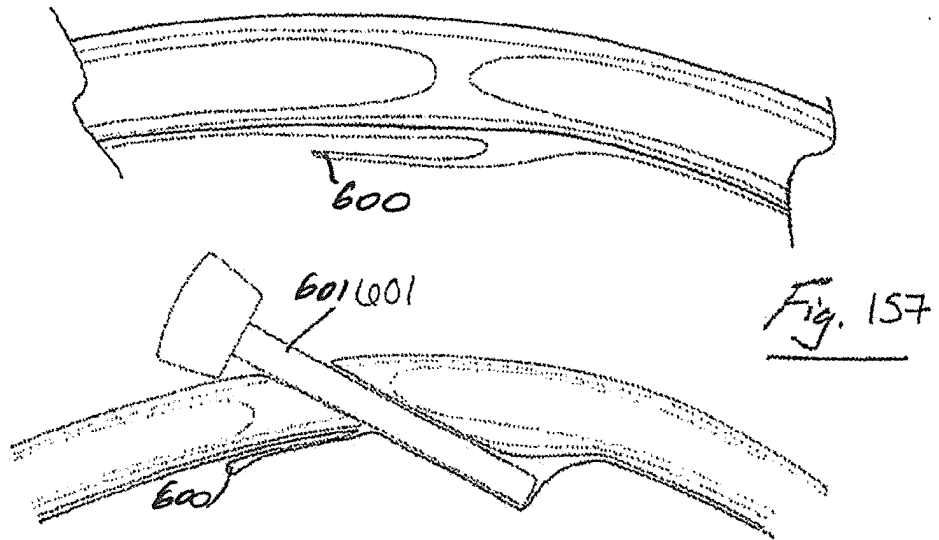
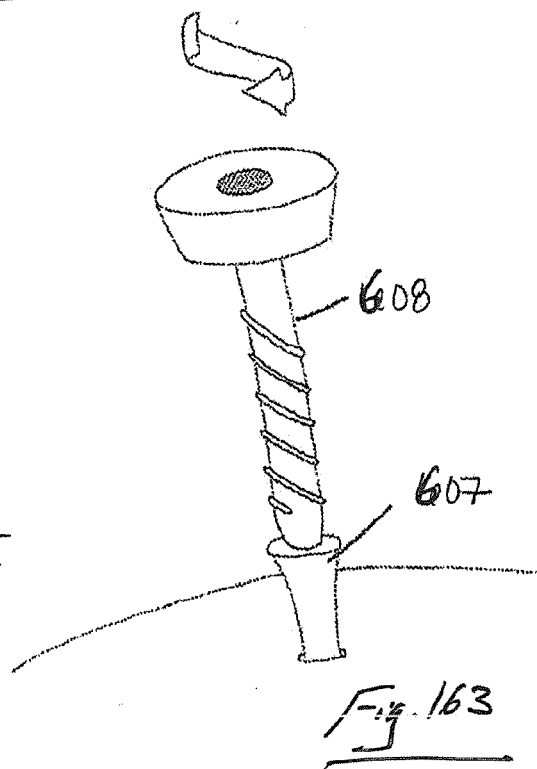
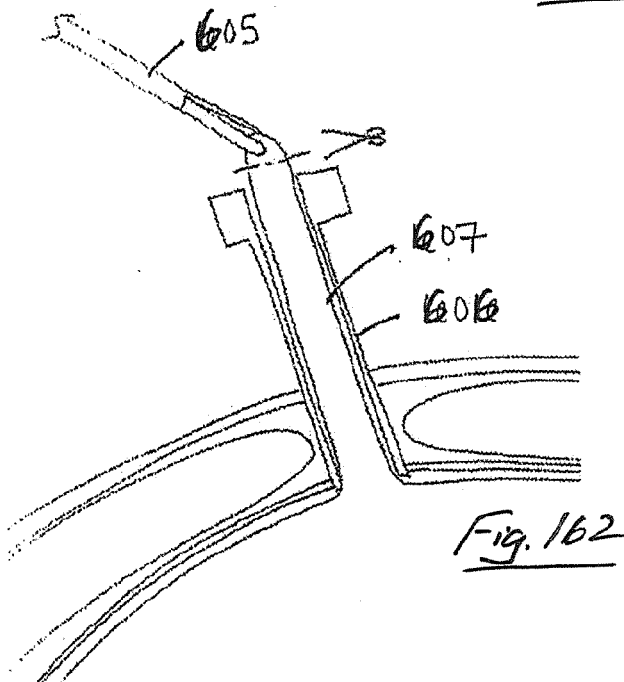
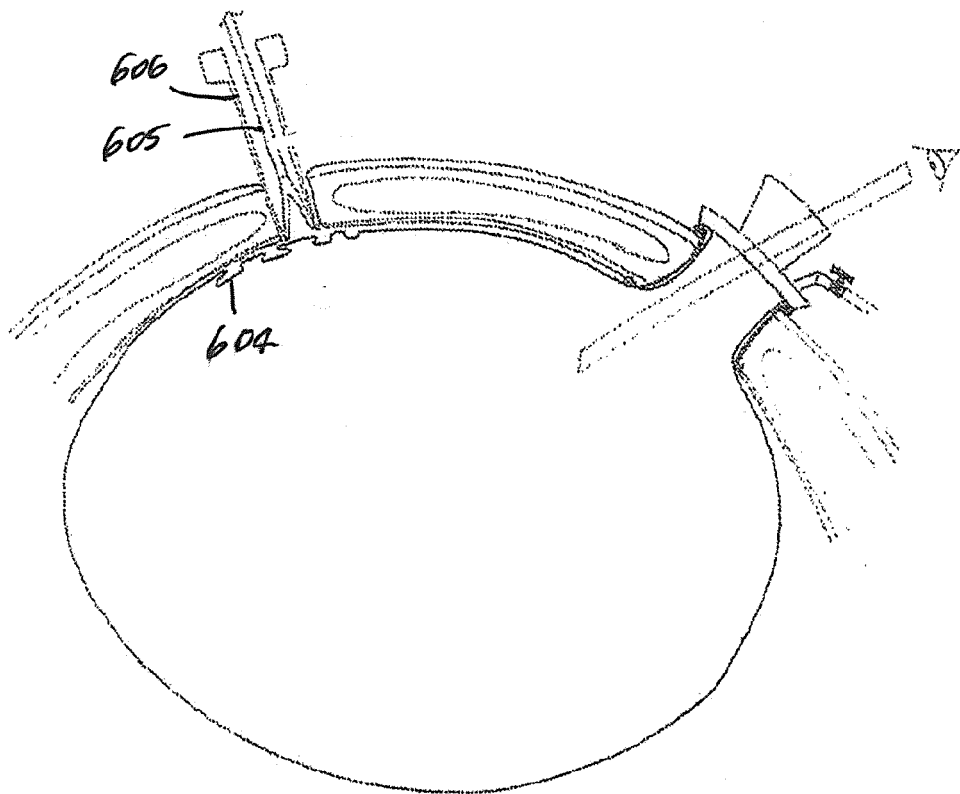
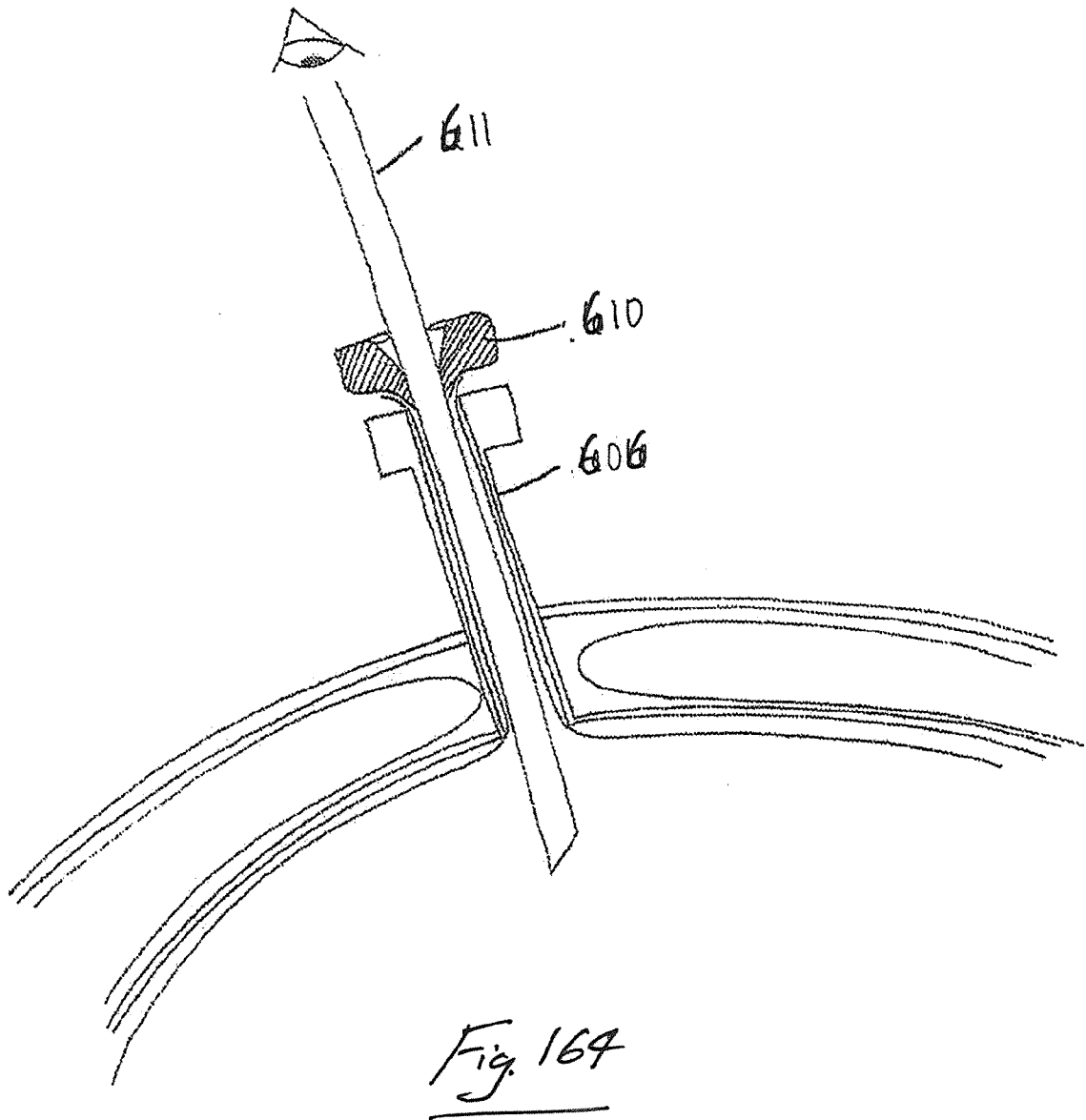


Fig. 155







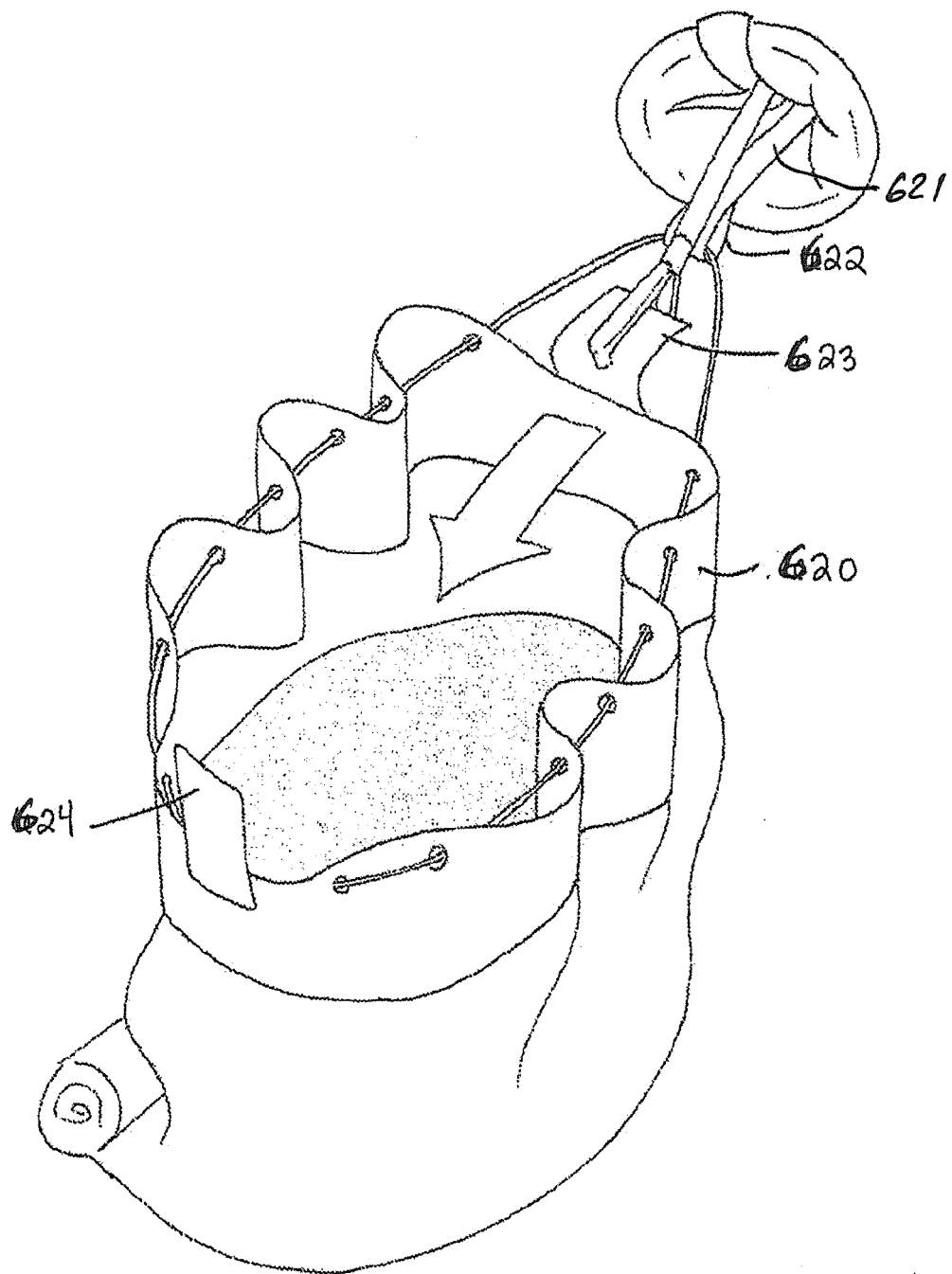


Fig. 165

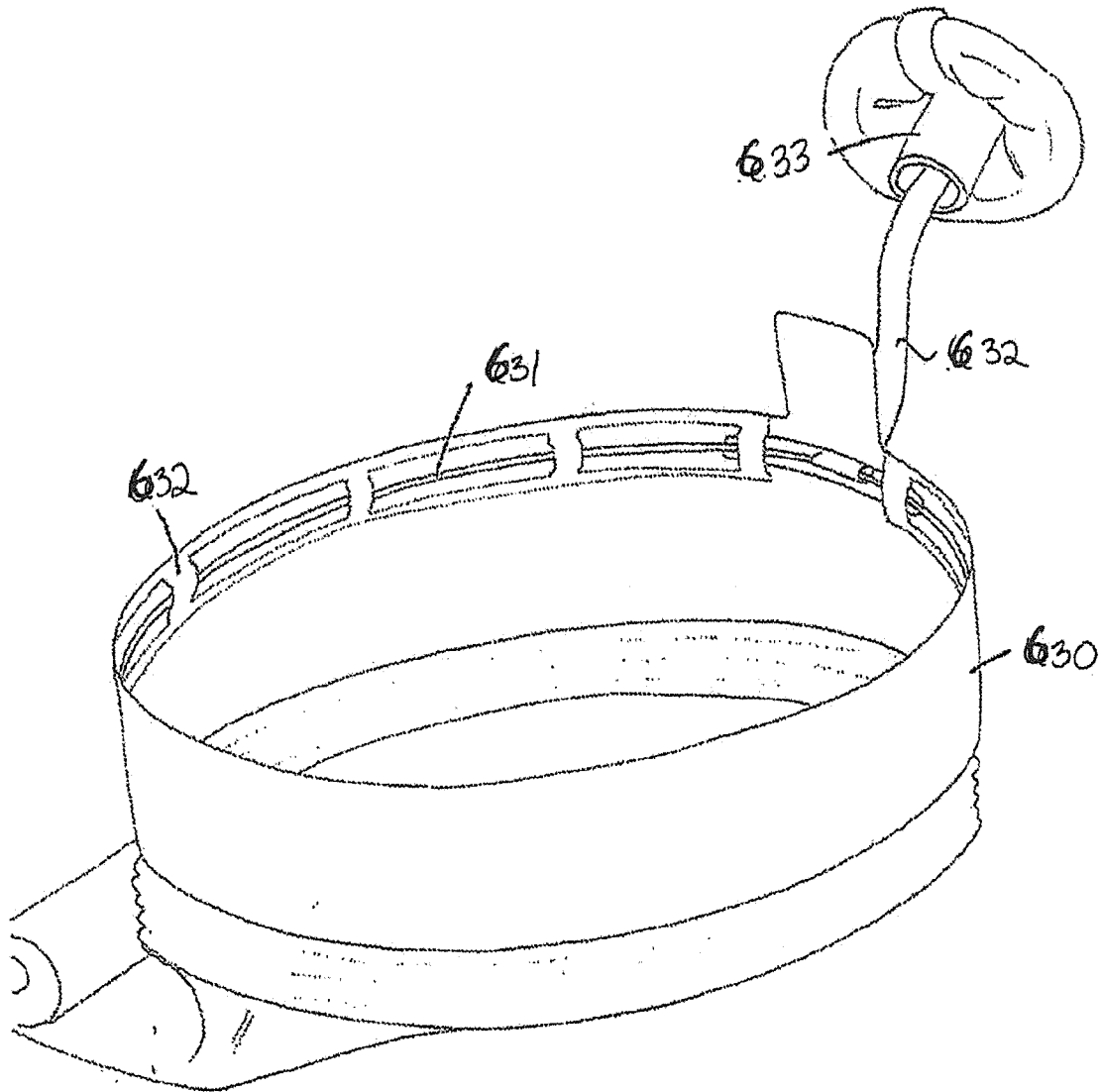


Fig. 166

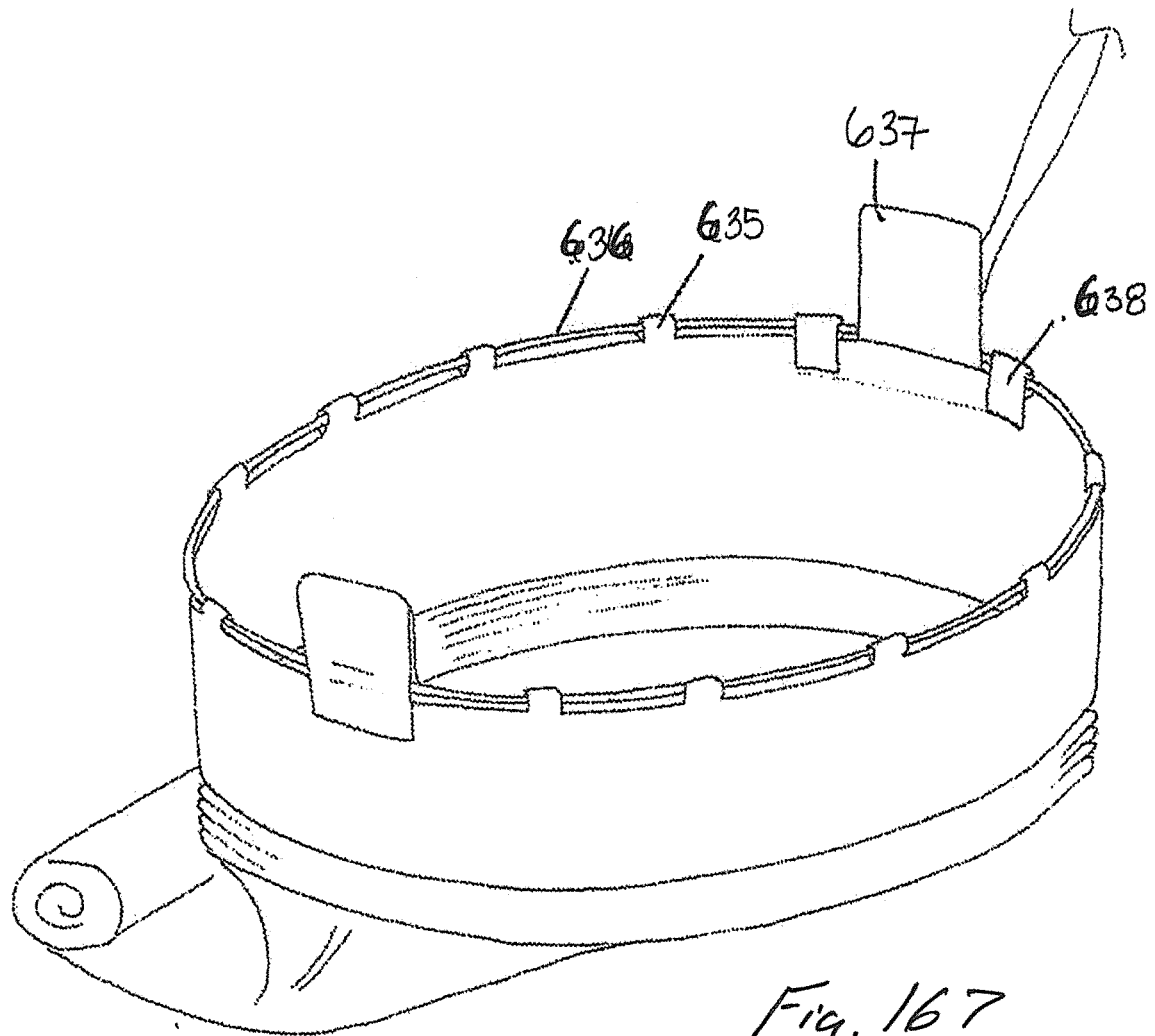


Fig. 167

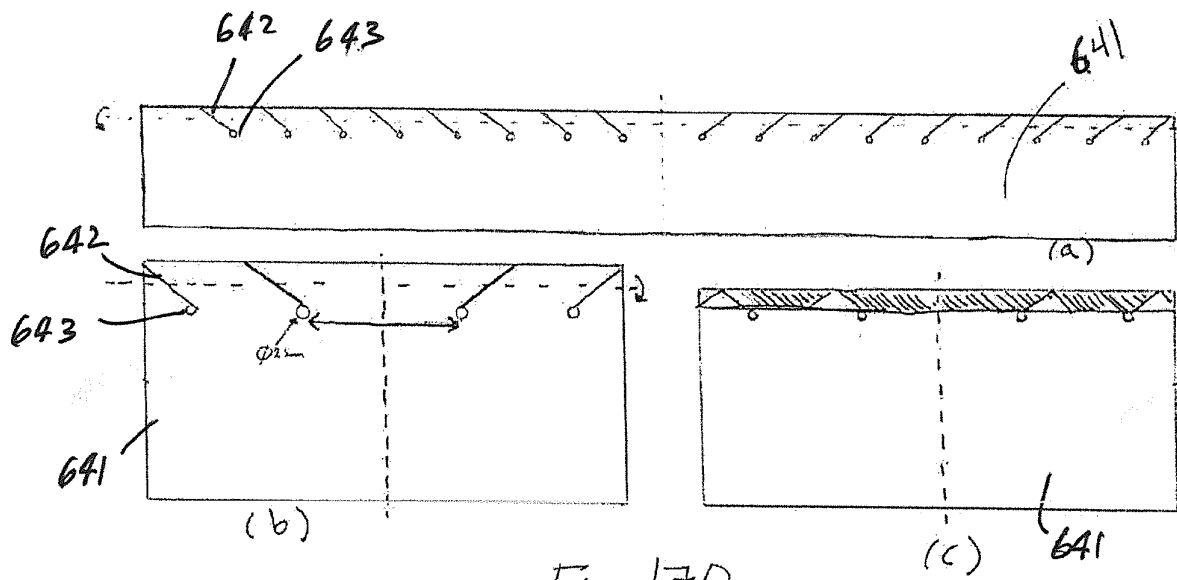
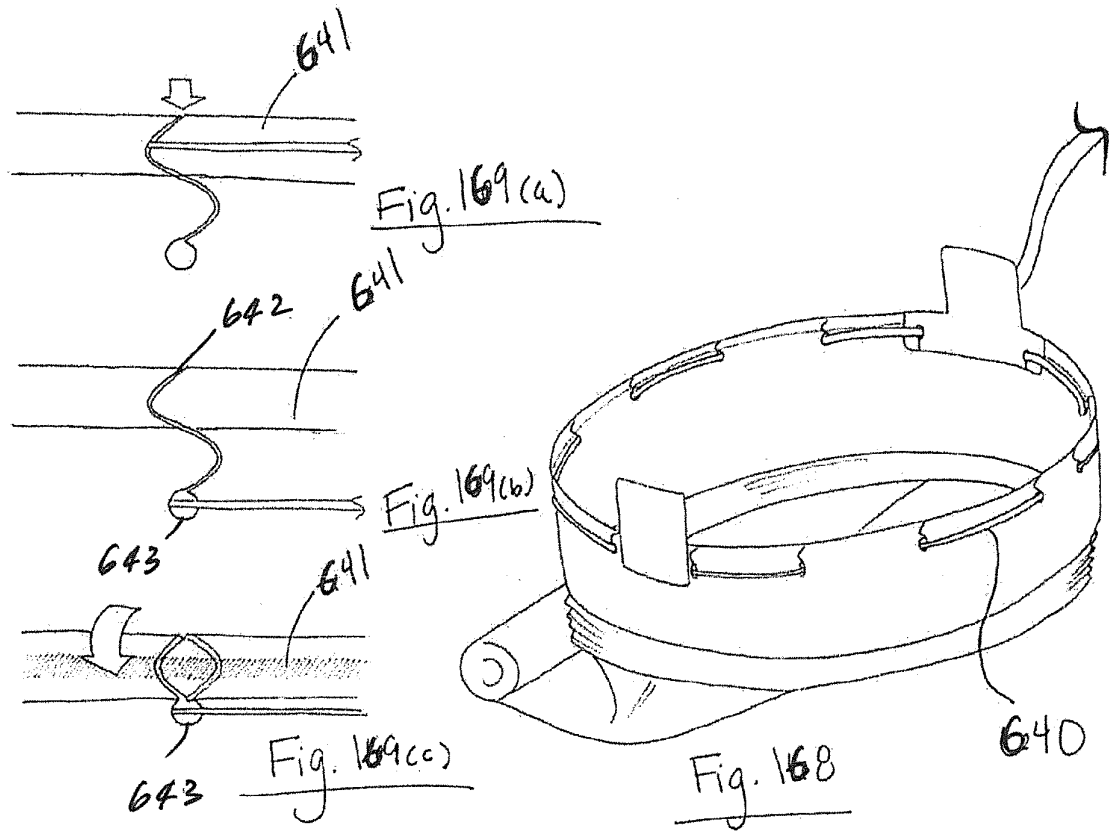


Fig. 170

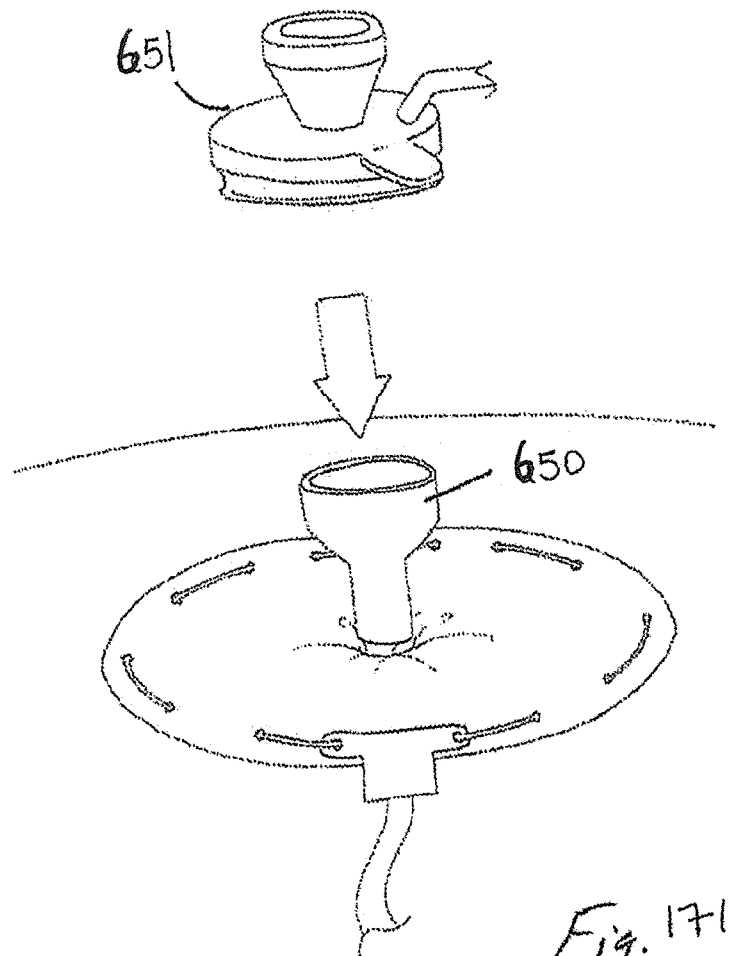


Fig. 171

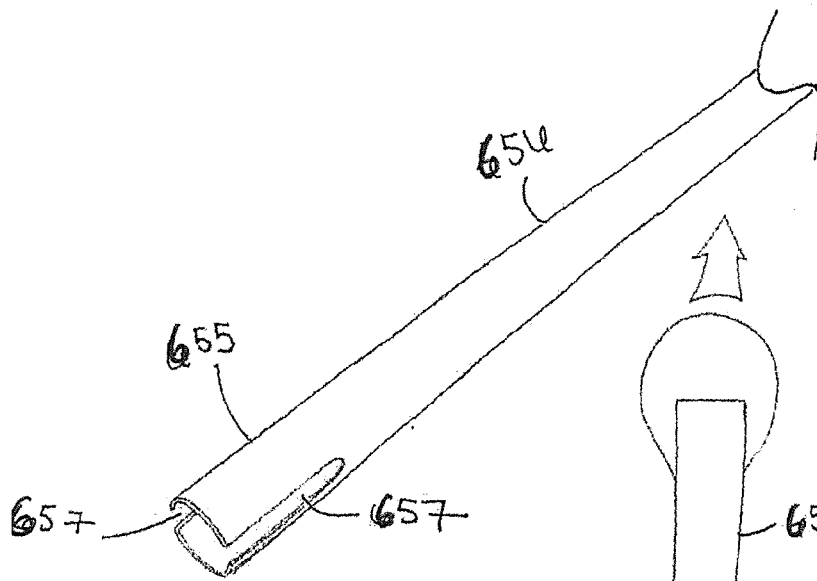


Fig. 172

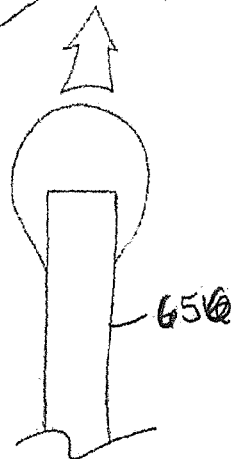


Fig. 173

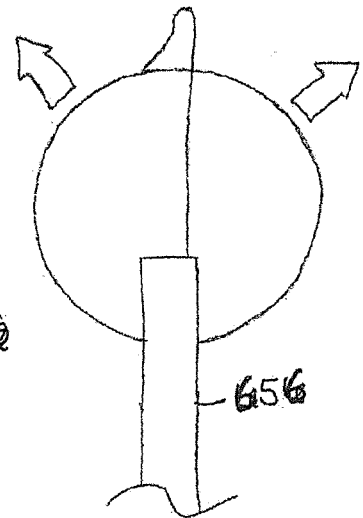


Fig. 174

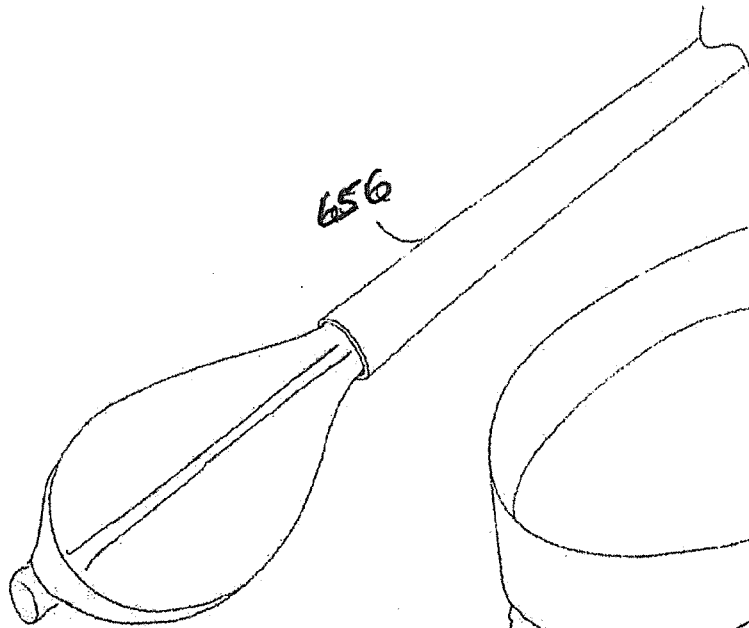


Fig. 175

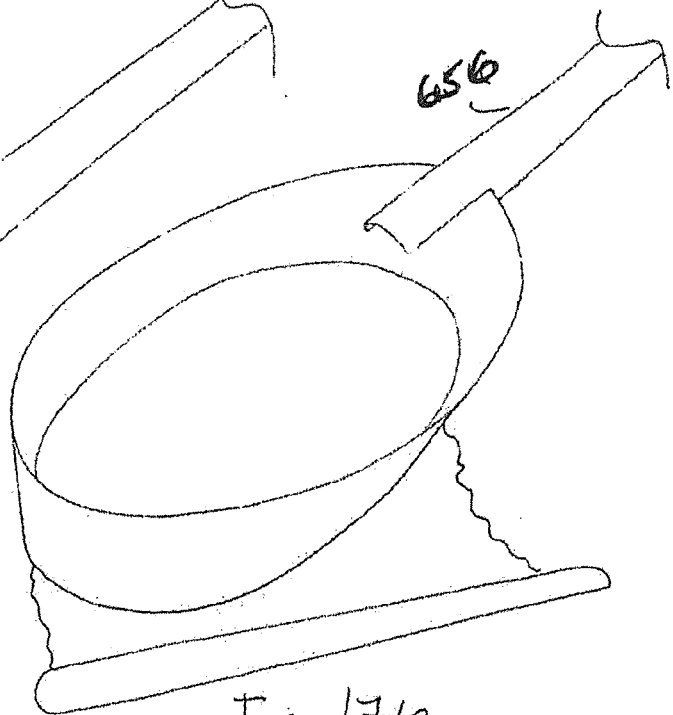


Fig. 176

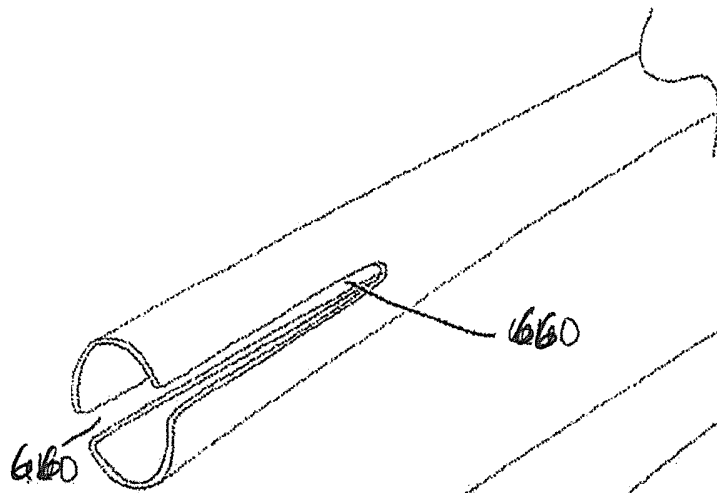


Fig. 177

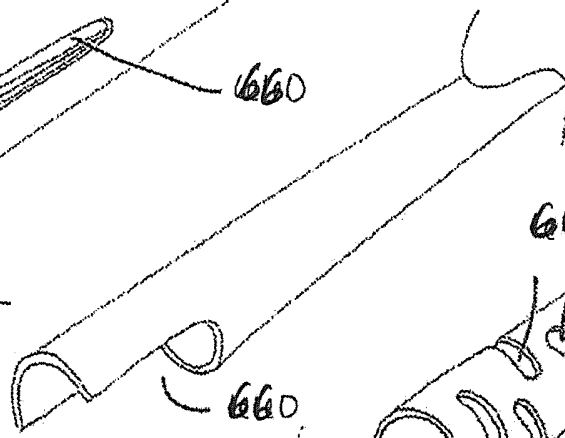


Fig. 178

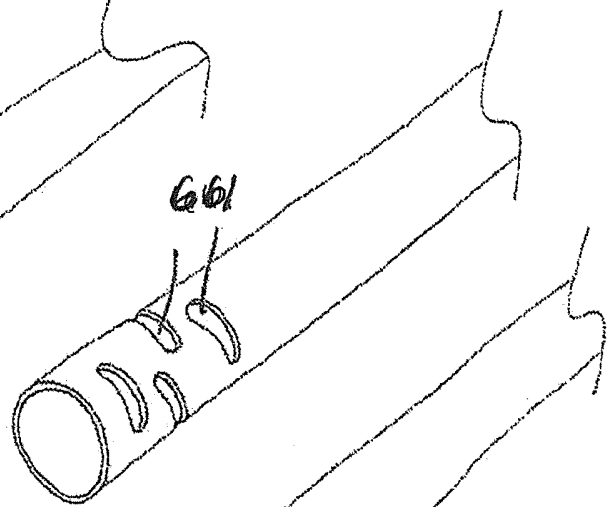


Fig. 179

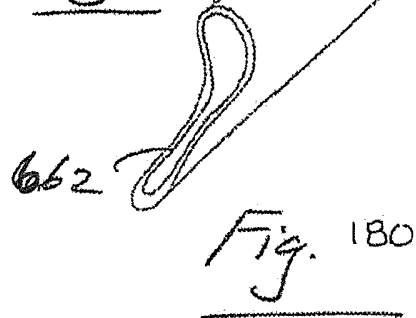


Fig. 180

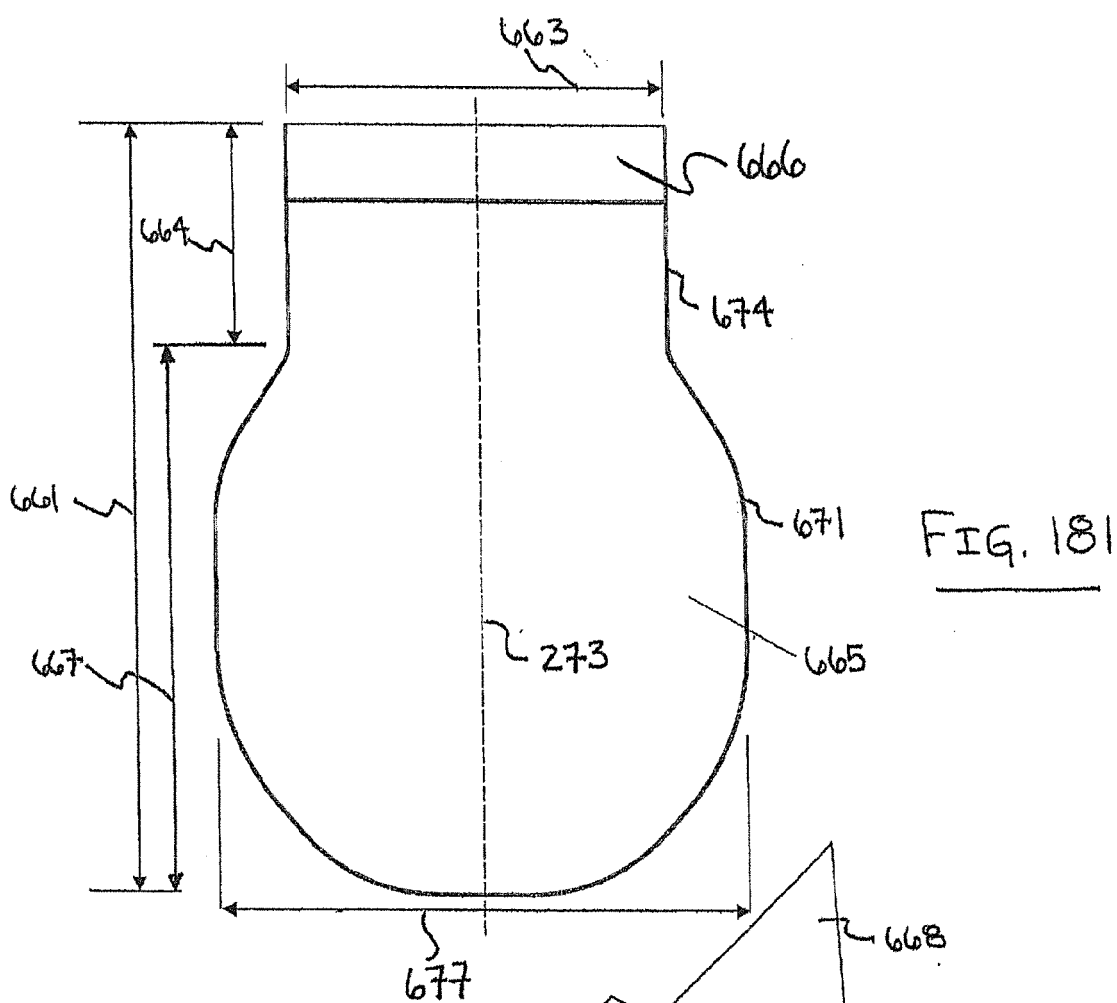


FIG. 181

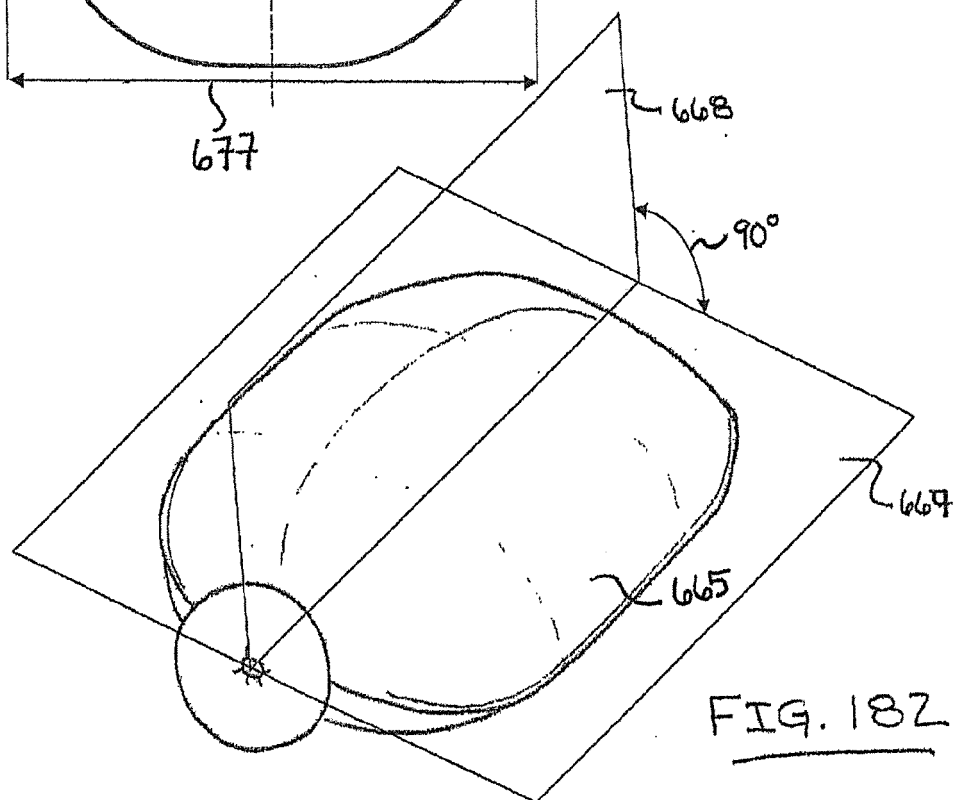


FIG. 182

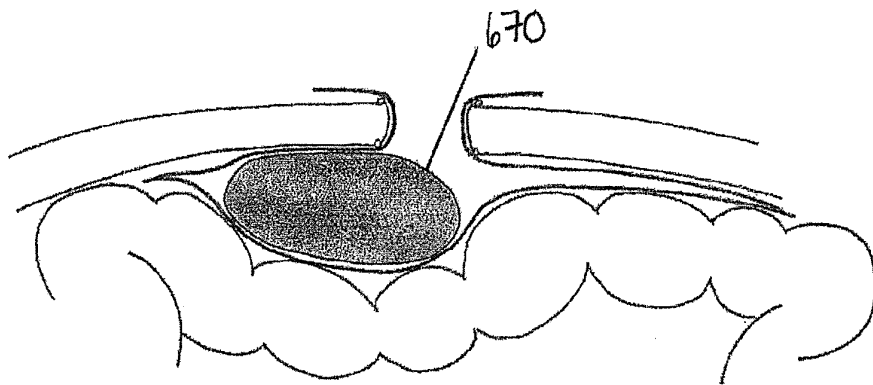


FIG. 183

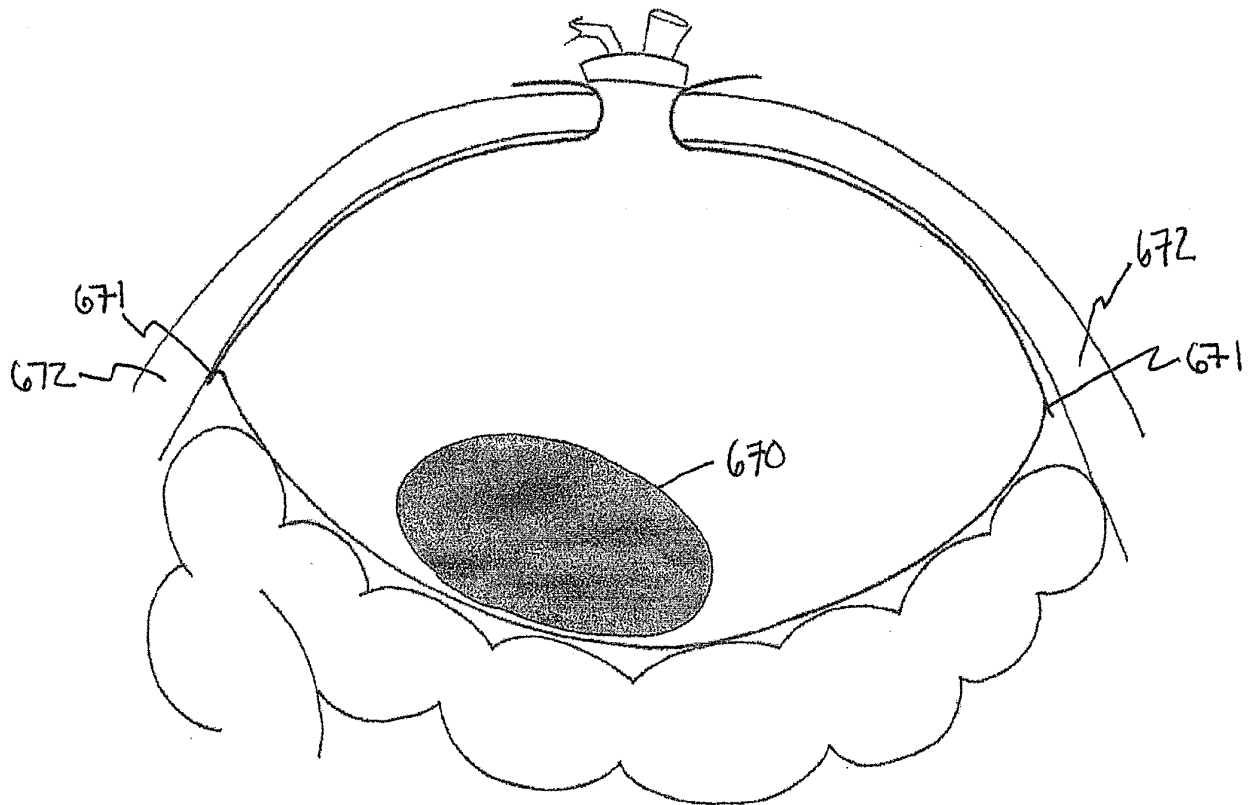
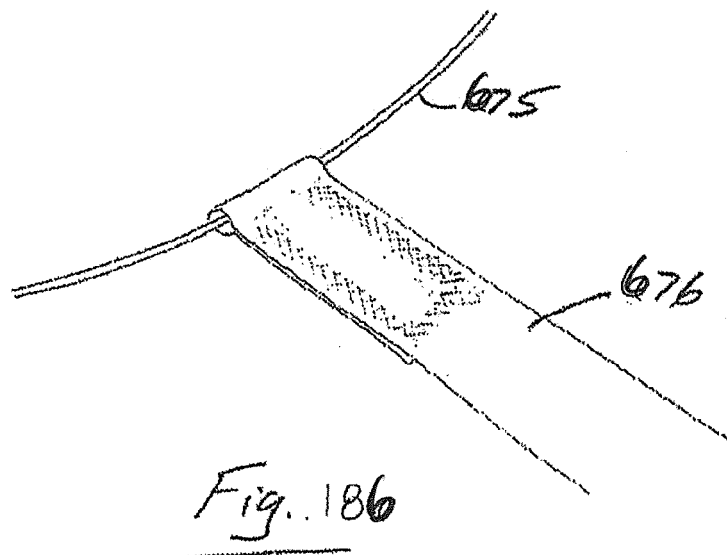
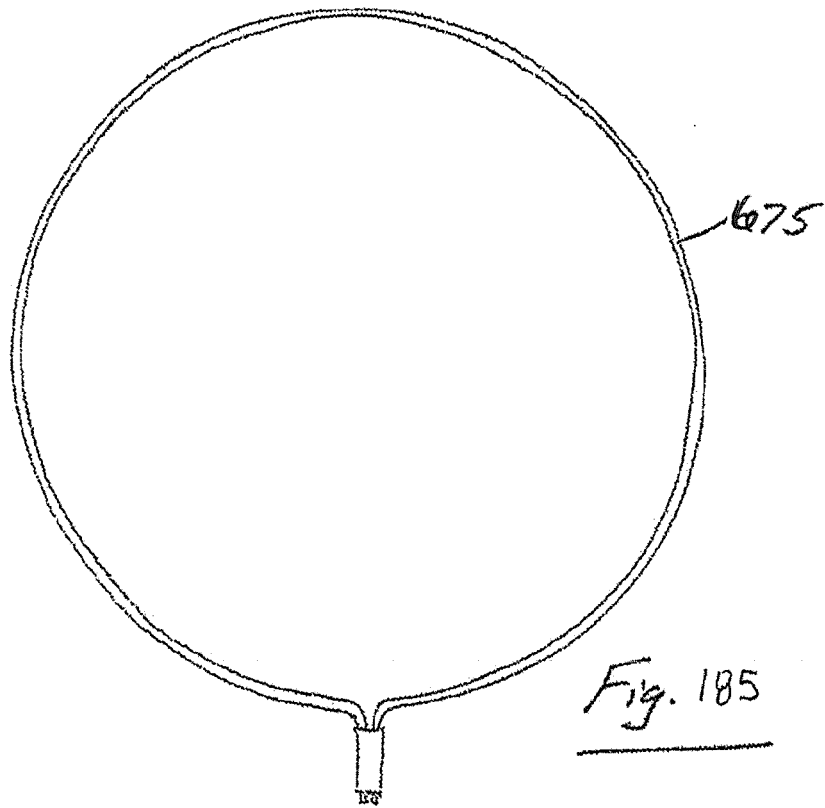
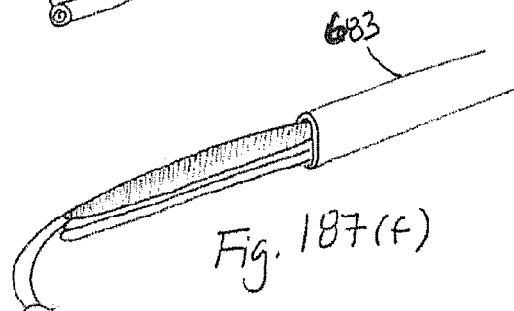
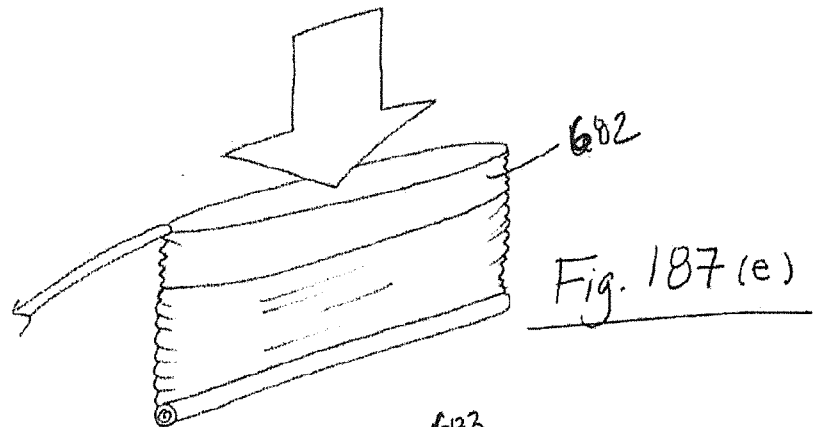
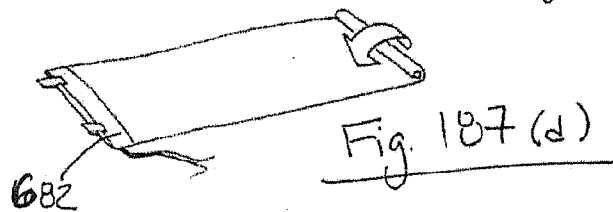
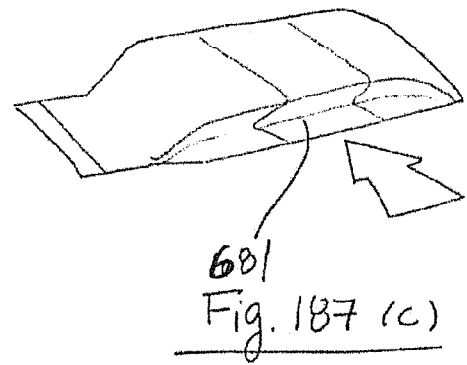
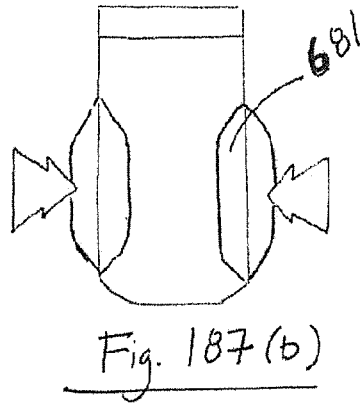
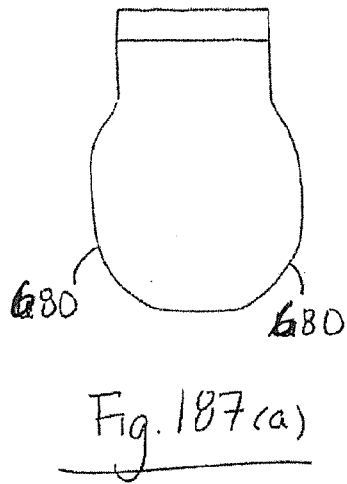


FIG. 184





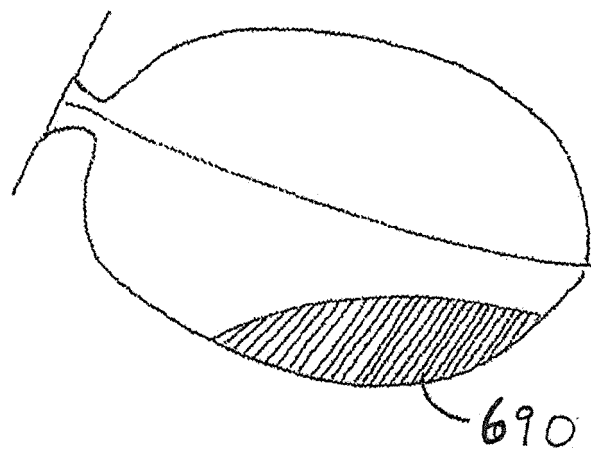


Fig. 188

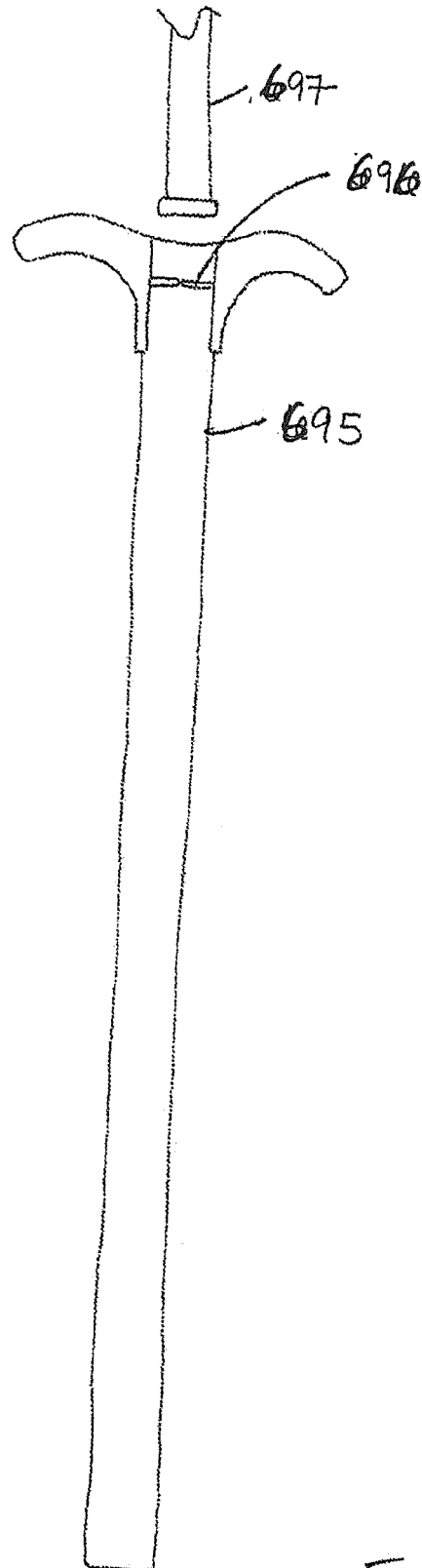


Fig. 189

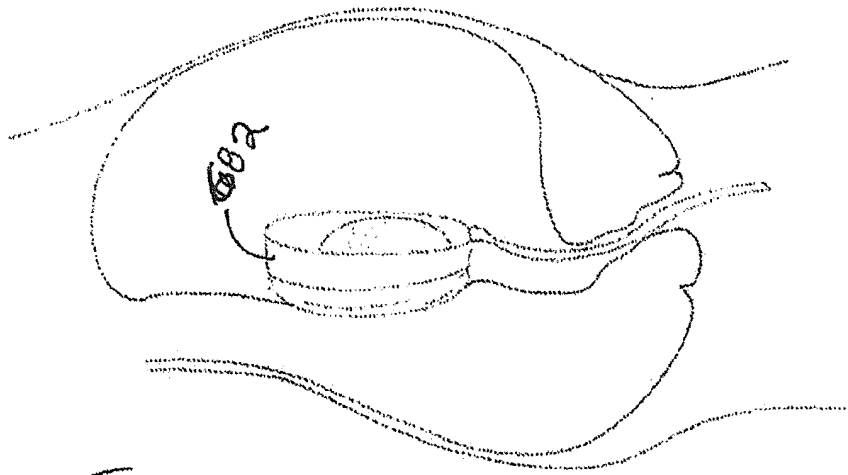


Fig. 190(a)

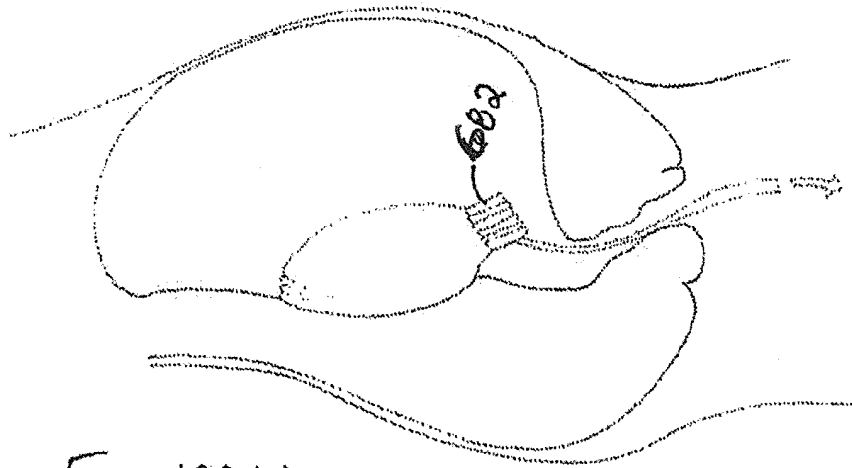


Fig. 190(b)

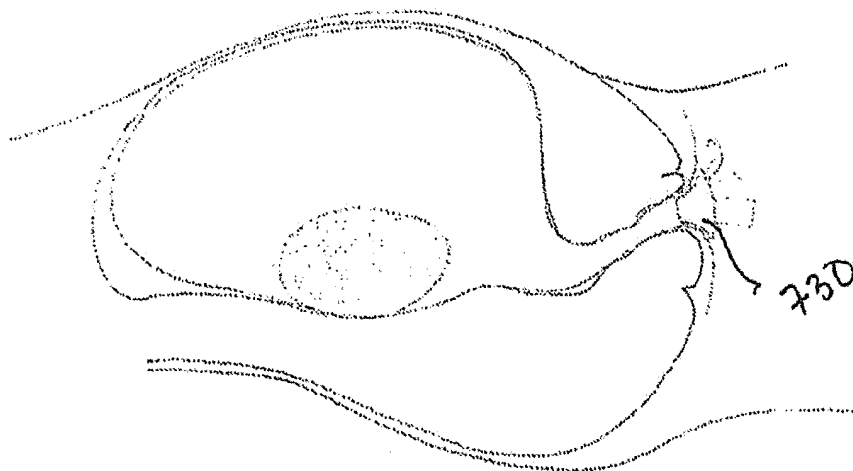


Fig. 190(c)

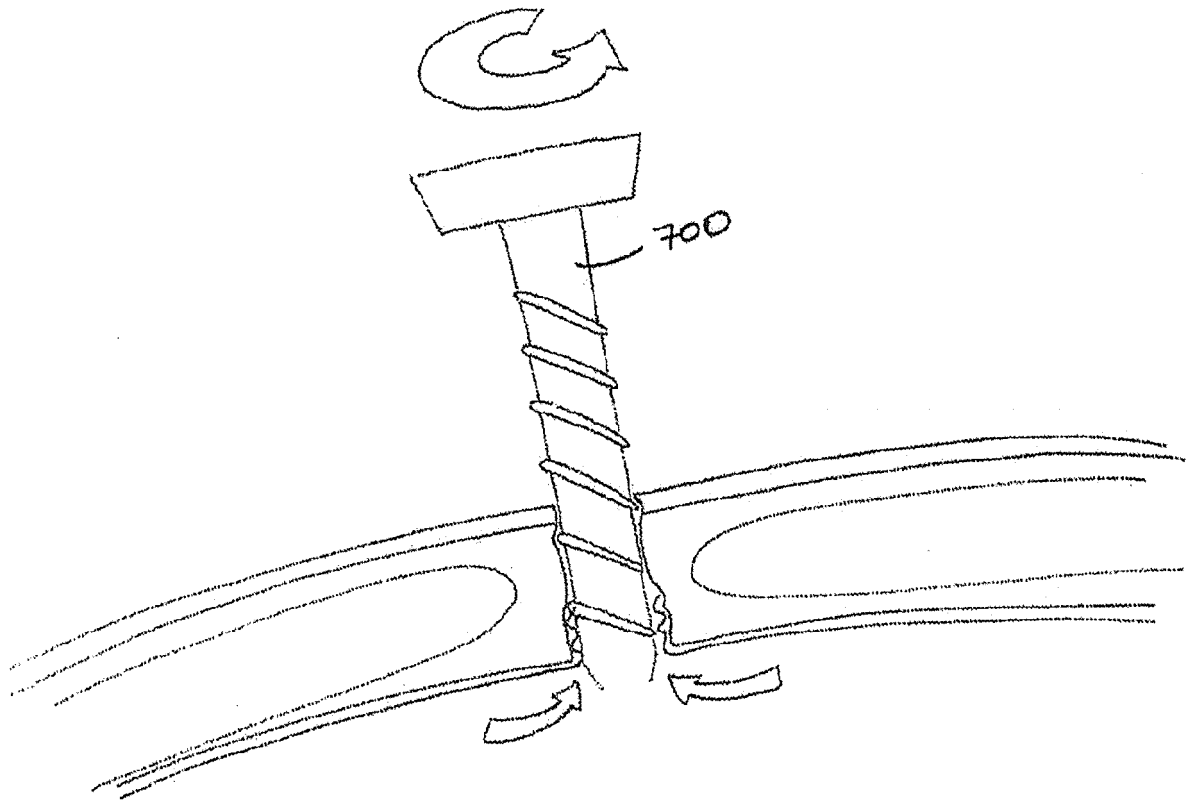
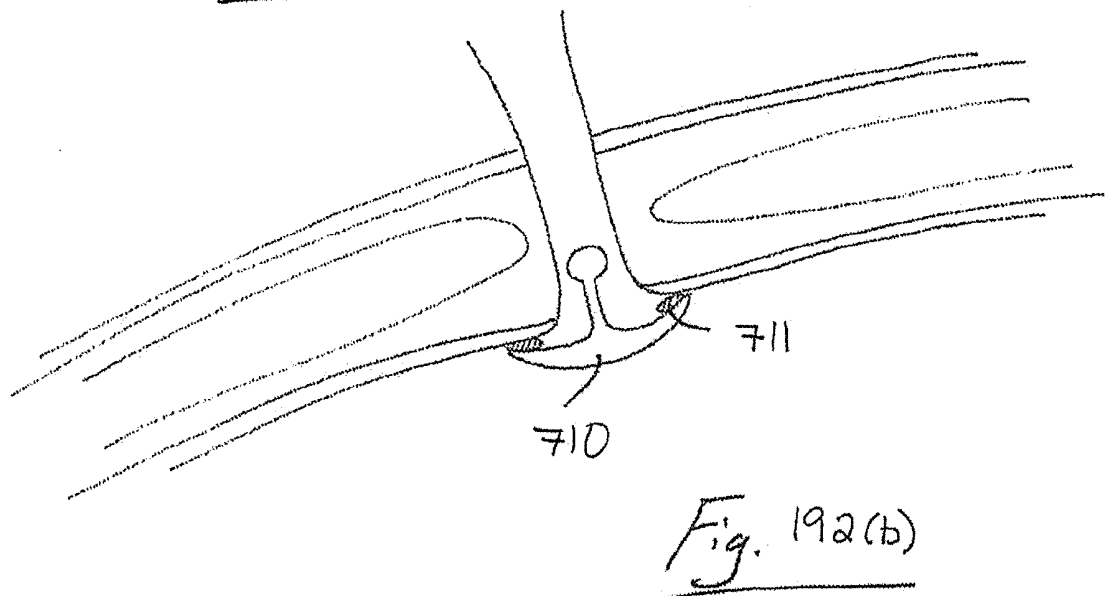
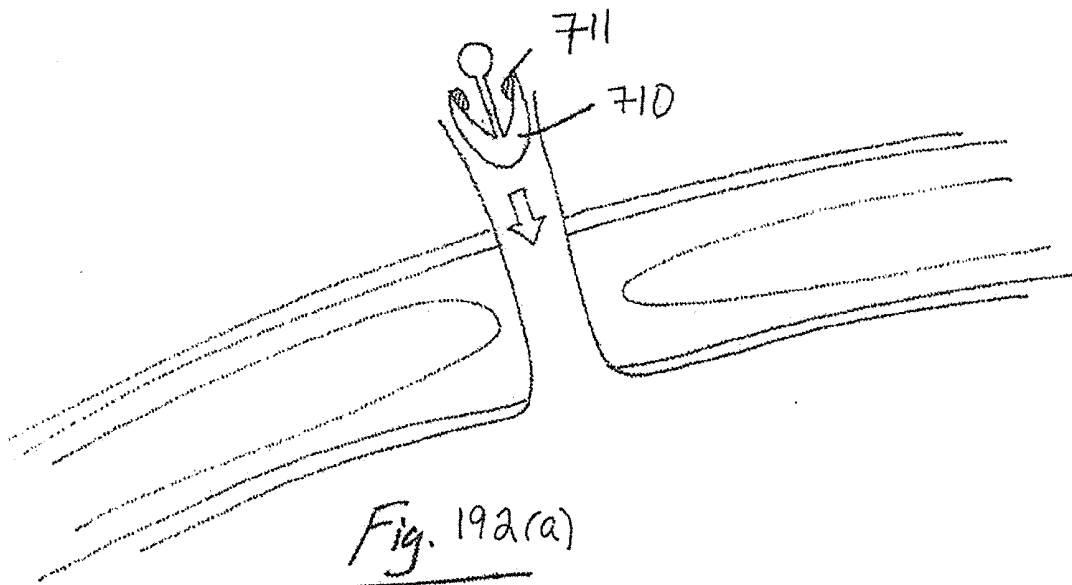


Fig. 191



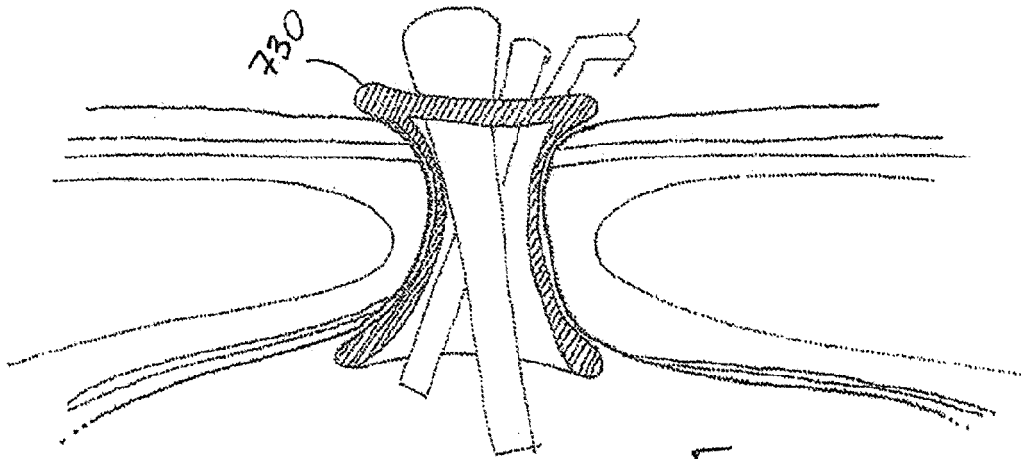


Fig. 193(a)

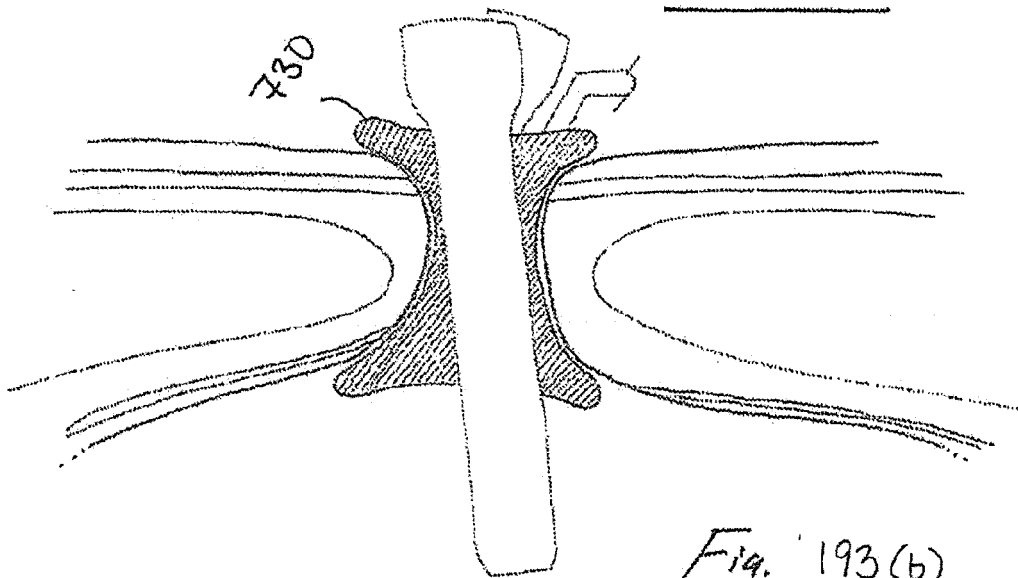


Fig. 193(b)

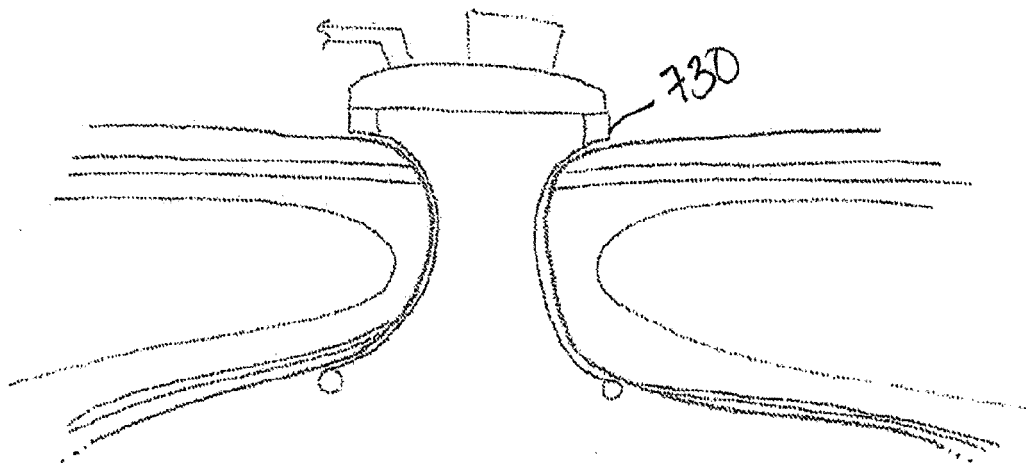
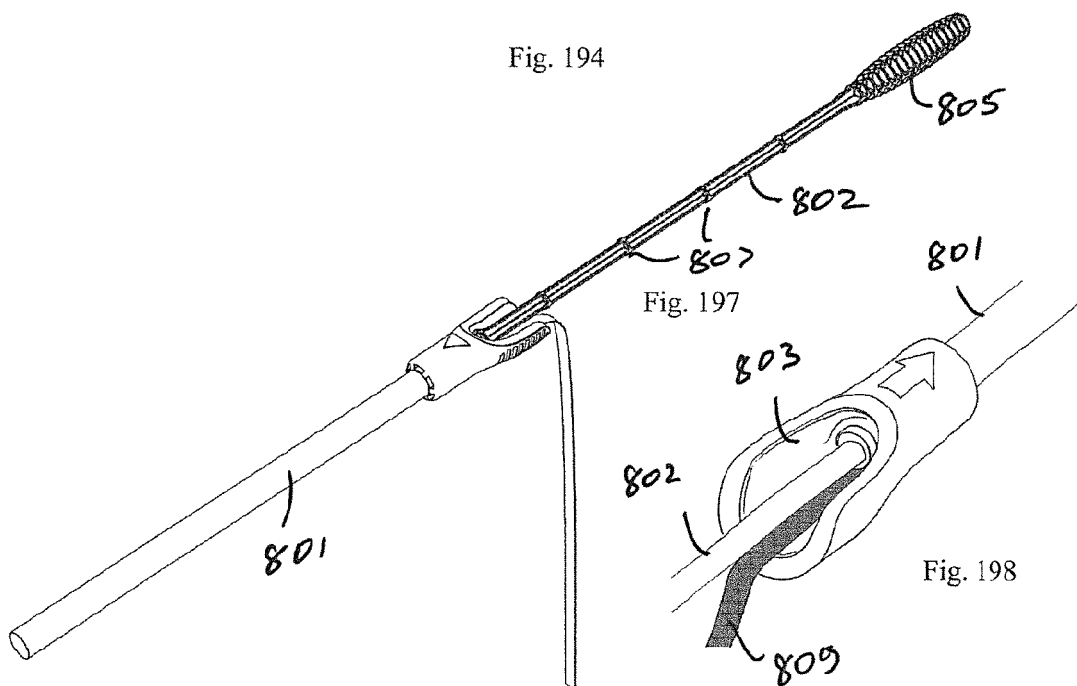
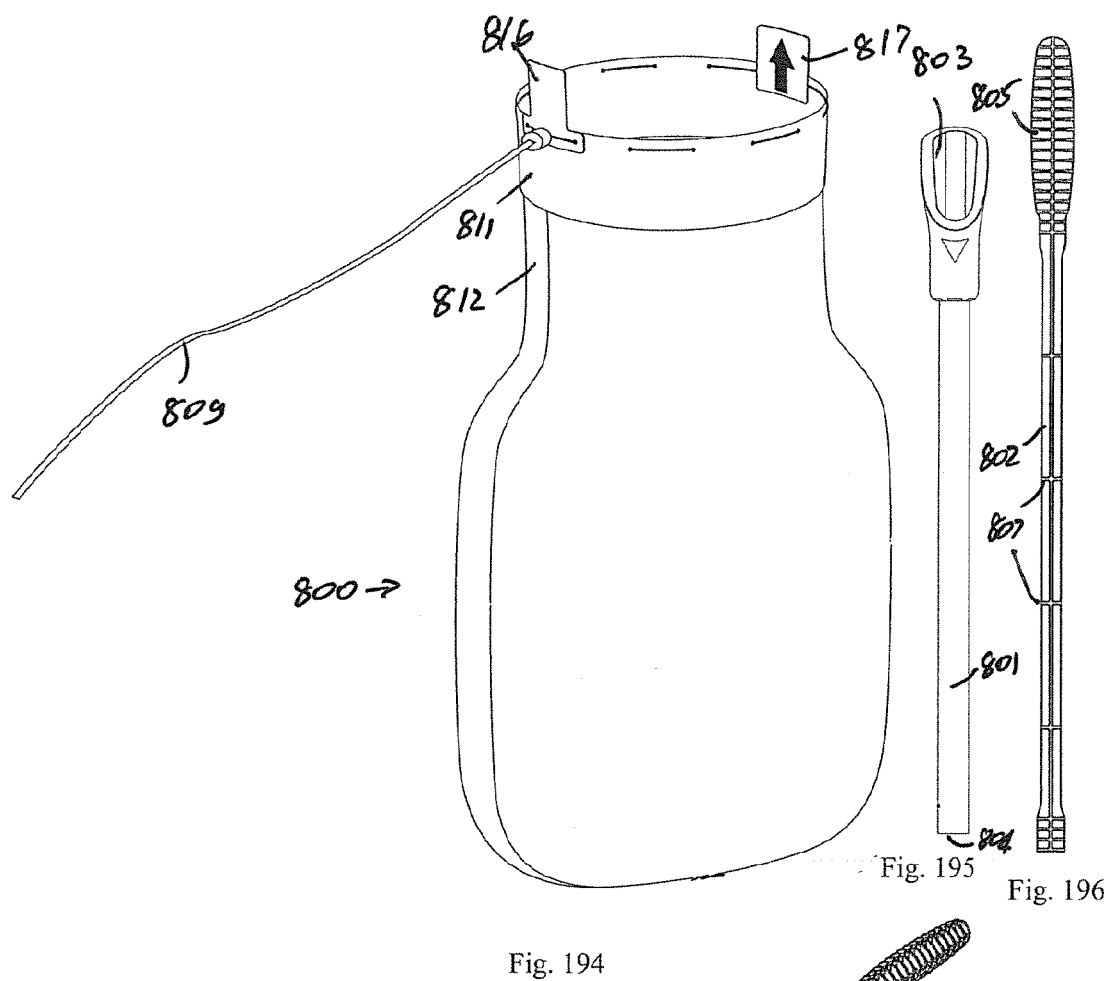


Fig. 193(c)



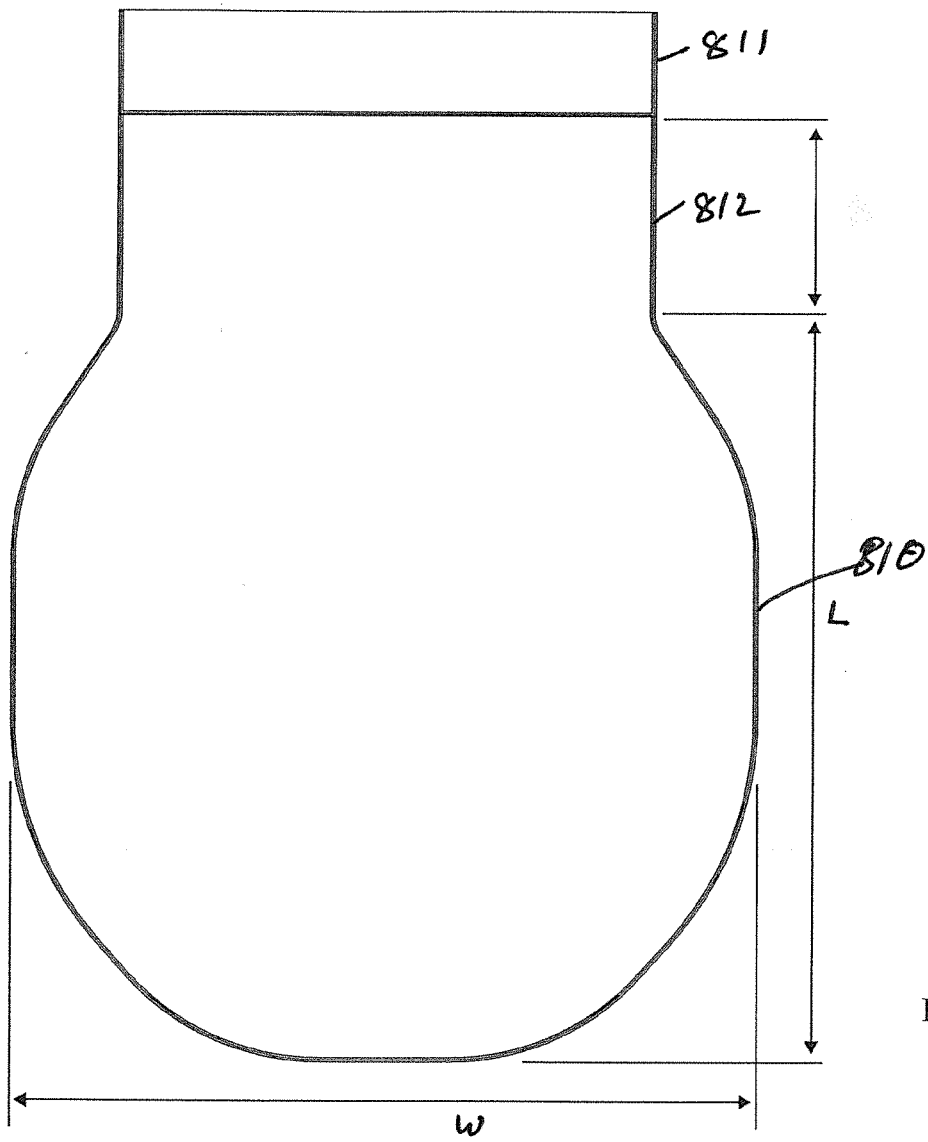


Fig. 199

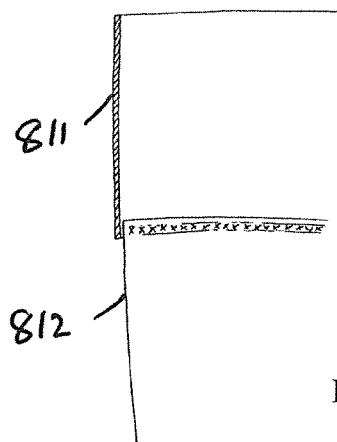
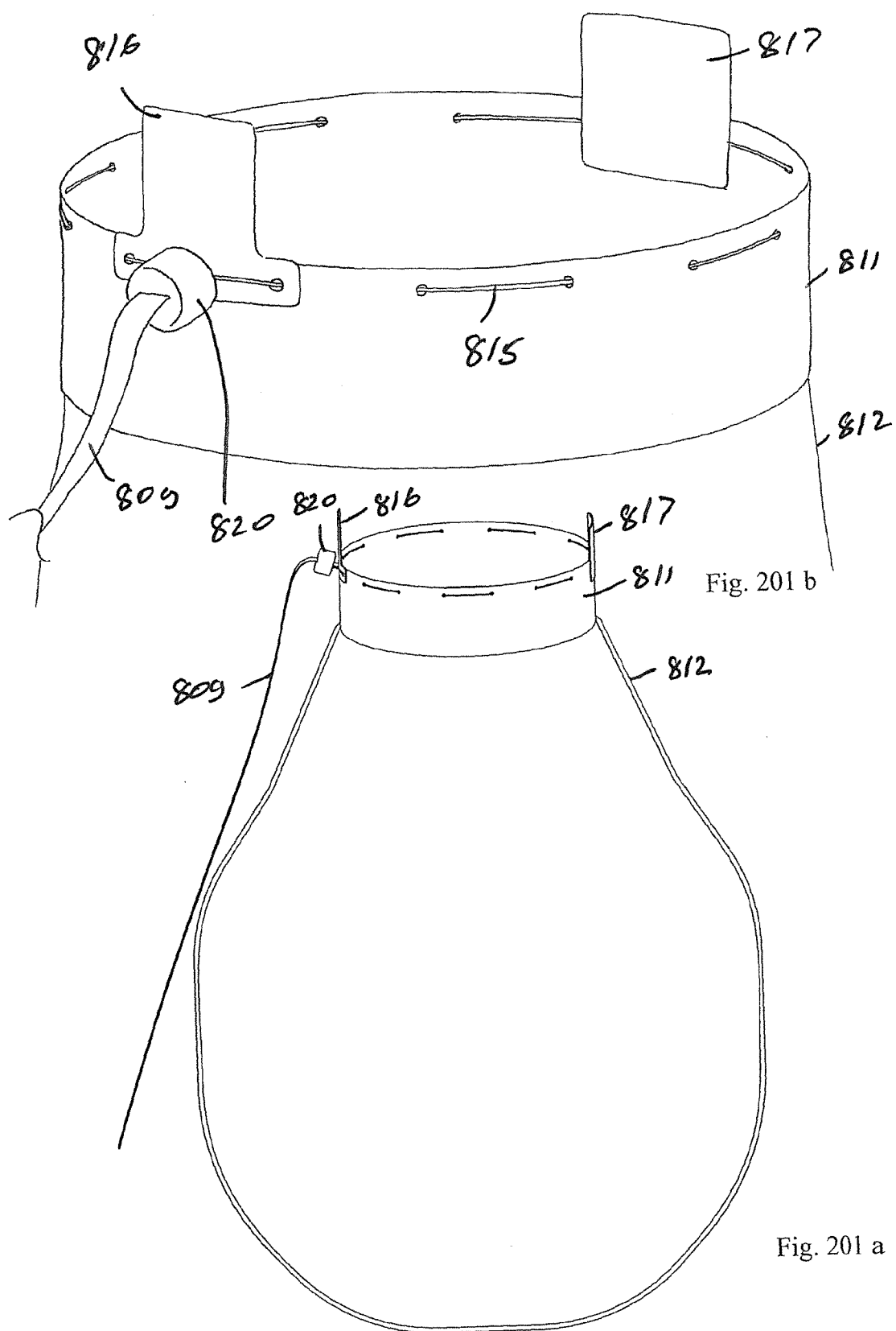
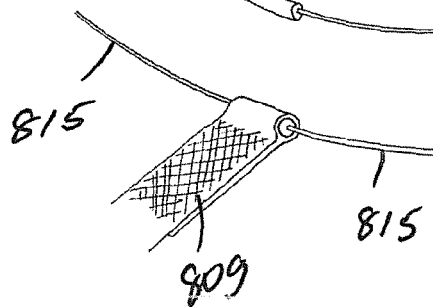
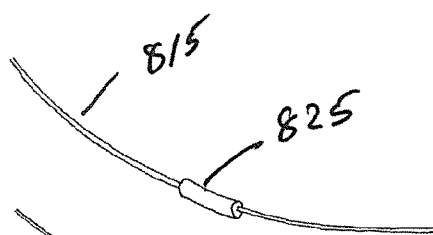
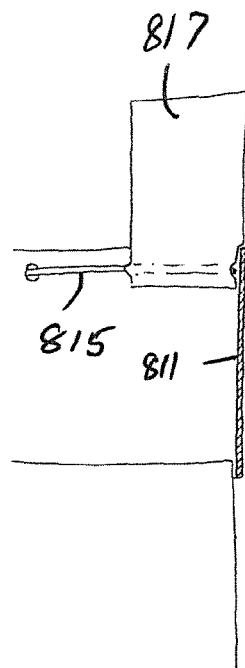
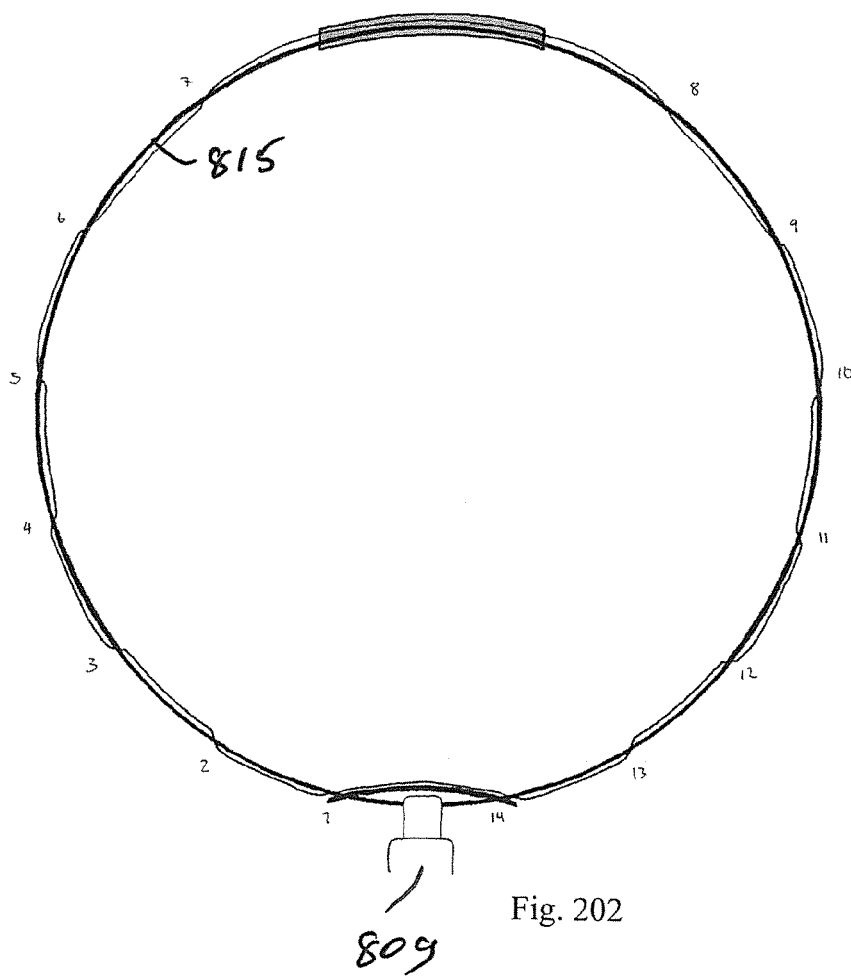


Fig. 200





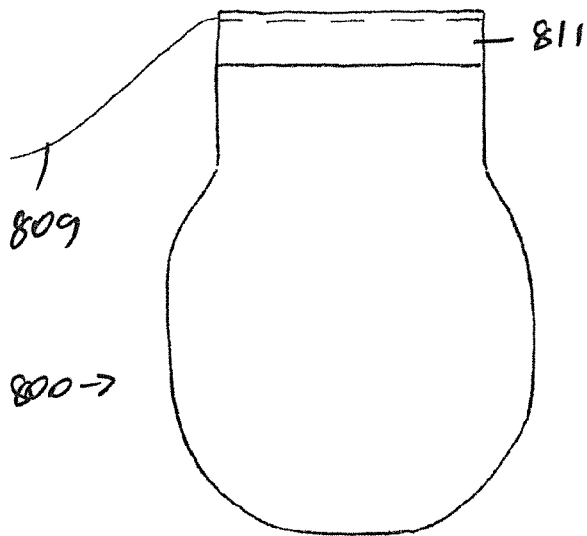


Fig. 206

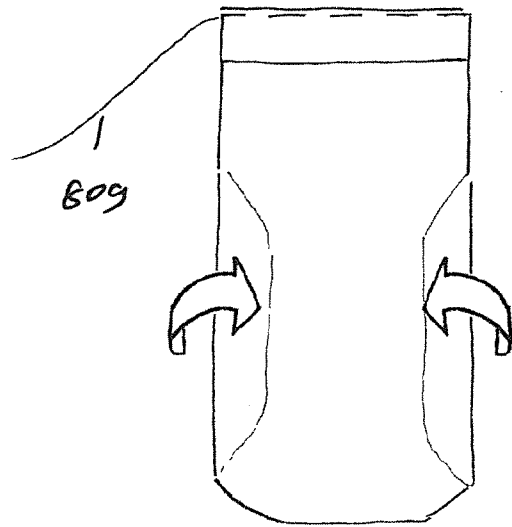


Fig. 207

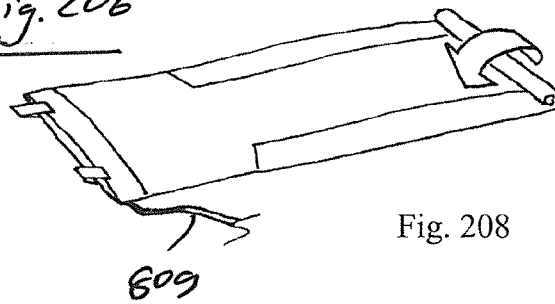


Fig. 208

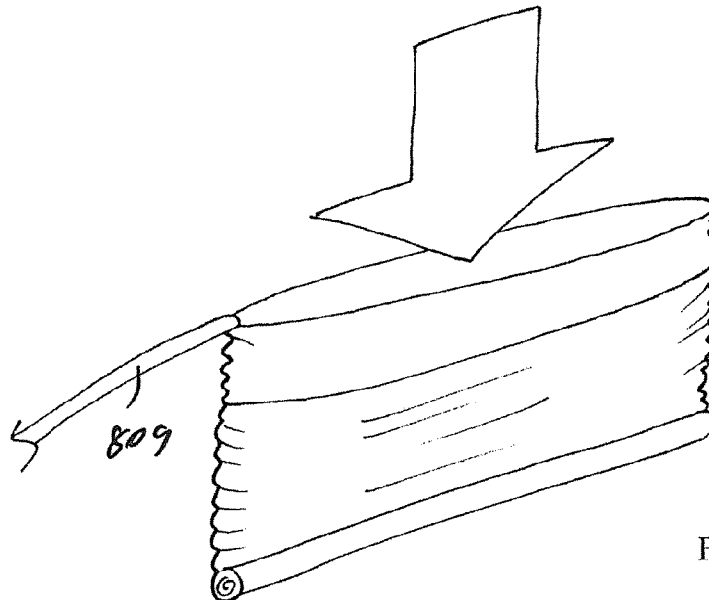


Fig. 209

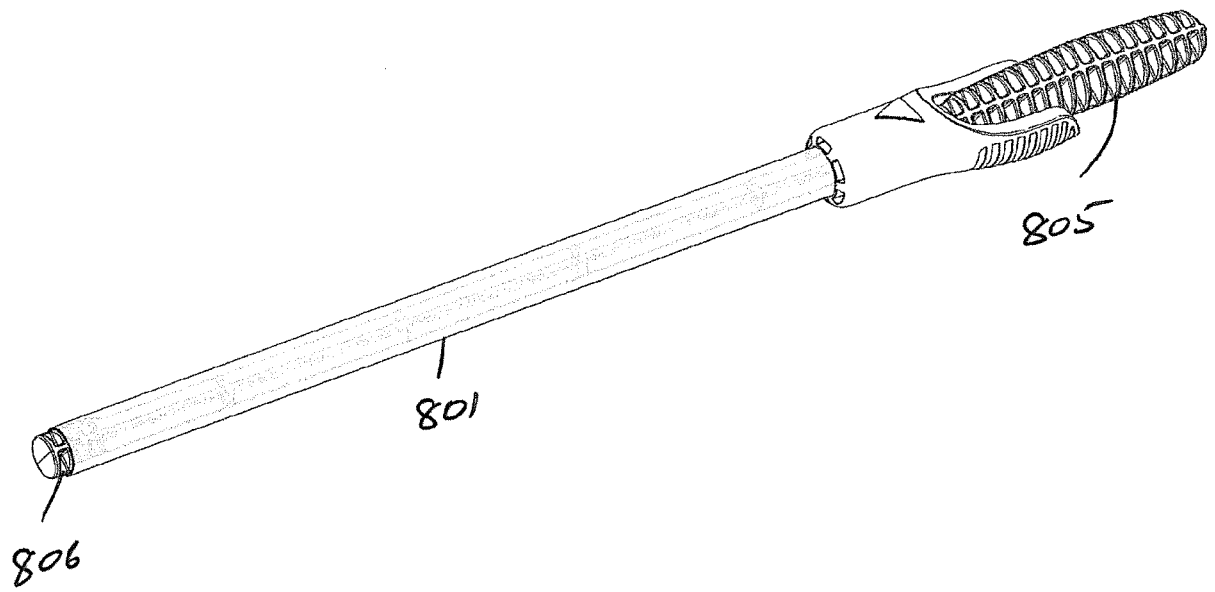


Fig. 210

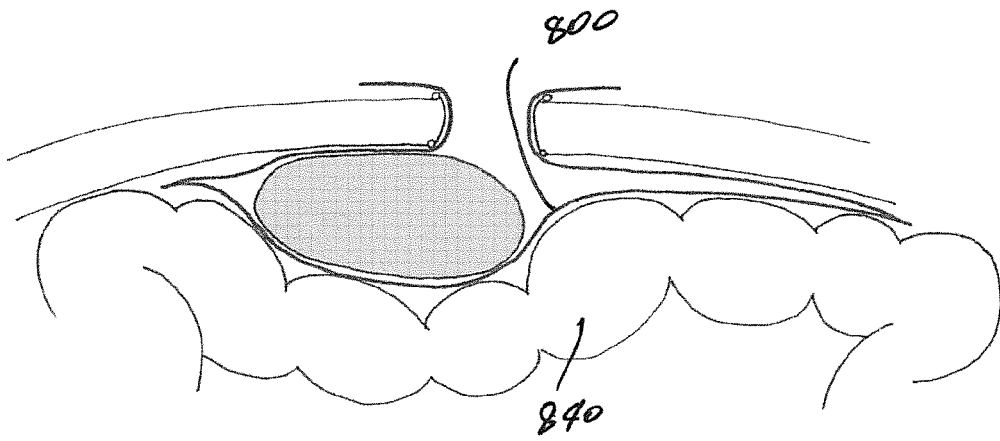


Fig. 211

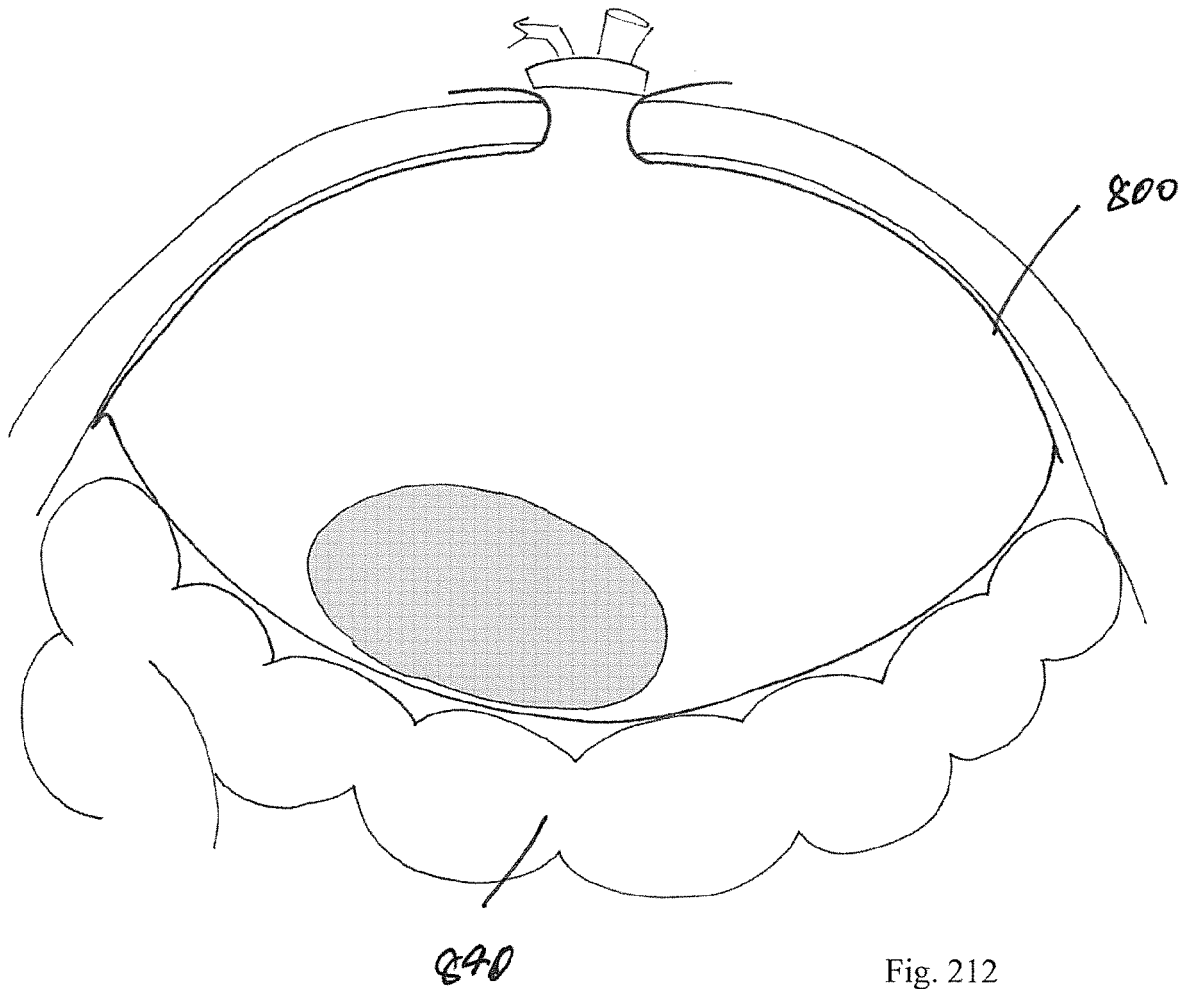


Fig. 212

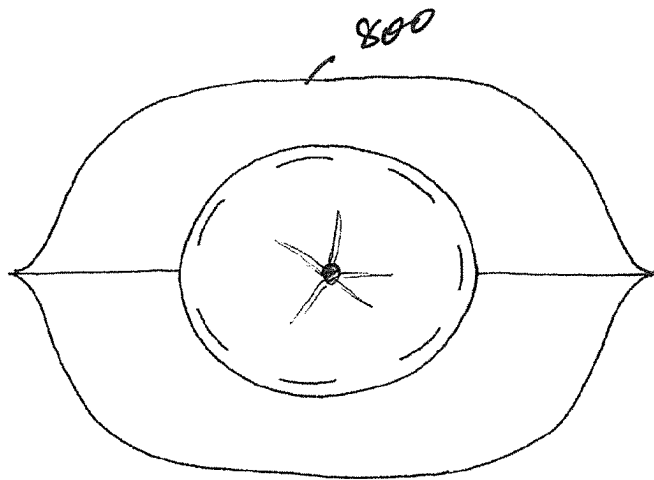


Fig. 213

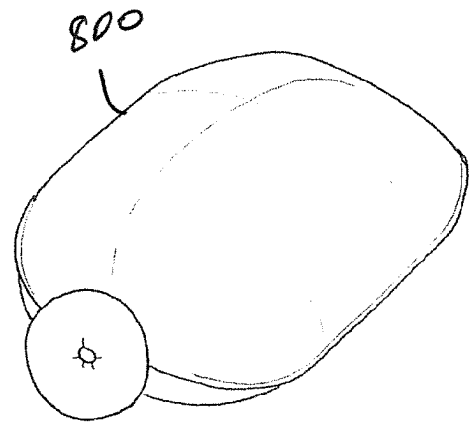


Fig. 214

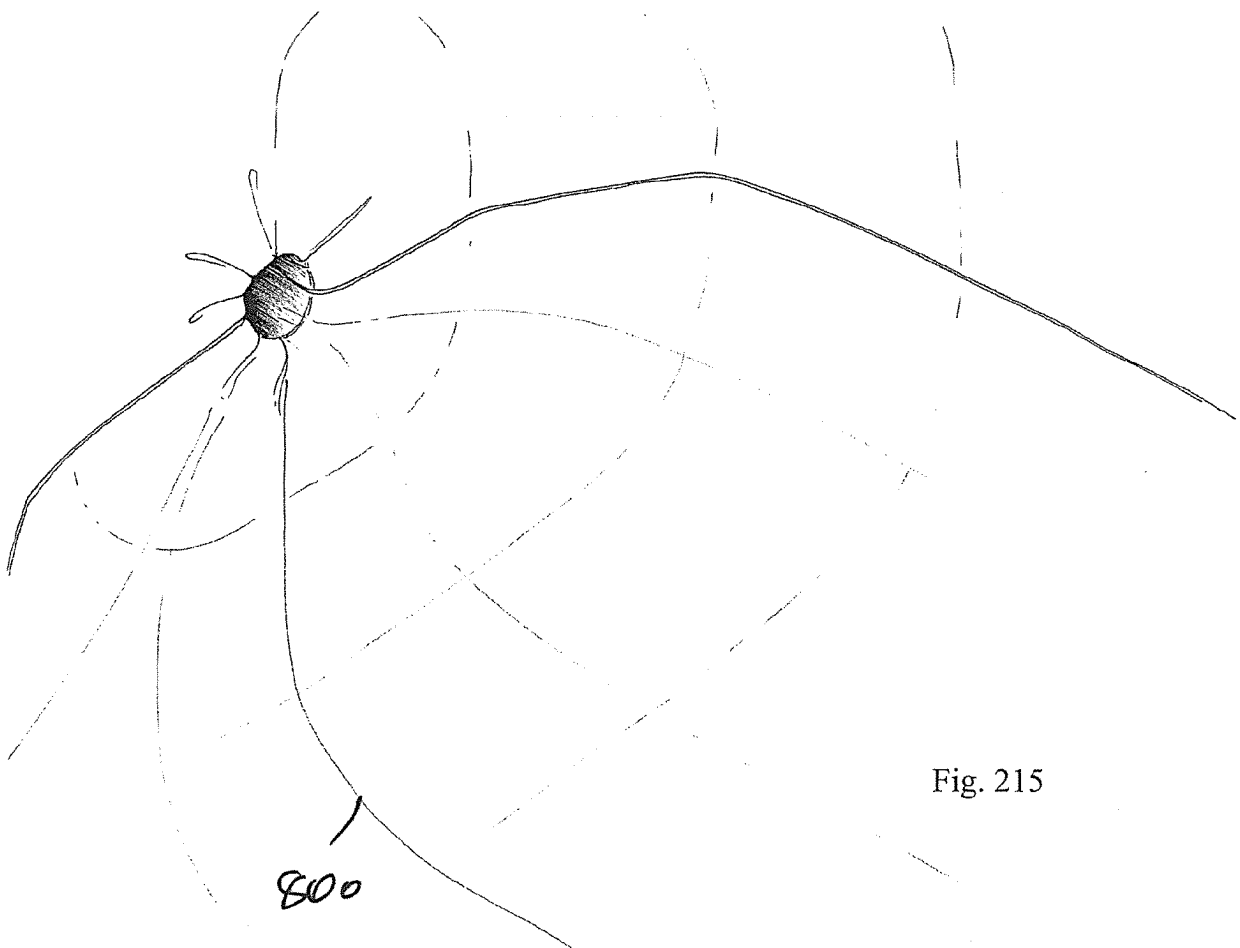


Fig. 215

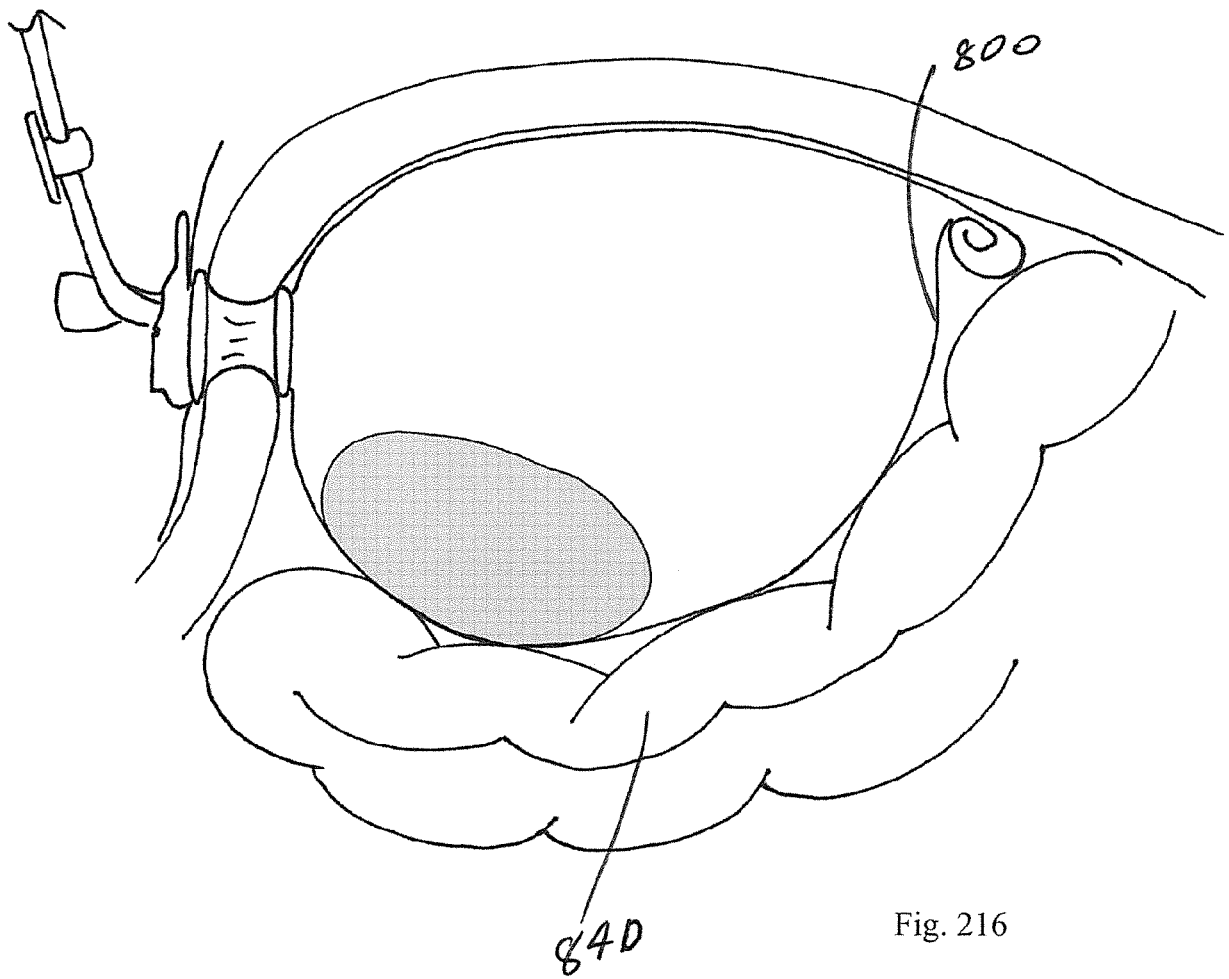


Fig. 216

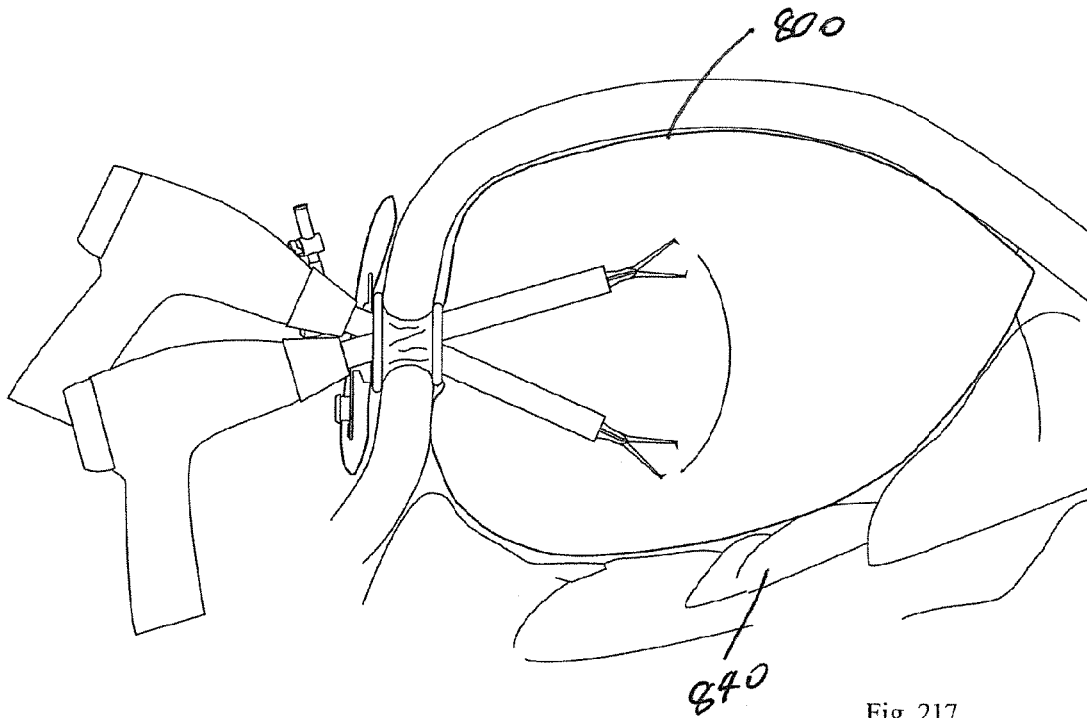


Fig. 217

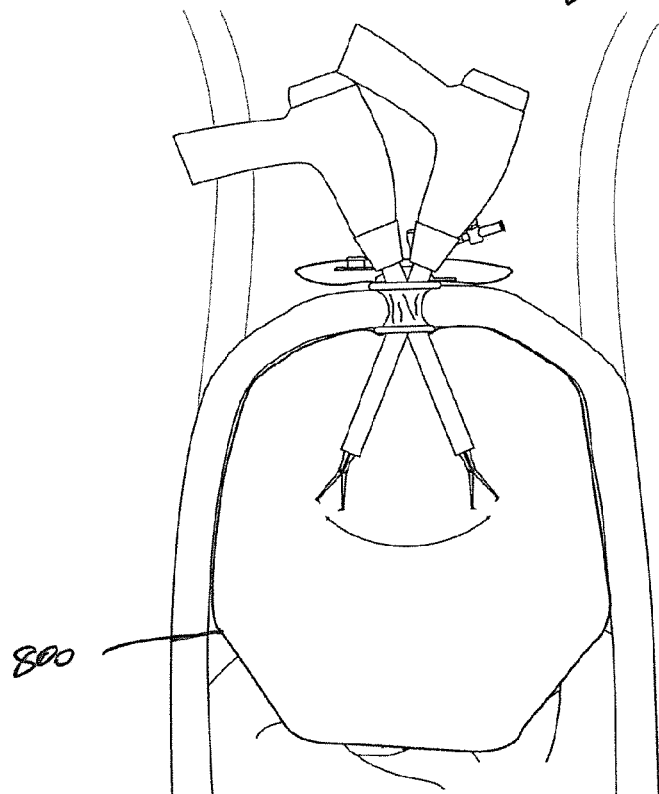


Fig. 218

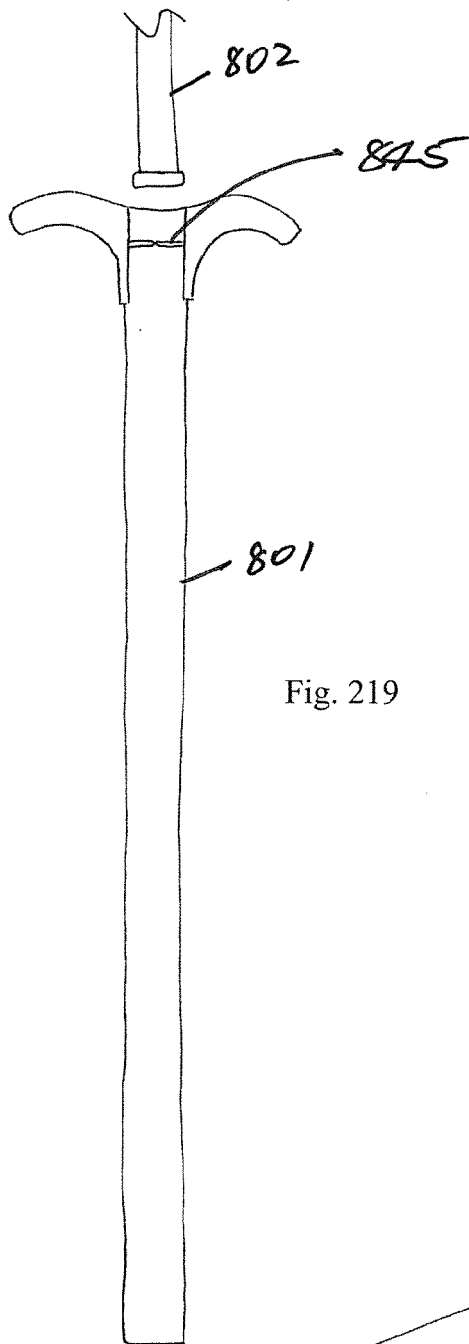


Fig. 219

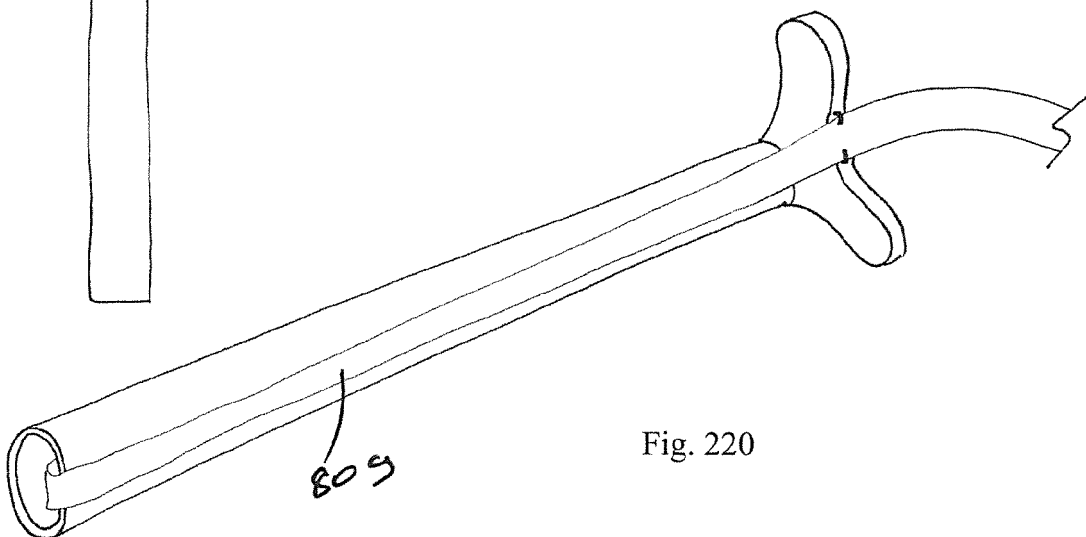


Fig. 220

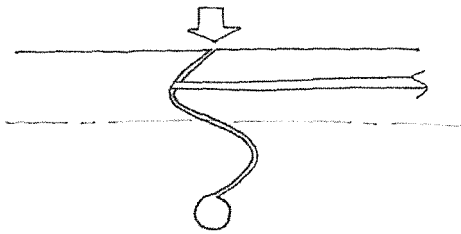


Fig. 222a

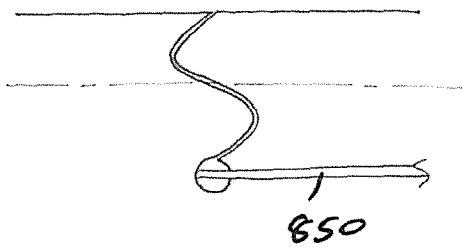


Fig. 222b

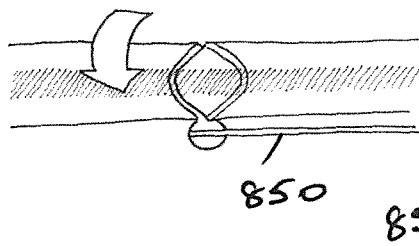


Fig. 222c

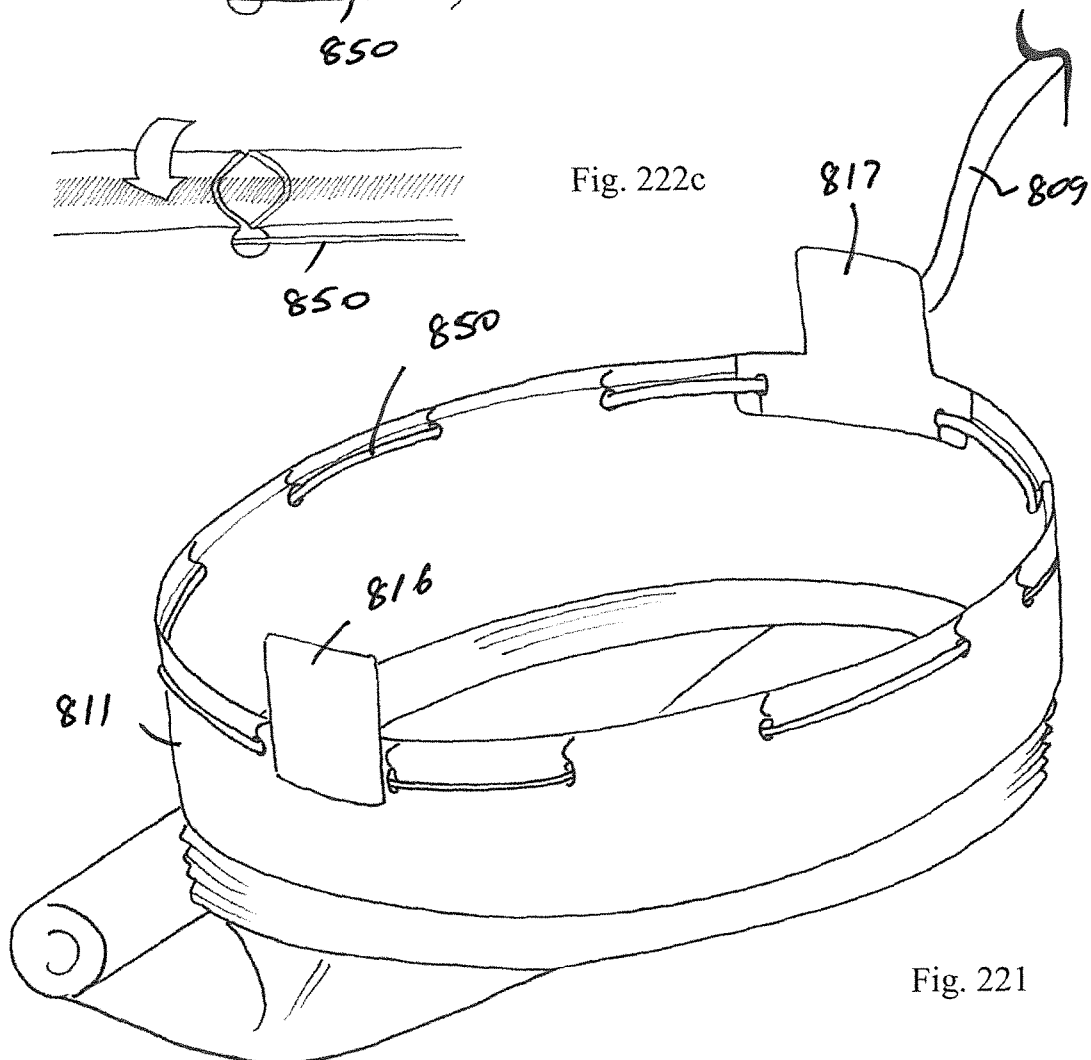


Fig. 221

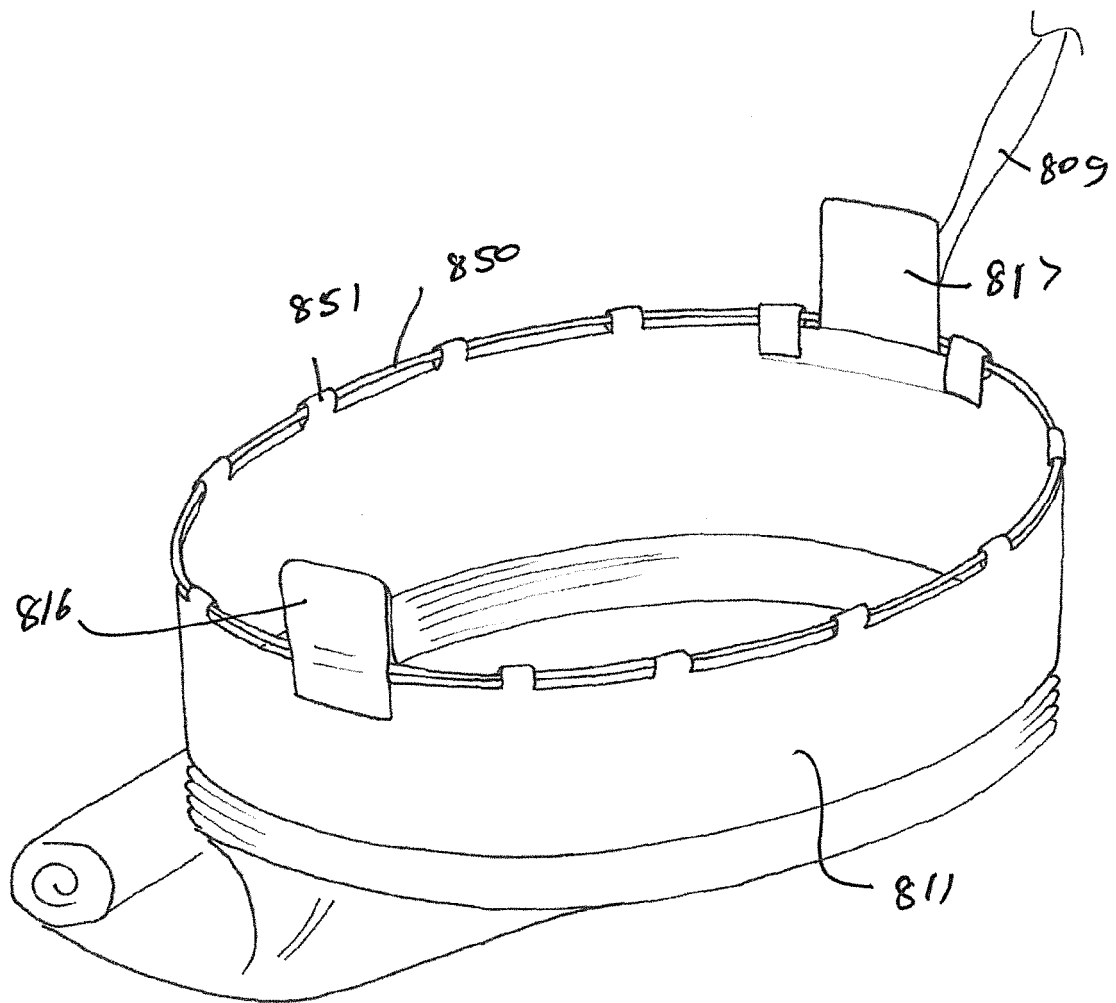
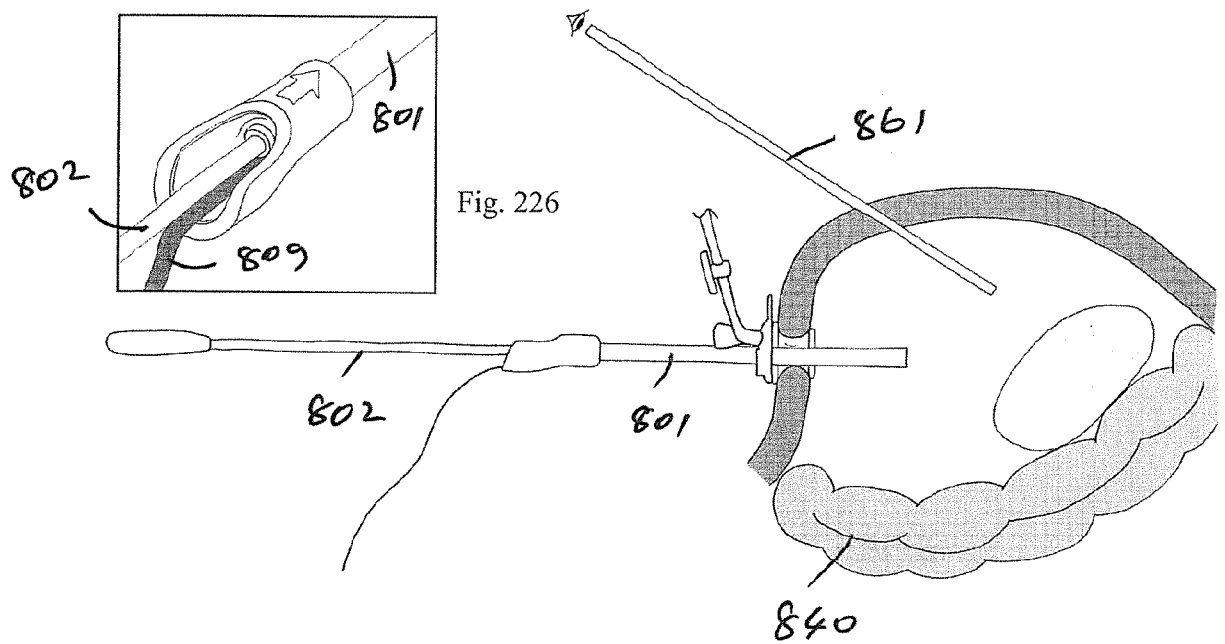
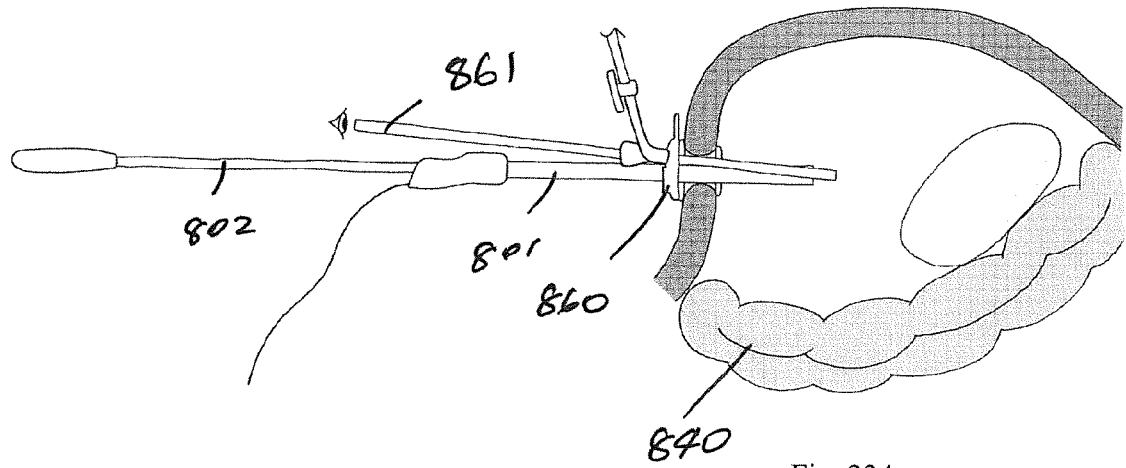


Fig. 223



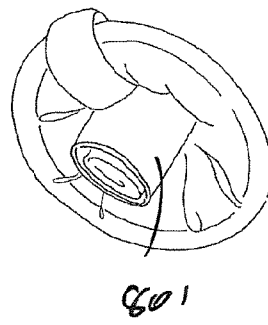


Fig. 227

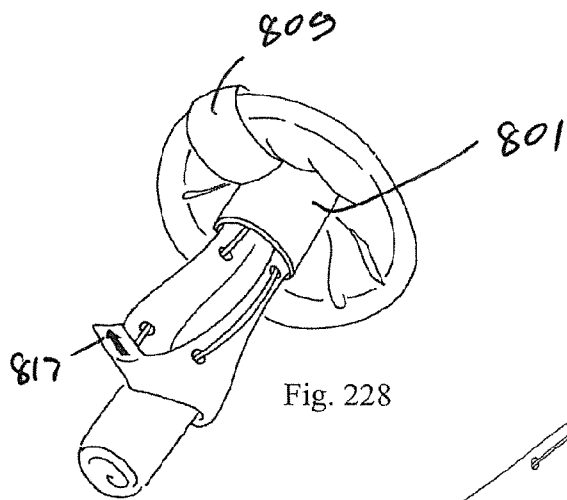


Fig. 228

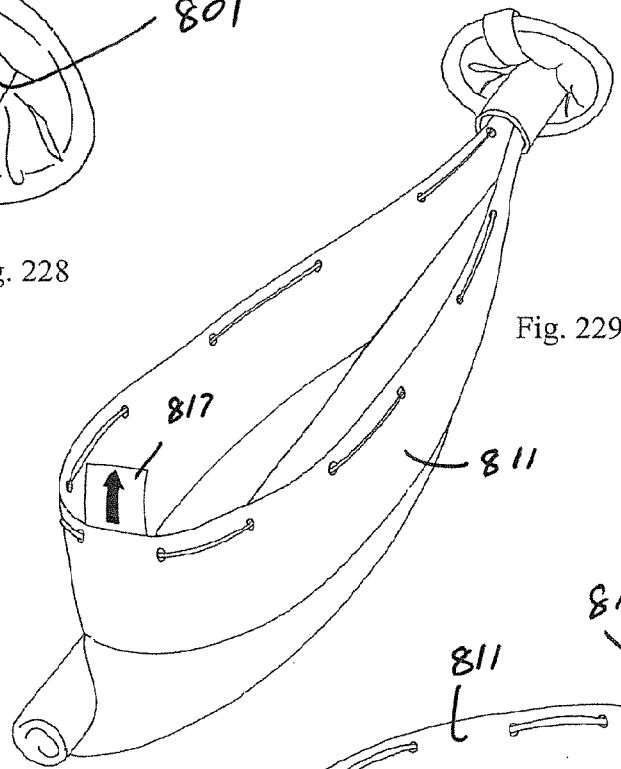


Fig. 229

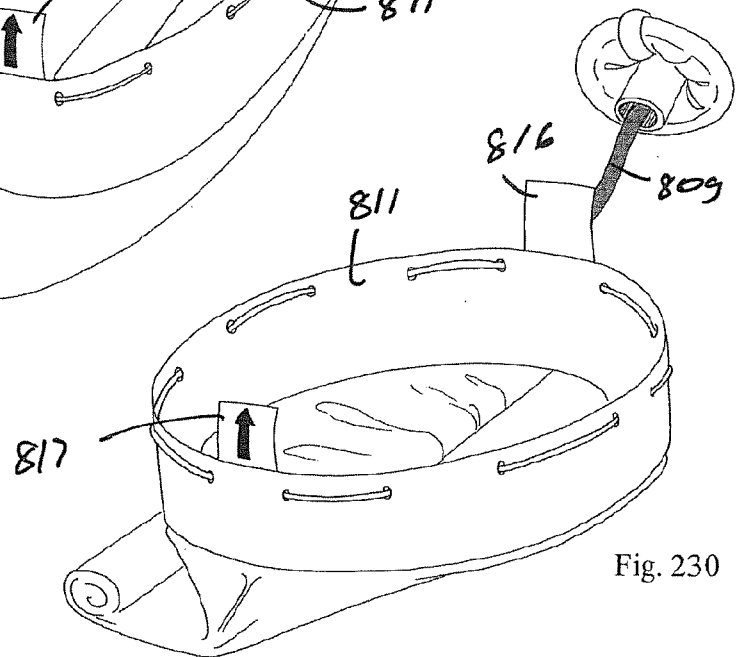
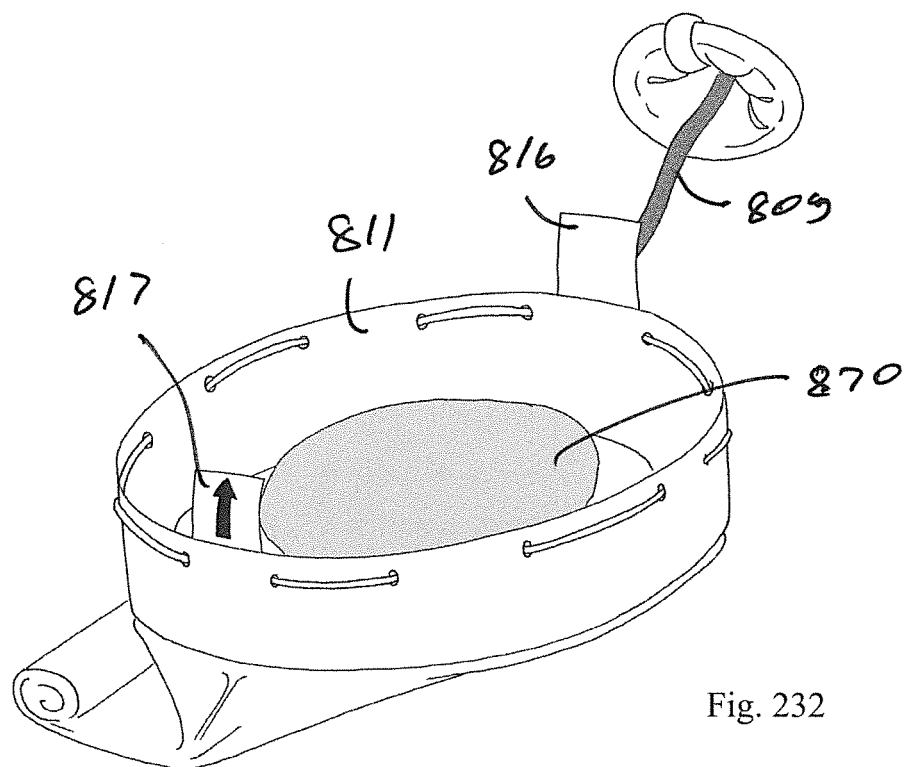
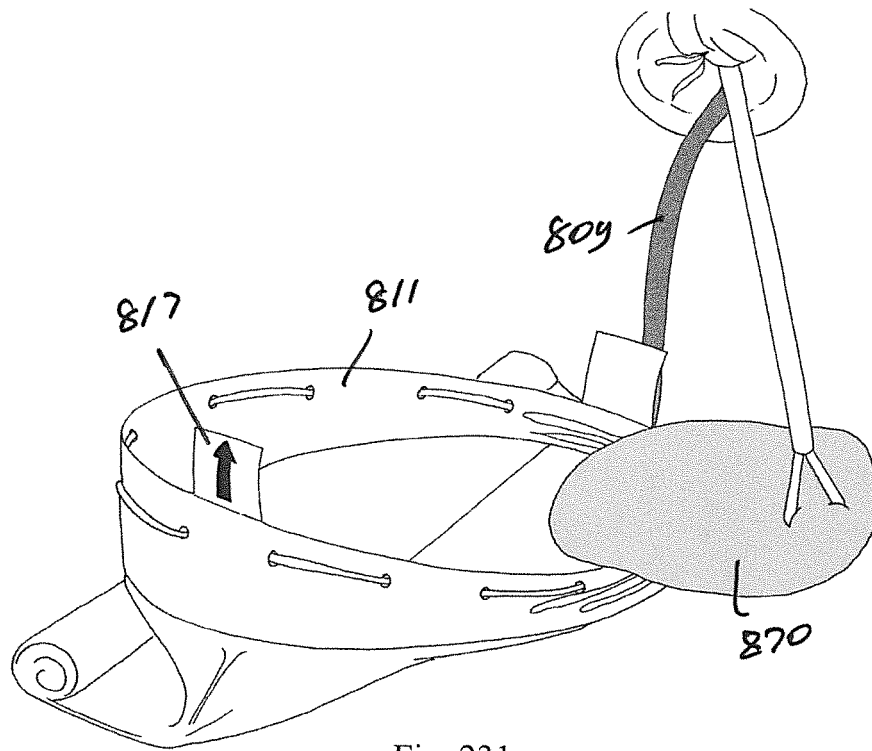


Fig. 230



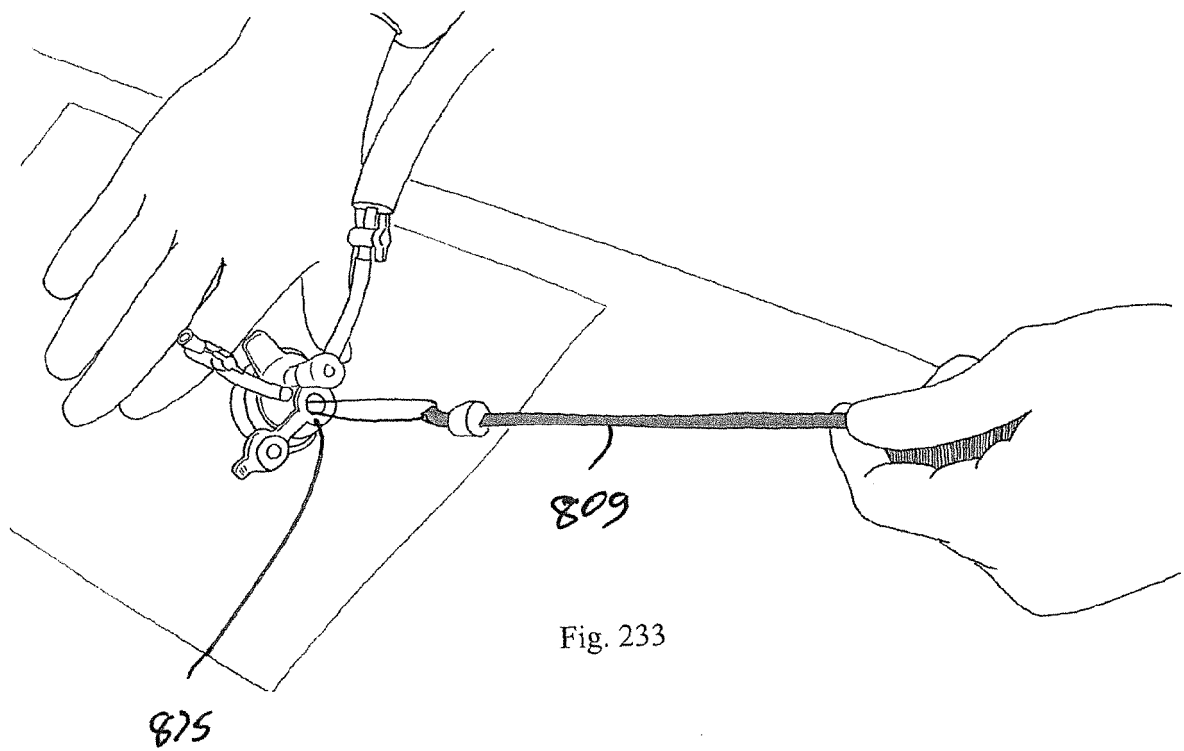


Fig. 233

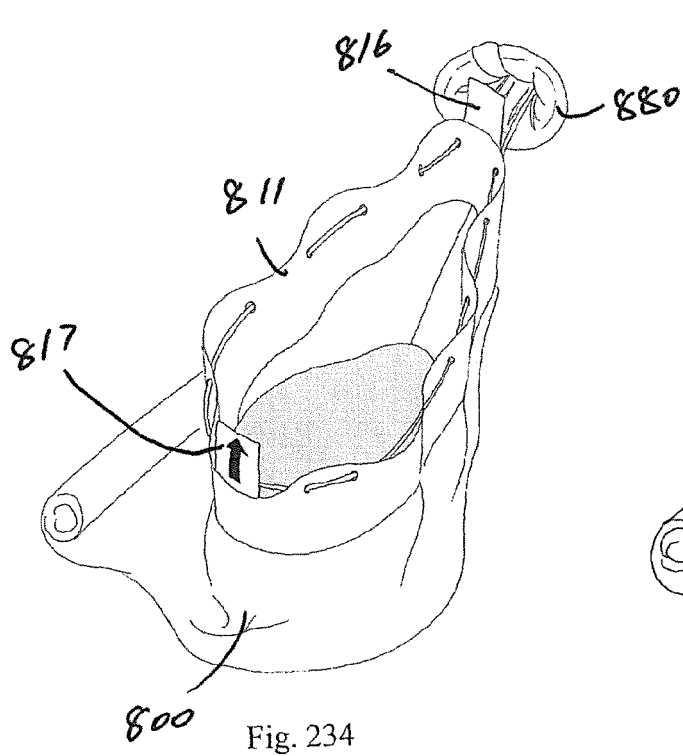


Fig. 234

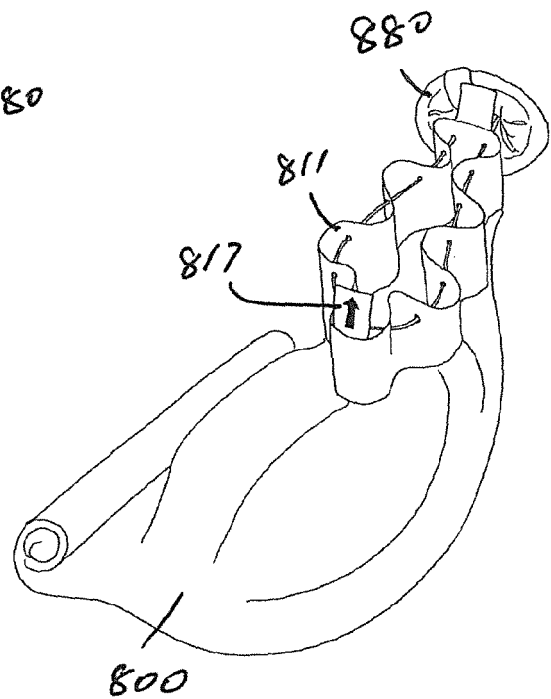


Fig. 235

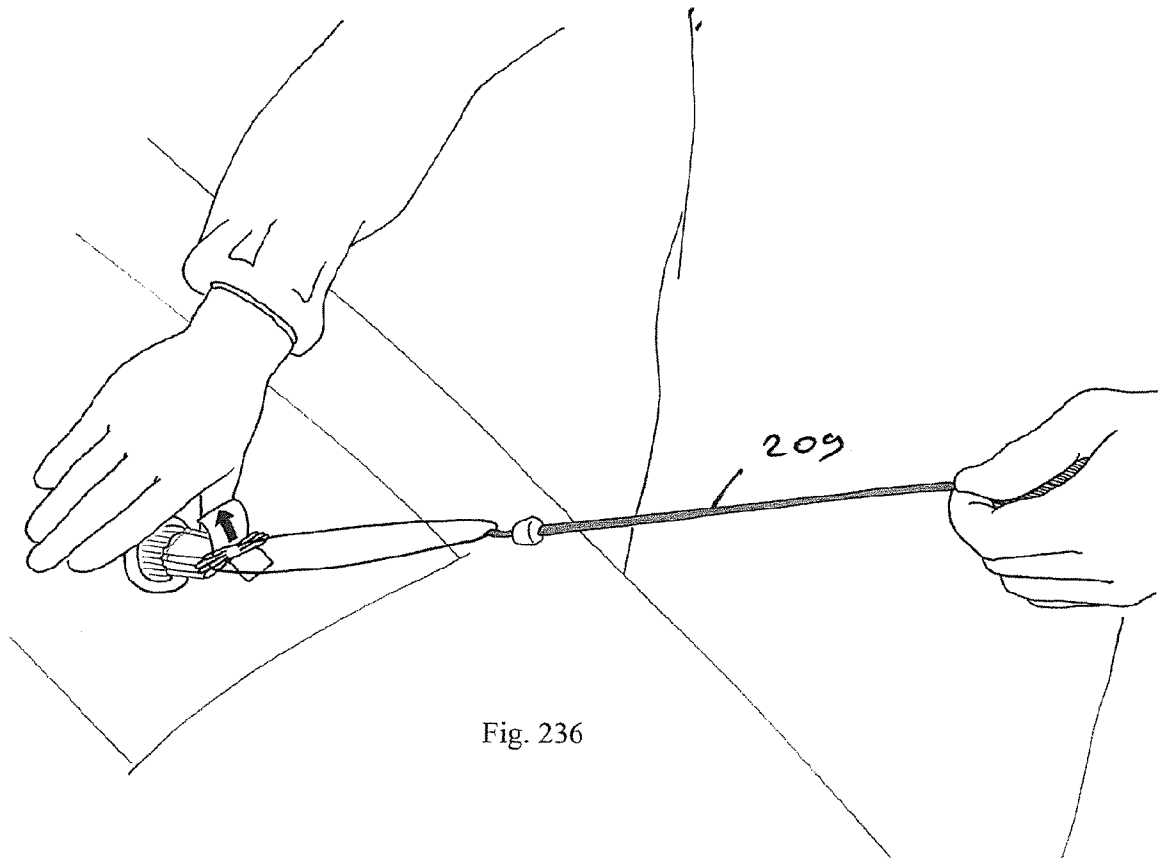


Fig. 236

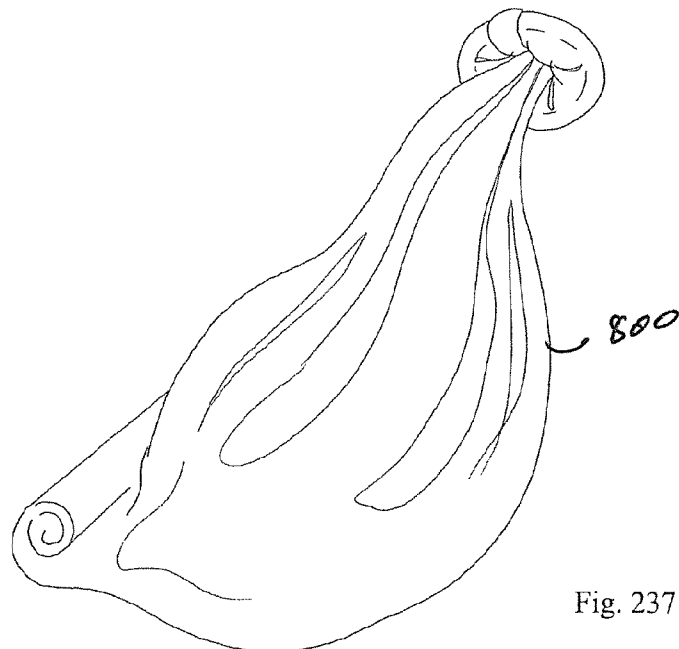


Fig. 237

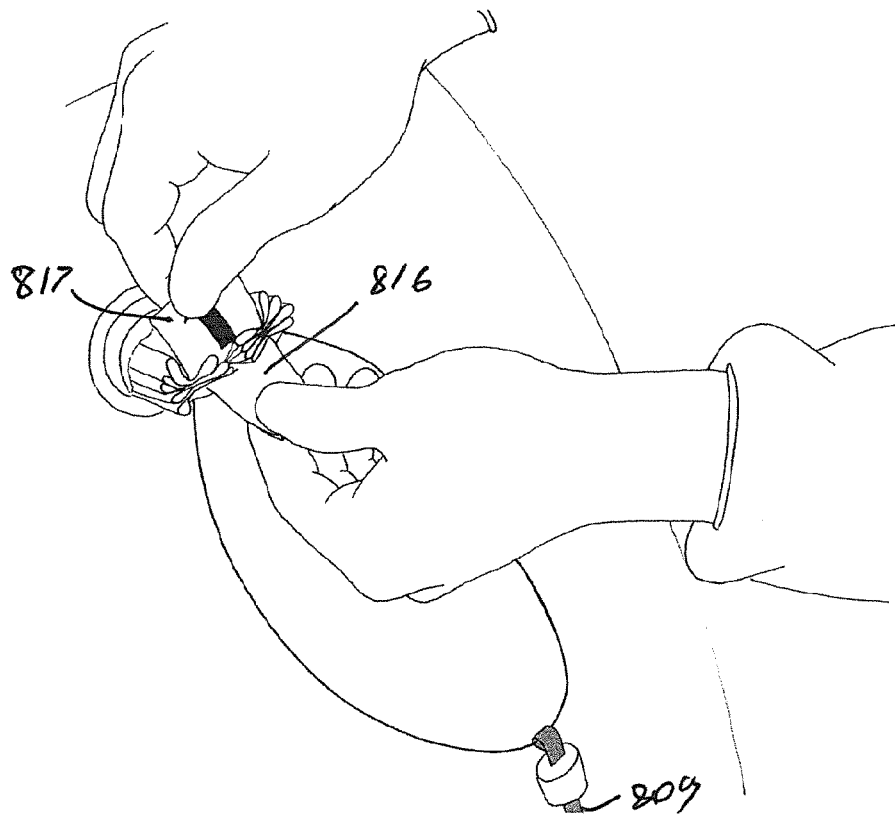


Fig. 238

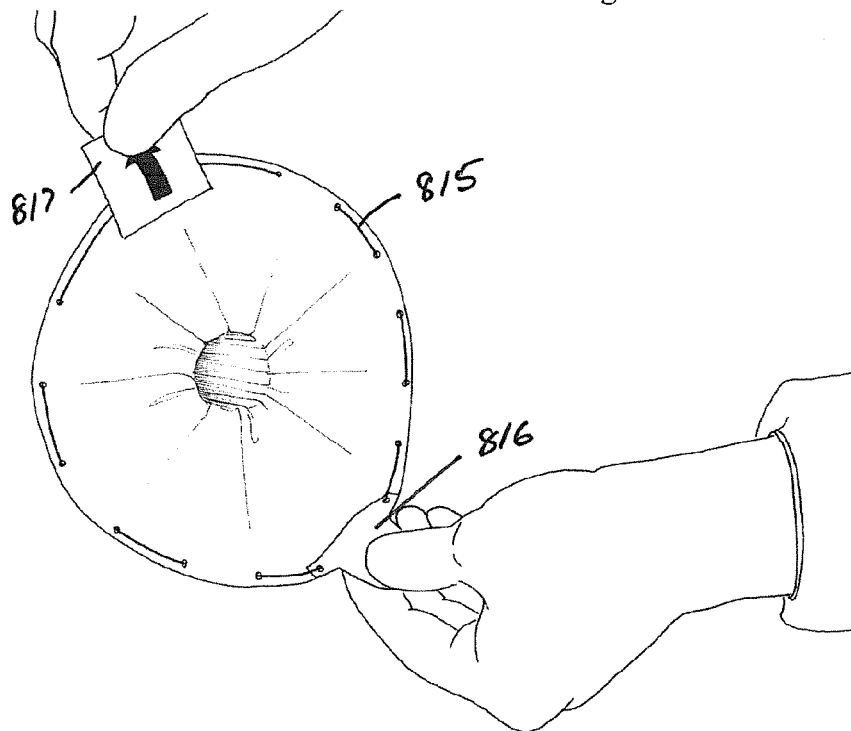
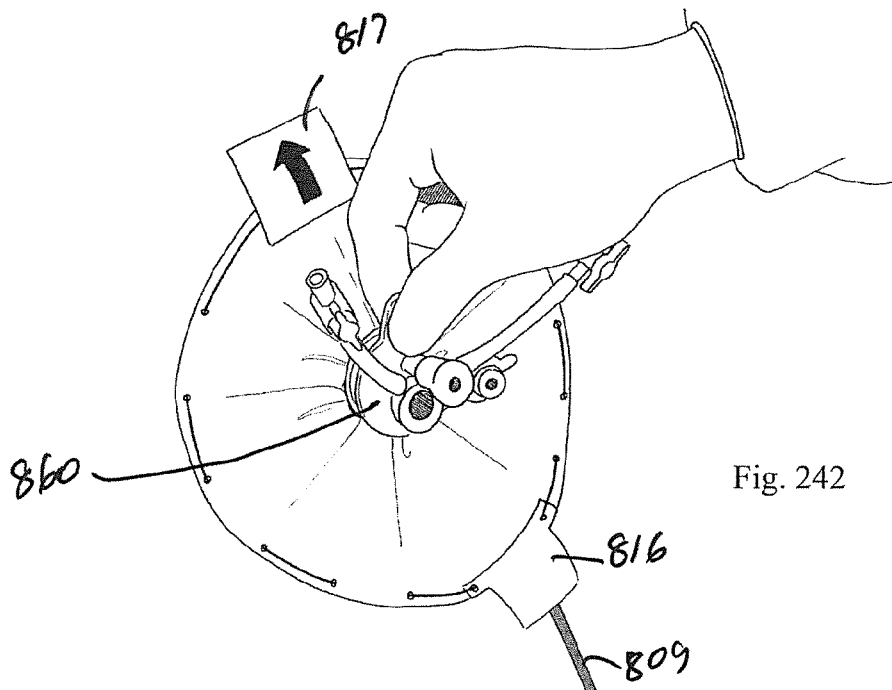
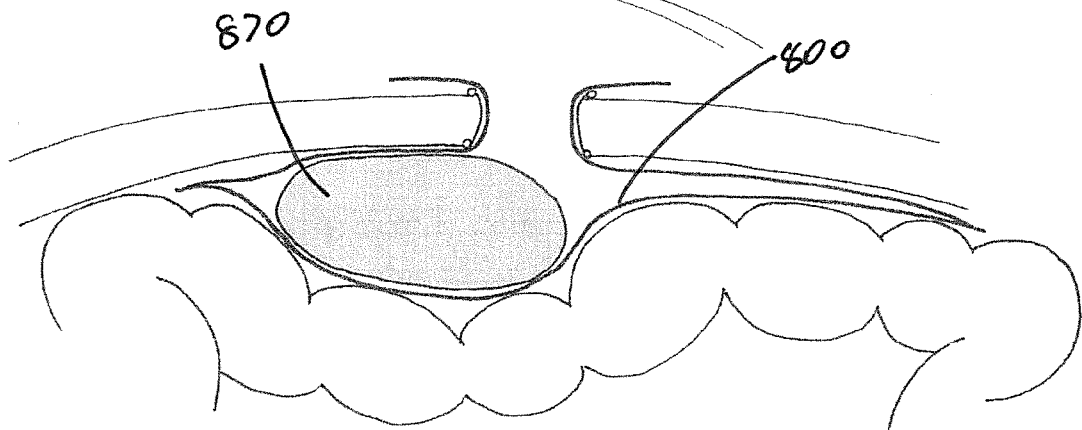
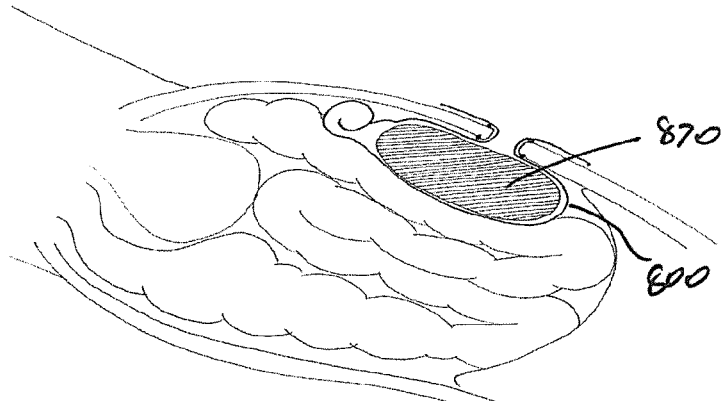
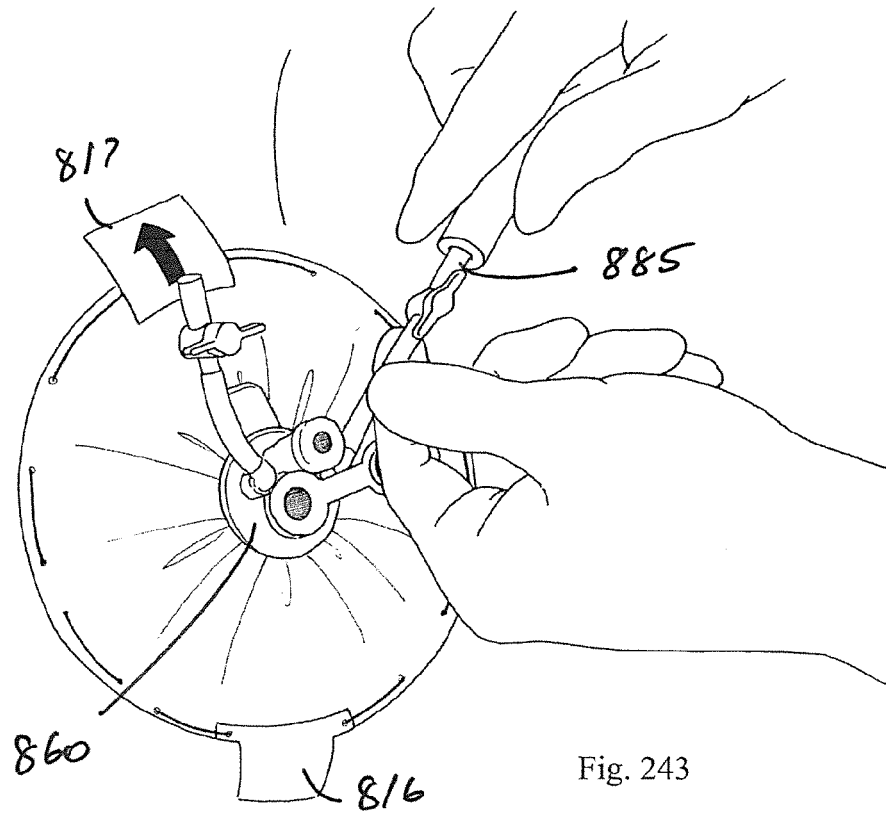


Fig. 239





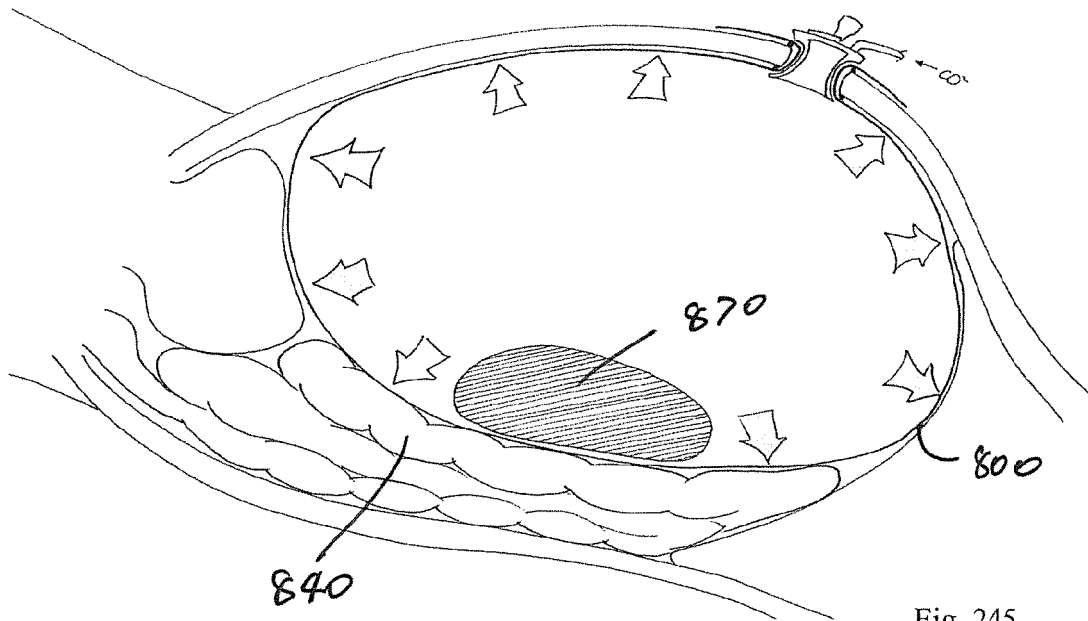


Fig. 245

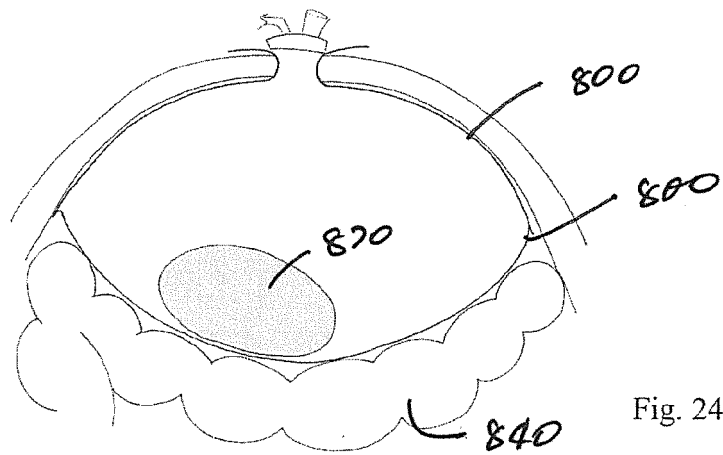


Fig. 246

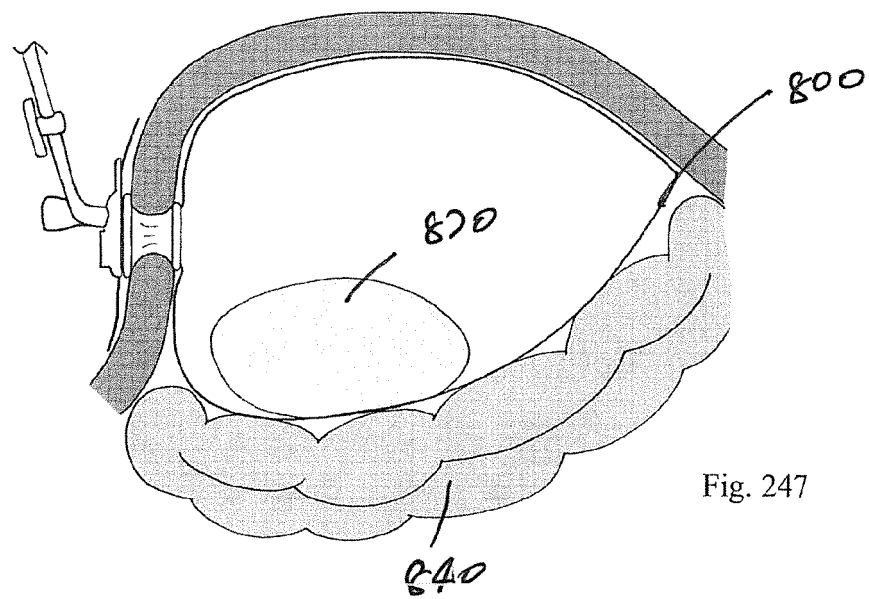


Fig. 247

Fig. 248

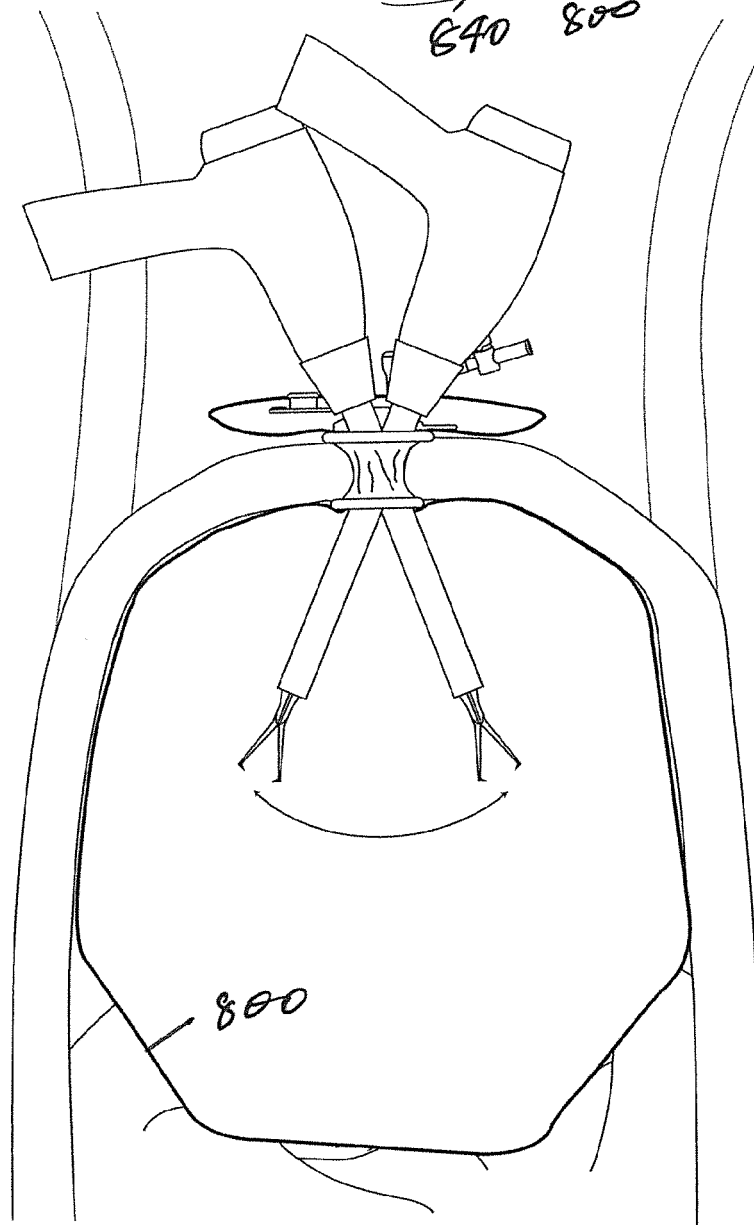
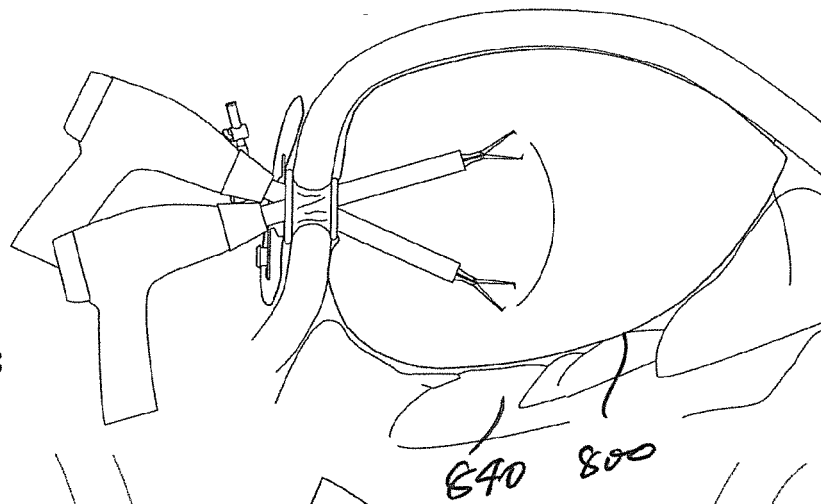
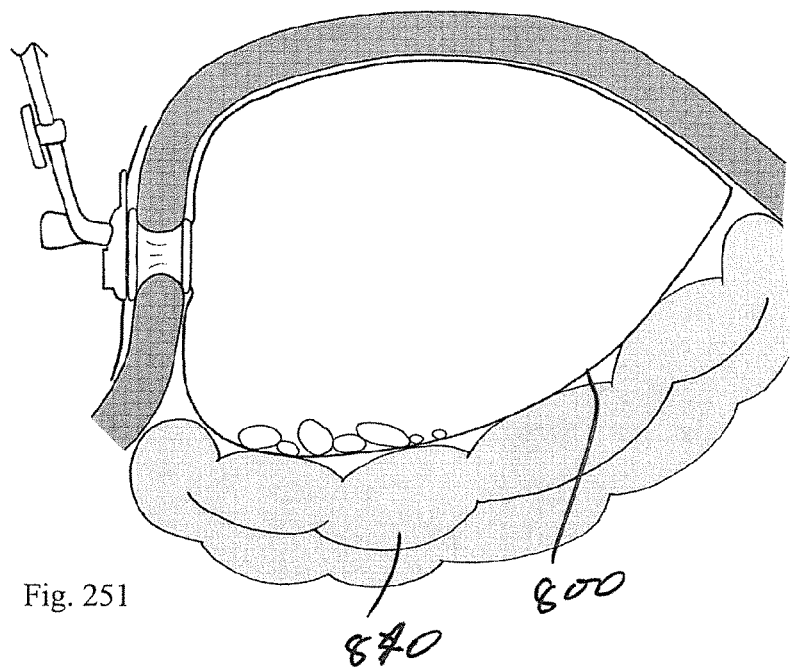
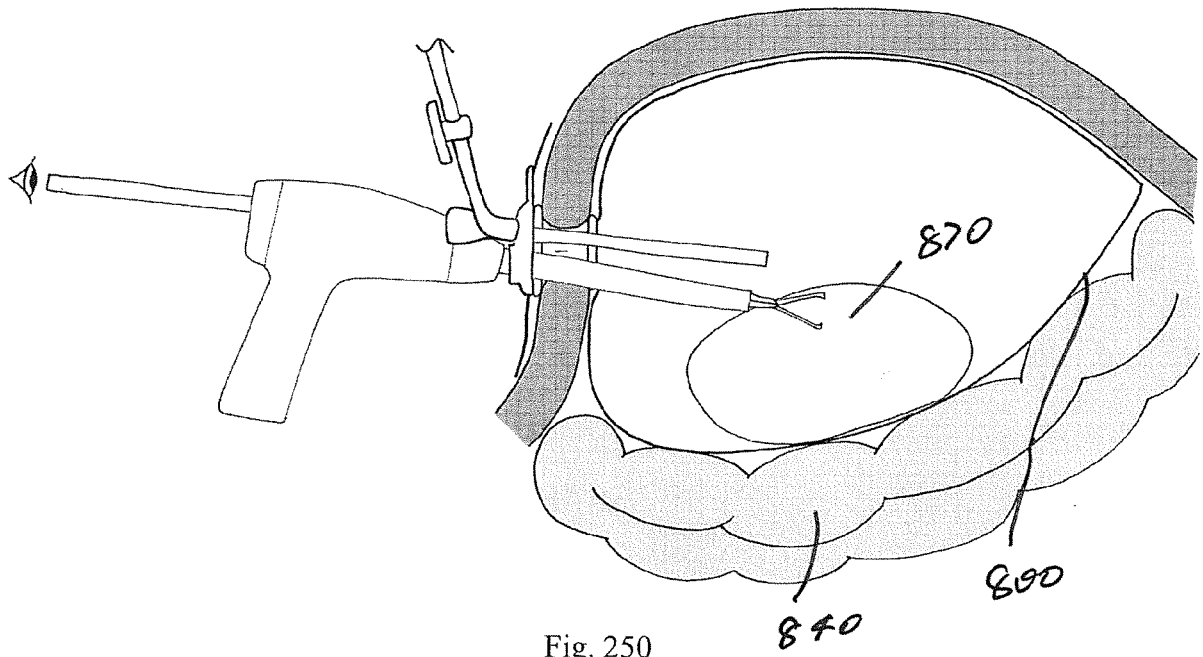


Fig. 249



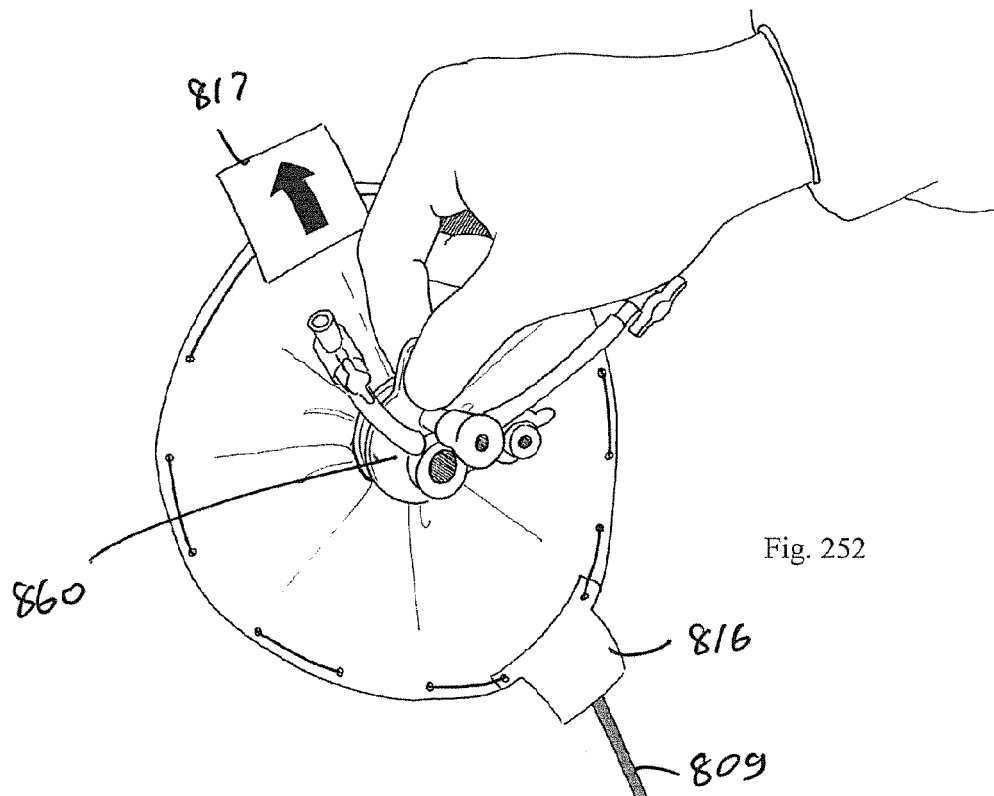


Fig. 252

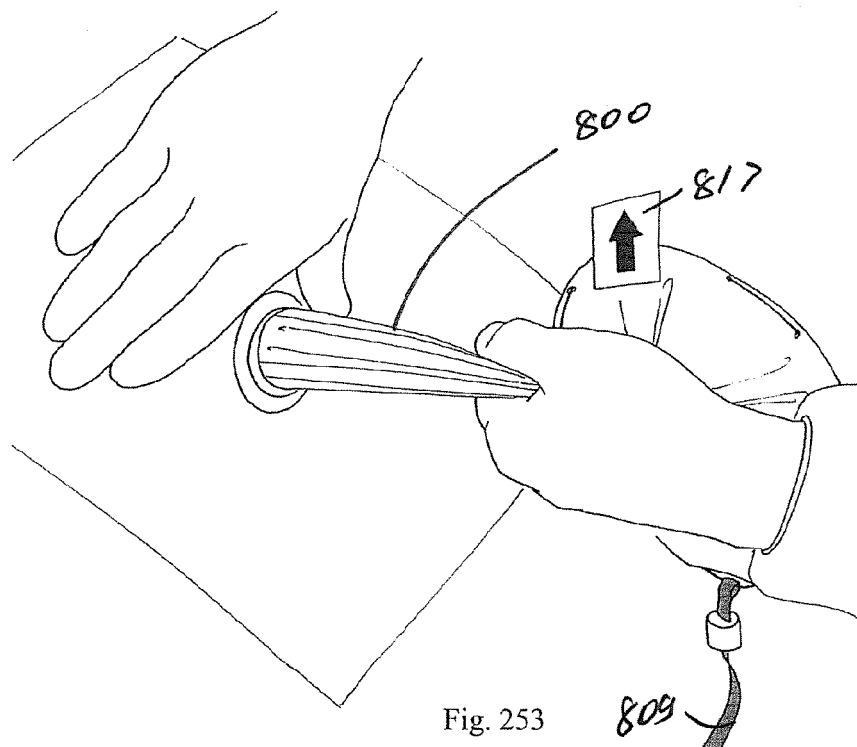


Fig. 253

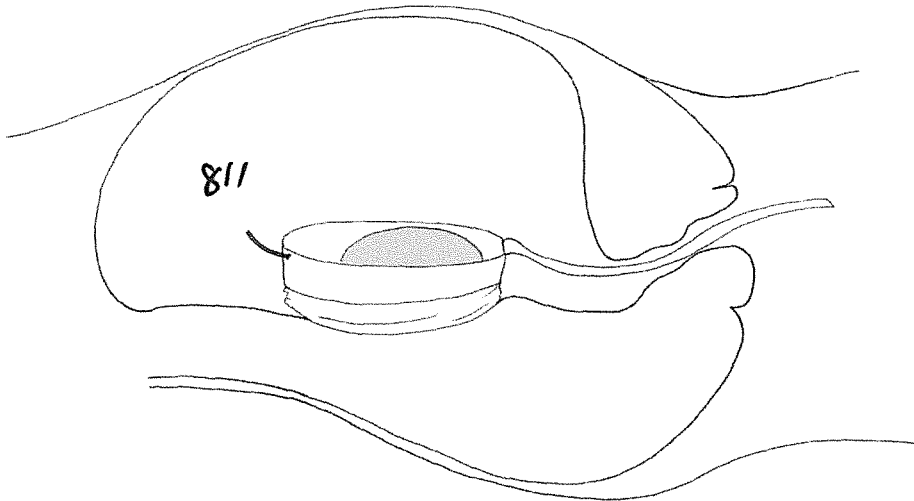


Fig. 254

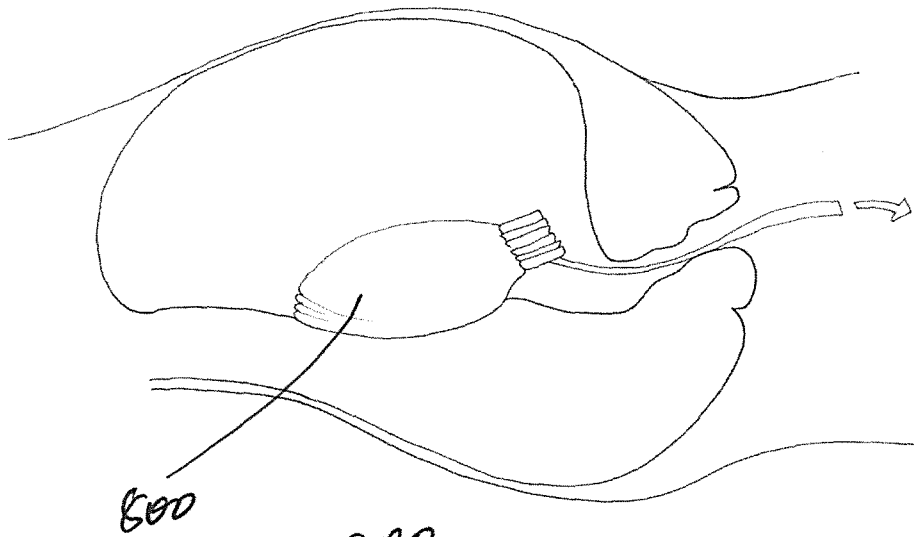


Fig. 255

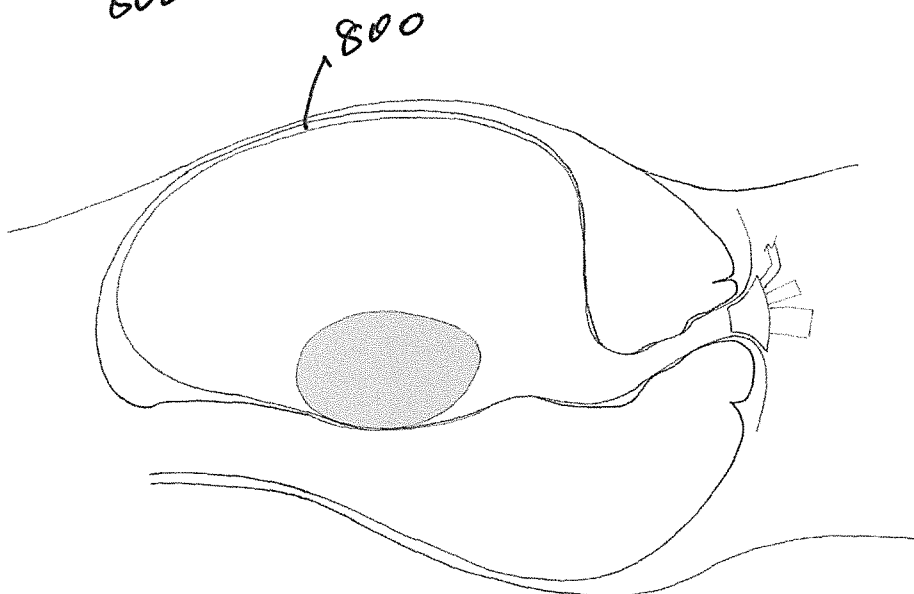


Fig. 256

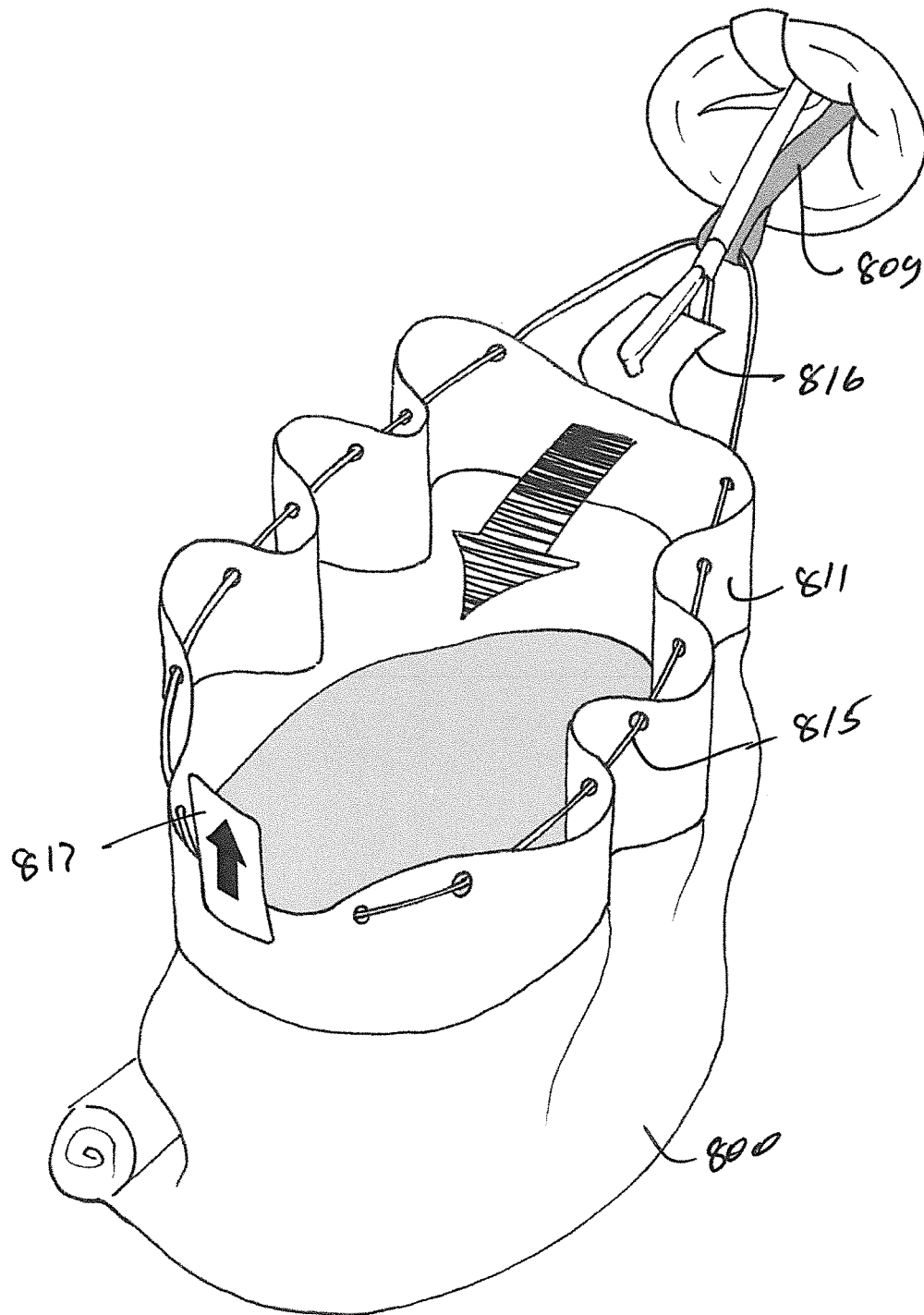


Fig. 257

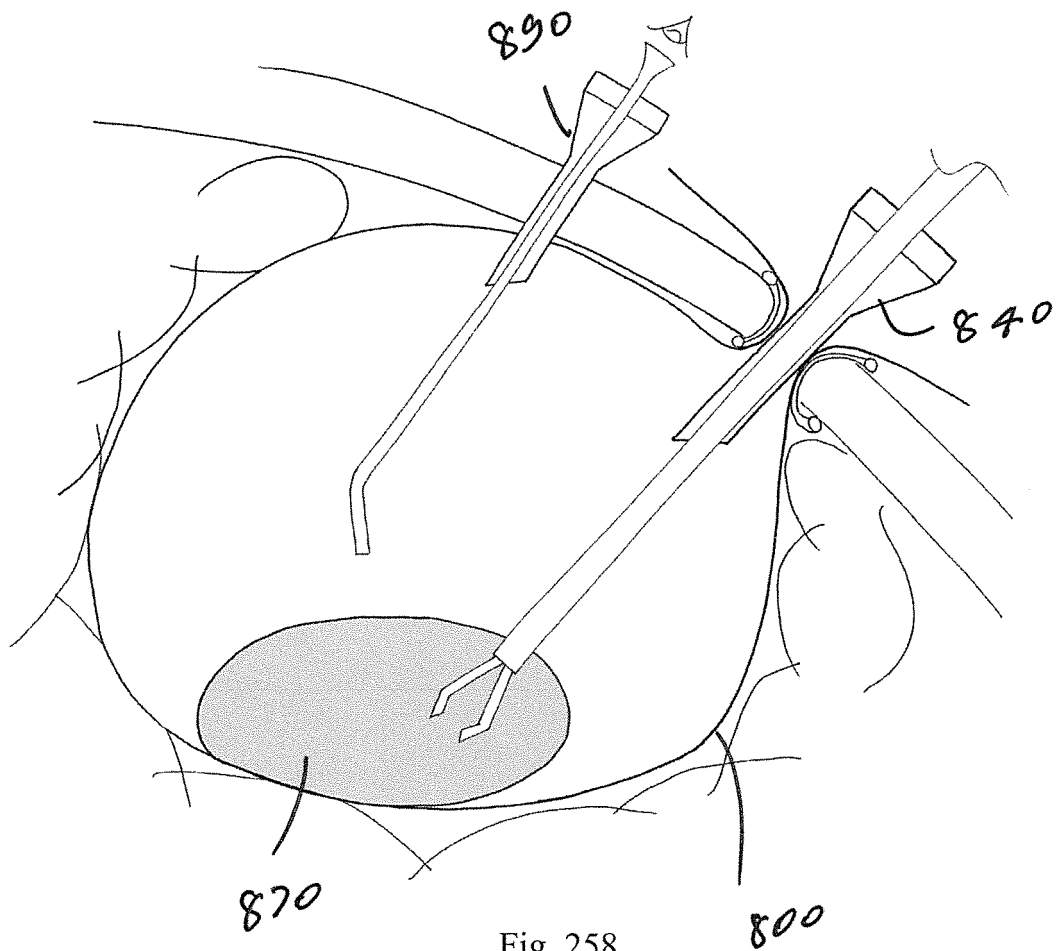


Fig. 258

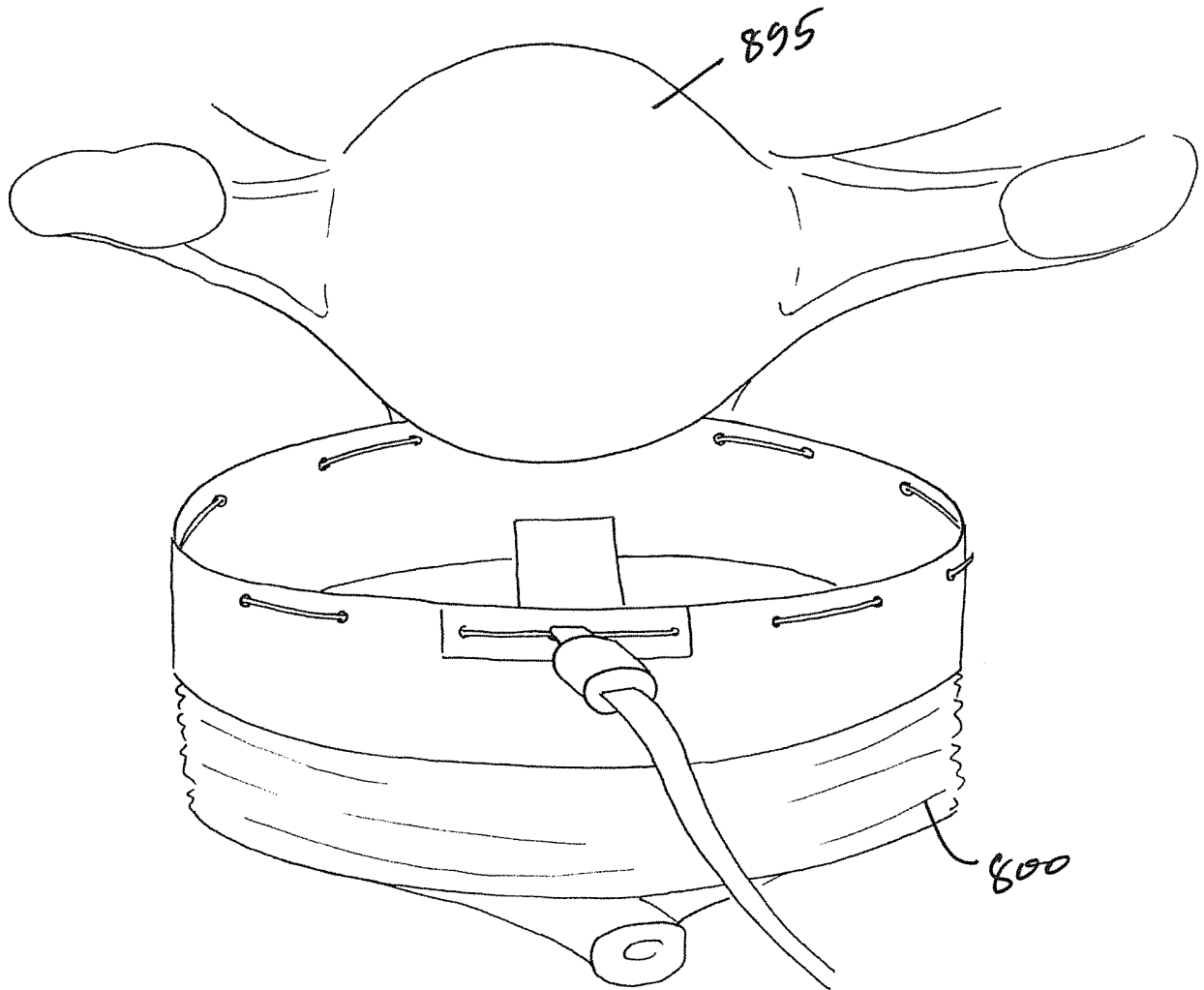


Fig. 259

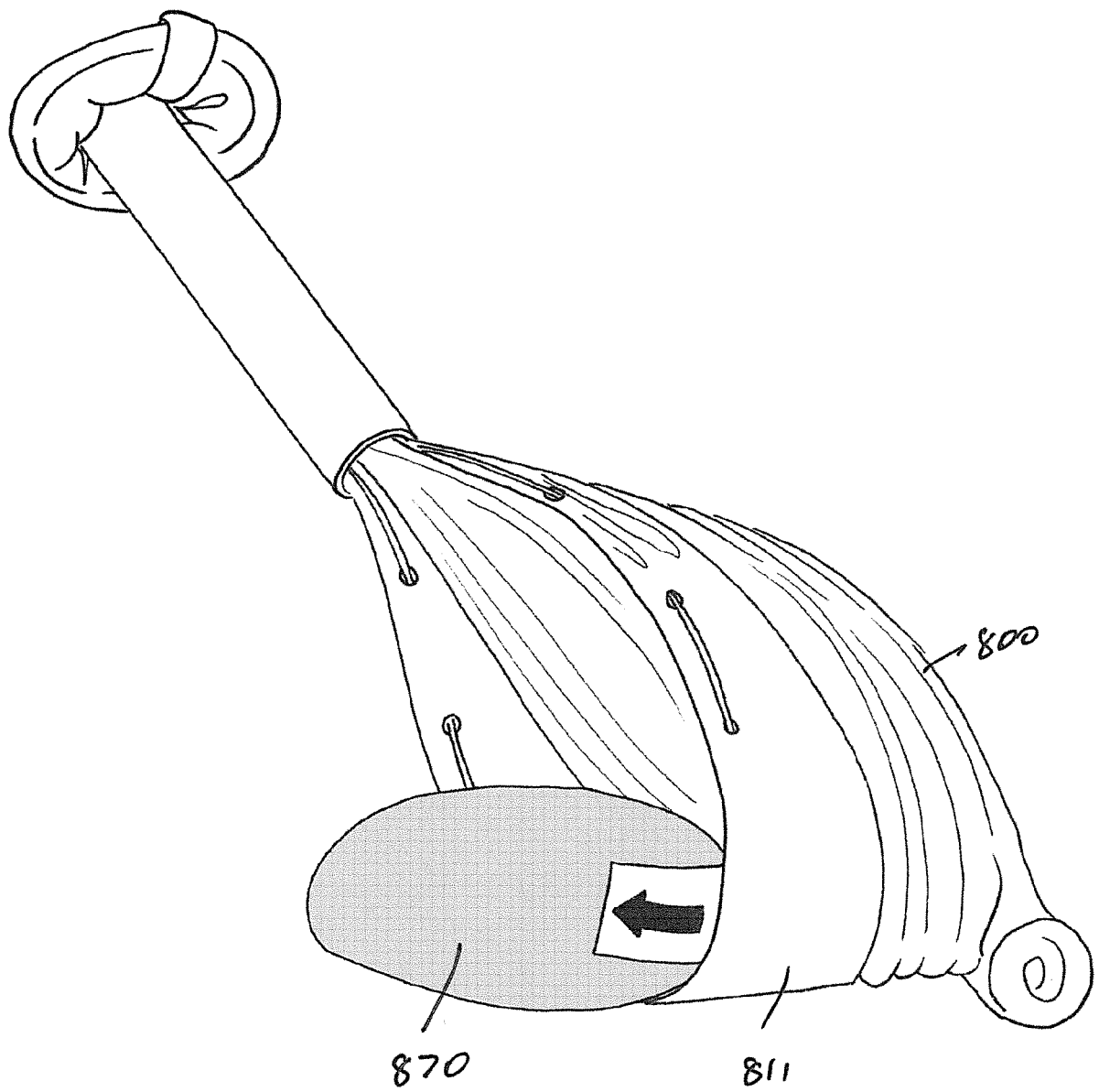


Fig. 260

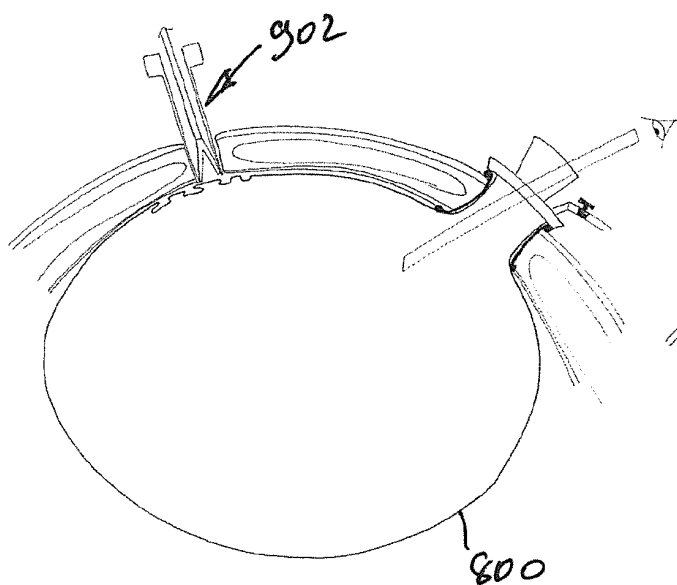


Fig. 261

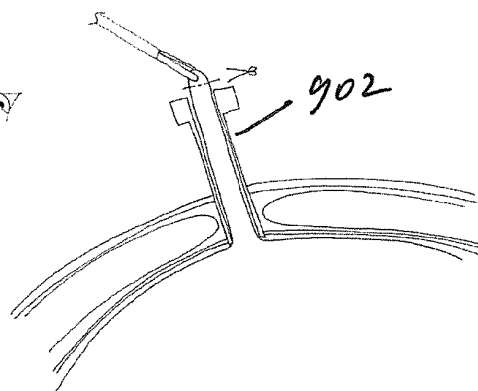


Fig. 262

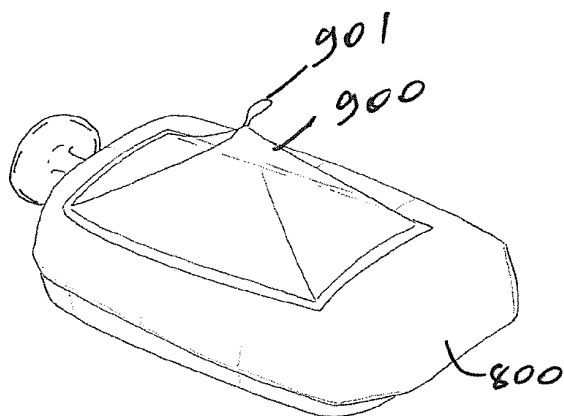


Fig. 263

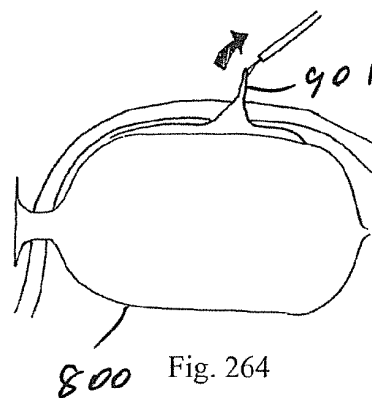


Fig. 264

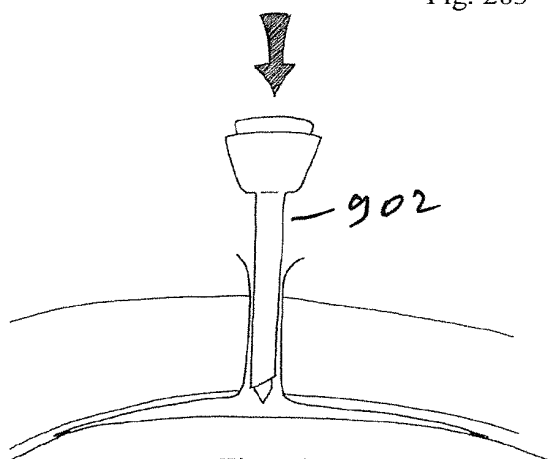


Fig. 265

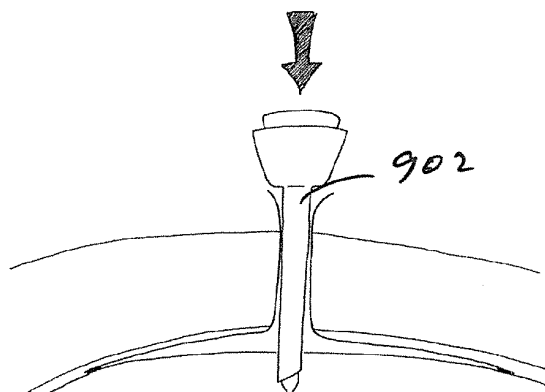


Fig. 266

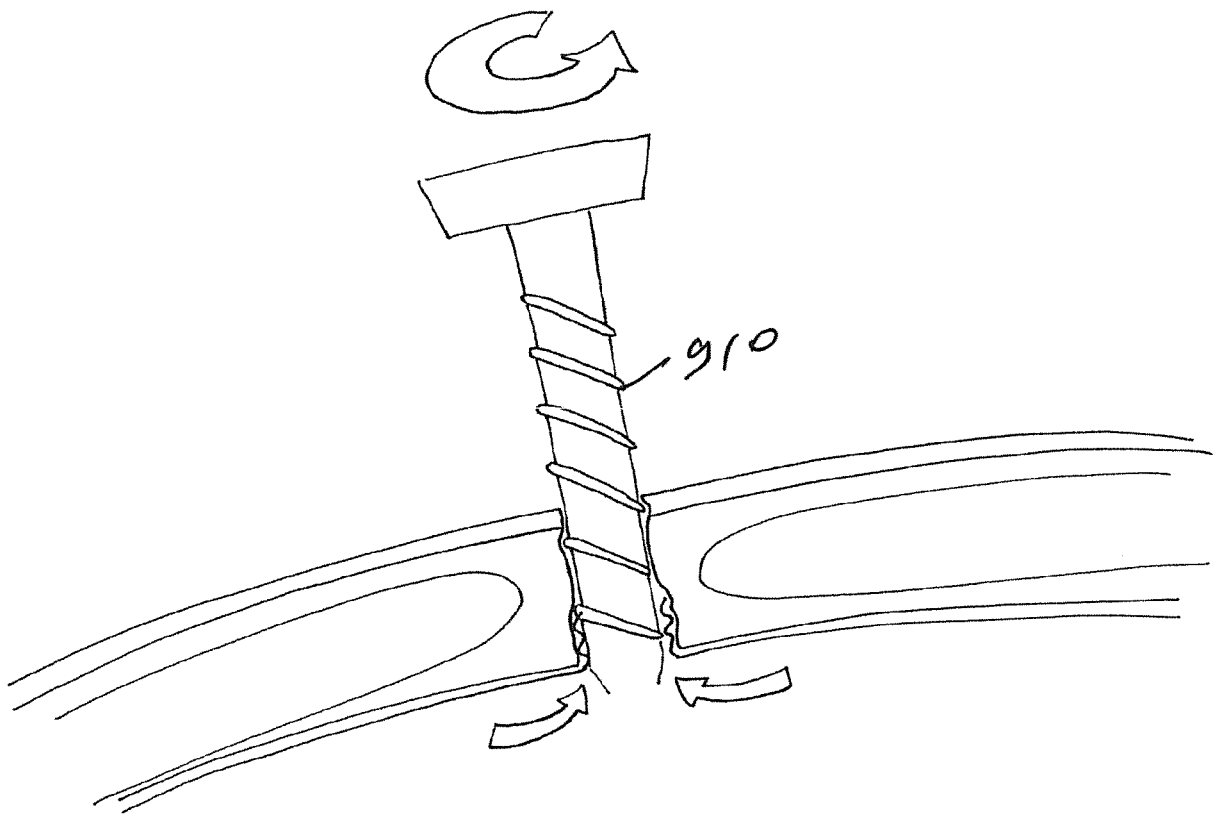
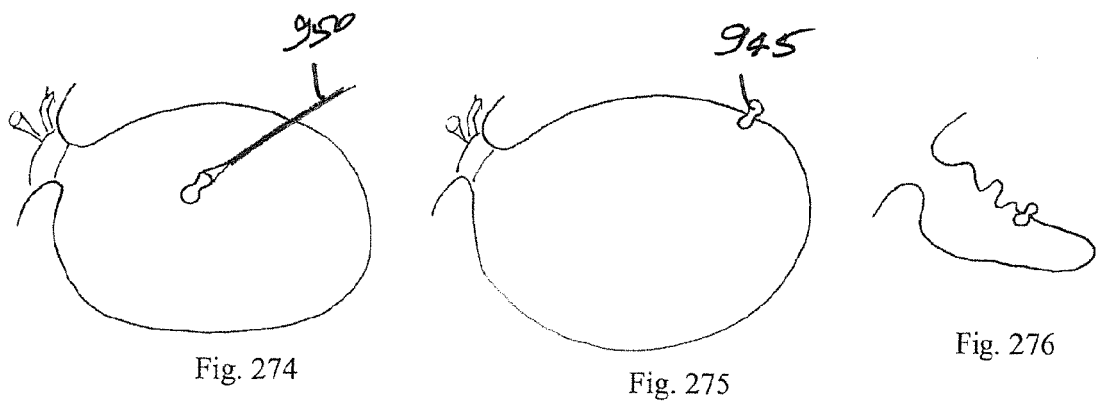
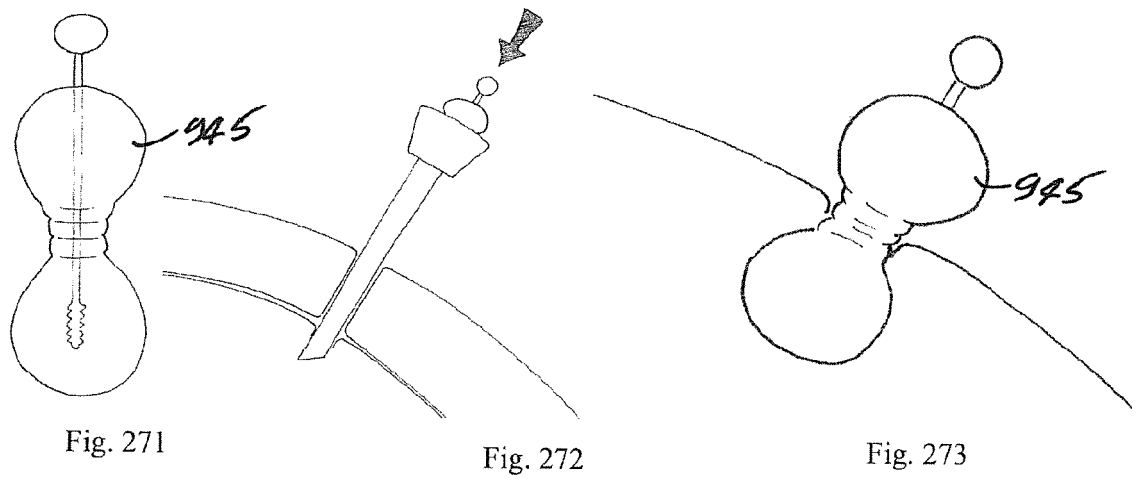
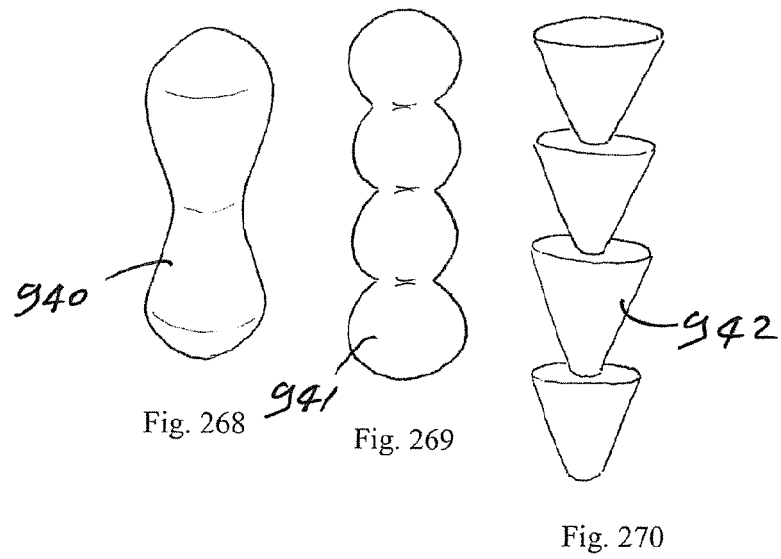


Fig. 267



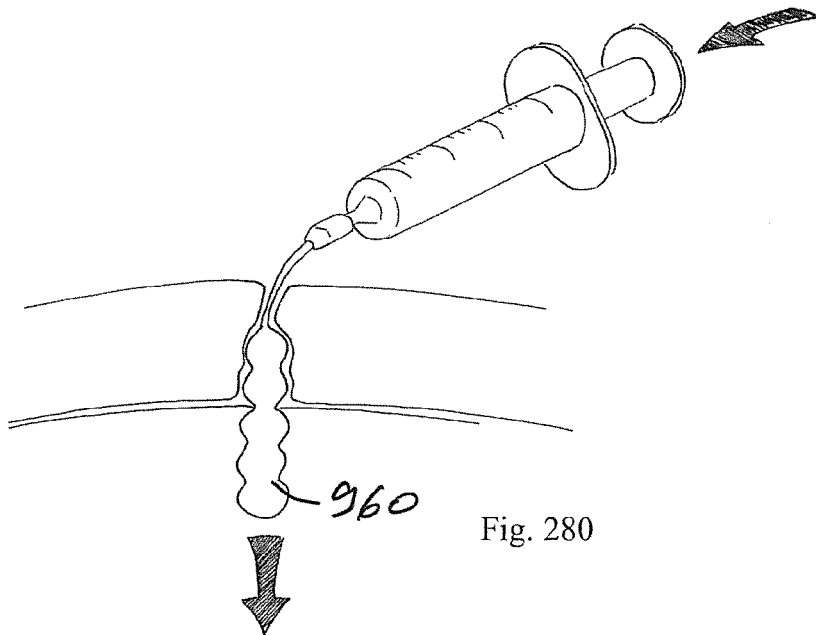
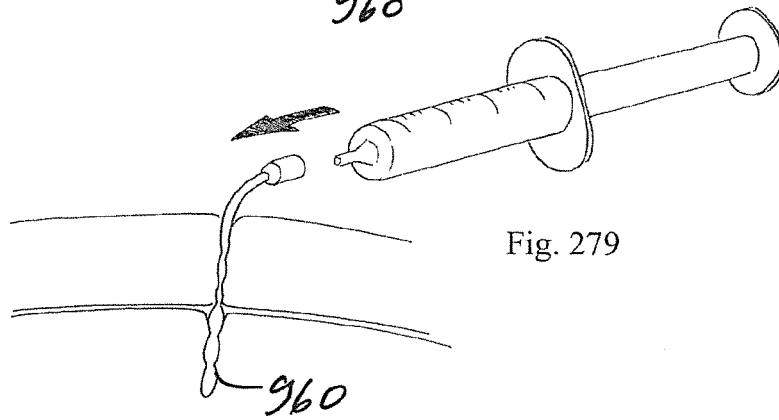
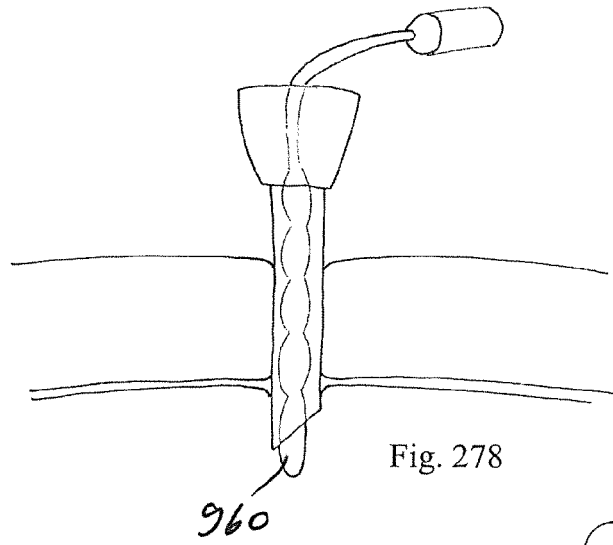
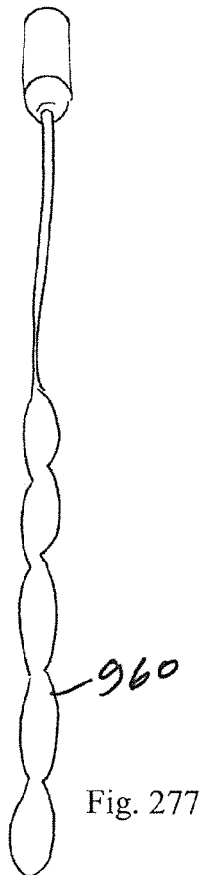


Fig. 281a

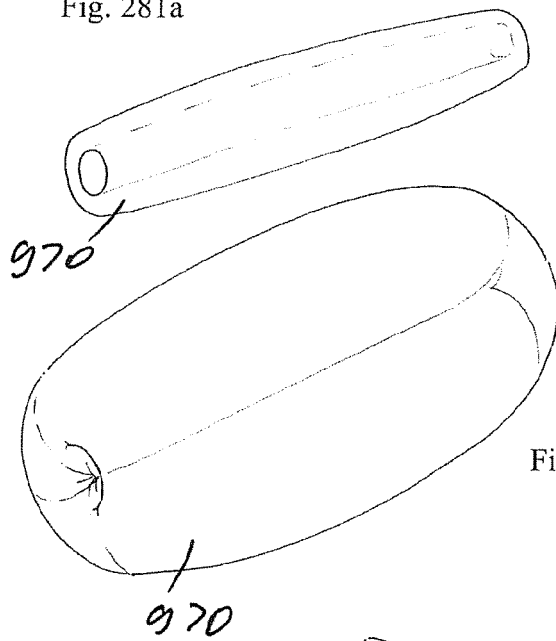


Fig. 281b

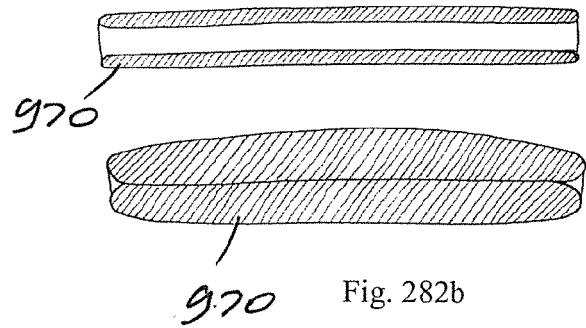


Fig. 282a

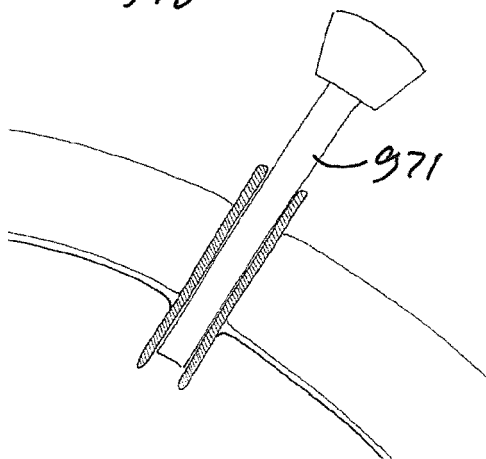


Fig. 283

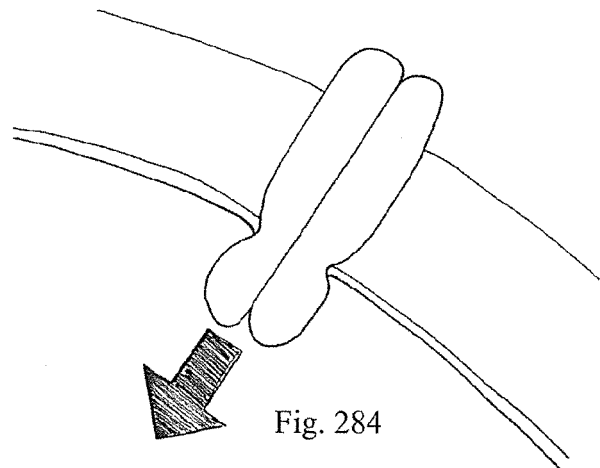


Fig. 284

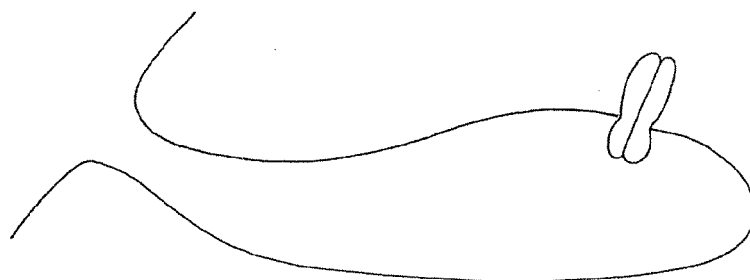


Fig. 285

REFERENCES CITED IN THE DESCRIPTION

This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

Patent documents cited in the description

- WO 2010099541 A [0001]
- WO 2013093030 A [0001]
- US 5618296 A [0001]
- WO 9509666 A [0002]
- WO 2011090866 A [0003]
- US 20110087235 A [0004]
- US 20050090717 A [0101] [0183]
- US 8187178 B [0121] [0184]
- US 20110071389 A [0121] [0184]

专利名称(译)	充气式气腹装置		
公开(公告)号	EP3013251B1	公开(公告)日	2018-02-07
申请号	EP2014733617	申请日	2014-06-25
[标]申请(专利权)人(译)	阿特波斯有限公司		
申请(专利权)人(译)	阿特洛波斯有限公司		
当前申请(专利权)人(译)	阿特洛波斯有限公司		
[标]发明人	BONADIO FRANK HALPIN LUCY DOLORES WILLIAMS STEPHEN		
发明人	BONADIO, FRANK HALPIN, LUCY DOLORES WILLIAMS, STEPHEN		
IPC分类号	A61B17/02 A61B17/00 A61B17/34 A61B90/40		
CPC分类号	A61B17/0218 A61B17/00234 A61B17/3423 A61B90/40 A61B2017/00287 A61B2017/00557 A61B2017/00637 A61B2017/00867 A61B2017/0225 A61B2017/320024 A61B2090/0801 A61B2090/08021		
优先权	61/839461 2013-06-26 US 14/251362 2014-04-11 US 61/968770 2014-03-21 US 61/940681 2014-02-17 US		
其他公开文献	EP3013251A1		
外部链接	Espacenet		

摘要(译)

用于在腹腔镜手术期间接收组织840的可充气人工气腹袋800在其近端具有组织接收开口，该组织接收开口由近端开口处的袖带811限定。袖带811具有用于递送和取出袋800的闭合构造和用于接收组织的打开构造。袖带811被偏置成打开配置。袋800具有从袖带延伸的主体，并且袋的主体比袖带811更柔韧。偏置元件815将袖带811偏压成打开构型。系绳809用于关闭袋子以便取回。袋800产生人造气腹，其隔离袋内的组织。

