



(51) International Patent Classification:

A61B 1/32 (2006.01) A61B 17/42 (2006.01)
A61B 17/3209 (2006.01)

(21) International Application Number:

PCT/IL2018/050934

(22) International Filing Date:

23 August 2018 (23.08.2018)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/549,097	23 August 2017 (23.08.2017)	US
62/549,078	23 August 2017 (23.08.2017)	US
62/558,469	14 September 2017 (14.09.2017)	US
62/558,460	14 September 2017 (14.09.2017)	US

(71) Applicant: MEMIC INNOVATIVE SURGERY LTD.

[IL/IL]; 6 Yonatan Netanyahu Street, 1st Floor, 6037604 Or-Yehuda (IL).

(72) Inventors: COHEN, Dvir; P.O. Box 51, 1924500 Ramot-

Menashe (IL). LEVINSON, Yaron; 25 Pinkas Street, Apt. 4, 6266211 Tel-Aviv (IL). MAIMON, Eyal; 1 Nofey Prat Street, 9061800 Kfar Edumim (IL). ROTEM, Idan; 63 J. L. Gordon Street, 6438817 Tel Aviv (IL).

(74) Agent: EHRLICH, Gal et al.; G. E. EHRLICH (1995)

LTD., 11 Menachem Begin Road, 5268104 Ramat Gan (IL).

(81) Designated States (unless otherwise indicated, for every

kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every

kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

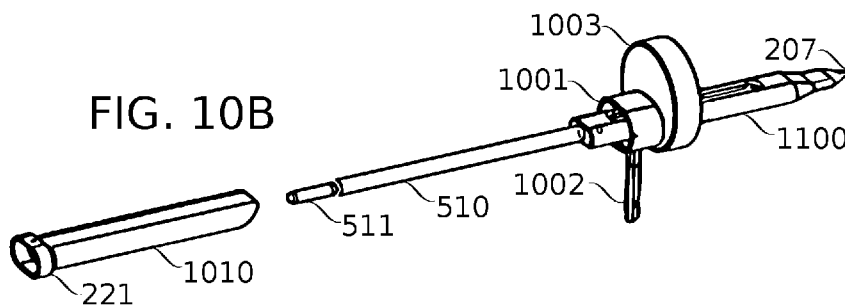
Declarations under Rule 4.17:

— of inventorship (Rule 4.17(iv))

Published:

— with international search report (Art. 21(3))
— before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))

(54) Title: TOOLS AND METHODS FOR VAGINAL ACCESS



(57) Abstract: Trocar components and methods of use are described, wherein the trocar components are configured to provide access to intraperitoneal space via the rectouterine pouch to surgical tools, which optionally include one or more surgical robot members. The surgical tools are optionally 5 mm or more in diameter. In some embodiments, a cannula part has a lumen sized to provide to a plurality of the surgical tools simultaneous transvaginal access to the intraperitoneal space via the rectouterine pouch. In some embodiments, an incision sized to receive a distal aperture of the cannula is created, optionally using one or two dilators. The dilators are sized to create (optionally starting from a puncture by a needle 2 mm in diameter or less) an oblong aperture. In some embodiments, the oblong aperture is at least twice as wide across a long diameter as across a short diameter.



TOOLS AND METHODS FOR VAGINAL ACCESS

RELATED APPLICATIONS

This application claims the benefit of priority under 35 USC § 119(e) of U.S. Provisional Patent Application No. 62/549,097 filed August 23, 2017; U.S. Provisional Patent Application
5 No. 62/549,078 filed August 23, 2017; U.S. Provisional Patent Application No. 62/558,460 filed September 14, 2017; and U.S. Provisional Patent Application No. 62/558,469 filed September 14, 2017; the contents of which are incorporated herein by reference in their entirety.

This application is also a part of a set of filings which are co-filed, co-pending and co-assigned:

10 U.S. Patent Application entitled "TOOLS AND METHODS FOR VAGINAL ACCESS"
(Attorney Docket No. **74919**);

PCT Patent Application entitled "TOOLS AND METHODS FOR VAGINAL ACCESS"
(Attorney Docket No. **74918**);

15 Canadian Patent Application entitled "TOOLS AND METHODS FOR VAGINAL
ACCESS" (Attorney Docket No. **74920**);

U.S. Patent Application entitled "TOOLS AND METHODS FOR VAGINAL ACCESS"
(Attorney Docket No. **74927**);

Canadian Patent Application entitled "TOOLS AND METHODS FOR VAGINAL
ACCESS" (Attorney Docket No. **74928**);

20 U.S. Patent Application entitled "TOOLS AND METHODS FOR VAGINAL ACCESS"
(Attorney Docket No. **74926**); and

U.S. Patent Application entitled "TOOLS AND METHODS FOR VAGINAL ACCESS"
(Attorney Docket No. **74924**);

the disclosures of which are all incorporated herein by reference.

FIELD AND BACKGROUND OF THE INVENTION

25 The present invention, in some embodiments thereof, relates to the field of intraperitoneal surgery and more particularly, to devices and methods for laparoscopic access to the intraperitoneal space.

Culdoscopy is an endoscopic procedure performed to examine the rectouterine pouch and pelvic
30 viscera by the introduction of a culdoscope through the posterior vaginal wall. The culdoscope may be a modified laparoscope. A trocar is first inserted through the vagina into the posterior cul-de-sac, the space behind the cervix, allowing then the entry of the culdoscope. Due to the

position of the patient, intestines fall away from the pelvic organs which can then be inspected. Conditions diagnosable by culdoscopy include tubal adhesions (causing sterility), ectopic pregnancy, and salpingitis. Culdoscopy allows the performance of minor procedures such as tubal sterilization.

5 SUMMARY OF THE INVENTION

There is provided, in accordance with some embodiments of the present invention, a stepped dilator for use with a trocar kit to provide intraperitoneal access via a body recess, comprising: a dilator body having a longitudinal axis in proximal to distal direction; a first tapered region of the dilator body tapering narrower toward a distal end of the dilator body; a second tapered region of the dilator body tapering narrower in a distal direction, located proximally to the first tapered region, and separated from the first tapered region by an isolating region.

In some embodiments, the isolating region comprises a region of constant cross-section perpendicular to the longitudinal axis extending between the first tapered region and the second tapered region.

15 In some embodiments, the isolating region comprises a region with no cross-section perpendicular to the longitudinal axis larger in any direction than the largest cross-section perpendicular to the longitudinal axis of the first tapered region.

In some embodiments, the isolating region is at least 3 mm long.

In some embodiments, the isolating region is no more than 20 mm long.

20 In some embodiments, the isolating region is between 5-15 mm long.

In some embodiments, a whole tapering extent of at least one of the first tapered region and the second tapered region tapers between its smallest cross-sectional area and its largest cross-sectional area perpendicular to the longitudinal axis over a longitudinal distance of 15 mm or less.

25 In some embodiments, through the tapering extent of at least one of the first tapered region and the second tapered region, a diameter of the stepped dilator perpendicular to the longitudinal axis increases by at least 7.5 mm.

In some embodiments, a cross-section with the largest cross-sectional area of the first tapered region perpendicular to the longitudinal axis has at least one axis of about 10 mm or longer.

30 In some embodiments, a distal tip of the first tapered region comprises a front surface aperture of 4 mm² or less, and expands in a distal direction from the front surface aperture through a radius of curvature of at least 2.5 mm.

In some embodiments, the front surface aperture is an aperture of an inner lumen sized to allow partial advancement of a trocar needle having a diameter less than or equal to about 2 mm in a longitudinal direction through the aperture.

In some embodiments, the cross-section of the first tapered region having the largest cross-sectional area perpendicular to the longitudinal axis has at least one axis of about 7.5 mm or less.

There is provided, in accordance with some embodiments of the present invention: a kit comprising the stepped dilator described above, along with a handle and a trocar needle; wherein the stepped dilator and handle together define a lumen sized to accept passage the trocar needle from a proximal end of the handle to a distal tip of the stepped dilator.

In some embodiments, the trocar needle is provided with a handle extending at least 5 cm past a proximal end of the handle when a distal tip of the trocar needle is advanced 5 mm past the distal tip of the stepped dilator.

In some embodiments, the trocar needle is provided with a dull-tipped, inner, spring-loaded stylet to act as a Veress needle having an extended position and a collapsed position; wherein the stylet comprises a blunt end extending past a sharp tip of the trocar needle in the extended position and preventing the sharp tip from injuring tissue; and wherein the stylet moves to the collapsed position upon sufficient longitudinal force being exerted so that it no longer extends past the sharp tip, allowing the sharp tip can operate to penetrate tissue.

In some embodiments, provided with a stopper device is configured to resist advancing the distal tip of the trocar needle more than 5 mm beyond the distal tip of the stepped dilator.

In some embodiments, a cross-section with the largest cross-sectional area of the second tapered region perpendicular to the longitudinal axis has at least one axis of about 21 mm or longer.

There is provided, in accordance with some embodiments of the present invention. A cannula to provide intraperitoneal access across a wall of a body recess, wherein a cross-section of an inner lumen of the cannula transverse to a longitudinal axis of the cannula has a long and a short axis; and wherein the long axis is at least twice as long as the short axis.

In some embodiments, the short cross-section axis is between 5 mm and 10 mm.

In some embodiments, the long cross-section axis is between 10 mm and 30 mm.

In some embodiments, a luminal wall defining the transverse cross-section of the cannula comprises straight sections on opposite sides of the transverse cross-section.

In some embodiments, the straight sections are interconnected through curved sections.

In some embodiments, the cannula is at least 5 cm long.

In some embodiments, an edge defining an aperture at a distal end of the cannula comprises a first edge portion extending along one side of the aperture, and a second edge portion extending along another side of the aperture, wherein the first edge portion extends along a side more distal along the longitudinal axis than the second edge portion.

5 In some embodiments, the first and second edge portions extend along opposite sides of the aperture.

In some embodiments, the first edge portion and the second edge portion extend along the long cross-section axis.

10 In some embodiments, the cannula comprises a handle extending at least 10 cm from a proximal end of the cannula.

There is provided, in accordance with some embodiments of the present invention, a cannula for use with a trocar to provide intraperitoneal access via a body recess, wherein an edge defining an aperture at a distal end of the cannula comprises a first edge portion extending along one side of the aperture, and a second edge portion extending along another side of the aperture, wherein the first edge portion extends along a side more distal along the longitudinal axis than the second edge portion.

In some embodiments, the first edge portion is positioned at least 5 mm more distally than the second edge portion.

20 In some embodiments, the first edge portion and the second edge portion each comprise respective a straight portion.

There is provided, in accordance with some embodiments of the present invention, a method of using a cannula to provide intraperitoneal access to a body cavity via a body recess, comprising: inserting a distal end of the cannula into a dilated aperture of a rectouterine pouch via transvaginal access; wherein an edge defining an aperture at the distal end of the cannula comprises a first edge portion extending along one side of the aperture, and a second edge portion extending along an opposite side of the aperture, and the first edge portion is positioned more distally along the cannula and from the dilated aperture than the second edge portion; and wherein the cannula is inserted so that the aperture at the distal end of the cannula is oriented to open toward the side of the second edge portion and facing toward a rectum adjacent to the rectouterine pouch.

30 In some embodiments, the method comprises inserting a flexible robotic arm through the cannula and into the rectouterine pouch, so that it exits the aperture in a direction oriented away from the rectum.

In some embodiments, the inserting a distal end of the cannula into the rectouterine pouch comprises sliding the cannula over an outer dilator; the outer dilator has a tapered distal insertion end sized and shaped to fittingly slide over an inner dilator having a tapered distal insertion end with a rounded tip; the cannula slides fittingly over the outer dilator; and at least the outer dilator is inserted into the rectouterine pouch via transvaginal access.

In some embodiments, the method comprises sliding the cannula over the outer dilator while the inner dilator remains within the outer dilator.

In some embodiments, the inserting a distal end of the cannula into the rectouterine pouch comprises sliding the cannula over a stepped dilator; the stepped dilator has a tapered distal insertion end with a rounded tip, a second tapered region, and an isolating region between the second tapered region and the tapered distal insertion end; the cannula slides fittingly over the stepped dilator; and the dilator is inserted into the rectouterine pouch via transvaginal access.

In some embodiments, the rounded tip has a hole sized to pass a trocar needle having a diameter less than or equal to about 2 mm.

There is provided, in accordance with some embodiments of the present invention, a kit for providing intraperitoneal access via a body recess, comprising: a cannula, wherein a transverse cross-section of an inner lumen of the cannula transverse to a longitudinal axis of the cannula has a long axis long enough to allow simultaneous insertion of at least two cylindrical members, each at least 8 mm in diameter; a stepped dilator having a dilator body with: a first tapered region of the dilator body tapering narrower toward a distal end of the dilator body, and a second tapered region of the dilator body tapering narrower in a distal direction, located proximally to the first tapered region, and separated from the first tapered region by an isolating region; and a trocar needle provided with a handle region extending past a proximal end of the inner dilator when a distal tip of the trocar needle is advanced to the distal tip of the inner dilator.

In some embodiments, the long axis of the cannula inner lumen transverse cross-section is at least 21 mm.

In some embodiments, the cannula has a short cross-sectional axis; and wherein the long cross-sectional axis is at least twice as long as the short cross-sectional axis.

In some embodiments, the kit comprises an arm sheath for a plurality of robotic arms, sized to fit within the cannula, having a minimum diameter of about 10 mm, and a maximum diameter of at least twice the minimum diameter.

There is provided, in accordance with some embodiments of the present invention, a method of gaining intraperitoneal access via a body recess, comprising: inserting a first stage of a stepped dilator into a rectouterine pouch to widen an aperture in a wall of the rectouterine

pouch; and inserting a second stage of the stepped dilator into the rectouterine pouch to widen the aperture; wherein the first and second stages of the stepped dilator each comprise a region which tapers narrower in a distal direction, and wherein the first and second stages of the dilator are separated by an isolating region at least 3 mm long.

5 In some embodiments, the method is preceded by: inserting the stepped dilator transvaginally to the wall of the rectouterine pouch; and advancing a trocar needle from within the stepped dilator to produce the aperture in the wall of the rectouterine pouch.

In some embodiments, an inner lumen of the cannula has at least one cross-sectional axis of at least 20 mm.

10 In some embodiments, the inner lumen of the cannula has at least one cross-sectional axis of less than about 12 mm.

There is provided, in accordance with some embodiments of the present invention, a method of gaining intraperitoneal access via a body recess, comprising: inserting a camera to an intraperitoneal space with a wall of a rectouterine pouch in a field of view of the camera; illuminating the wall of the rectouterine pouch using an intraperitoneally positioned illumination device; selecting a position for an aperture in the wall of the rectouterine pouch, based on light from the illumination device visible from outside the rectouterine pouch; advancing a trocar needle from outside the rectouterine pouch to press against the selected position in the wall of the rectouterine pouch; verifying the position of the trocar needle, based on one or more images from the camera within the intraperitoneal space; and piercing the rectouterine pouch to form the aperture, using the trocar needle.

There is provided, in accordance with some embodiments of the present invention, a kit for setting a position of a robotic arm system along a longitudinal axis of a cannula inserted to a body orifice, wherein the robotic arm system comprises a motor unit and at least one robotic arm extending, when positioned, distally from the motor unit along the longitudinal axis, the kit comprising: the cannula, including a cannula body configured for insertion to the body orifice; a mounting block, configured for attachment to the cannula; and an assembly attached to the mounting block and comprising a spacing arm and an aligning arm, and movable between a stowed position and a deployed position; wherein the deployed position of the assembly places elements of the aligning arm where they indicate a predetermined position along the longitudinal axis.

30 In some embodiments, the mounting block attaches to the cannula by connecting to an access device having a lumen sized to fittingly accept the cannula therewithin.

In some embodiments, the spacing arm and the aligning arm deploy by hinging around a plurality of stopped hinges, each stopped hinge defining at least a stopped deployed position, and a stopped stowed position.

In some embodiments, the kit further comprises: the motor unit and the at least one robotic arm extending distally from the motor unit to a predetermined distance from a stopper-receiving portion of the motor unit; wherein a distal end of the at least one robotic arm aligns with a distal end of the cannula when the at least one robotic arm is inserted to the cannula, and a stopper portion of the aligning arm contacts a stopper-receiving portion of the motor unit to prevent longitudinal advance of the motor unit.

In some embodiments, the kit comprises an arm sheath with a lumen sized to accept the at least one robotic arm, and an outer surface sized to fit within the cannula.

There is provided, in accordance with some embodiments of the present disclosure, an inner dilator for use with a trocar kit to provide intraperitoneal access via a body recess, having a distal insertion end tapered over a longitudinal distance of 15 mm or less between a distal tip and a fully dilating cross-section of the inner dilator, wherein: the fully dilating cross-section of the inner dilator has at least one axis of about 10 mm or longer; the distal tip of the insertion end comprises a front surface aperture of 4 mm² or less, and the distal tip of the insertion end expands in a direction along the taper from the front surface aperture through a radius of curvature of at least 2.5 mm; and the front surface aperture is an aperture of an inner lumen sized to allow partial advancement of a trocar needle having a diameter less than or equal to about 2 mm in a longitudinal direction through the aperture.

In some embodiments, the fully dilating cross-section of the inner dilator has at least one axis of about 7.5 mm or less.

There is provided, in accordance with some embodiments of the present disclosure, a kit comprising the inner dilator described above, along with the trocar needle, wherein the inner dilator is at least 17 cm long, and the trocar needle is provided with a handle extending at least 5 cm past a proximal end of the inner dilator when a distal tip of the trocar needle is advanced 5 mm past the distal tip of the inner dilator.

In some embodiments, the trocar needle is provided with a dull-tipped, inner, spring-loaded stylet to act as a Veress needle, wherein the stylet, in its extended position prevents a sharp tip of the needle from injuring tissue, but collapses upon sufficient longitudinal force being exerted so that the sharp tip can operate to penetrate tissue.

In some embodiments, provided with a stopper device is configured to resist advancing the distal tip of the trocar needle more than 5 mm beyond the distal tip of the inner dilator.

There is provided, in accordance with some embodiments of the present disclosure, a kit comprising the inner dilator described above, along with an outer dilator, wherein the outer dilator has a distal insertion end tapered over a longitudinal distance of 15 mm or less between a distal opening and a fully dilating cross-section of the outer dilator, wherein the fully dilating cross-section of the inner dilator has at least one axis of about 21 mm or longer.

In some embodiments, the distal opening has an inner lumen sized to fittingly enclose the fully dilating cross-section of the inner dilator.

In some embodiments, the kit is provided with a stopper configured to resist advancing the distal tip of the outer dilator more than 15 mm beyond the distal tip of the inner dilator.

There is provided, in accordance with some embodiments of the present disclosure, a trocar kit for providing intraperitoneal access via a body recess, comprising: a cannula, wherein a transverse cross-section of an inner lumen of the cannula transverse to a longitudinal axis of the cannula has a long axis long enough to allow simultaneous insertion of at least two cylindrical members, each at least 8 mm in diameter; an inner dilator at least long enough to leave an external handling region of about 10 cm while inserted fully into a body aperture 7 cm long, and having a distal insertion end tapered over a longitudinal distance short enough to reach a complete first-stage dilation within 15 mm or less of movement between a distal tip of 4 mm² area or less and a fully dilating cross-section of the inner dilator, wherein the fully dilating cross-section of the inner dilator has at least one axis about half as long as the long axis of the cannula cross-section; a trocar needle provided with a handle region extending past a proximal end of the inner dilator when a distal tip of the trocar needle is advanced to the distal tip of the inner dilator; and an outer dilator, wherein the outer dilator has a distal insertion end tapered over a longitudinal distance of 15 mm or less between a distal opening sized to fittingly surround the fully dilating cross-section of the inner dilator, and a fully dilating cross-section of the outer dilator, wherein the fully dilating cross-section of the outer dilator is sized to be fittingly surrounded by the inner lumen of the cannula.

In some embodiments, the long cross-sectional axis of the cannula inner lumen is at least 21 mm.

In some embodiments, the cannula has a short cross-sectional axis; and wherein the long cross-sectional axis is at least twice as long as the short cross-sectional axis.

There is provided, in accordance with some embodiments of the present disclosure, a trocar kit for providing intraperitoneal access via a body recess, comprising: an inner dilator, an outer dilator, and a cannula; wherein: the outer dilator is sized and shaped to fittingly insert over the inner dilator; the cannula is sized and shaped to fittingly insert over the outer dilator; the

inner dilator is provided with a rounded distal tip having a hole sized for the longitudinal pass of a trocar needle portion having a diameter less than or equal to about 2 mm; and an inner lumen of the cannula has at least one cross-sectional axis of at least 20 mm; and wherein the inner lumen of the cannula has at least one cross-sectional axis of less than about 12 mm.

5 In some embodiments, the inner dilator and the outer dilator are each tapered from a respective narrower distal insertion end to a respective full-size cross-section within 15 mm along a longitudinal axis.

There is provided, in accordance with some embodiments of the present disclosure, a method of gaining intraperitoneal access via a body recess, comprising: inserting an inner dilator
10 transvaginally to a wall of a rectouterine pouch; advancing a trocar needle from within the inner dilator to produce an aperture in the wall of the rectouterine pouch; inserting the inner dilator no more than 15 mm into the rectouterine pouch to widen the aperture; inserting an outer dilator no more than 15 mm into the rectouterine pouch by sliding the outer dilator over the inner dilator and across the aperture while the aperture is held open by the inner dilator; and inserting a distal
15 end of a cannula into the rectouterine pouch by sliding the cannula over the outer dilator and across the aperture while the aperture is held open by the outer dilator; wherein an inner lumen of the cannula has at least one cross-sectional axis of at least 20 mm.

In some embodiments, the inner lumen of the cannula has at least one cross-sectional axis of less than about 12 mm.

20 There is provided, in accordance with some embodiments of the present disclosure, a method of gaining intraperitoneal access via a body recess, comprising: inserting a camera to an intraperitoneal space with a wall of a rectouterine pouch in a field of view of the camera; illuminating the wall of the rectouterine pouch using an intraperitoneally positioned illumination device; selecting a position for an aperture in the wall of the rectouterine pouch, based on light
25 from the illumination device visible from outside the rectouterine pouch; advancing a trocar needle from outside the rectouterine pouch to press against the selected position in the wall of the rectouterine pouch; verifying the position of the trocar needle, based on one or more images from the camera within the intraperitoneal space; and piercing the rectouterine pouch to form the aperture, using the trocar needle.

30 There is provided, in accordance with some embodiments of the present disclosure, a kit for setting a longitudinal position of a robotic arm system along a longitudinal axis of a cannula inserted to a body orifice, wherein the robotic arm system comprises a motor unit and at least one robotic arm extending distally from the motor unit along the longitudinal axis, the kit comprising: a cannula, including a cannula body configured for insertion to the body orifice and

a cannula handle extending proximally along a longitudinal axis of the cannula; a mounting block, including a block body and a clamp configured to clamp the cannula handle at a selected longitudinal position relative to the block body; and a motor unit stopper, including a longitudinally extended member attached to the block body, and movable between: a first position extending a predetermined length from the block body to a proximal end of the motor unit stopper, and a second position; wherein the proximal end of the motor stopper unit in the first position is positioned to contact and prevent longitudinal advance of the motor unit upon insertion of the at least one robotic arm to the cannula, thereby defining a predetermined longitudinal position of the robotic arm system relative to the cannula; and wherein the second position of the motor unit stopper removes the motor unit stopper proximal end from a position preventing the longitudinal advance of the motor unit from the predetermined longitudinal position.

In some embodiments, the motor unit stopper is attached to the block body by a hinge, the first position comprises orientation of the motor unit stopper along the longitudinal axis of the cannula, and movement between the first position and the second position comprises rotation of the motor unit stopper on the hinge.

In some embodiments, the motor unit stopper is movable between the first position and the second position without disturbing the position of either the cannula or the motor unit when the at least one robotic arm is inserted to the cannula.

In some embodiments, the kit further comprises: the motor unit and the at least one robotic arm extending distally from the motor unit to a predetermined distance from a stopper-receiving portion of the motor unit; wherein a distal end of the at least one robotic arm aligns with a distal end of the cannula when the at least one robotic arm is inserted to the cannula, and the motor unit stopper contacts the stopper-receiving portion of the motor unit to prevent longitudinal advance of the motor unit.

In some embodiments, the kit comprises a plurality of extenders, each comprising a tube with a luminal cross-section sized to receive a robotic arm having a cross-sectional axis of at least 7 mm, and a length sized to extend longitudinally from the proximal end of the stopper to a position distal to the longitudinal position of the block body.

In some embodiments, the block body is slotted to receive the plurality of extenders at a position and orientation allowing guidance of robotic arms along the longitudinal axis to an aperture of the cannula body.

There is provided, in accordance with some embodiments of the present disclosure, a kit comprising an inner dilator having a distal tip sized to partially dilate an incision, and an outer

dilator sized to slide distally over the inner dilator to further dilate the incision with a distal tip of the outer dilator, wherein: the overall dilation of the inner and outer dilators is at least enough to allow simultaneous insertion of at least two cylindrical members each having a diameter of at least about 8 mm, while a longitudinal distance along each of the inner and outer dilators over which dilation occurs is less than about 20 mm; at least one of the inner dilator and the outer dilator are marked near a proximal end to indicate a relative position at which the two dilators are positioned, including at least a mark indicating alignment of the distal ends of the two dilators, and a mark indicating a longitudinal position difference of one dilator relative to the other of the longitudinal distance over which dilation occurs.

In some embodiments, both the inner and outer dilators are marked with a distance scale indicating distance along each dilator to its distal end.

In some embodiments, the distance scales of the inner and outer dilators are numerically aligned when the distal ends of each are aligned.

In some embodiments, the kit comprises an indicating indexer configured to change the force needed to translate the inner and outer dilators longitudinally over one another, depending on the relative longitudinal positions of the inner and outer dilators.

Unless otherwise defined, all technical and/or scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which the invention pertains. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of embodiments of the invention, exemplary methods and/or materials are described below. In case of conflict, the patent specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and are not intended to be necessarily limiting.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

Some embodiments of the present disclosure are described, by way of example only, with reference to the accompanying drawings. With specific reference now to the drawings in detail, it is stressed that the particulars shown are by way of example, and for purposes of illustrative discussion. In this regard, the description taken with the drawings makes apparent to those skilled in the art how embodiments of the present disclosure may be practiced.

In the drawings:

FIG. 1 is a schematic representation of portions of a human female pelvic anatomy, referenced by descriptions herein according to some embodiments of the present disclosure;

FIG. 2A schematically represents a kit of trocar components, according to some embodiments of the present disclosure;

FIG. 2B schematically represents distal portions of components in the kit of trocar components, according to some embodiments of the present disclosure;

5 FIGs. 3A–3H schematically represent a method of using trocar components to establish intraperitoneal access through rectouterine pouch wall or another body wall, according to some embodiments of the present disclosure;

FIG. 3I schematically represents the cannula configure of *Figure 3H*, along with inserted tools, according to some embodiments of the present disclosure;

10 FIG. 3J is a schematic flowchart outlining preparation for a laparoscopic procedure using trocar, according to some embodiments of the present disclosure;

FIG. 3K is a schematic flowchart representing a method of dilating and cannulating an access incision into a rectouterine pouch, according to some embodiments of the present disclosure;

15 FIG. 3L schematically represents a wider view (compared to *Figure 3H*) of the positioning of cannula relative to anatomical structures of a female lower abdomen/pelvic region, according to some embodiments of the present disclosure;

FIG. 4 schematically represents a dual-verification method of locating an incision for providing transvaginal access to a rectouterine pouch, according to some embodiments of the present disclosure;

20 FIGs. 5A–5C schematically represent different stopper mechanisms for use with trocar components, according to some embodiments of the present disclosure;

FIGs. 5D–5F schematically illustrate a needle, needle holder, and needle handle, according to some embodiments of the present disclosure;

25 FIGs. 5G–5I schematically illustrate mechanisms for controlling the relative positioning of an inner dilator, outer dilator, and needle according to some embodiments of the present disclosure;

FIG. 5J is a flowchart schematically outlining a method of using indicators to establish and maintain known penetration depths of the trocar needle, dilator, and/or cannula parts, according to some embodiments of the present disclosure;

30 FIG. 5K illustrates a manufactured example of a dilation and cannulation kit comprising members described and illustrated in, for example, *Figures 3A–3I* and *5D–5I*, according to some embodiments of the present disclosure;

FIGs. 6A–6E schematically represent dilation using a single dilator trocar kit, according to some embodiments of the present disclosure;

FIGs. 7A–7B are images taken from inside an insufflated abdomen of robotic arms inserted through the cannula in a configuration similar to that of *Figure 3I*, according to some
5 embodiments of the present disclosure;

FIGs. 8A–8B illustrate scale features of outer dilator and inner dilator, according to some embodiments of the present disclosure;

FIGs. 9A–9D comprise views representing an instrument holder for cannula and its configuration for use, wherein instrument holder includes a motor unit stopper for use in setting
10 an initial robotic arm position, according to some embodiments of the present disclosure;

FIGs. 10A–10E schematically illustrate views representing a collapsing instrument holder for cannula and its configuration for use in setting an initial robotic arm position relative to cannula, according to some embodiments of the present disclosure;

FIGs. 10F–10J schematically represent components of collapsing instrument holder,
15 according to some embodiments of the present disclosure;

FIGs. 11A–11E schematically represent a stepped dilator, dilator handle, and trocar needle, according to some embodiments of the present disclosure; and

FIGs. 12A–12C schematically represent a duck-bill gasket used to seal access to the proximal aperture of an access device, according to some embodiments of the present disclosure.

20 DESCRIPTION OF SPECIFIC EMBODIMENTS OF THE INVENTION

The present invention, in some embodiments thereof, relates to the field of intraperitoneal surgery and more particularly, to devices and methods for laparoscopic access to the intraperitoneal space.

Overview

25 A broad aspect of some embodiments of the present invention relates to trocar components (provided individually and/or in kits) configured to provide access to intraperitoneal space via the rectouterine pouch for surgical tools, optionally including one or more surgical robot members (herein, “robotic arms”). Compared, for example, to umbilicus entry to the intraperitoneal space, rectouterine pouch access to the peritoneal space provides potential
30 advantages for reduced invasiveness, reduced patient trauma, reduced visible scarring, and/or speed of the operation.

An aspect of some embodiments of the present invention relates to a cannula having a lumen with an oblong cross-section, and configured to simultaneously accept two or more substantially cylindrical tools extending side-by-side along the lumen.

In some embodiments, a cannula part is provided which has sufficient longitudinal length to extend between a wall of the rectouterine pouch and a position near to the entrance to the vagina, for example, about 7–15 cm long. Optionally, an intravaginal length of cannulation is selectably extended (*e.g.*, between from 7 up to about 15 cm) by use of the cannula together with an additional part; for example, a trocar part, with which it may be telescopically mated. Alternatively, in some embodiments, a selection of different cannula lengths is provided (*e.g.*, in a length range of from 7 to 15 cm, for example, at least three cannulas in this range; and optionally cannulas of sizes in about 1 cm or 2 cm length increments from each other). This provides a potential advantage for avoiding a possibility of pinching between two telescoping cannula portions.

Optionally, the cannula lumen cross-section is sized to provide simultaneous, side-by-side intraperitoneal access to two substantially cylindrical (*e.g.*, tubular) tools having a diameter of about 8.6 mm. In some embodiments, the substantially cylindrical tools comprise tube-shaped robotic arms. The cannula lumen cross-section, in some embodiments, has a longest axis at least twice as long as a shortest axis. Optionally, the cannula cross-section is sized so that there is also room for a third tool having a cross-section with a maximum axis length of about 6 mm or less. Compared to a circular cross-section sized to pass two or more such cylindrical tools, such an oblong cross-section provides a potential advantage for allowing a smaller overall cannula perimeter, with a correspondingly smaller incision needed to accept the cannula.

In some embodiments, the cannula has a distal aperture which is slanted relative to the longitudinal axis of the cannula so that it opens toward the direction of intraperitoneal space when inserted into the rectouterine pouch. Potentially, this helps to provide space for robotic members and/or other tools to bend to enter the intraperitoneal space.

Optionally, the cannula is constructed from stainless steel or another material which can be sterilized and re-sterilized to surgical use standards. Optionally, the cannula is disposable and provided in a sterilized condition.

An aspect of some embodiments of the present invention relates to geometries of dilators configured to achieve incision dilation while advancing to longitudinal distances kept short and/or controlled to avoid trauma to delicate tissues near the rectouterine pouch; and to methods of dilation adjusted for the geometries of the dilators.

A significant potential complication of opening a rectouterine pouch incision to the intraperitoneal space is damage to the rectum. The damage may be due for example, to over-penetration causing puncture, scraping, and/or crushing during initial puncture and/or dilation of the incision. In some embodiments, a constraint on maximum longitudinal advance is set by the width of the rectouterine pouch, and a need to reduce the potential for accidental damage to nearby internal organs, for example, the rectum.

In some embodiments of the invention, features of a trocar kit and/or its method of use potentially act to reduce a risk of injury due to over-penetration during cannulation. In particular, target insertion depth (*e.g.*, minimum depth at full dilation) is kept low in some embodiments (*e.g.*, dilation of up to about 7.5–15 mm occurs over about 10–20 mm of longitudinal insertion depth, for example, 13 mm, 15 mm, 17 mm, 19 mm, or 20 mm). In some embodiments, dilation occurs over about up to 30 mm, 40 mm, or 50 mm of insertion depth.

Overall insertion depth by a dilator after dilation itself is complete is optionally somewhat larger than this (*e.g.*, up to about 2–5.5 cm). However, application of potentially injurious force is particularly likely during a dilating phase in which the expanding portion of the dilator is being advanced through an incision—since this is the phase of dilator operation during which overcoming resistance by use of additional force is expected and normal.

Optionally, insertion in a dilating phase of operation to an excessive depth is treated as itself having a higher risk of causing injury (*e.g.*, due to the internal proximity of delicate tissues) than the elevation of maximum peak insertion force that may result from the lowered mechanical advantage of a shortened dilation depth.

In some embodiments, dilators are provided as a pair of dilators. In some embodiments, the pair of dilators comprises a first dilator and a second dilator, wherein the first dilator is smaller in cross-section than the second dilator. Optionally, the first dilator is provided as an “inner” dilator relative to the second, larger and “outer” dilator. The inner and outer dilators are configured to slide longitudinally relative to one another.

In some embodiments, a dilator is provided as a single dilator which dilates using a stepwise plurality of staged dilator expansions (*e.g.*, two or three).

In some embodiments, each dilator or dilator step provides expansion from an initial incision width (smallest cross-sectional area of the dilator stage) to a final incision width (largest cross-sectional area of the dilator stage) within about 15 mm of longitudinal advance per stage. Optionally the expansion occurs within another longitudinal distance of advance, for example, a distance from within the range of about 10–20 mm, for example, 13 mm, 15 mm, 17 mm, 19 mm, or 20 mm.

The amount of widening over the travel of the dilator stage is optionally itself in the range of about 7.5–15 mm, for example, about 7.5 mm, 10 mm, 12 mm, 12.5 mm, or 15 mm. This fairly rapid rate of dilation as a function of longitudinal advance accepts loss of mechanical advantage in exchange for a reduced required insertion depth to complete dilation. Total insertion depth during dilation can be about the length of a single step (optionally plus a few millimeters past the expanding part of the dilator, *e.g.*, plus 5–10 millimeters) when a plurality of dilators are used; wherein a subsequent dilator is inserted over the previous dilator. When a single (stepwise expanding) dilator is used, insertion depth during dilation may be the sum of the lengths of the individual dilation stages; plus an optional isolating region between the dilation steps having a length of, for example, about 5–15 mm; and optionally plus a few more millimeters (*e.g.*, 5–10 mm) past the expanding part of the dilator. For example, the total insertion depth may be about 50 mm. In some embodiments, the isolating region is at least 3 mm long. Additionally or alternatively, the isolating region is less than about 20 mm long.

Related to this, the inventors have realized that mechanical properties of the pouch wall tissue related to resisting dilation (*e.g.*, resistance to tearing and/or stretching) are potentially more permissive of a lowered mechanical advantage than the mechanical properties of the walls of other intraperitoneal access positions, for example, the mechanical properties of the skin, fat, and/or muscle layers of the umbilical region. This has allowed use of a dilator design which has reduced mechanical advantage (is blunter), in exchange for such potential advantages as a shorter dilator insertion depth and/or a smaller number of dilation steps.

Dividing the dilation into stages (*e.g.*, by using a plurality of dilators and/or a plurality of isolated dilation steps) potentially gives greater control over dilation by providing a stopping point mid-dilation. This potentially reduces a chance of uncontrolled tearing during dilation, and/or allows inspection of initial dilation to ensure that there is no unexpected damage (*e.g.*, excessive bleeding) which might be aggravated by further dilation.

In some embodiments, the first dilator and/or dilator step has a blunted distal-most portion. The distal-most portion optionally has a port through which a trocar needle can be extended. Optionally, the distal-most portion curves proximally, widening in both width and height through a radius of at least about 2.5 mm, then expanding primarily in width to form a wide oblong cross-section about 15 mm proximal to the distal-most portion (or another distance, for example in the range of about 10–20 mm).

In some embodiments, the second dilator and/or dilator step has a distal-most portion defining a lumen sized to fittingly slide over the first dilator. The outer perimeter of the distal-most cross-section is only slightly larger than the oblong cross-section at the proximal end of the

expanding cross-section region of the first dilator. From there, the second dilator's cross-section also expands going proximally for about 15 mm (or another distance, for example in the range of about 10–20 mm). The maximum of the further expansion is by about, for example, about 5 mm, 7.5, mm, 10 mm, or 12.5 mm. Optionally, there is a larger expansion along one axis of the
5 incision cross-section than along another axis; for example, there may be a relative factor of expansion of about 1:1.5, 1:2, or 1:3.

In some embodiments, the cannula is sized to fittingly slide over the second dilator to reach a position with its distal aperture inserted within the rectouterine pouch.

In some embodiments, the trocar needle used with the first dilator is provided together
10 with a holder and/or handle which are sized so that the maximum distal advance of the trocar needle is limited by interference between the handle and/or the holder. In some embodiments, the dilators are provided with a stopper and/or indicating indexer which allow tracking of their relative position, and/or resist, indicate, and/or prevent over-advancement of one dilator relative to the other.

Optionally, the dilator, dilators, handle, holder, and/or trocar needle are constructed from
15 stainless steel or another suitable material which can be sterilized and re-sterilized to surgical use standards (*e.g.*, by autoclaving). Optionally, one or more of these parts is disposable and provided in a sterilized condition.

An aspect of some embodiments of the present invention relates to dilator safety performance
20 maintained and/or enhanced by feedback features and/or methods which help monitor dilator advancement.

In some embodiments, a potential for loss of control of position (*e.g.*, sudden accidental over-advancement as tissue gives way, and/or as the end of the expanding region of the dilator is reached) is reduced by moving each dilator with respect to a fixed (*e.g.*, clamped to the patient
25 table) reference. For example, the first dilator is moved with reference to its initial position and/or an already inserted needle; and/or the second dilator is moved relative to the inserted position of the first dilator. Monitoring of position relative to a fixed reference position potentially encourages a user of the dilators to ease back on force when nearing a dilator's target position.

In some embodiments, a stopper arrangement changes (*e.g.*, increases) a sliding
30 resistance between two components in relative motion as a target dilator advancement limit is approached and/or reached. The change may indicate reaching a target position to a user, and/or mechanically resist advance beyond the target position.

An aspect of some embodiments of the present invention relates to methods of cannulizing a rectouterine pouch wall while monitoring the penetration using information communicated across the rectouterine wall. In some embodiments, initial rectouterine pouch penetration (*e.g.*, using a trocar needle) is visualized from using a camera and/or light source already inserted to the intraperitoneal space from another location, for example, the umbilical. Upon needle contact, a region of indentation may be interiorly observed before actual puncture. Alternatively or additionally, transillumination of the rectouterine pouch wall by an intraperitoneally located light source is observed from outside the pouch in order to help position a needle used for initial penetration. The method has a potential advantage insofar as the rectouterine wall is located both in a difficult region to directly access (due to its position deep within the vagina), and nearby sensitive internal structures which could lead to surgical complications if damaged during cannulization. Dual inside-to-outside and outside-to-inside needle position verification allows seeing from the outside (by the illumination) that the targeted port position (aimed at by a needle which is to create an initial opening) is in a reasonable location relative to internal structures which are to be targeted/avoided, and then confirming that the actual port position which the needle will create really is at the position aimed at.

It is noted that the method described with respect to the rectouterine pouch may be adapted to cannulization of other areas, wherein a first introduction of a camera and light source is from a first port into an intrabody space, and cannulization is to be performed to create another port in a region which perhaps provides some advantage (*e.g.*, better suited to receive a larger incision which is needed for larger tools, and/or provides a preferred direction and/or position of access by the tools), but may also be at greater risk of complication during its creation (*e.g.*, because it is in a region which is more difficult to target externally, and/or because it is associated with certain safety risks if performed incorrectly). More generally, in some embodiments, where a plurality of ports are to be used, second and subsequent ports may be opened under two-sided observation after camera and lighting are established within a first port.

An aspect of some embodiments of the present invention relates to achieving dependable and preferably rapid initial positioning of robotic arms relative to a cannulated surgical access-way. The cannula helps to provide access to a region of surgical activity which is not only internal, but also positioned at the end of a restrictive tunnel. The robotic arms themselves may be articulated along their length in such a way that the result of a commanded motion is different depending on exactly what their starting position is relative to the potential restrictions on motion presented by the cannula and/or the geometry of the internal body space in which they are operated.

Two parameters of particular importance are the distance of longitudinal advance of one or more robotic arms through the lumen of the cannula, and the angle of approach of the one or more robotic arms. An incorrect distance of longitudinal advance potentially leads to unexpectedly restricted motion (*e.g.*, because an articulated arm portion has not advanced out of the cannula as much as expected), or even injury (*e.g.*, a collision with body tissue due to over-advancing). An incorrect angle of approach potentially leads to torquing of the robotic arms and/or cannula due to mutual interference as the robotic arms advance. In some more extreme cases, this could lead to difficulty with robotic arm advance, and/or to disturbance of the cannula position. Even if the alignment is correct enough to achieve safe robotic arm introduction to the region of surgical activity, robotic arms may not perform fully as expected, because of lateral interference forces. Such forces are potentially hard to judge by visual inspection to allow correction or compensation.

These concerns potentially applies to one or both of robotic arm movements fully under the direct guidance of a surgeon, and robotic arm movements which are at least partially under automatic control. Moreover, it can be difficult to judge the angle of approach and initial distance of longitudinal advance required for intended device operation, potentially leading to an iterative and/or painstaking setup period before the surgery can begin.

In some embodiments of the present invention, apparatus elements attached to the cannula are deployed to provide indications of where a robotic arm device should be placed. In some embodiments, these elements include spacing devices and/or guides which, once deployed, provide clear indications of whether robotic arm-cannula alignment is correct, and/or help to prevent incorrect alignments.

Before explaining at least one embodiment of the invention in detail, it is to be understood that the invention is not necessarily limited in its application to the details of construction and the arrangement of the components and/or methods set forth in the following description and/or illustrated in the drawings. The invention is capable of other embodiments or of being practiced or carried out in various ways.

Dilator and Cannula for Rectouterine Pouch Access

Reference is now made to *Figure 1*, which is a schematic representation of portions of a human female pelvic anatomy, referenced by descriptions herein according to some embodiments of the present disclosure.

Referenced in particular by descriptions herein is rectouterine pouch **19**, including a portion of rectouterine pouch wall **3** accessible from vagina **17**. Also shown in *Figure 1* are

bladder **15**, uterus **13**, and rectum **11**. As shown, uterus **13** is in an anteverted position (*i.e.*, tilted forward toward the bladder). In a significant number of patients, uterus **13** may be retroverted (*i.e.*, tilted posteriorly), or in another position. In some embodiments, a retroverted uterus can be manipulated into a position which improves intraperitoneal access through the rectouterine pouch **3**; for example, by use of a uterine manipulator.

Reference is now made to *Figure 2A*, which schematically represents a kit of trocar components **200**, according to some embodiments of the present disclosure.

In some embodiments, the trocar components **200** are sized and shaped to open an intraperitoneal aperture in the rectouterine pouch **19**, transvaginally via vagina **17**.

In some embodiments, trocar components **200** comprise a trocar needle **207**, an inner dilator **205**, an outer dilator **203**, and/or a cannula **201**, **1010**. The components are optionally sized and shaped to be nested one within the next in the order listed. Optionally, a distance of longitudinal advancement of trocar components **200** along one another is indicated and/or limited by use of a stopper and/or indicating indexer, for example, leaf spring device **503** or another device, for example as described in relation to *Figures 5A–5C* herein.

Reference is now made to *Figure 2B*, which schematically represents distal portions of components in the kit of trocar components **200**, according to some embodiments of the present disclosure. Reference is also made to *Figures 3A–3H*, which schematically represent a method of using trocar components **200** to establish intraperitoneal access through rectouterine pouch wall **3** or another body wall, according to some embodiments of the present disclosure.

While *Figures 3A–3H* illustrate a two-dilator expansion procedure, it is to be understood that more dilators (*e.g.*, three, four, five, or more) are optionally used. Using more dilators is optionally coupled to steeper-sloped dilator tip designs (*i.e.*, less expansion per mm of advance), which can help to reduce resistance to insertion. Optionally, only one dilator is used (for example as described in relation to *Figures 6A–6E* herein). The inventors have found that two dilators, each expanding along a longitudinal distance of about 15 mm, are apparently enough to reach a fully dilated size of about 30 mm x 10 mm, without undue use of insertion force and/or elevated risk of patient injury. In particular, 15 mm appears to be a safe distance of direct penetration through a vaginal wall **3** into rectouterine pouch **19**, which does not carry a significant risk of accidental injury to the adjacent rectum **19**.

In *Figure 3A*, a distal end of inner dilator **205** is shown advanced to wall **3** of rectouterine pouch **19**, *e.g.*, with blunt tip **215** positioned in contact with wall **3**. The advance is made transvaginally, in some embodiments. Vagina **17** is not shown in the sequence of *Figures 3A–3H*,

but may be understood to surround distal portions of the trocar components near the rectouterine pouch wall **3**.

In *Figure 3B*, a sharp tip **217** of trocar needle **207** is advanced out of a distal port **216** of inner dilator **205** sufficiently to puncture wall **3** and enter rectouterine pouch **19**.

5 Optionally, trocar needle **207** is no more than 2 mm in diameter (distal port **216** is sized large enough to pass trocar needle **207**; for example, distal port **216** may be about 2.1 mm in diameter to pass a trocar needle **207** having a 2 mm diameter). Potentially, this limitation on diameter helps to reduce a risk of serious complications developing in the case of accidental penetration into rectum **11**. Optionally, trocar needle **207** comprises a Veress needle having a
10 blunt, spring-loaded center stylet which in its extended position prevents the sharp tip of the needle from injuring tissue, but collapses upon sufficient longitudinal force being exerted so that the sharp tip can operate to penetrate tissue. Such a needle potentially serves to prevent unintended injury (*e.g.*, penetration to the rectum **11**) during penetration of the wall **3** of the rectouterine pouch **19**.

15 In some embodiments, trocar needle **207** is restrained by a stopper device from protruding more than a few millimeters (*e.g.*, no more than about 3 mm, 5 mm, 8 mm, or 10 mm) from the distal tip of inner dilator **205** by a stopper and/or indicating indexer. Potentially, the restriction on protrusion reduces opportunity for the needle cause injury by over-penetrating the outer tissue wall to be dilated and injuring an internal tissue surface. A penetration distance
20 chosen, *e.g.*, 5 mm, may be enough to stretch and puncture with the trocar needle **207** an outer tissue wall having tissue pressed against the inner dilator **205**, while being short enough that puncture of any deeper tissue layer beyond the outer tissue wall is prevented. A method of positioning trocar needle **207** for penetration is described, for example, in relation to *Figure 4*, herein.

25 In the position of *Figure 3C*, inner dilator **205** is advanced through the hole opened by trocar needle **207**, up to about the wide cross-section **219** of the distal region of inner dilator **205**. During the advance, blunt tip **215** of inner dilator **205** is first pushed into the hole in wall **3** made by needle **207**. Further advance of dilator **205** widens the hole in wall **3** according to the expansion of the tip through tapering region **218** of dilator **205** between blunt tip **215** and wide
30 cross-section **219**. In some embodiments, a first (distal) dilating stage of stepped dilator **1100** (that is, a portion of dilator **1100** comprising distal tapering region **1121**) is used for these operations.

In some embodiments, the overall distance between wide cross-section **219** and the distal-most profile of blunt tip **215** (at distal port **216**) is about 15 mm. Potentially, this distance

is short enough to prevent injury to the wall of the rectouterine pouch **19** opposite to the wall **3** which inner dilator **205** penetrates (*e.g.*, short enough to prevent injury to the rectum). However, providing some distance of expansion allows the widening slope of the dilator tip to provide mechanical advantage during insertion, so that tissue at the puncture is widened gradually. In some embodiments, another dilator tip length is used, for example about 10 mm, 12 mm, 14 mm, 16 mm, or 18 mm.

In some embodiments, blunt tip **215** rounds back from a substantially flat distal-most profile with a radius of curvature of about 2.5 mm. While a blunt profile potentially results in initially higher resistance to the advance of inner dilator **205**, the blunt tip profile has the potential advantage of reducing a likelihood of injury to the wall of the rectouterine pouch **19** opposite to the wall **3** which inner dilator **205** penetrates (*e.g.*, short enough to prevent injury to the rectum).

Optionally, cross-section **219** is about 20 mm across its longest axis, and about 10 mm across its shortest axis. Optionally, the largest axis of the inner dilator **205** at cross-section **219** is about, for example, 15 mm, 16 mm, 17 mm, 18 mm, 19 mm, or 20 mm. Optionally, the shortest axis of cross-section **219** is, for example, about 5 mm, 7 mm, 8 mm, 10 mm, 11 mm or 13 mm. Optionally, the ratio between the longest axis and the shortest axis is, for example, about 1.5, 2, 2.5, 3, or 3.5. In some embodiments, the expansion of one or both of the longest axis and the shortest axis of cross-section **219** uses the whole available length of the dilator tip. In some embodiments, proximal-going expansion across the shortest cross-sectional axis occurs over a distance of, for example, about 2.5 mm, 3 mm, or 4 mm of longitudinal travel, then levels out. Optionally, proximal-going expansion across the longest cross-sectional axis occurs over a distance of, for example, about 8 mm, 10 mm, 12 mm, 13 mm, 15 mm, or 18 mm.

Optionally, the expansion of the longest cross-sectional axis through tapering region **218** is substantially linear as a function of longitudinal distance for a portion of inner dilator **205** leading up to wide cross-section **219**. Optionally the expansion is non-linear, *e.g.*, curved to expand faster and/or slower as the overall perimeter of the entry hole into wall **3** grows larger. For example, a relatively blunt tip potentially takes advantage of tissue compliance due to elasticity around an initially small entrance incision, while a more gradual cross-section expansion is used where non-elastic expansion (*e.g.*, by tearing) dominates.

Optionally, there is a gradual increase in expansion rate (slope) moving still further proximally. Potentially, this allows percentage stretch of a hole perimeter as a function of longitudinal advance is maintained at a lower initial value than would be produced by a more linear expansion over the same distance. This may reduce resistance to insertion, potentially

reducing a risk of trauma. Optionally, either linear or non-linear expansion is used through tapering region **213** of outer dilator **203**.

In some embodiments, needle **207** is retracted before and/or as inner dilator **205** is advanced, potentially reducing a risk of injury to the opposite wall. In some embodiments, the distance of advance of inner dilator **205** is controlled by recording the position of inner dilator **205** upon needle penetration, and comparing that position to more advanced positions. Optionally, a stopper is positioned after initial penetration by needle **207** and locked into place relative to the patient (*e.g.*, locked to the table by a positioning arm) so that no more than a predetermined total advance distance (*e.g.*, 15 mm) is allowed.

In *Figure 3D*, outer dilator **203** is shown longitudinally advanced over inner dilator **205** until it reaches wall **3** of rectouterine pouch **19** (*e.g.*, so that distal edge **212** of outer dilator **203** contacts wall **3**). In *Figure 3E*, outer dilator **203** is advanced further, up to the level of the wide profile **214** of outer dilator **203**. It is noted that the advance in *Figure 3E* shows the distal edge **212** of outer dilator **203** brought to the same longitudinal position as the distal-most profile of inner dilator **205**. In some embodiments, a second (proximal) dilating stage of stepped dilator **1100** (that is, a portion of dilator **1100** of *Figures 11A–11E* comprising proximal tapering region **1117**) is used for these operations; except that instead of advancing over an inner dilator, the proximal tapering region **1117** follows after distal tapering region **1121**, and optionally after an intervening isolating region **1119**, as described herein in relation to *Figures 11A–11E*.

Figures 3A–3D show first insertion of inner dilator **205**, then outer dilator **203** over inner dilator **205**. In some embodiments, outer dilator **203** is optionally used as the *first* inserted dilator (or insertion along with inner dilator **205**), followed by puncture using trocar needle **203**, dilation with the inner dilator **205** sliding distally from within outer dilator **203**, and finally dilation with outer dilator **203** sliding distally over inner dilator **205**. This provides a potential advantage, for example, in the location of a puncture target location for needle **207**, which can be seen through the relatively open lumen of outer dilator **203** upon transillumination of the wall **3** of the rectouterine pouch **19**. The difference in insertion order can also affect the arrangement of stoppers/indicating indexers (for example as described in relation to *Figures 5A–5I*) used to determine the relative distances of the different dilation components during insertion and/or dilation.

Optionally, the distal aperture **220** of outer dilator **203** is sized to fittingly accommodate the wide cross-section **219** of inner dilator **205**. The outer perimeter at distal edge **212** may be just slightly larger than distal aperture **220** (*i.e.*, distal edge **212** is optionally sharp). Optionally,

distal edge **212** is blunt and/or rounded, for example, with an initial wall thickness (or diameter, in the case of a rounded edge) of, for example, about 100 μm , 200 μm , 500 μm , or 1 mm.

In some embodiments, the distance between distal edge **212** and wide profile **214** is about 15 mm. In particular, the distance is optionally any distance described in relation to the distance
5 between the distal-most profile of inner dilator **205** and wide cross-section **219**. Optionally, these two distances are about equal. Optionally, the distance for outer dilator **203** is slightly shorter (*e.g.*, about 0.5 mm, 1 mm, or 1.5 mm shorter), which potentially reduces a possibility of injury due to tissue contact with distal edge **212**. Optionally, a distance of advance of outer dilator **203** relative to inner dilator **205** is controlled by use of a stopper and/or indicating indexer, for
10 example as described in relation to *Figures 5A–5C* herein.

Optionally, the longest axis of wide cross-section **214** is, for example, about 20 mm, 23 mm, 25 mm, 27 mm, 33 mm, or 35 mm. Optionally, the shortest axis of wide cross-section **214** is, for example, about 5 mm, 7 mm, 8 mm, 10 mm, 11 mm, or 13 mm. Optionally, the ratio
15 between the longest axis and the shortest axis is, for example, about 1.5, 2 2.5, 3, or 3.5. The dimensional descriptions apply as well to slanted distal aperture **209** of cannula **201**, **1010** and the internal lumen of cannula **201**, **1010** which is sized to slide over outer dilator **203** in a fitting association (*e.g.*, with a relative size tolerance of about 1 mm or less). The internal lumen of cannula **201**, **1010** is sized to accept a plurality of (typically tubular, tube-sheathed and/or cylindrical) tools positioned side-by side, for example, two tools of 8 mm diameter or more and
20 one tool of about 5–6 mm diameter or more. In some embodiments, the cross-sections of the cannula **201**, **1010** and dilators **203**, **205** are (for example as shown in *Figures 2B* and *5K*) substantially rectangular with rounded ends (*e.g.*, rounded so that the short sides are formed as sections of substantially circular arcs).

A rounded-end shape has a potential advantage for enclosing a plurality of side-by-side
25 cylindrical tools, since it allows packing of the outermost tool sides toward the outside of the cross-section, while minimizing wasted corner space (the “waste” is not so much with respect to the cannula interior as it is with respect to the creation of a potentially larger-than-necessary incision hole). Round corners (as opposed to sharp corners) also provide a potential advantage during insertion, by helping to distribute forces which might otherwise tend to focus cutting
30 toward the corners and lead, *e.g.* to less predictable dilation and/or incisions which heal more unpredictably. Between the arced ends, a straight line section (used in some embodiments) has the potential advantage of maintaining an outer profile suitable for staying in full peripheral contact with the dilated tissue surrounding it (*e.g.*, to maintain a tension seal), without extra widening which would create space that the cylindrical tools within do not need to use. In some

embodiments, there is a slight outward bowing introduced along the longer sides of the cross-sections (*e.g.*, less than 1 mm of bowing per 5 mm of perimeter), potentially enhancing tension contacts between the dilators and/or cannula and the aperture edges of the tissue wall they penetrate.

5 In the examples illustrated herein (*e.g.*, *Figures 2B, 5K*), the cross-sections of the dilators and cannula shown display mirror symmetry around both a long axis and a short axis. However, the cross-sectional shape, in some embodiments, need not have any particular symmetry arrangement, and can be otherwise shaped (*e.g.*, as a round-cornered irregular triangle) to suit the accommodation of different-sized arrangements of tools.

10 In *Figure 3F*, vaginal cannula **201** is shown with its distal-most edge **211** brought up to the position of wall **3**. In *Figure 3G*, cannula **201** is shown advanced into the rectouterine pouch **19**, so that distal-most edge **211** is about even with the distal-most portion of outer dilator **203**. *Figure 3H* shows cannula **201** with inner dilator **203** and outer dilator **205** removed. Optionally, inner dilator **205** is removed at any time after outer dilator **203** is in place. Optionally, the two
15 dilators **203, 205** are removed together.

In some embodiments, an inner lumen cross-section of cannula **201** is sized to fittingly slide over section of outer dilator **203** having the size of wide cross-section **214**. Optionally advance of cannula **201** is performed by use of a handle **202**. Optionally, the maximum advance of cannula **201** relative to outer dilator **203** is controlled by the use of a stopper and/or indicating
20 indexer.

In some embodiments, the lumen of cannula **201** is about 7 cm, or another length long enough to reach between a distal-most position within the rectouterine pouch, and a proximal position at or outside the vaginal orifice (*e.g.*, up to about 15 cm). Optionally, a flange **221** is provided at a proximal end of the lumen of cannula **201**. While cannula **201** (with a handle **202**,
25 as next described) is shown in *Figure 3F–3I*, it should be understood that the foregoing descriptions of cannula **201** also apply, in some embodiments, to the use of cannula **1010** of *Figure 10B*.

Handle **202** of cannula **201** is long enough, in some embodiments, to extend past either of the dilators **203, 205** while they are inserted, and enough past to provide a grasping region (*e.g.*,
30 a grasping region of about 10 cm; overall length is optionally at least about 37 cm). In some embodiments, inner dilator **205** is at least long enough to reach its fully inserted position while providing a handle (*e.g.*, about 17 cm overall), and outer dilator **203** is at least 10 cm longer again (*e.g.*, about 27 cm overall).

Reference is now made to *Figure 3L*, which schematically represents a wider view (compared to *Figure 3H*) of the positioning of cannula **201**, **1010** relative to anatomical structures of a female lower abdomen/pelvic region, according to some embodiments of the present disclosure. Among the anatomical structures shown are the uterus **15**, vagina **17**, bladder **15**, rectum **11**, and rectouterine pouch **19**, also shown as in *Figure 1*. Not shown is the handle of cannula **201**, **1010**, and optional associated devices such as insufflation sealing which may be provided for, *e.g.*, at the vaginal orifice, and/or a trocar part which may be telescopically fitted to extend the length of cannula **201**, **1010**.

In some embodiments, slanted distal aperture **209** of cannula **201**, **1010** is slanted at an angle between a leading distal-most edge portion **211**, and a following, more proximal edge portion **210**. The longitudinal distance between distal-most edge portion **211** and proximal-most edge portion **210**, in some embodiments, is about 15 mm. In some embodiments, the distance is, for example, about 10 mm, 12 mm, 14 mm, 16 mm, or 18 mm. A potential advantage of the slanting of distal aperture **209** is to allow the cannula edge to be relatively retracted on an unprotected side of the cannula which could otherwise be accidentally positioned to scrape the rectum **11**. As for the more-protruding leading-edge side of the aperture **211**: (1) upon insufflation, tissue is generally lifted away from the rectum, reducing contact risk posed by this side, and (2) the robotic arms, where they exit the cannula, will generally be curved across the plane of the leading edge as they reach deeper into the peritoneal space. This potentially prevents contact of the leading edge with delicate internal tissues. Robotic arm positions can be seen, for example, in *Figures 7A–7B*, herein.

Reference is now made to *Figure 3I*, which schematically represents the cannula configuration of *Figure 3H*, along with inserted tools, according to some embodiments of the present disclosure. Reference is also made to *Figures 7A–7B*, which are images (from a viewpoint inside an insufflated abdomen) of robotic arms inserted through the cannula **201** in a configuration similar to that of *Figure 3I*, according to some embodiments of the present disclosure. Descriptions in relation to *Figures 3I* and *7A–7B* of cannula **201** also apply to cannula **1010** of *Figure 10B*.

In *Figure 3I*, the lumen of cannula **201** is shown occupied by two substantially cylindrical members, for example, robotic arms **305** (having diameter, for example, of about 8.6 mm), and another cylindrical (*e.g.*, tubular) member comprising a tool **307** (for example, a laparoscopic illuminator, camera, clamp, cutter, and/or another tool). In some embodiments, the luminal cross section of cannula **201** is sized to accommodate a plurality of substantially

cylindrical members (*e.g.*, robotic arms) each having a diameter of at least about 8 mm, and optionally another tool operated through a tube having a diameter of at least about 5 mm.

As shown in *Figure 3I*, tools exiting the slanted distal aperture **209** of cannula **201** positioned within rectouterine pouch **19** are optionally oriented to curve downward from the aperture, into the rest of the peritoneal space where operations of a laparoscopic procedure are to be performed (for example, clipping ligaments in preparation for hysterectomy. Optionally, this orientation is assisted by the angled slot of distal aperture **209** described in relation to *Figures 3F–3H*, herein.

In *Figures 7A–7B*, robotic arms **305** include surgical tools **704** positioned on a distal end of each robotic arm **305**. The robotic arms **305** are shown entering into an insufflated abdominal cavity from cannula **201** through slanted distal aperture **209**. Also illustrated in the image is the dilated incision **702** through which cannula **201** has entered the abdominal cavity.

Reference is now made to *Figure 3J*, which is a schematic flowchart outlining preparation for a laparoscopic procedure using trocar **200**, according to some embodiments of the present disclosure. The procedure shown represents a setup portion of a larger procedure (for example, a hysterectomy), including elements of the method described in relation to *Figures 3A–3I*.

Optionally, at block **110**, in some embodiments, a laparoscopically mounted camera is inserted into the peritoneal space, for example from the umbilical. Optionally, the camera is provided together with an illumination source. Optionally, a separately mounted illumination source is also inserted from the umbilical.

At block **112**, in some embodiments, the peritoneal space is insufflated, *e.g.*, by inflating with CO₂. Insufflation is performed to enhance access and/or visibility during the procedure.

At block **114**, in some embodiments, a sleeve is inserted transvaginally. The sleeve potentially helps to maintain insufflation pressure (pseudo-peritoneum) used for visualization and/or access during the rest of the procedure. The sleeve may be provided, for example, as an Alexis® nylon sleeve (Applied Medical Resources Corporation), or as part of a GelPOINT® path transanal access platform (Applied Medical Resource Corporation). In some embodiments, the sleeve is inserted at a later stage; for example, just before attachment of a sealing unit at block **122**.

At block **116**, in some embodiments, a speculum is inserted to the vagina to assist in visualization (optionally or additionally, a tenaculum is used).

At block **118**, in some embodiments, a uterine manipulator is inserted transvaginally to the uterus. The uterine manipulator may be, for example, a Karl Storz uterine manipulator. The uterine manipulator is used to move the uterus during the procedure, for example, to help provide

maneuvering room for other instruments, and/or to help move the uterus away from the rectum to provide increased safety. If a tenaculum was used, the tenaculum may be removed at this point.

At block **120**, in some embodiments, dilation and cannula introduction is performed, for example, as described in relation to *Figures 3A–3G*, *Figure 3K*, *Figures 6A–6E*, and/or *Figures 11A–11E* herein. The speculum may be removed during dilation, for example if it begins to interfere with dilation, or afterward.

At block **122**, in some embodiments, a sealing unit is attached to cannula **201** (or cannula **1010**, for example as described in relation to *Figures 10A–10C*, herein). The sealing unit optionally comprises elements of a GelPOINT® system; optionally in combination with elements specifically adapted for use with cannula **201**. Optionally, the uterine manipulator is re-positioned to pass through the sealing unit.

At block **124** in some embodiments, fixation is performed. Fixation comprises securing of the cannula and/or sealing unit to a platform which is stationary relative to the patient (*e.g.*, an operating table).

At block **125**, in some embodiments, one or more robotic arms are aligned to cannula **201**, **1010** in preparation for introduction into cannula **201**, **1010**, for example as described in relation to *Figures 9A–9D* or *Figures 10C–10J*, herein.

At block **126**, in some embodiments, one or more robotic arms and/or other tools are introduced through the cannula **201**, **1010**, for example as described in relation to *Figure 3H* herein.

Reference is now made to *Figure 3K*, which is a schematic flowchart representing a method of dilating and cannulating an access incision into a rectouterine pouch, according to some embodiments of the present disclosure. In some embodiments, the method of *Figure 3K* details operations of block **120** of *Figure 3J*.

The method is outlined briefly. Additional details for the operations of *Figure 3K* are described in relation to *Figures 3A–3G*, herein.

At block **130**, in some embodiments, inner dilator **205** (alternatively, in some embodiments, stepped dilator **1100** of *Figures 11A–11E*) is inserted transvaginally to reach a wall **3** of the rectouterine pouch **19** (for example as described in relation to *Figure 3A*). At block **132**, in some embodiments, a tip of a needle (inserted through inner dilator **207**) is located at the site where an access incision is to be made (for example, as detailed in relation to *Figure 4*). At block **134**, in some embodiments, the rectouterine pouch is punctured (for example, as described in relation to *Figure 3B*). At block **136**, in some embodiments, the pouch wall **3** is crossed with the

first dilator stage, for example, a tip of inner dilator **205** (for example as described in relation to *Figure 3C*). Alternatively, in some embodiments, pouch wall **3** is crossed with a first (distal) dilating stage of stepped dilator **1100**. At block **138**, in some embodiments, the pouch wall **3** is crossed with a second dilator stage, for example, the outer dilator **203** (for example, as described in relation to *Figures 3D–3E*). Alternatively, in some embodiments, pouch wall **3** is crossed with a second (proximal) dilating stage of stepped dilator **1100**. At block **140**, in some embodiments, the pouch wall is crossed with cannula **201**, **1010** (for example, as described in relation to *Figures 3F–3G*). At block **142**, in some embodiments, the dilators **203**, **205** and/or trocar needle **207** are removed, and the flowchart ends.

Reference is now made to *Figure 4*, which schematically represents a dual-verification method of locating an incision for providing transvaginal access to a rectouterine pouch **19**, according to some embodiments of the present disclosure.

Shown in *Figure 4* is a schematic representation of rectouterine pouch **19**, including a vaginal wall **3** of the rectouterine pouch **19**, approached by an inner dilator **205**, with a trocar needle **207** partially extended. Also shown are an intraperitoneally inserted (*e.g.*, from the umbilical) camera **402** and an illumination source **401**. Camera **402** and illumination source **401** are shown on the same laparoscopic instrument **400**; optionally they are separately provided. Camera **402** has an associated field of view **402A**, while illumination source **401** has an associated illumination field **401A**.

In some embodiments, the tip of needle **207** is positioned before puncture against a region of wall **3**, wherein the region selected based on external observation of light from illuminator **401** visible from outside (*e.g.*, as viewed transvaginally, optionally using the speculum and/or uterine manipulator to increase visibility). Optionally, the region selected is a region through which trans-illumination light intensity is observed to be relatively large compared to surrounding regions. Such well trans-illuminated wall areas are potentially among the thinnest, most easily penetrated portions of wall **3** accessible to needle **207**.

In some embodiments, as needle **207** is pressed against wall **3** to puncture it, camera **402** is used to visualize the results. In the camera images, for example, there may be initially a protrusion, other tissue distortion, and/or other change (*e.g.*, a color change due to pressure on the tissue) visible at the site of penetration (*e.g.*, a distortion visible from a side of wall **3** opposite a side contacting the needle), and afterwards the needle itself may become visible. Optionally, the visualization helps to verify that the intended region is being penetrated (*e.g.*, that the region being penetrated is suitable to provide intraperitoneal access), and/or to help identify and/or prevent imminent rectal puncture or another insertion mistake. By using visual cues that

communicate across the rectouterine wall **3**, even before the wall **3** itself has been punctured, complications may potentially be avoided.

Reference is now made to *Figures 5A–5C*, which schematically represent different stopper and/or movement interference devices for use with trocar components **200**, according to some
5 embodiments of the present disclosure.

Figure 5A illustrates a stopper-and-shoulder type of stopper device. In this embodiment, outer dilator **203** is inserted first, or along with inner dilator **205**. The view of *Figure 5A* cuts away a portion of outer dilator **203** so that the connection between inner dilator **205** and proximal stopper **501** can be seen.

Proximal stopper **501** is positioned along inner dilator **205** at a position where surface **501A** is brought into abutment with a proximal surface (cut-away in *Figure 5A*) of outer dilator **203** at the position where inner dilator **203** is as far distal relative to outer dilator **205** as it should be allowed to go (*e.g.*, with a distal-most portion of inner dilator **205** positioned 15 mm in advance of a distal-most portion of outer dilator **203**). Then, when it is the turn of outer dilator
10 **203** to be advanced, the distance between surface **501A** and the proximal surface of outer dilator **203** can be measured to determine the advance distance. Optionally another stopper device (for example, that of *Figure 5B* and/or *5C*) prevents and/or indicates distal over-advancement of outer dilator **203** relative to inner dilator **205**.

Figure 5B illustrates a leaf-spring based stopper and/or motion interference device. The
20 device comprises a bracket **504** attached to a distal portion of outer dilator **203**, which in turn supports a leaf spring **505** configured to press down onto the body of inner dilator **205**. This interaction optionally is set to a force sufficient to retard free sliding (*e.g.*, due to gravity) of the inner dilator **205** relative to the outer dilator **203**.

Optionally, inner dilator **205** comprises a receiving shape **505A** (hidden by leaf spring
25 **505**) which is positioned to contact leaf spring **505** and interfere with further longitudinal movement of outer dilator **203** relative to inner dilator **205**. The stopper device may be used to prevent over-advancement of either or both of inner dilator **205** and outer dilator **203** with respect to one another. There may be a plurality of different receiving shapes **505A**, allowing different positioned to be noted. Though referred to as a “stopper”, the stopper device of
30 *Figure 5B* is optionally configured using shapes and/or surface friction that interfere to resist, rather than completely prevent further longitudinal motion. The stopper device of *Figure 5B* in such embodiments may be alternatively described as an indicating indexer. Optionally, the indicating indexer indicates relative dilator positions by the positions from which further

longitudinal movement is resisted, and/or positions at which the two dilators click into place (as indicated through audible and/or tactile feedback).

The interfering occurs at one or more relative longitudinal positions of the two dilators **203**, **205**, for example, when the two are positioned with their distal-most portions in alignment, as shown in *Figure 3E*. Receiving shape **505A** may comprise, for example, an inset portion into which leaf spring **505** falls when the two parts are aligned. Additionally or alternatively, receiving shape **505A** comprises a raised portion. In this case, a lumen of outer dilator **203** may be shaped to pass over the raised portion of receiving shape **505A** until leaf spring **505** encounters it. Alternatively, lumen of outer dilator **203** may be shaped to *prevent* passage of the receiving shape, so that it also acts as a type of stopper-and-shoulder arrangement. It should be understood that although the device is illustrated with respect to two nested dilators, it may alternatively or additionally be implemented between a cannula and a dilator.

Figure 5C represents a different leaf-spring based stopper device. In this example, there are optionally provided be a plurality of leaf springs **503** mounted to the body of outer dilator **203**. Somewhere along its length, inner dilator **205** has a receiving shape (not shown), which comprise an indentation and/or a raised portion that contacts leaf spring **503** at a certain relative longitudinal position of the two dilators **203**, **205**; for example, when the two are positioned with their distal-most portions in alignment, as shown in *Figure 3E*. The leaf spring device of *Figure 5C* can be placed in any suitable location along the body of outer dilator **203**, but preferably in a location that remains outside body cavities during use. There may be a plurality of such devices provided on outer dilator **203**, and/or a plurality of receiving shapes provided on inner dilator **205**. Optionally, the leaf spring device is provided on inner dilator **205**, and the outer dilator provided with the appropriate receiving shape(s).

In some embodiments, a stopper and/or motion interference device replaces the leaf spring with another mechanism. For example, in some embodiments, a plunger is provided (*e.g.*, a spring-loaded pressing member such as a ball bearing, for example as described in relation to *Figures 8A–8B*). The plunger presses down to hold two nested parts in relative position (*e.g.*, two dilators; dilator and cannula), and/or indicate that a particular relative position has been reached, by force of friction and/or by interference between the plunger and one or more stopper/indicating indexing shapes—and/or by release of such interference and/or friction.

Reference is now made to *Figures 5D–5F*, which schematically illustrate a needle **207**, needle-passing dilator handle **510**, and needle handle **511**, according to some embodiments of the present disclosure.

In *Figure 5D*, needle **207** is shown partially inserted within the hollow body of dilator handle **510**. Optionally, dilator handle **510** serves as a handle for inner dilator **205**; *e.g.*, attached via a screw thread or other attachment means. Proximal end **512** of needle **207** is adapted to attach to needle handle **511** of *Figure 5E*, for example as shown in *Figure 5F*. In some 5 embodiments, the relative lengths of needle **207** and dilator handle **510** (and the relative position of handle **511** on needle **207**) are configured to help control the maximum distance of distal advance of needle **207** relative to inner dilator **205**, for example as described in relation to *Figures 5G–5I*.

Reference is now made to *Figures 5G–5I*, which schematically illustrate devices for 10 controlling the relative positioning of inner dilator **205**, outer dilator **203**, and needle **207** according to some embodiments of the present disclosure. Reference is also made to *Figures 8A–8B*, which illustrate scale features of outer dilator **203** and inner dilator **205**, according to some embodiments of the present disclosure.

In *Figure 5G* and *Figures 8A–8B*, outer dilator **203** is shown provided with an integrated 15 scale window **515**. Scale window **515** optionally comprises at least one reference mark alongside a longitudinal window formed in a portion (*e.g.*, cut into a proximal end) of outer dilator **203**. In some embodiments, outer dilator **203** is used to define a reference position allowing inference of the current position of the wall **3** of rectouterine pouch **19**. Outer dilator can be inserted as far as it will easily pass. Since it is too blunt to puncture wall **3**, it will normally be brought to a halt 20 with its distal-most end abutting wall **3**. This can be verified by viewing using a speculum, for example.

In some embodiments, a distance scale is marked (*e.g.*, in centimeters from the distal end) on one or both of the inner dilator **205** and outer dilator **203**, for example, scale **811** of the outer dilator **203**, and scale **813** of the inner dilator **205** (*Figures 8A–8B*).

Also illustrated in *Figures 8A–8B* is ball-stop device **815**, which is an example of an 25 indicating indexer. In some embodiments, ball stop device **815** comprises a spring-loaded ball bearing, configured to protrude outwards from near a proximal end of inner dilator **205** under elastic pressure. The lumen of outer dilator **203** is sized so that it can be pushed over and then slide forward along ball-stop device **815**. The ball of ball-stop device **815** is pushed inward 30 during this motion. It acts (by continuing to press outward) to help center outer dilator **203** over inner dilator **205**, and optionally to resist spontaneous relative translation (sliding) of the two dilators (*e.g.*, due to the weight of outer dilator **205**). Upon sufficient advance of outer dilator **203**, ball-stop device **815** is freed from the proximal side of outer dilator **203**, potentially inducing a tactile and/or audible click, and/or causing changes to the mechanical handling of the

dilators **203**, **205** which indicate to a user that outer dilator **203** has been fully advanced. Optionally, outer dilator **203** comprises one or more indentations and/or bumps along its lumen at positions which change the force with which the ball-stop device **815** interferes with relative longitudinal translation of the dilators **203**, **205**. It should be understood that the ball-stop device **815** may alternately be provided on the outer dilator **203** at a position where it interacts while sliding with the inner dilator **205** (and optionally bumps and/or indentations thereof). An indicating indexer is optionally provided for indication and/or control of relative longitudinal positions of other pairs of elements of the trocar kit; for example, between inner dilator **205** and needle **207**, and/or between outer dilator **203** and cannula **201**, **1010**. In some embodiments, a ball stop is provided which controls relative motion between stepped dilator **1100** and cannula **201**, **1010** when locked. In some embodiments, a ball stop is provided which controls relative motion between cannula **1010** and an access device **1001** (*Figure 10A*) when locked.

In *Figure 8B*, the ball stop device **815** is shown still compressed by outer dilator **203**. In *Figure 8A*, the ball stop device is free of compression. Once the ball-stop device **815** is free of compression, the ball is pressed outward to the full extent of its travel, resulting in an increased resistance (upon contacting a proximal surface **817** of outer dilator **203**) to accidentally pushing inner dilator **205** distally relative to dilator **203**.

In *Figure 5H*, inner dilator **205** is shown partially inserted into outer dilator **203**. Scale markings **517** on inner dilator **205** allow monitoring of the distal advance of inner dilator **205** relative to outer dilator **203**, *e.g.*, to a position distally even with the distal end of outer dilator **203** (in preparation for needle puncture), and/or to a position a few millimeters (*e.g.*, 15 mm) distally in advance of outer dilator **205** during initial dilation.

Also in *Figure 5H*, needle **207** is shown still only partially advanced with respect to handle **510** and the dilators **203**, **205**. In some embodiments, dilator handle **510** fits (*e.g.*, screws into) to a socket in inner dilator **205**, holding it at a predetermined distance from the distal end of inner dilator **205**.

In *Figure 5I*, the relative positions of components shown is as may occur immediately after needle **207** punctures the wall **3** of the rectouterine pouch **19**.

The distal-most end of inner dilator **205** has been brought forward even with the distal-most end of outer dilator **203**, as monitored from the relative positions of scale window **515** and markings **517**. Handle **511** has been pushed forward so that it abuts a proximal end of dilator handle **510**, preventing further advancement of needle **207**. Lengths of needle **207**, inner dilator **203**, and dilator handle **510** are set so that needle **207** now protrudes from the front of inner dilator **203** by a predetermined amount which has been determined to fall within the range of

distances which are enough to penetrate a wall **3** of a rectouterine pouch **19**, but avoid risk of also penetration a wall of the rectum **11**.

Optionally, the distal-most end of inner dilator **205** remains longitudinally offset from (e.g, distal to) the distal-most end of outer dilator **203** by some distance to control the maximum advance of needle **207**. Optionally, maximum advance of needle **207** relative to inner dilator **205** is performed first, before advance to penetrate wall **3**. This provides a potential advantage for allowing marked relative positions of window **515** and scale **517** to indicate the distance of needle advance in detail.

During dilation, in some embodiments, distal advance of outer dilator **203** relative to inner dilator **205** potentially disrupts the longitudinal frame of reference that outer dilator **203** initially establishes. In some embodiments, the frame of reference is maintained by clamping inner dilator **205** into place once it has advanced into the rectouterine pouch.

Reference is now made to *Figure 5K*, which illustrates a manufactured example of a dilation and cannulation kit comprising members described and illustrated in, for example, *Figures 3A–3I* and *5D–5I*, according to some embodiments of the present disclosure. Items are to scale with each other, and shown along their total length, with the exception of needle **207**, which has been truncated on the right hand (sharpened, distal) side. Illustrated, from top to bottom, are:

- Needle **207** including handle **511**,
- Cannula **201**, including handle **202** and slanted distal aperture **209**, comprising proximal edge portion **210** and distal-most edge portion **211**.
- Outer dilator **203**, including scale window **515**, distal edge **212**, tapering region **213**, and the wide profile **214** at the proximal side of tapering region **213**.
- Inner dilator **205**, including distal port **216** positioned in blunt tip **215**, tapering region **218**, and the wide cross-section **219** at the proximal side of tapering region **218**.
- Dilator handle **510**, which may optionally be attached to (e.g., screwingly attached to) inner dilator **205** to serve as a proximal-side handle thereof.

Reference is now made to *Figures 6A–6E*, which schematically represent dilation using a single dilator trocar kit, according to some embodiments of the present disclosure.

Figure 6A shows a schematic representation of a body cavity (e.g., a vagina **17**), together with an access device **601** which is optionally used, e.g., to help maintain insufflation pressure. In *Figure 6B*, a uterine manipulator **603** is optionally added (shown incompletely inserted), potentially allowing maneuvering to increase visibility of the trocar target region (which may be a wall **3** of a rectouterine pouch **19**, not shown). In *Figure 6C*, cannula **605** (optionally

constructed as described herein in relation to cannula **201** or cannula **1010**) is inserted to the target region. Cannula **605** comprises a handle **605A** to allow maneuvering of the lumen region of the cannula **605**.

In *Figure 6D*, a dilator **607** is being inserted to cannula **605**, optionally using handle **607A**. The tip of dilator **607** is optionally shaped to begin at its distal-most longitudinal position with a cutting tip. Alternatively, the tip of dilator **607** is blunted (shaped, for example, like that of inner dilator **205**). Optionally, dilator **607** allows for passage distally of a trocar needle via an aperture. Optionally, initial puncture is made separately by a needle, *e.g.*, passed into the lumen of cannula **605** (not shown), and dilator **607** inserted afterward.

In some embodiments, the tip of dilator **607** expands to any appropriate cross-sectional size; for example, any cross-section described in relation to wide cross-section **219**. Optionally, the longitudinal distance between distal-most position and the tip cross-section of greatest expansion is about 15 mm or any other suitable distance; for example, as described in relation to dilator tips of dilators **203**, **205**. Relative to use of two or more dilators, one-step dilation over the same maximum dilator tip distance is potentially simpler from the point of view of component exchanges and manipulations. There is a potential tradeoff, however, of increased penetration length (*e.g.*, if expansion angle is maintained), and/or (*e.g.*, if expansion angle is increased) of increased resistance to penetration.

In *Figure 6E*, the tip of dilator **607** is shown maximally advanced, with dilation complete. To complete trocarization, the cannula **605** may now be advanced through the dilated aperture, and the dilator **607** removed.

Reference is now made to *Figure 5J*, which is a flowchart schematically outlining a method of using indicators to transitively establish and maintain known insertion depths of the trocar needle **207**, dilator **203**, **205**, and/or cannula **201**, **1010** parts, according to some embodiments of the present disclosure.

At block **551**, in some embodiments, a first dilator (*e.g.*, inner dilator **205**) is brought into position with its blunt tip against the outer wall of the rectouterine pouch. This position may be established, for example, by direct visualization (*e.g.*, using a speculum), by noting where insertion resistance is encountered, and/or indirectly, *e.g.*, by shining a light through the dilator lumen, and monitoring the projected light spot (*e.g.*, the position where the spot reaches its smallest, most sharply defined shape) through the rectouterine pouch wall using a camera positioned inside the intraperitoneal space. Optionally, the first dilator is held clamped in position, *e.g.*, by a table-attached clamping arm. Optionally, at block **552**, the depth of penetration relative to the natural orifice opening is noted, providing a total vaginal length (TVL)

which may be used separately from the transitive method of establishing trocar component positions, and/or to verify insertion depths set by the transitive method.

At block **553**, in some embodiments, a trocar needle **207** is inserted into dilator **205**, *e.g.*, via holder **210**. Insofar as holder **210** is itself sized to insert to a predetermined longitudinal position relative to the first dilator, the amount of visible shaft of trocar needle **207** optionally provides an indication of where the needle tip is positioned relative to the distal end of the first dilator. This can be used to control an advancing distance of trocar needle **207**, optionally along with a stopper device (such as a shoulder stopper) that prevents over-advancement of the trocar needle **207**. Optionally, the needle remains in place at least until the first dilator is advanced over it.

At block **555**, in some embodiments, a second dilator (*e.g.*, outer dilator **203**) is advanced over the first dilator until a part of the second dilator (*e.g.*, its proximal end, or an index mark) it is suitably aligned to a part of the second dilator (*e.g.*, an index marking on a scale). If an index scale is used, it may be on either or both of the first and second dilator. For the sake of description of the method, the distal ends of the first and second dilators are assumed to be aligned to one another in the aligned position; optionally, they are offset by some known amount. Additionally or alternatively, the insertion depth of the second dilator relative to the total vaginal length is used to longitudinally position and/or verify the position of the second dilator.

At this stage, the longitudinal positions of the distal ends of the first dilator and second dilator are both known relative to that of the rectouterine pouch wall which is to be dilated. Optionally, either dilator can be advanced or retracted relative to the other in any suitable sequence, and so long as the sequence of movements and their distances are tracked, their positions relative to the rectouterine pouch wall will remain known.

For example, at block **557**, in some embodiments, the first dilator is advanced into the rectouterine pouch while the second dilator remains fixed (*e.g.*, clamped). The relative motion is optionally monitored by looking at scale marking motions on the proximal ends of the dilators. At block **559**, *e.g.*, once the first dilator is sufficiently advanced (for example, 15 mm) to achieve a full first-stage dilation, the second dilator is advanced (*e.g.*, until the original relative alignment of the two is restored). Additionally or alternatively, the changing insertion depths are controlled/monitored relative to the total vaginal length.

Additionally or alternatively (at blocks **558** and/or **560**), advancing distance is controlled by making reference to the TVL determined at block **552**.

From this position, at block **561**, the first dilator (and trocar needle **207**, if not yet withdrawn) may be removed; and the second dilator may remain as a longitudinal positioning

reference for positioning the cannula **201** at block **563**. Optionally, the cannula **201** has a handle which is long enough to support a scale and/or reference mark that aligns with some visible part of the second dilator (a scale mark, distal end, or indicator mark) when the cannula **201** is in place. Additionally or alternatively, the cannula insertion depth is controlled/monitored relative to the total vaginal length.

At block **565**, the second dilator is removed. Cannula **201** is now positioned crossing the rectouterine wall, and at a known longitudinal depth relative to the rectouterine wall.

Optionally, the stepped dilator **1100** of *Figures 11A–11E* is used by this method, with the insertion of *Figure 6E* optionally occurring in two stages, one for each of distal tapering region **1121**, and proximal tapering region **1117**, optionally with a pause between the two stages enabled by allowing the physician to sense a change in insertion resistance upon transitioning from distal tapering region **1121** to isolating region **1119**.

Cannula Fixation and Robot Alignment to Cannula

Free-Positioned Alignment System

Reference is now made to *Figures 9A–9D*, which comprise views representing an instrument holder **900** for cannula **201** and its configuration for use, wherein instrument holder **900** includes a motor unit stopper **902** for use in setting an initial robotic arm position, according to some embodiments of the present disclosure.

Figure 9A schematically shows an instrument holder **900**, comprising a mounting block **901** and a stopper arm **902**. In some embodiments, stopper arm **902** is hinged upon hinge **904** to rotate to different orientations relative to mounting block **901**.

Figure 9B shows cannula **201** and associated handle **202** mounted to mounting block **901**. Mounting, in some embodiments, comprises positioning a proximal end of handle **202** in a well-defined position relative to mounting block **901**; for example, flush with an aperture **903** of a lumen within block **901** sized and shaped to receive handle **202**. Fixation of cannula **201** relative to block **901** is optionally assured by tightening a tightening handle **905**.

Figure 9C shows an overall mounting arm assembly **910**, comprising a table-mounting block **911** (configured, *e.g.*, with clamps and tightening handles supporting firm attachment to a table or other stabilizing surface), and jointed arm **912** extending distally from the mounting block **911** to attachment at its distal side to mounting block **901**.

Figure 9D shows the instrument holder **900** with motor unit stopper **902** configured to set an initial working distance of motor unit **930**. Motor unit stopper **902** is sized so that when positioned to horizontally protrude from block **901**, its distal end marks the distance at which

motor unit **930** should be set so that its arms (which themselves have a well-known predetermined length) reach a defined (and safe) initial distal-most position within cannula **201** (e.g., at the distal end of the cannula) when cannula **201** positioned relative to mounting block **901** as shown in *Figure 9B* (the cannula is not shown in *Figure 9D*). Robotic arms **305** (not seen) pass distally from motor unit **930** within extenders **920**. Extenders **920** comprise tubes having a lumen sized to pass the robotic arms **305** thereinto, for example, a lumen of, for example, at least 7 mm, 8 mm, 9 mm, or 10 mm. Extenders **920** may be aligned (e.g., set within slots **921** of *Figure 9B*) to guide the robotic arms **305** to cannula **201**.

By appropriate positioning of its parts at marked and predetermined positions, a kit comprising cannula **201** and handle **202**, and instrument holder **900** with motor unit stopper **902** (optionally including extenders **920** and/or motor unit **930** and associated arms **305**) potentially helps to achieve a rapid, reproducible initial setup providing a well-determined initial relationship between the distal end of cannula **201**, and the distal ends of robotic arms **305**, for example, an alignment of the two ends.

To allow initiation of robotic arm movement in a distal direction, motor unit stopper **902** is allowed to swing away from the horizontal position (e.g., downward), so that motor unit **930** can advance distally without interference. Preferably, attachment of motor unit stopper **902** to block **901** is configured to allow conversion between a first position that prevents advance of motor unit **930**, and a second position that allows it, without disturbing the position of structures on either side of it—for example, without disturbing positions of the cannula **201** or the motor unit **930**. The attachment is not necessarily by a hinge (for example, stopper **902** may be telescoping, slideable within block **901**, or otherwise moveable). A hinge provides a potential advantage by allowing a reproducible longitudinal stopper position to be obtained when the stopper is oriented in the horizontal (oriented to the longitudinal axis of the cannula) position, together with ready conversion to a non-stopping position without having to exert either longitudinal force to slide the stopper **902**, or torque on a fastener to release the stopper **902**.

Lock-Positioned Alignment System

Reference is now made to *Figures 10A–10E*, which schematically illustrate views representing a collapsing instrument holder **1000** for cannula **1010** and its configuration for use in setting an initial robotic arm position relative to cannula **1010**, according to some embodiments of the present disclosure. Reference is also made to *Figures 10F–10J*, which schematically represent components of collapsing instrument holder **1000**, according to some embodiments of the present disclosure. Further reference is made to *Figures 12A–12C*, which

schematically represent a duck-bill gasket **1050** used to seal access to the proximal aperture of access device **1001**, according to some embodiments of the present disclosure.

Figure 10A shows an access device **1001**, shaped for insertion into a gel seal **1003** (in a position indicated, for example, in *Figures 10B–10C*). In use, gel seal **1003** is positioned at the vaginal entrance to provide protected access; thus, elements positioned to the right of gel seal **1003** would be positioned intra-vaginally during a procedure, and elements positioned to the left would be positioned extra-vaginally.

Trans-seal region **1001A** of access device **1001** is flanged on either side, and seats within the gel membrane of gel seal **1003**. External (proximal) side **1001B** of access device **1001** (also referred to herein as a "trocar"), in some embodiments, is provided with a mounting projection **1002**. Lumen **1004** of access device **1001** is sized to allow insertion of stepped dilator **1100** (*e.g.*, as shown in *Figure 10B*), or optionally another dilator/dilator system, for example, a two-piece dilator comprising inner dilator **205** and outer dilator **203**. Also shown in *Figure 10B* assembled together with stepped dilator **1100** are needle handle **511** of needle **207** and dilator handle **510**.

Lumen **1004** of access device **1001** is also sized to allow insertion of a cannula **1010** over stepped dilator **1100** (or other dilatory system). When inserted to lumen **1004**, in some embodiments, cannula **1010** is fittingly contained by access device **1001**, and optionally locked thereto. Cannula **1010**, in some embodiments, comprises a body with an elongated (*e.g.*, oval) cross-section, having a slanted distal aperture **209**, for example as described in relation to *Figure 3L*, and a flange **221** at its proximal side. In some embodiments, flange **221** includes a receiving indentation for a ball stop **1006** provided on access device **1001** which controls relative motion between cannula **1010** and access device **1001** when locked.

In some embodiments, a two-seal "duck bill" gasket **1050** (shown in *Figure 10C*, and *Figures 12A–12C*) is inserted into the proximal aperture of access device **1001** after positioning of cannula **1010**. A first sealing member **1050A** of gasket **1050** is normally sealed (two opposite sides pushed together) when there is nothing inserted into cannula **1010**. Upon insertion of robotic arm guide **1032** (explained further in relation to *Figures 10D–10E*), first sealing member **1050A** is forced open, while a second sealing member **1050B**, which is normally open, is shaped so that it seals around the robotic arm guide **1032**. The sealing members **1050A**, **1050B** are made of a soft elastic material, for example, a silicone rubber. In some embodiments, gasket body **1050C** comprises a rigid polymer or metal material that provides support to sealing members **1050A**, **1050B** to help maintain the overall shape of gasket **1050**. In some embodiments, each of the sealing members **1050A**, **1050B** is shaped so that it has a long axis and a short axis, with the long axis being at least twice as long as the short axis.

The placement of the above-described elements generally corresponds, in some embodiments, to the operations of block **120** (dilate and introduce cannula) and block **122** (sealing unit) of *Figure 3J*.

Fixation corresponding to operation of block **124** of *Figure 3J*, in some embodiments, comprises attaching mounting projection **1002** to mounting block **1020**. Mounting block **1020** may itself be affixed, for example, to a platform stationary relative to the patient, such as an operating table. In some embodiments, attachment comprises insertion of mounting projection **1002** into a receiving aperture **1061** (shown in *Figure 10J*) of mounting block **1020**. In some embodiments, notch **1005** or another shape on mounting projection **1002** engages to a lock within receiving aperture **1061**. Optionally, button **1021** is pressed to assist and/or activate engagement and/or release of mounting projection **1002** from the lock.

Mounting block **1020** is a component of instrument holder **1000**. Attached to mounting block **1020**, instrument holder **1000** additionally comprises spacing arm **1024** and aligning arm **1030**. In some embodiments, spacing arm **1024** is attached to mounting block **1020** by stopped hinge **1022**, and aligning arm **1030** is in turn attached to spacing arm **1024** by stopped hinge **1028**. In some embodiments, spacing arm **1024** is telescoping (reversibly extendible and retractable). Optionally, release and/or locking of telescoping is controlled by a button **1026** or other control member.

The use of stopped hinges **1022**, **1028** and button **1026** to position spacing arm **1024** and aligning arm **1030** are further illustrated in *Figures 10D–10E*.

In some embodiments of the invention, cannula **1010** is used to provide intraperitoneal access to one or more robotic arms, for example robotic arms **305** as described, for example, in relation to *Figures 3I* and *7A–7B*. As also described in the *Overview* herein, it is a potential advantage to be able to position these robotic arms so that they begin at a well-known distance of longitudinal advance through cannula **1010**, and at a well-known angle of approach (generally an angle of approach axially aligned with cannula **1010**).

In *Figures 10D–10E*, spacing arm **1024** and aligning arm **1030** comprise an assembly shown fully deployed (in a deployed position) in order to help set a starting position for robotic arms being introduced into cannula **1010**. In some embodiments, this comprises:

- Rotating spacing arm **1024** 90° from its vertically downward stowed position (seen in *Figure 10C*) to a horizontal deployment position.
- Extending spacing arm **1024** to set the spacing distance by pressing on button **1026**, and pulling to expose telescoping arm portion **1024A**.

- Rotating aligning arm **130** from its stowed position against spacing arm **1024** (as in *Figure 10C*) to a vertical deployment position.

Optionally, spacing arm **1024** and aligning arm **1030** are returned to their stowed positions after use to position robotic arms correctly. A potential advantage of post-use stowing is so that they do not interfere with further movements of the robotic arms (*e.g.*, advancing deeper into the intraperitoneal cavity).

Figures 10F–10J illustrate these components in more detail. *Figures 10F–10H* show spacing arm **1024** in a collapsed (unextended) configuration. Stopped hinges **1022**, **1028** can be seen, including details of an optional embodiment thereof. For example, in the case of stopped hinge **1022**, each of (optionally four) projections **1063** is spaced around a circumference of stopped hinge **1022**. Each projection **1063** extends into a respective notch of plate **1022A**. The projections **1063** are fixed to the orientation of block **1020** (*Figure 10J*), while plate **1022A** is affixed to spacing arm **1024**. Accordingly, as long as the projections **1063** engage the notches of plate **1022A**, spacing arm **1024** is fixed in orientation. Upon actuating button **1062**, projections **1063** are recessed out of their notches, allowing spacing arm **1024** to rotate freely. Optionally, the projections **1063** are held in place by plate **1022A** once the notches are moved out of alignment with them, so that button **1022A** can be released while spacing arm **1024** continues to move freely.

Once spacing arm **1024** is deployed by a full 90°, the notches of plate **1022A** and projections **1063** come back into alignment, allowing projections **1063** to spring back into place, locking spacing arm **1024** in a new orientation. This mechanism has potential advantages of (1) only allowing one deployed orientation of spacing arm **1024**, and (2) being strong enough to hold the horizontally deployed weight of spacing arm **1024** and aligning arm **1030** without collapsing.

Stopped hinge **1028**, in some embodiments, comprises a similar mechanism, comprising projections **1065**, button **1067**, and notched plate **1066**. Notched plate **1066** is again affixed to spacing arm **1024**, while the projections **1065** are fixed to the orientation of aligning arm **1030**.

In the fully deployed position of *Figures 10D–10E*, horizontal bar **1068** (*Figure 10I*) of aligning arm **1030** acts to set an elevation across which robotic arms should be horizontally passed while aiming at cannula **1010** in order to be level with the longitudinal axis of cannula **1010**. Similarly, the robotic arms should pass through the space between vertical bars **1064** in order to approach cannula **1010** along its central longitudinal axis. Furthermore, in some embodiments, a motor unit or other housing holding a robotic arm which is being positioned is in position when some designated portion of it (a "stopper-receiving portion") is pressed up against

a portion of aligning arm **1030** that acts as a stopper, for example, pressed against the vertically extending main bar **1069** of aligning arm **1030**.

Returning now to *Figures 10D–10E*: in some embodiments, robotic arms **305** are sheathed (*e.g.*, two round arms side-by-side) using arm sheath **1032** before being passed into cannula **1010** (*e.g.*, in performing the operations of block **126** of *Figure 3J*). In some
5 embodiments, arm sheath **1032** has an inner lumen sized to allow at least two robotic arms having a minimum outer diameter of about 8.6 mm to pass therethrough. In some embodiments, for example, a minimum diameter of the inner lumen of arm sheath **1032** is about 9 mm, 10 mm, or 11 mm.

A potential advantage of sheath **1032** is to ensure that the arms are straight, and moreover
10 to ensure that no part of them will be accidentally snagged during passage into and through cannula **1010**. In some embodiments, moreover, there is provided a gasket **1034**, comprising a fitted hole for each of two robotic arms that may be passed therethrough. Gasket **1034** fittingly attaches to a proximal end of arm sheath **1032**, to act as another protective seal. Optionally, arm
15 sheath **1032** is constructed from stainless steel. Optionally gasket **1034** is manufactured from a flexible polymer such as a silicone rubber.

Stepped Dilator

Reference is now made to *Figures 11A–11E*, which schematically represent a stepped dilator **1100**, dilator handle **510**, and trocar needle **207**, according to some embodiments of the
20 present disclosure. Reference

Dilator handle **510** and trocar needle **207** are, in some embodiments, substantially as described, for example, in relation to *Figures 5D–5F, 5H, 5I, and 5K*.

In some embodiments, stepped dilator **1100** comprises, along a distal working end **1115** of a single dilator, distal and proximal tapering regions **1121** and **1117** (*Figures 11B, 11C*). Distal
25 and proximal tapering regions **1121, 1117** are, in some embodiments, substantially shaped as described for embodiments of tapering regions **213** and **218** of the two-dilator combination which is described herein, for example, in relation to *Figures 2A–3I and 5K*. Nevertheless, some specific details of tapering region shapes are repeated here for clarity.

In some embodiments, distal (first) tapering region **1121** has a blunted distal-most
30 portion. The distal-most portion optionally has a port through which a trocar needle **207** can be extended. Optionally, the distal-most portion curves proximally, widening in both width and height through a radius of about 2.5 mm, then expanding primarily in width to form a wide oblong cross-section about 15 mm proximal to the distal-most portion (or another distance, for example in the range of about 10–20 mm).

In some embodiments, a non-dilating isolating region **1119** extends longitudinally between the distal and proximal tapering regions. Preferably, isolating region **1119** is long enough to allow an inserting physician to sense a reduction in insertion resistance upon passing the proximal side of distal tapering region **1121**, and reduce insertion force in response, so that dilation is paused. This distance is optionally in the range of about 5–15 mm. In some 5 embodiments, isolating region **1119** is of a constant cross-section extending proximally from the distal side of isolating region **1119** until reaching the distal side of proximal tapering region **1117**. Alternatively, isolating region **1119A** (*Figure 11C*), in some embodiments, comprises a constriction (*e.g.*, a tapering constriction) relative to the proximal-side cross-section of distal tapering region **1121**, which potentially acts as a detente that accentuates for a physician the 10 sensation of reduced insertion resistance. The constriction may also allow a physician to detect by feel a difference between an initial dilation after which the dilated tissue perimeter retains elasticity (tending to re-collapse the dilated opening), and a dilation where a tear has been induced, potentially reducing the tendency of the dilated opening to exert inward force on the 15 dilator.

In some embodiments, proximal (second) tapering region **1117** has a distal-most cross-section arising from the proximal-most portion of isolating region **1119**. From there, the second dilator's cross-section expands going proximally for about 15 mm (or another distance, for example in the range of about 10–20 mm). The maximum of the further expansion is an 20 expansion by about, for example, about 5 mm, 7.5 mm, 10 mm, or 12.5 mm. Optionally, there is a larger expansion along one axis of the incision cross-section than along another axis; for example, there may be a relative factor of expansion of about 1:1.5, 1:2, or 1:3.

In some embodiments, a body **1111** of dilator **1100** is constructed of a sterilizable and re-sterilizable (*e.g.*, autoclave compatible) material, *e.g.* stainless steel. Optionally, an inset region 25 **1113** is provided on one or both sides of body **1111**. This potentially reduces weight of dilator **1100** (*e.g.*, in embodiments where body **1111** is constructed of a solid piece of metal).

In some embodiments, dilator **1100** is at least 17 cm in overall length without handle **510**. In some embodiments, threaded region **1102A** is threaded to accept a thread of handle **510**. Handle **510** is optionally at least 20 cm long.

30 **General**

As used herein with reference to quantity or value, the term “about” means “within $\pm 10\%$ of”.

The terms “comprises”, “comprising”, “includes”, “including”, “having” and their conjugates mean: “including but not limited to”.

The term “consisting of” means: “including and limited to”.

The term “consisting essentially of” means that the composition, method or structure may include additional ingredients, steps and/or parts, but only if the additional ingredients, steps and/or parts do not materially alter the basic and novel characteristics of the claimed composition, method or structure.

As used herein, the singular form “a”, “an” and “the” include plural references unless the context clearly dictates otherwise. For example, the term “a compound” or “at least one compound” may include a plurality of compounds, including mixtures thereof.

The words “example” and “exemplary” are used herein to mean “serving as an example, instance or illustration”. Any embodiment described as an “example” or “exemplary” is not necessarily to be construed as preferred or advantageous over other embodiments and/or to exclude the incorporation of features from other embodiments.

The word “optionally” is used herein to mean “is provided in some embodiments and not provided in other embodiments”. Any particular embodiment may include a plurality of “optional” features except insofar as such features conflict.

As used herein the term “method” refers to manners, means, techniques and procedures for accomplishing a given task including, but not limited to, those manners, means, techniques and procedures either known to, or readily developed from known manners, means, techniques and procedures by practitioners of the chemical, pharmacological, biological, biochemical and medical arts.

As used herein, the term “treating” includes abrogating, substantially inhibiting, slowing or reversing the progression of a condition, substantially ameliorating clinical or aesthetical symptoms of a condition or substantially preventing the appearance of clinical or aesthetical symptoms of a condition.

Throughout this application, embodiments of this disclosure may be presented with reference to a range format. It should be understood that the description in range format is merely for convenience and brevity and should not be construed as an inflexible limitation on the scope of the disclosure. Accordingly, the description of a range should be considered to have specifically disclosed all the possible subranges as well as individual numerical values within that range. For example, description of a range such as “from 1 to 6” should be considered to have specifically disclosed subranges such as “from 1 to 3”, “from 1 to 4”, “from 1 to 5”, “from 2 to 4”, “from 2 to 6”, “from 3 to 6”, *etc.*; as well as individual numbers within that range, for example, 1, 2, 3, 4, 5, and 6. This applies regardless of the breadth of the range.

Whenever a numerical range is indicated herein (for example “10-15”, “10 to 15”, or any pair of numbers linked by these another such range indication), it is meant to include any number (fractional or integral) within the indicated range limits, including the range limits, unless the context clearly dictates otherwise. The phrases “range/ranging/ranges between” a first indicate number and a second indicate number and “range/ranging/ranges from” a first indicate number “to”, “up to”, “until” or “through” (or another such range-indicating term) a second indicate number are used herein interchangeably and are meant to include the first and second indicated numbers and all the fractional and integral numbers therebetween.

Although the invention has been described in conjunction with specific embodiments thereof, it is evident that many alternatives, modifications and variations will be apparent to those skilled in the art. Accordingly, it is intended to embrace all such alternatives, modifications and variations that fall within the spirit and broad scope of the appended claims.

All publications, patents and patent applications mentioned in this specification are herein incorporated in their entirety by reference into the specification, to the same extent as if each individual publication, patent or patent application was specifically and individually indicated to be incorporated herein by reference. In addition, citation or identification of any reference in this application shall not be construed as an admission that such reference is available as prior art to the present invention. To the extent that section headings are used, they should not be construed as necessarily limiting.

It is appreciated that certain features, which are, for clarity, described in the context of separate embodiments within the present disclosure, may also be provided in combination in a single embodiment. Conversely, various features, which are, for brevity, described in the context of a single embodiment, may also be provided separately or in any suitable subcombination or as suitable in any other described embodiment of the present disclosure. Certain features described in the context of various embodiments are not to be considered essential features of those embodiments, unless the embodiment is inoperative without those elements.

WHAT IS CLAIMED IS:

1. A stepped dilator for use with a trocar kit to provide intraperitoneal access via a body recess, comprising:

a dilator body having a longitudinal axis in proximal to distal direction;

a first tapered region of the dilator body tapering narrower toward a distal end of the dilator body;

a second tapered region of the dilator body tapering narrower in a distal direction, located proximally to the first tapered region, and separated from the first tapered region by an isolating region.

2. The stepped dilator of claim 1, wherein the isolating region comprises a region of constant cross-section perpendicular to the longitudinal axis extending between the first tapered region and the second tapered region.

3. The stepped dilator of claim 1, wherein the isolating region comprises a region with no cross-section perpendicular to the longitudinal axis larger in any direction than the largest cross-section perpendicular to the longitudinal axis of the first tapered region.

4. The stepped dilator of claim 1, wherein the isolating region is at least 3 mm long.

5. The stepped dilator of claim 4, wherein the isolating region is no more than 20 mm long.

6. The stepped dilator of claim 5, wherein the isolating region is between 5-15 mm long.

7. The stepped dilator of claim 1, wherein a whole tapering extent of at least one of the first tapered region and the second tapered region tapers between its smallest cross-sectional area and its largest cross-sectional area perpendicular to the longitudinal axis over a longitudinal distance of 15 mm or less.

8. The stepped dilator of claim 7, wherein, through the tapering extent of at least one of the first tapered region and the second tapered region, a diameter of the stepped dilator perpendicular to the longitudinal axis increases by at least 7.5 mm.

9. The stepped dilator of claim 1, wherein a cross-section with the largest cross-sectional area of the first tapered region perpendicular to the longitudinal axis has at least one axis of about 10 mm or longer.

10. The stepped dilator of claim 1, wherein a distal tip of the first tapered region comprises a front surface aperture of 4 mm^2 or less, and expands in a distal direction from the front surface aperture through a radius of curvature of at least 2.5 mm.

11. The stepped dilator of claim 10, wherein the front surface aperture is an aperture of an inner lumen sized to allow partial advancement of a trocar needle having a diameter less than or equal to about 2 mm in a longitudinal direction through the aperture.

12. The stepped dilator of claim 1, wherein the cross-section of the first tapered region having the largest cross-sectional area perpendicular to the longitudinal axis has at least one axis of about 7.5 mm or less.

13. A kit comprising the stepped dilator of claim 1, along with a handle and a trocar needle; wherein the stepped dilator and handle together define a lumen sized to accept passage the trocar needle from a proximal end of the handle to a distal tip of the stepped dilator.

14. The kit of claim 13, wherein the trocar needle is provided with a handle extending at least 5 cm past a proximal end of the handle when a distal tip of the trocar needle is advanced 5 mm past the distal tip of the stepped dilator.

15. The kit of claim 13, wherein the trocar needle is provided with a dull-tipped, inner, spring-loaded stylet to act as a Veress needle having an extended position and a collapsed position; wherein the stylet comprises a blunt end extending past a sharp tip of the trocar needle in the extended position and preventing the sharp tip from injuring tissue; and wherein the stylet moves to the collapsed position upon sufficient longitudinal force being exerted so that it no longer extends past the sharp tip, allowing the sharp tip can operate to penetrate tissue.

16. The kit of claim 13, provided with a stopper device configured to resist advancing the distal tip of the trocar needle more than 5 mm beyond the distal tip of the stepped dilator.

17. The stepped dilator of claim 1, wherein a cross-section with the largest cross-sectional area of the second tapered region perpendicular to the longitudinal axis has at least one axis of about 21 mm or longer.

18. A cannula to provide intraperitoneal access across a wall of a body recess, wherein a cross-section of an inner lumen of the cannula transverse to a longitudinal axis of the cannula has a long and a short axis; and wherein the long axis is at least twice as long as the short axis.

19. The cannula of claim 18, wherein the short cross-section axis is between 5 mm and 10 mm.

20. The cannula of claim 18, wherein the long cross-section axis is between 10 mm and 30 mm.

21. The cannula of claim 18, wherein a luminal wall defining the transverse cross-section of the cannula comprises straight sections on opposite sides of the transverse cross-section.

22. The cannula of claim 21, wherein the straight sections are interconnected through curved sections.

23. The cannula of claim 8, wherein the cannula is at least 5 cm long.

24. The cannula of claim 18, wherein an edge defining an aperture at a distal end of the cannula comprises a first edge portion extending along one side of the aperture, and a second edge portion extending along another side of the aperture, wherein the first edge portion extends along a side more distal along the longitudinal axis than the second edge portion.

25. The cannula of claim 24, wherein the first and second edge portions extend along opposite sides of the aperture.

26. The cannula of claim 24, wherein the first edge portion and the second edge portion extend along the long cross-section axis.

27. The cannula of any one of claims 18-26, comprising a handle extending at least 10 cm from a proximal end of the cannula.

28. A cannula for use with a trocar to provide intraperitoneal access via a body recess, wherein an edge defining an aperture at a distal end of the cannula comprises a first edge portion extending along one side of the aperture, and a second edge portion extending along another side of the aperture, wherein the first edge portion extends along a side more distal along the longitudinal axis than the second edge portion.

29. The cannula of claim 28, wherein the first edge portion is positioned at least 5 mm more distally than the second edge portion.

30. The cannula of any one of claims 28-29, wherein the first edge portion and the second edge portion each comprise respective a straight portion.

31. A method of using a cannula to provide intraperitoneal access to a body cavity via a body recess, comprising:

inserting a distal end of the cannula into a dilated aperture of a rectouterine pouch via transvaginal access;

wherein an edge defining an aperture at the distal end of the cannula comprises a first edge portion extending along one side of the aperture, and a second edge portion extending along an opposite side of the aperture, and the first edge portion is positioned more distally along the cannula and from the dilated aperture than the second edge portion; and

wherein the cannula is inserted so that the aperture at the distal end of the cannula is oriented to open toward the side of the second edge portion and facing toward a rectum adjacent to the rectouterine pouch.

32. The method of claim 31, comprising inserting a flexible robotic arm through the cannula and into the rectouterine pouch, so that it exits the aperture in a direction oriented away from the rectum.

33. The method of claim 31, wherein:
- the inserting a distal end of the cannula into the rectouterine pouch comprises sliding the cannula over an outer dilator;
 - the outer dilator has a tapered distal insertion end sized and shaped to fittingly slide over an inner dilator having a tapered distal insertion end with a rounded tip;
 - the cannula slides fittingly over the outer dilator; and
 - at least the outer dilator is inserted into the rectouterine pouch via transvaginal access.
34. The method of claim 33, comprising sliding the cannula over the outer dilator while the inner dilator remains within the outer dilator.
35. The method of claim 31, wherein:
- the inserting a distal end of the cannula into the rectouterine pouch comprises sliding the cannula over a stepped dilator;
 - the stepped dilator has a tapered distal insertion end with a rounded tip, a second tapered region, and an isolating region between the second tapered region and the tapered distal insertion end;
 - the cannula slides fittingly over the stepped dilator; and
 - the dilator is inserted into the rectouterine pouch via transvaginal access.
36. The method of claim 35, wherein the rounded tip has a hole sized to pass a trocar needle having a diameter less than or equal to about 2 mm.
37. A kit for providing intraperitoneal access via a body recess, comprising:
- a cannula, wherein a transverse cross-section of an inner lumen of the cannula transverse to a longitudinal axis of the cannula has a long axis long enough to allow simultaneous insertion of at least two cylindrical members, each at least 8 mm in diameter;
 - a stepped dilator having a dilator body with: a first tapered region of the dilator body tapering narrower toward a distal end of the dilator body, and a second tapered region of the dilator body tapering narrower in a distal direction, located proximally to the first tapered region, and separated from the first tapered region by an isolating region; and

a trocar needle provided with a handle region extending past a proximal end of the inner dilator when a distal tip of the trocar needle is advanced to the distal tip of the inner dilator.

38. The kit of claim 37, wherein the long axis of the cannula inner lumen transverse cross-section is at least 21 mm.

39. The kit of claim 37, wherein the cannula has a short cross-sectional axis; and wherein the long cross-sectional axis is at least twice as long as the short cross-sectional axis.

40. The kit of claim 37, comprising an arm sheath for a plurality of robotic arms, sized to fit within the cannula, having a minimum diameter of about 10 mm, and a maximum diameter of at least twice the minimum diameter.

41. A method of gaining intraperitoneal access via a body recess, comprising:

inserting a first stage of a stepped dilator into a rectouterine pouch to widen an aperture in a wall of the rectouterine pouch; and

inserting a second stage of the stepped dilator into the rectouterine pouch to widen the aperture;

wherein the first and second stages of the stepped dilator each comprise a region which tapers narrower in a distal direction, and

wherein the first and second stages of the dilator are separated by an isolating region at least 3 mm long.

42. The method of claim 41, preceded by:

inserting the stepped dilator transvaginally to the wall of the rectouterine pouch; and

advancing a trocar needle from within the stepped dilator to produce the aperture in the wall of the rectouterine pouch.

43. The method of claim 42, continued by:

inserting a distal end of a cannula into the rectouterine pouch by sliding the cannula over the stepped dilator and across the aperture while the aperture is held open by the stepped dilator;

wherein an inner lumen of the cannula has at least one cross-sectional axis of at least 20 mm.

44. The method of claim 43, wherein the inner lumen of the cannula has at least one cross-sectional axis of less than about 12 mm.

45. A method of gaining intraperitoneal access via a body recess, comprising:

inserting a camera to an intraperitoneal space with a wall of a rectouterine pouch in a field of view of the camera;

illuminating the wall of the rectouterine pouch using an intraperitoneally positioned illumination device;

selecting a position for an aperture in the wall of the rectouterine pouch, based on light from the illumination device visible from outside the rectouterine pouch;

advancing a trocar needle from outside the rectouterine pouch to press against the selected position in the wall of the rectouterine pouch;

verifying the position of the trocar needle, based on one or more images from the camera within the intraperitoneal space; and

piercing the rectouterine pouch to form the aperture, using the trocar needle.

46. A kit for setting a position of a robotic arm system along a longitudinal axis of a cannula inserted to a body orifice, wherein the robotic arm system comprises a motor unit and at least one robotic arm extending, when positioned, distally from the motor unit along the longitudinal axis, the kit comprising:

the cannula, including a cannula body configured for insertion to the body orifice;

a mounting block, configured for attachment to the cannula; and

an assembly attached to the mounting block and comprising a spacing arm and an aligning arm, and movable between a stowed position and a deployed position;

wherein the deployed position of the assembly places elements of the aligning arm where they indicate a predetermined position along the longitudinal axis.

47. The kit of claim 46, wherein the mounting block attaches to the cannula by connecting to an access device having a lumen sized to fittingly accept the cannula therewithin.

48. The kit of claim 46, wherein the spacing arm and the aligning arm deploy by hinging around a plurality of stopped hinges, each stopped hinge defining at least a stopped deployed position, and a stopped stowed position.

49. The kit of claim 46, further comprising:

the motor unit and the at least one robotic arm extending distally from the motor unit to a predetermined distance from a stopper-receiving portion of the motor unit;

wherein a distal end of the at least one robotic arm aligns with a distal end of the cannula when the at least one robotic arm is inserted to the cannula, and a stopper portion of the aligning arm contacts a stopper-receiving portion of the motor unit to prevent longitudinal advance of the motor unit.

50. The kit of any one of claims 46-49, comprising an arm sheath with a lumen sized to accept the at least one robotic arm, and an outer surface sized to fit within the cannula.

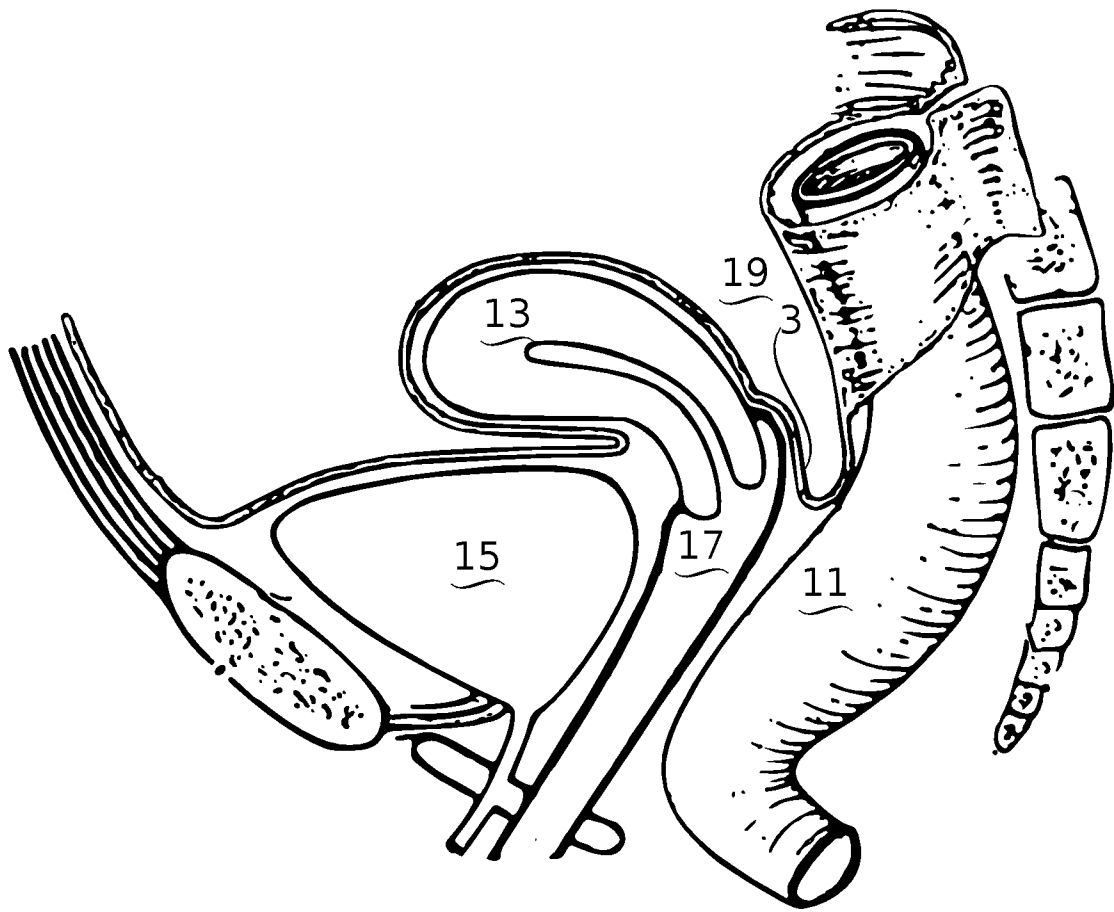


FIG. 1

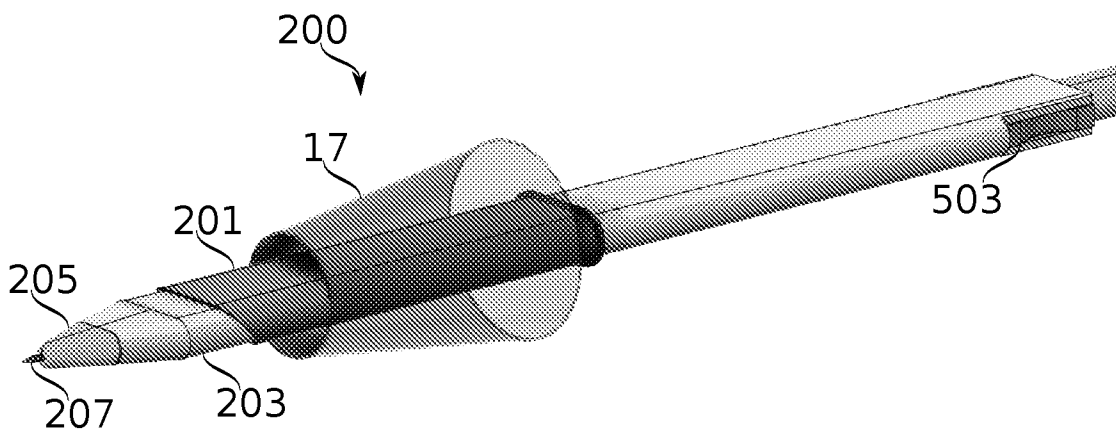


FIG. 2A

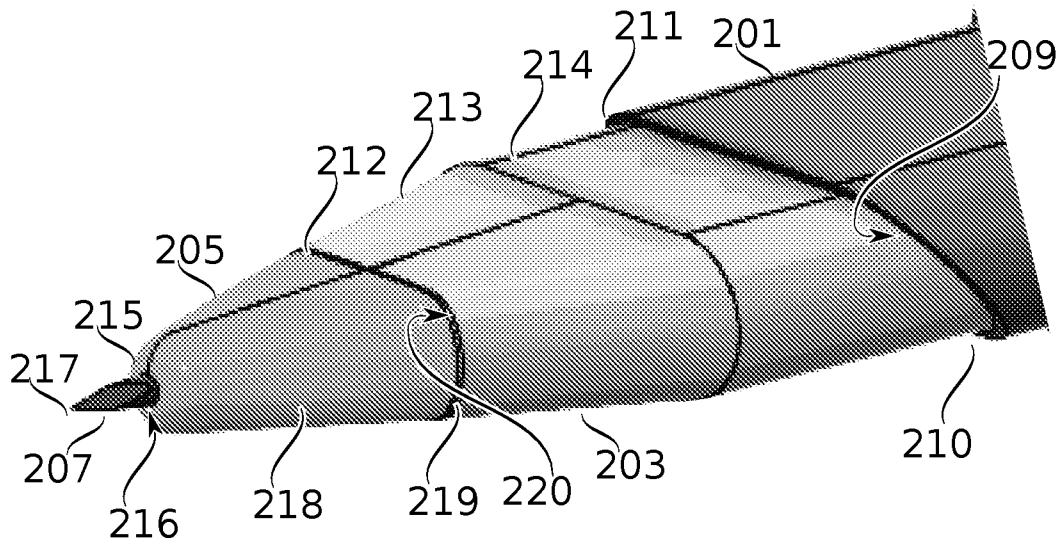
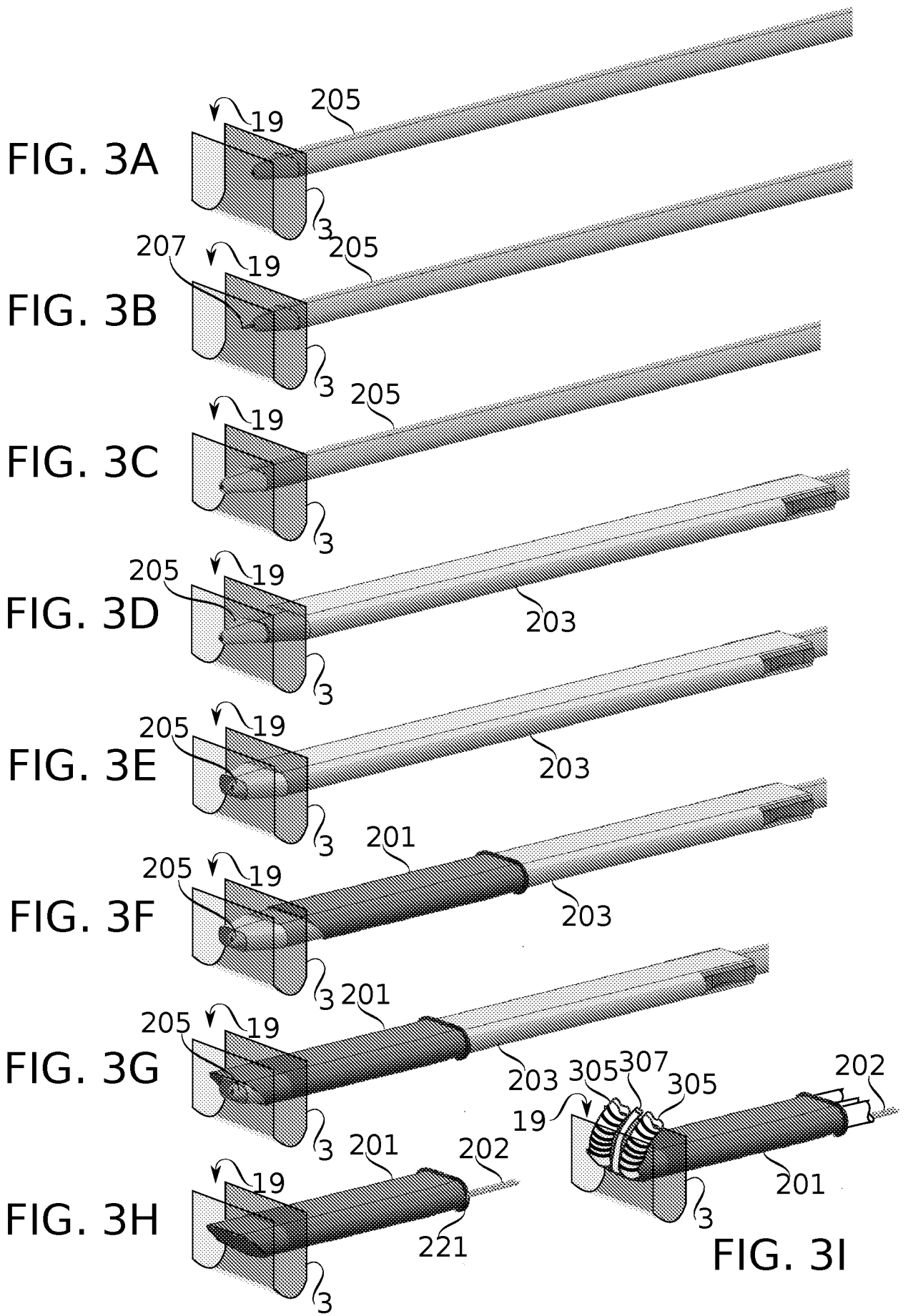
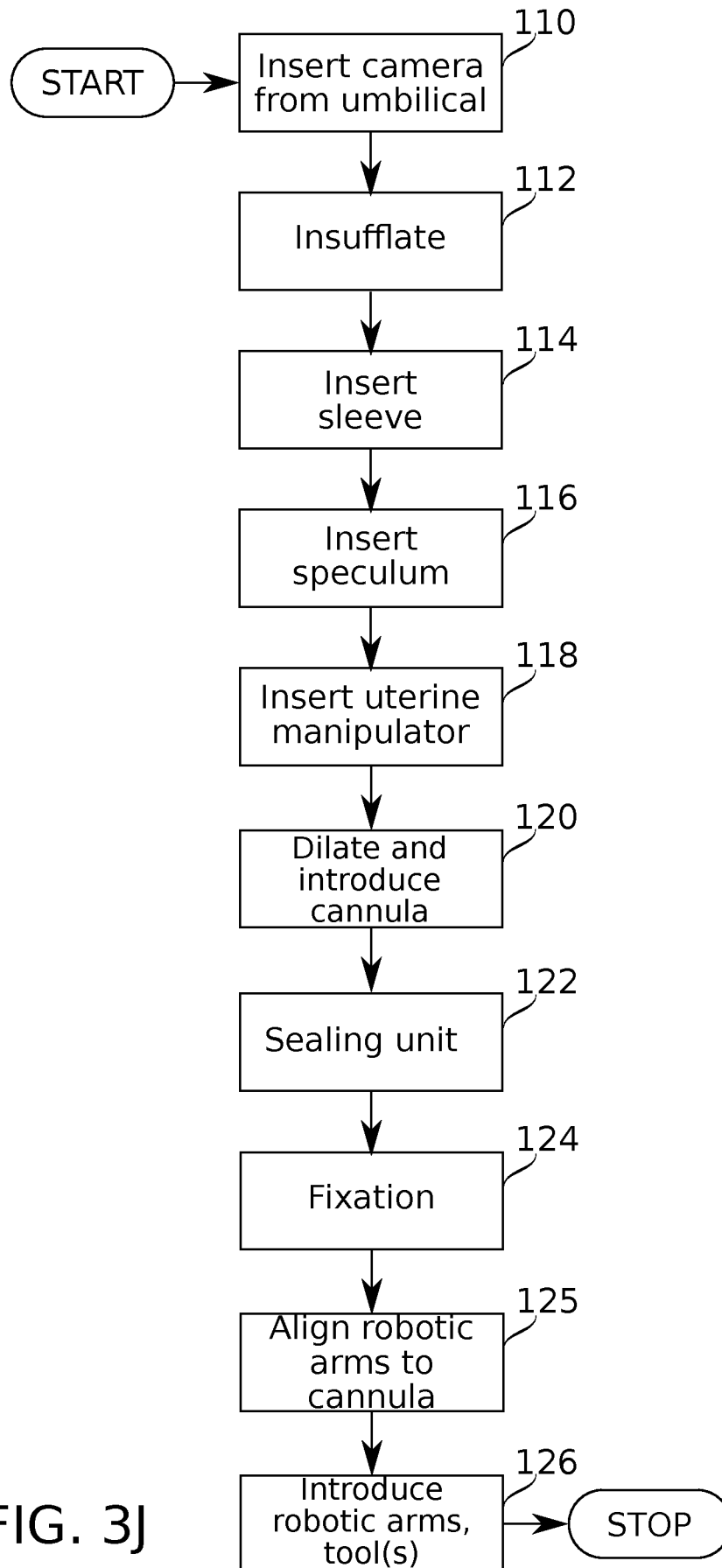


FIG. 2B





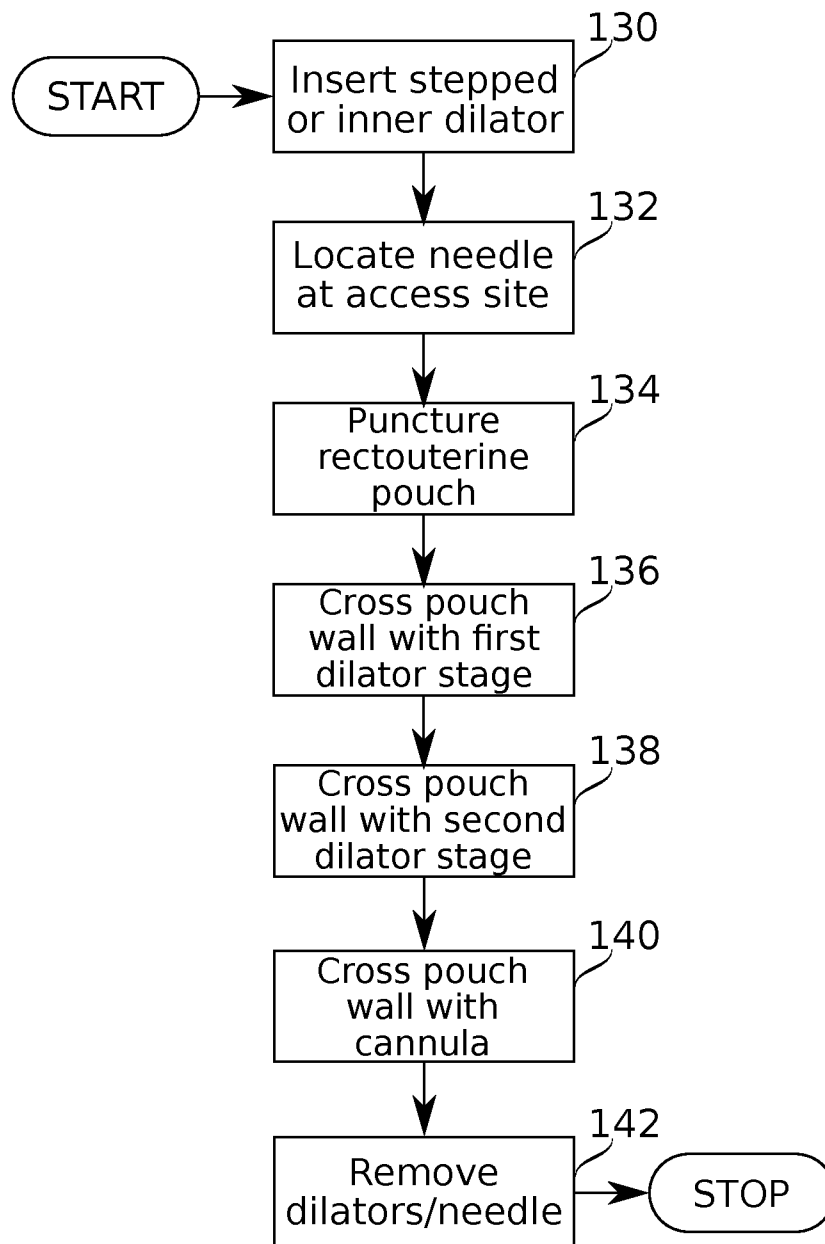


FIG. 3K

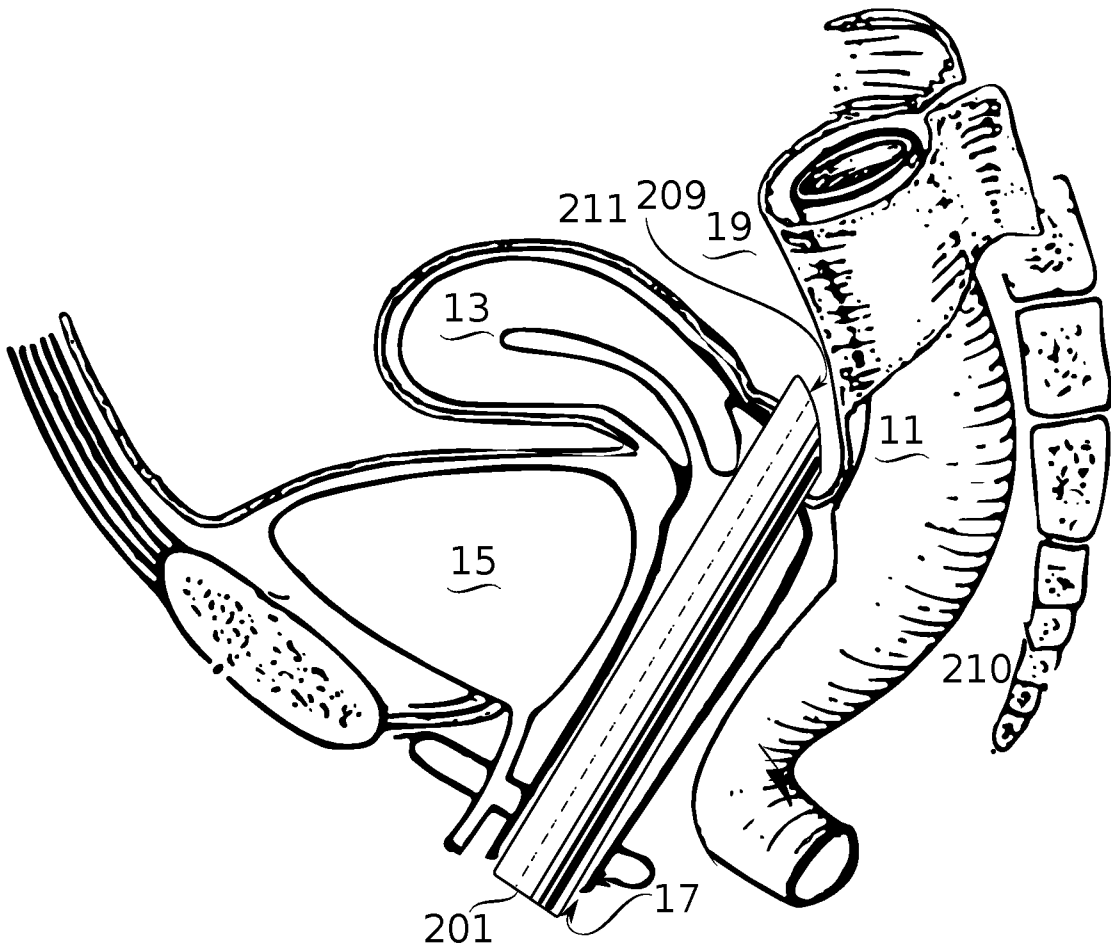
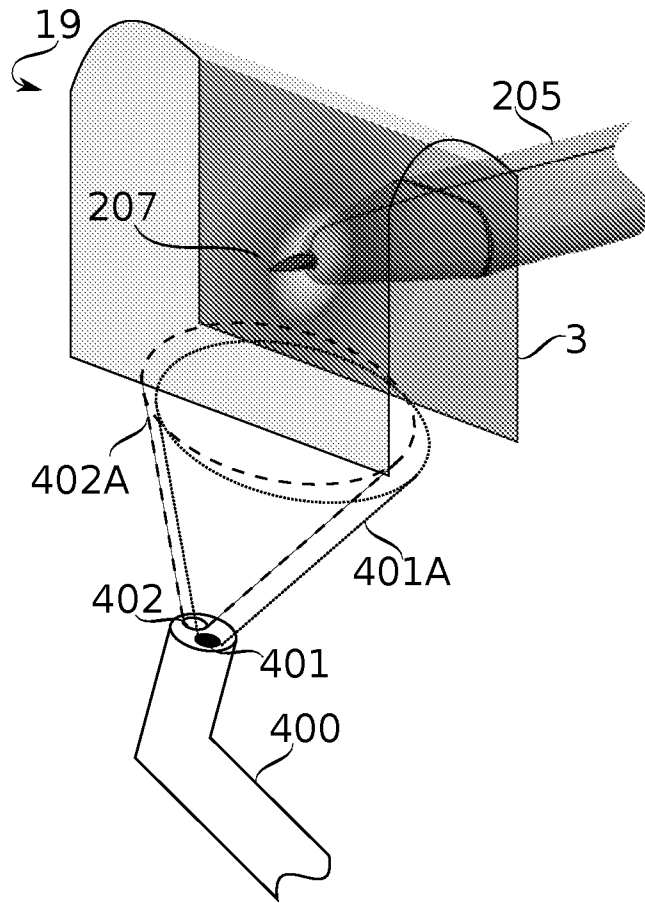
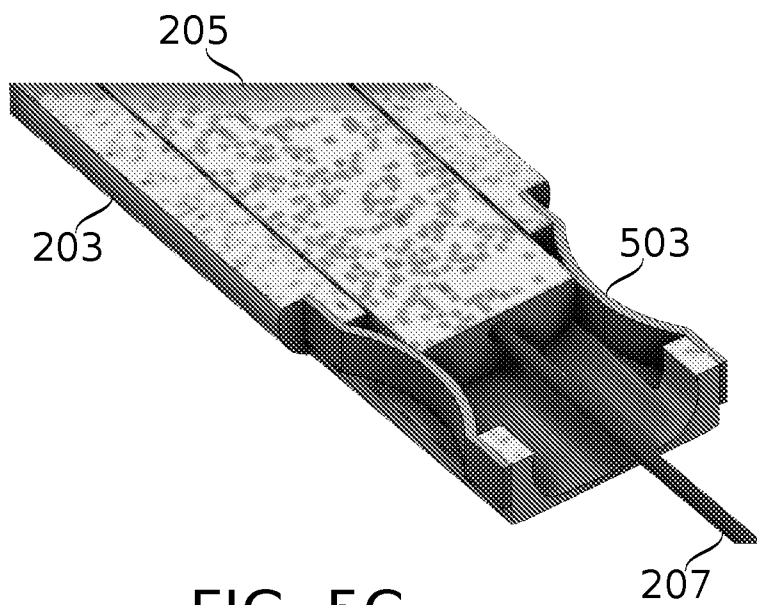
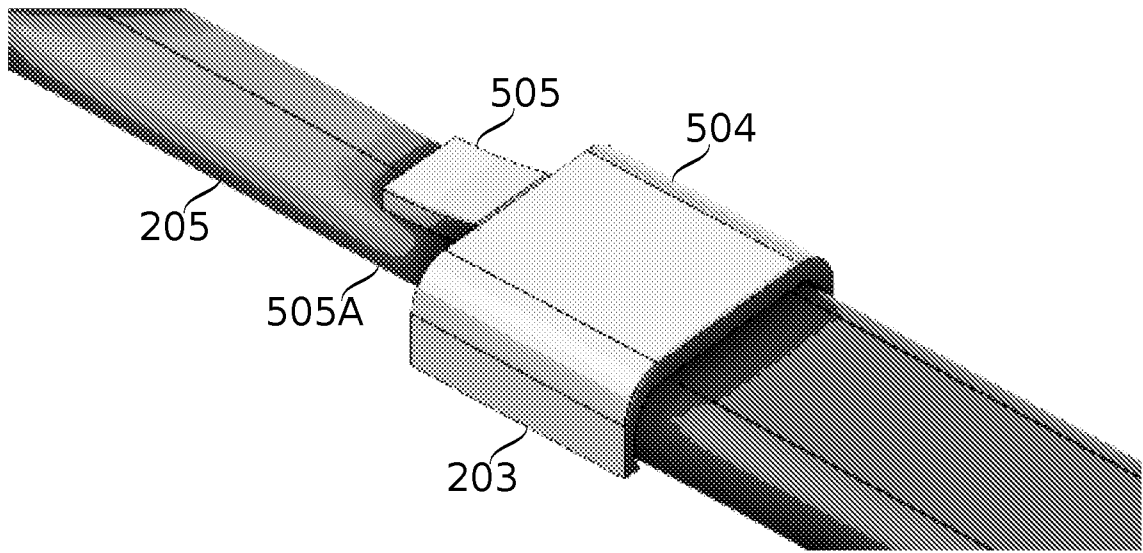
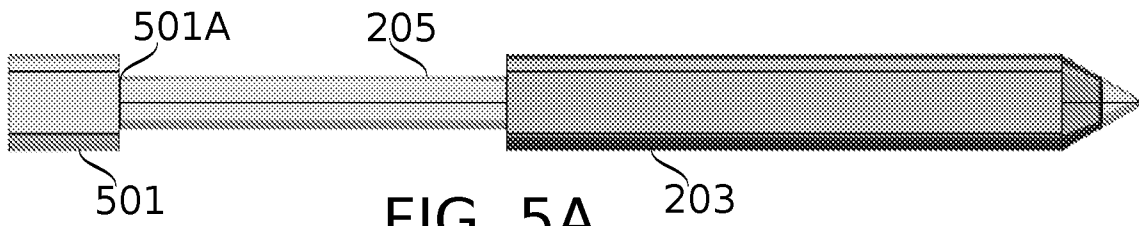
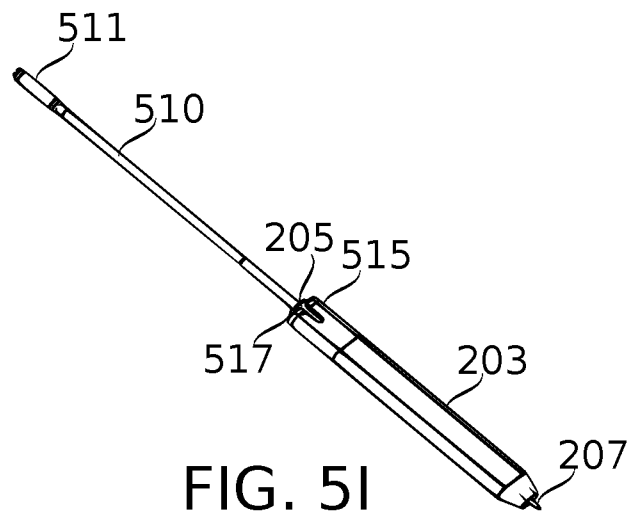
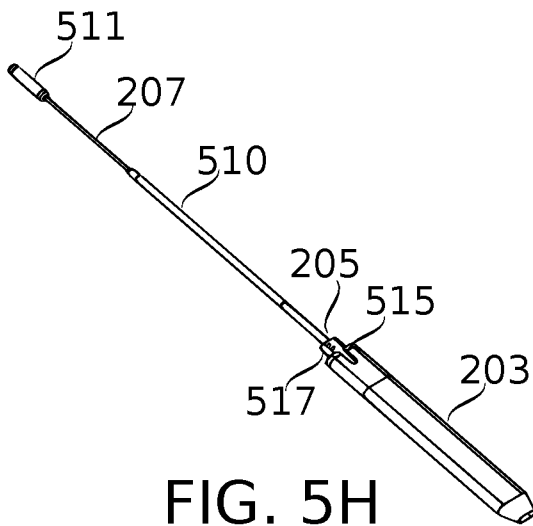
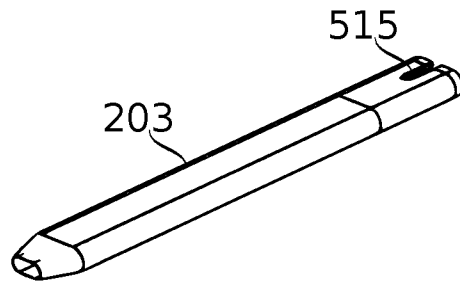
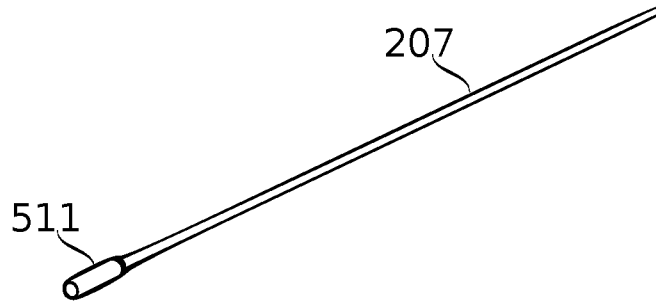
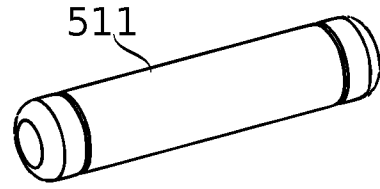
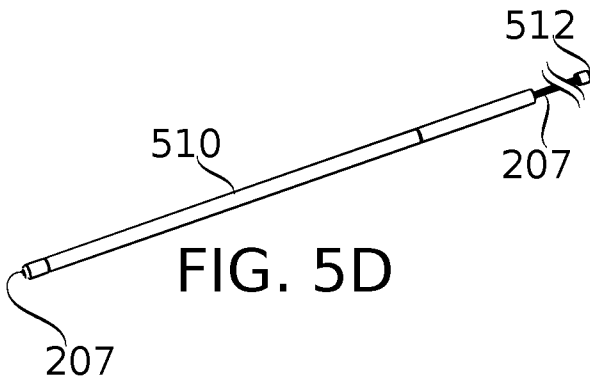


FIG. 3L

FIG. 4







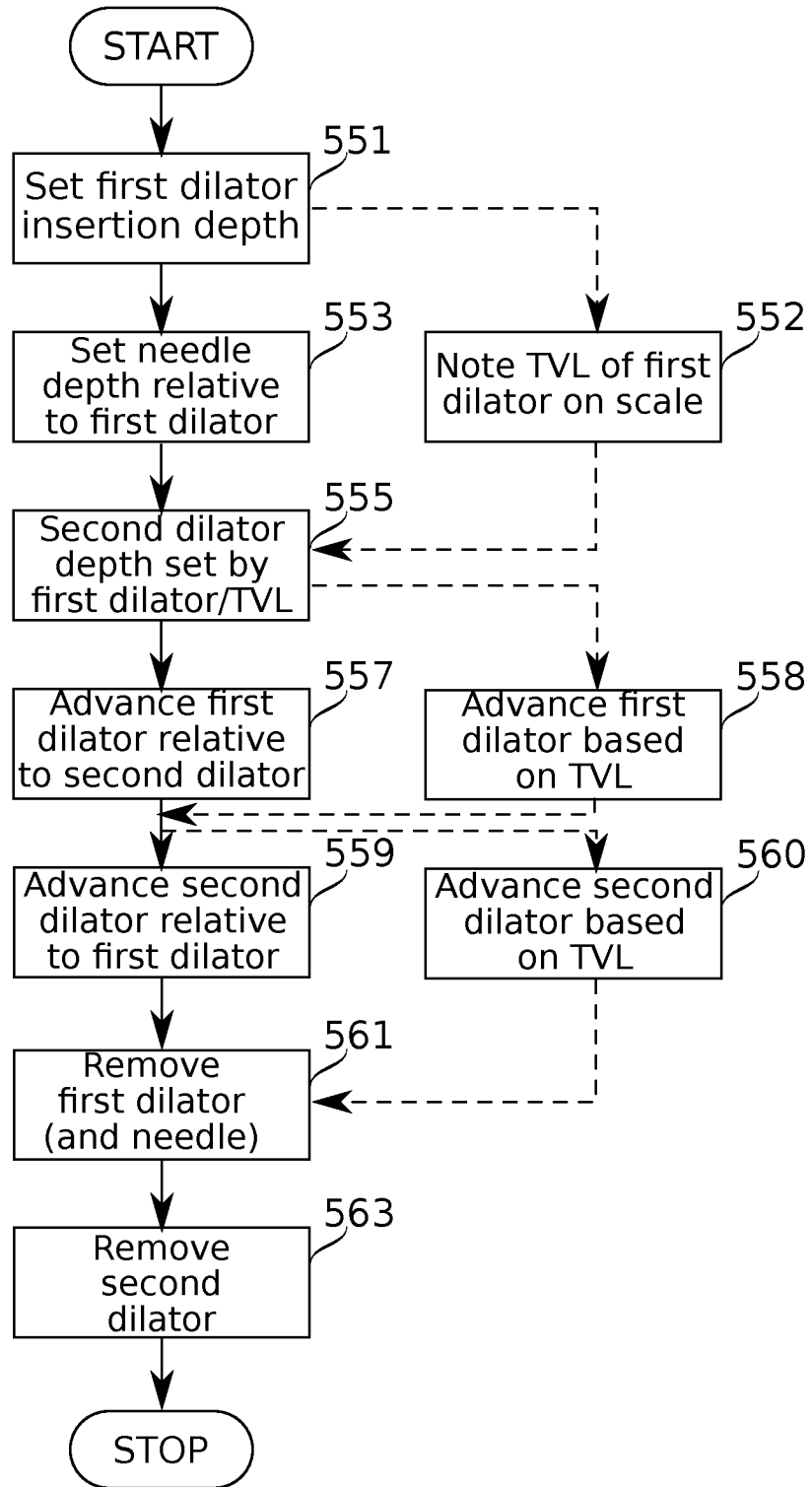


FIG. 5J

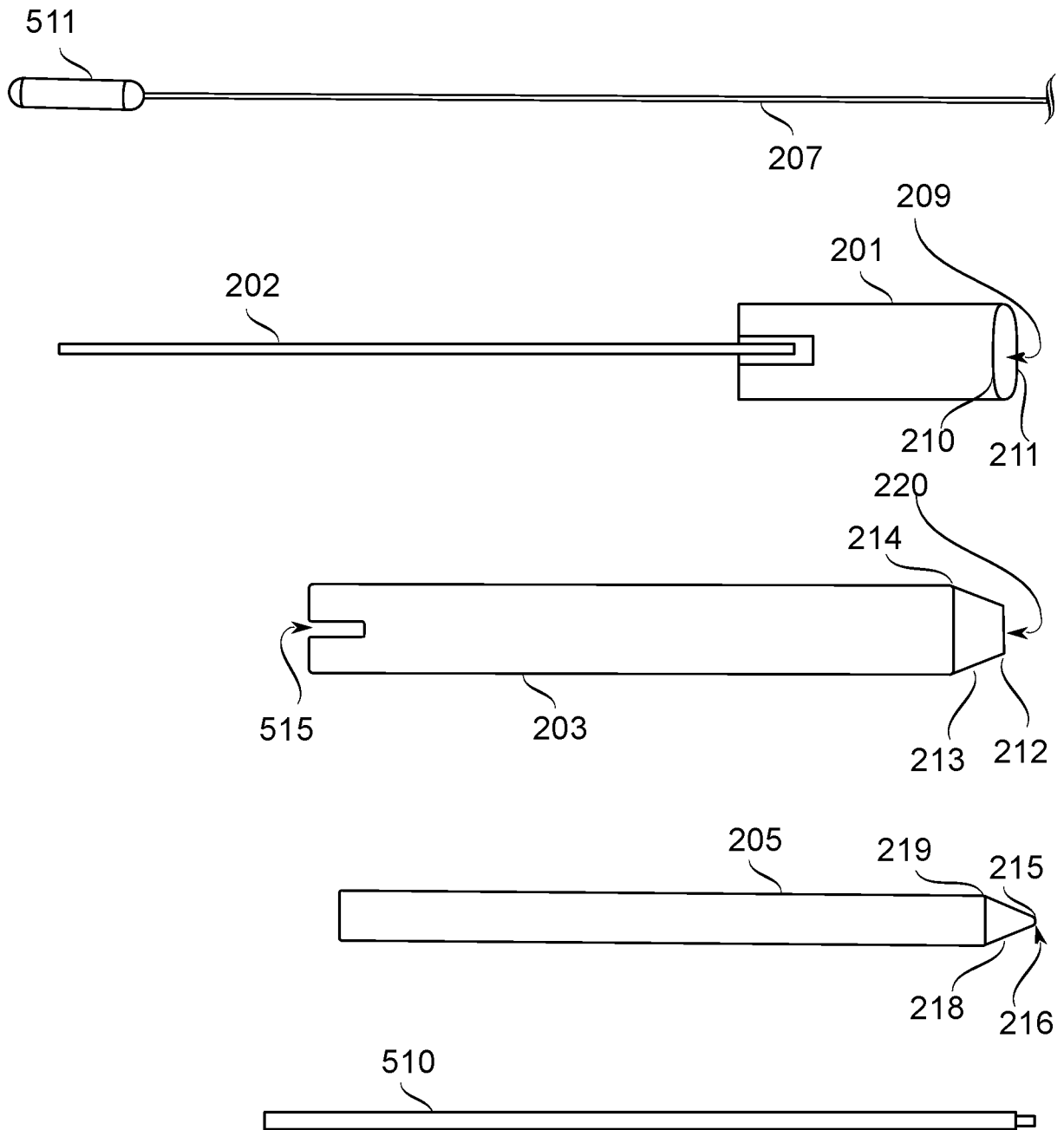
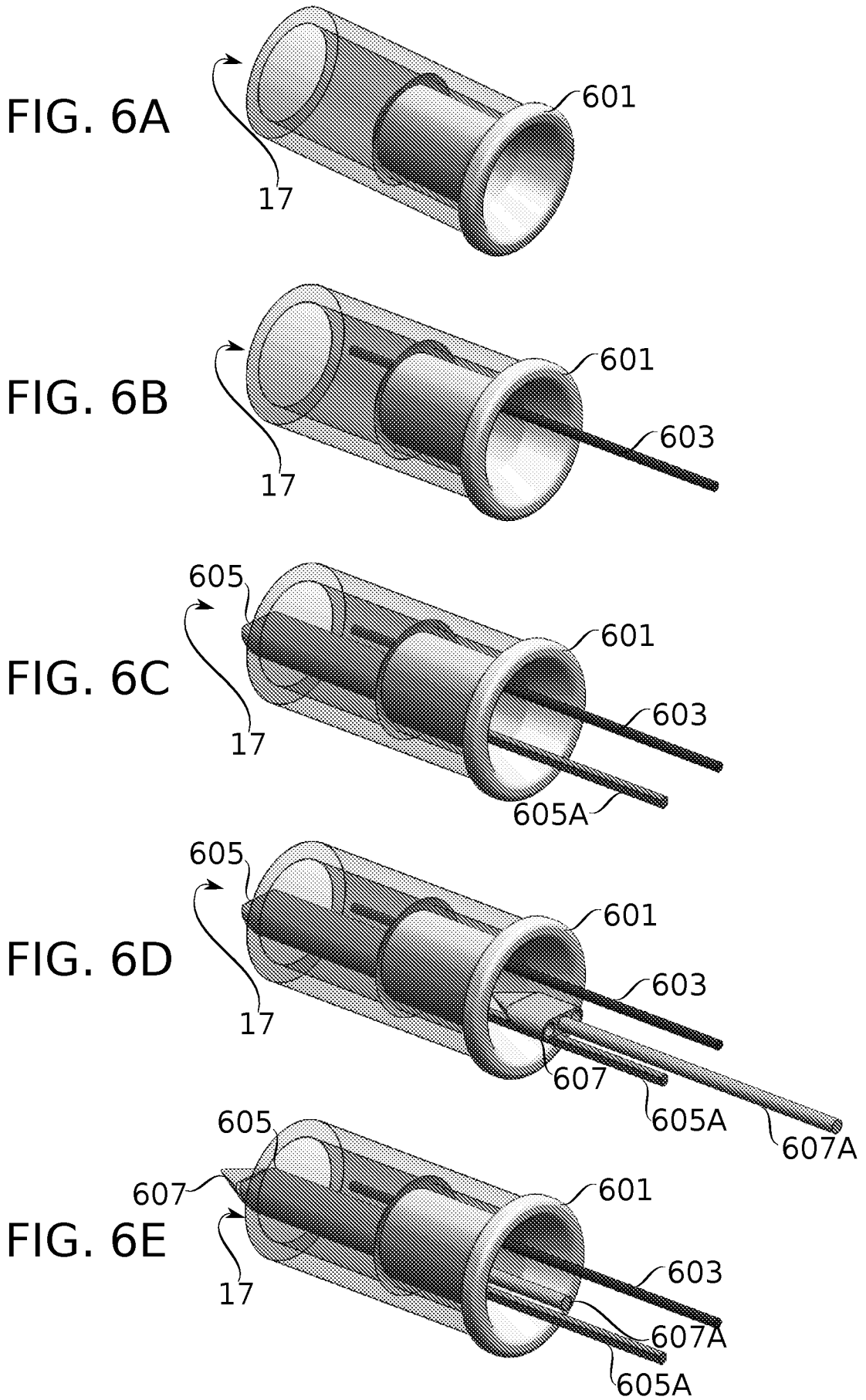


FIG. 5K



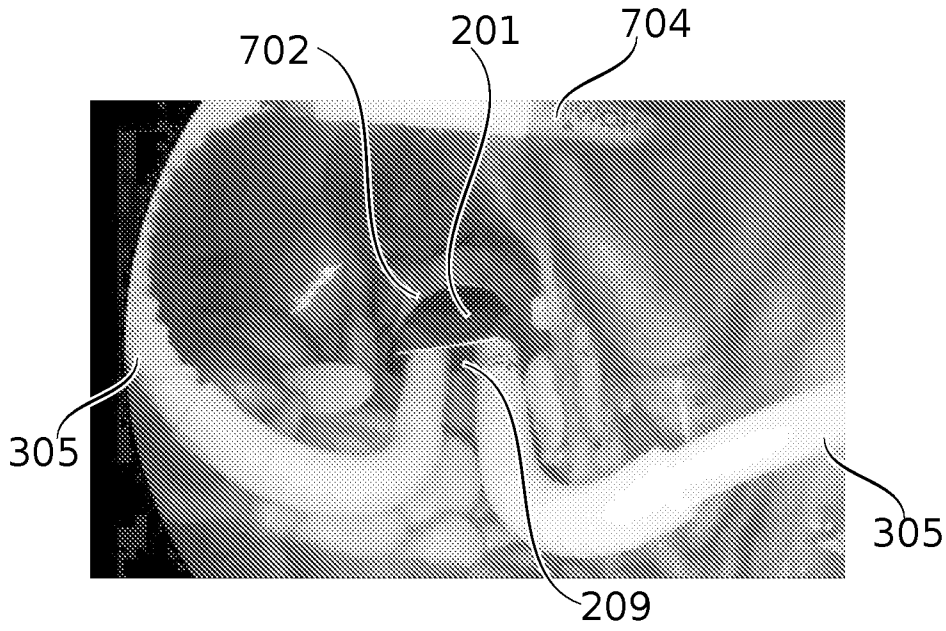


FIG. 7A

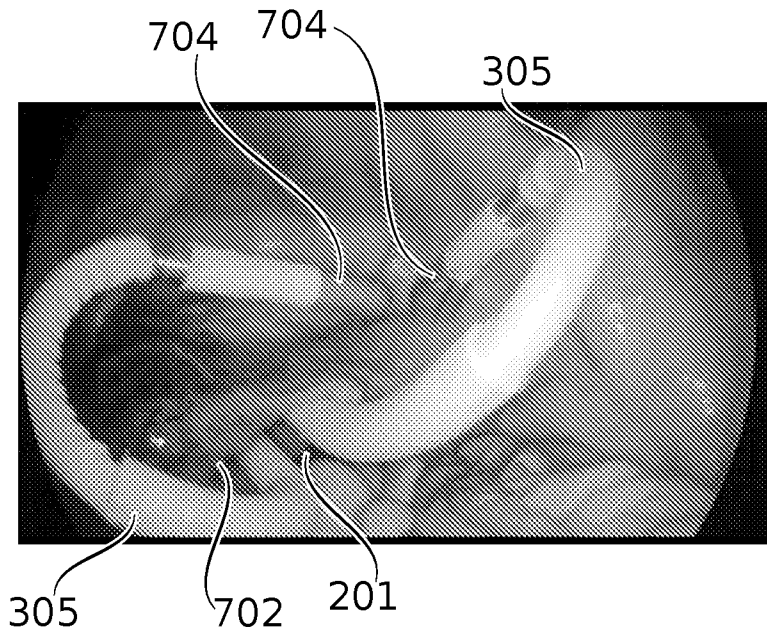


FIG. 7B

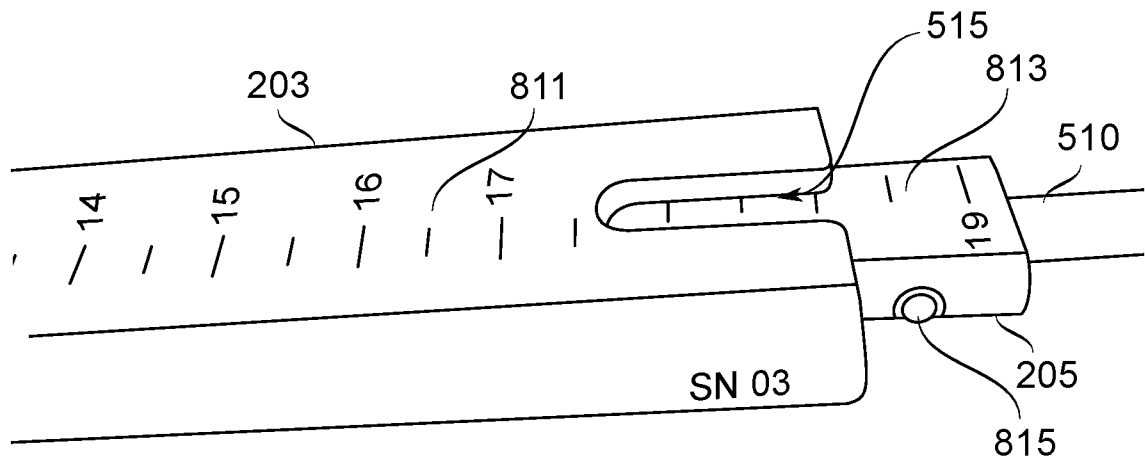


FIG. 8A

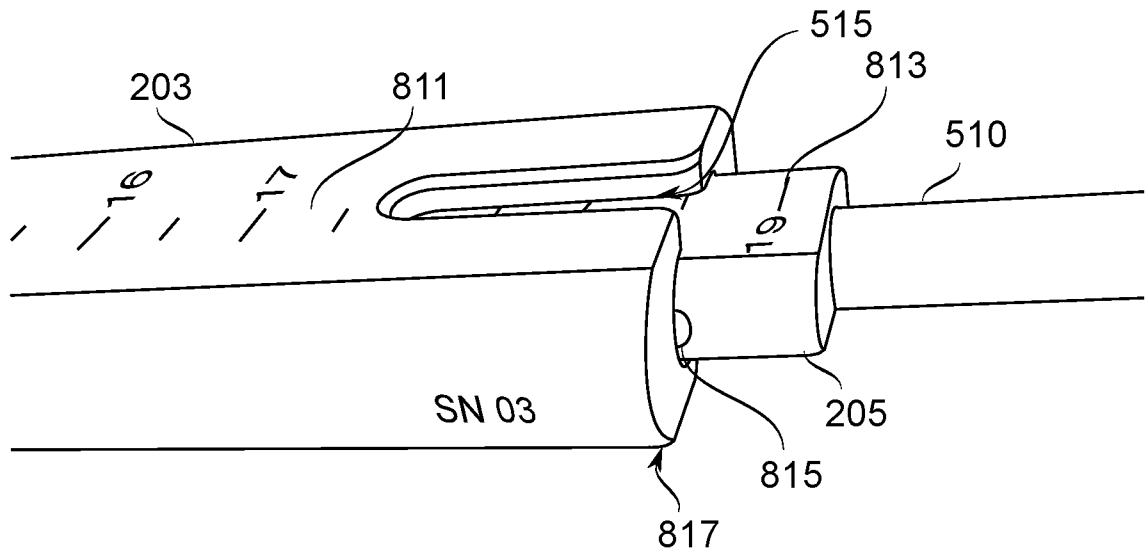
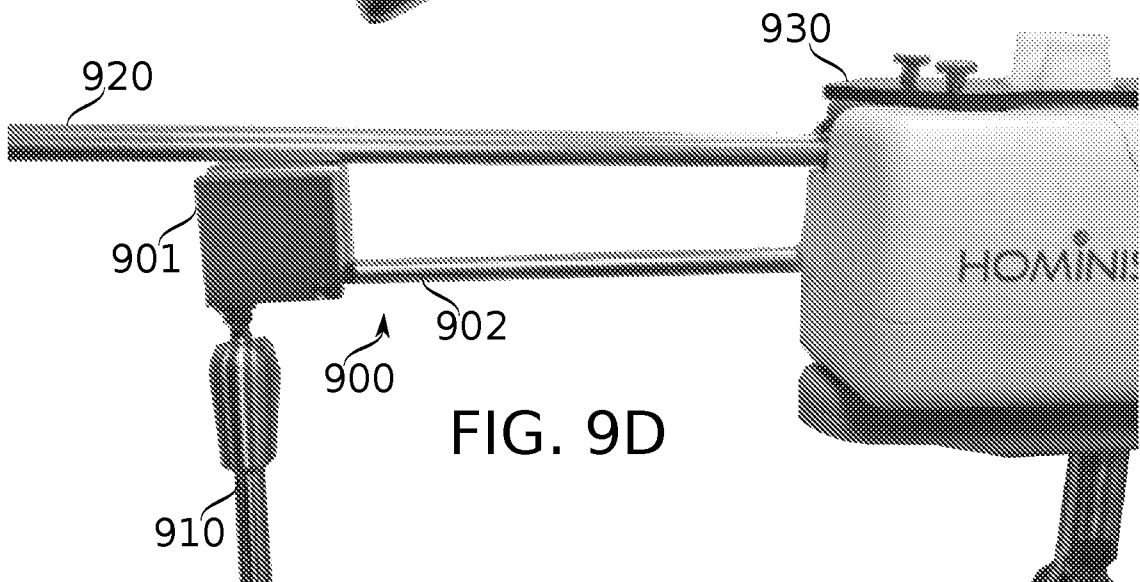
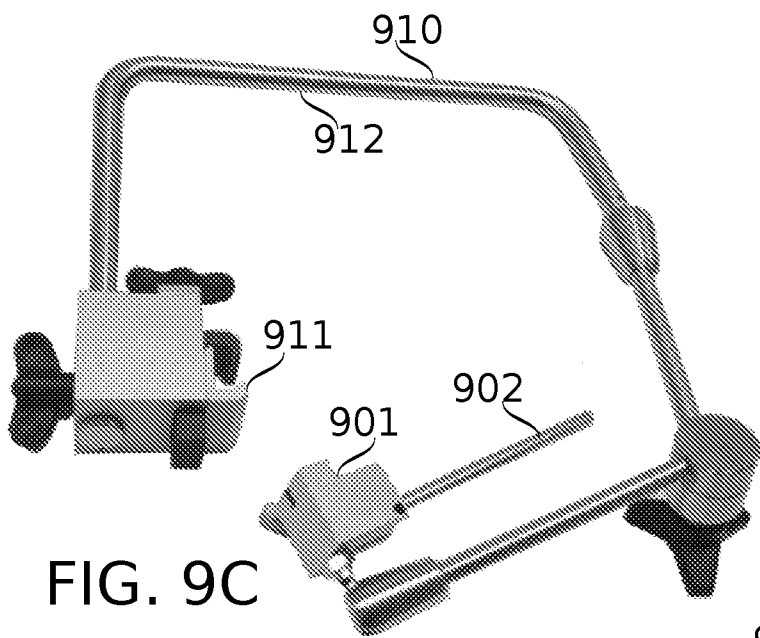
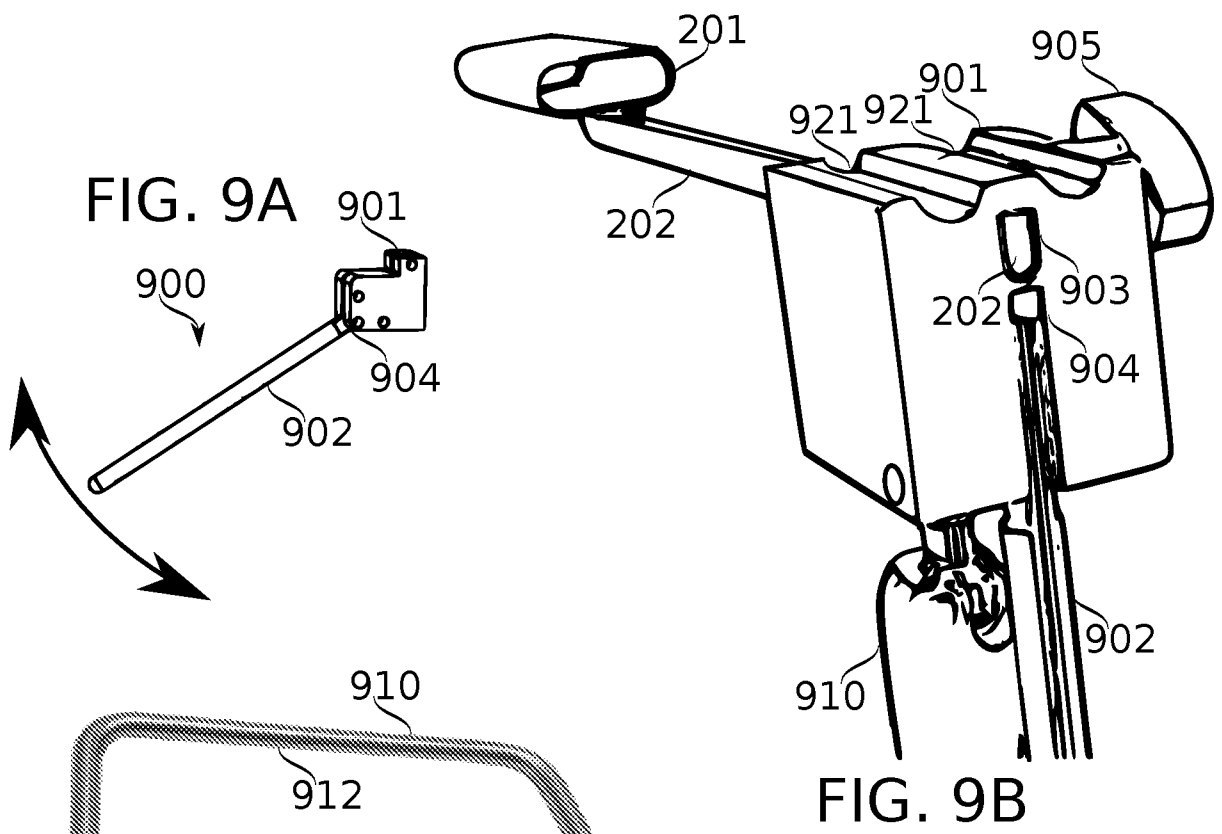
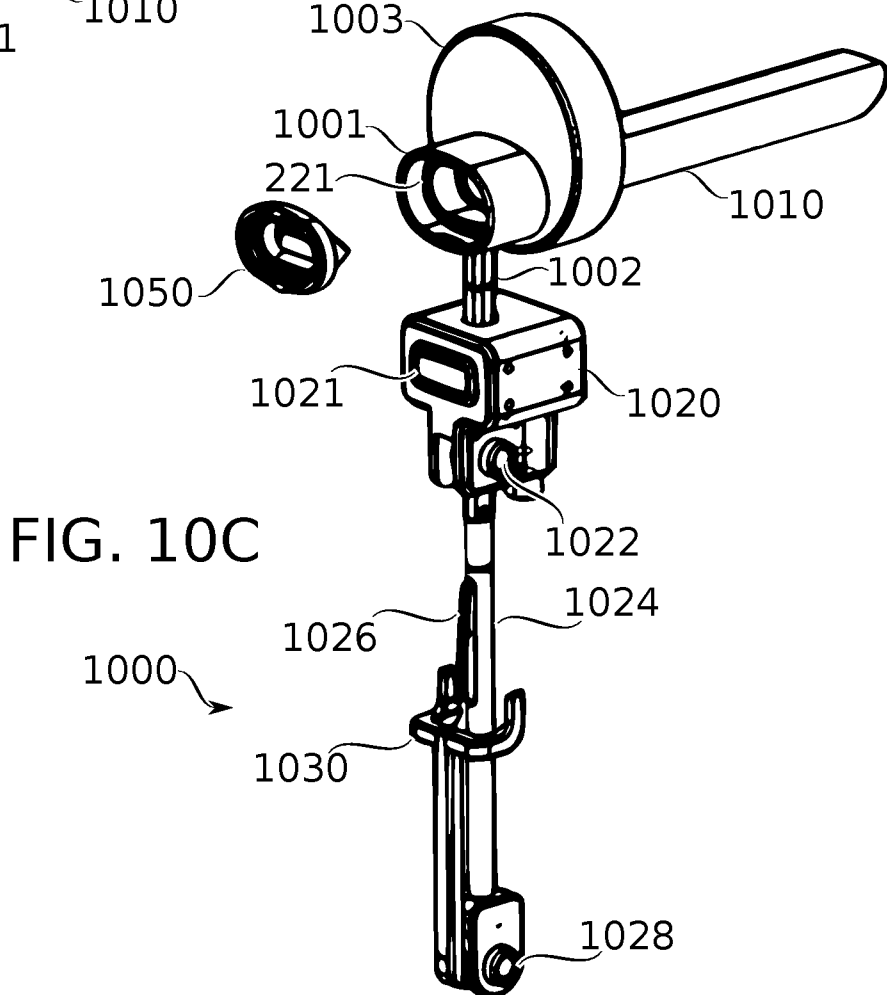
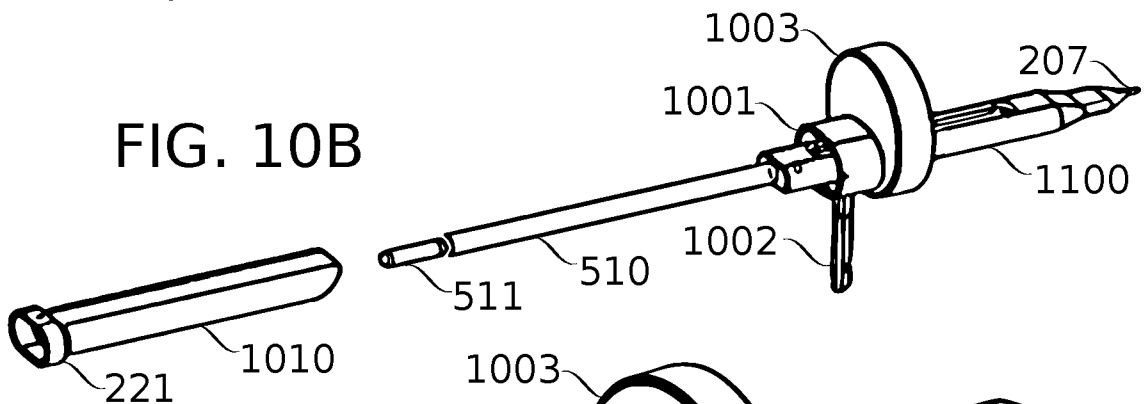
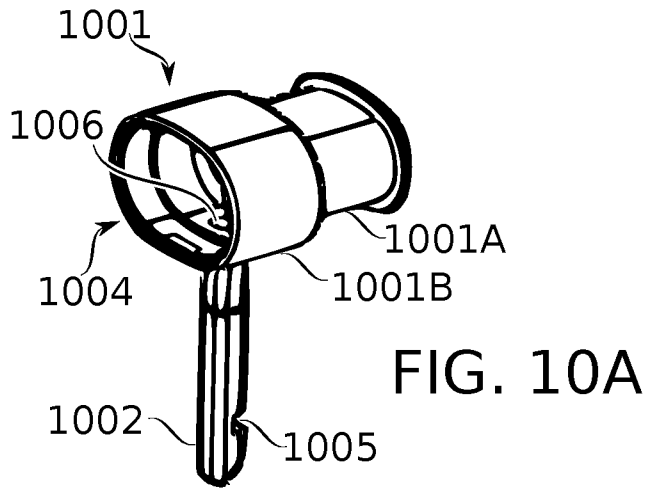
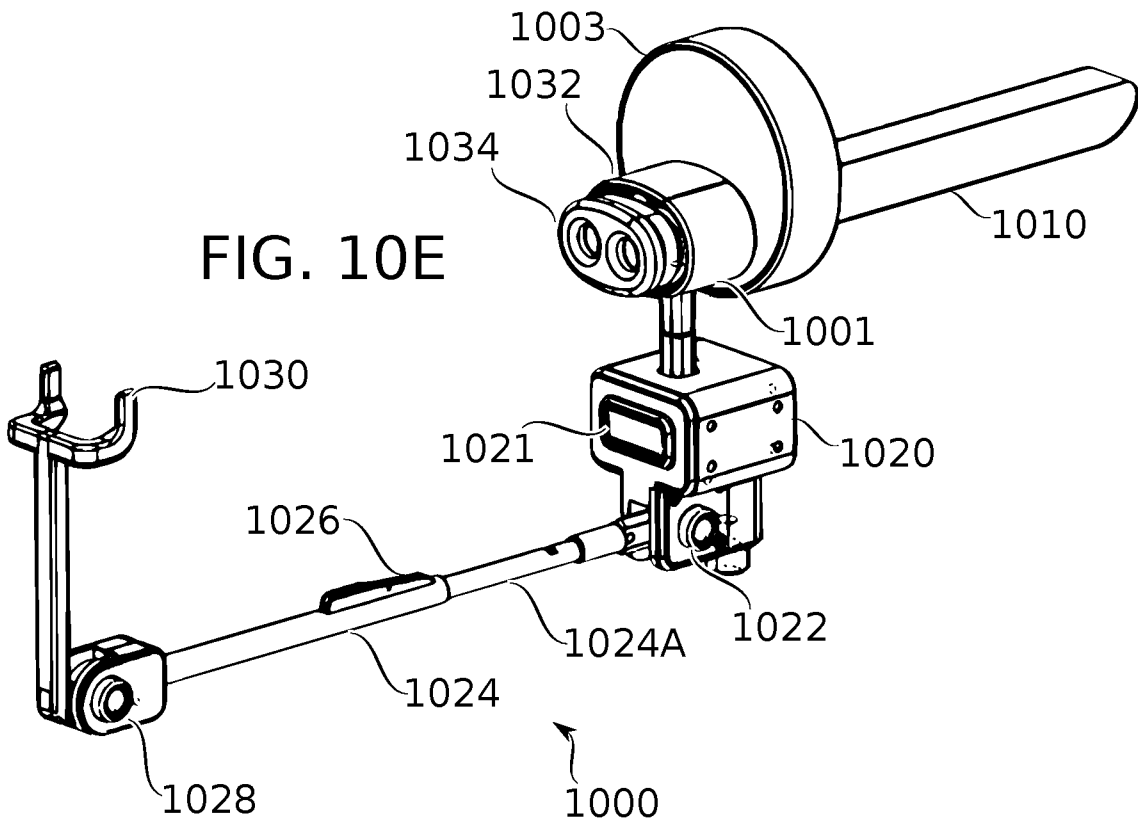
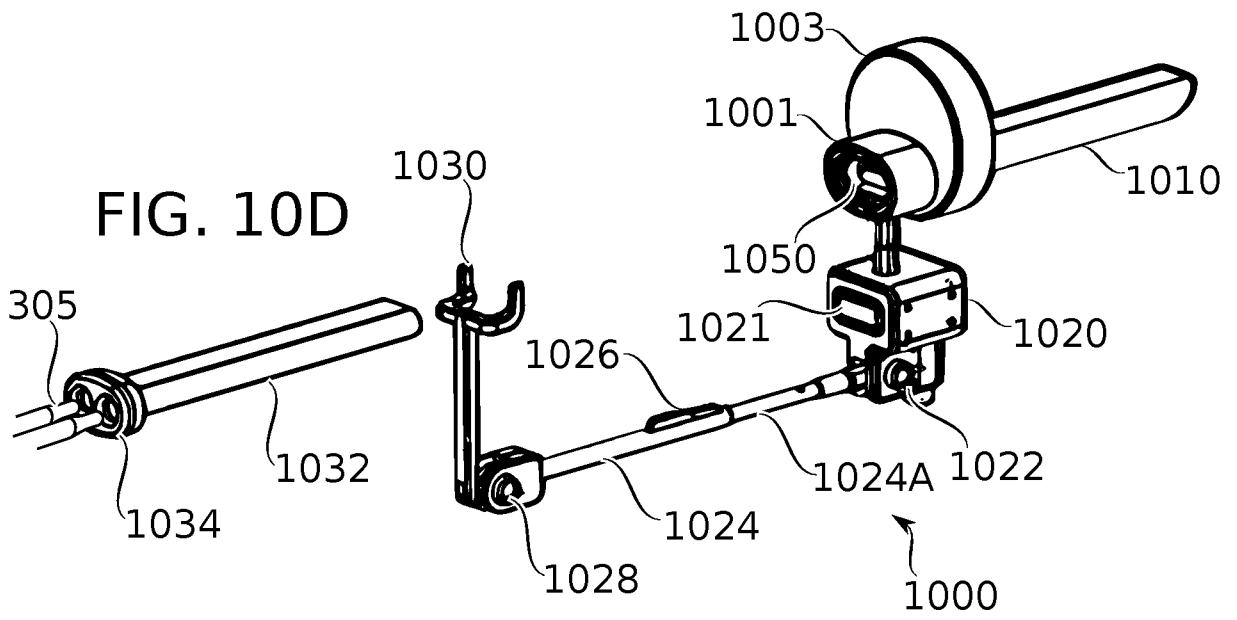
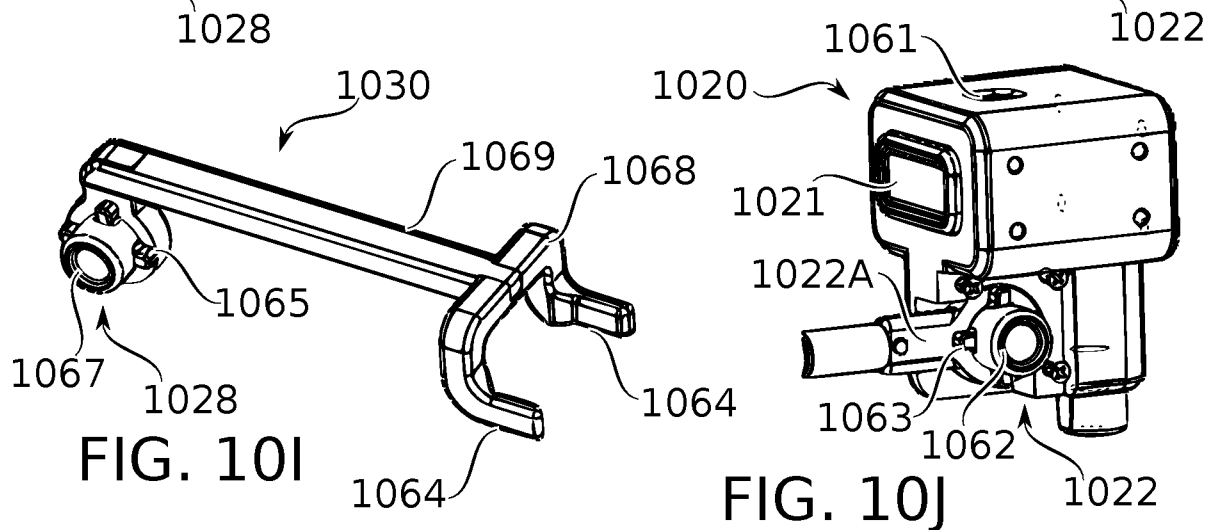
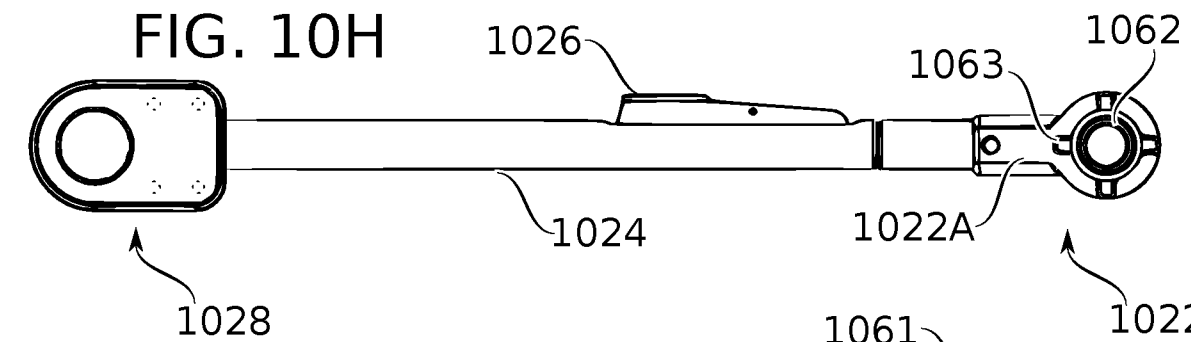
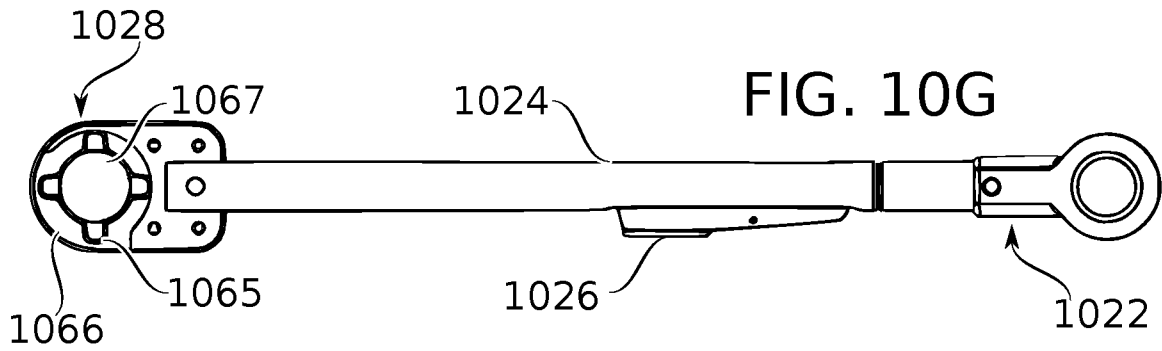
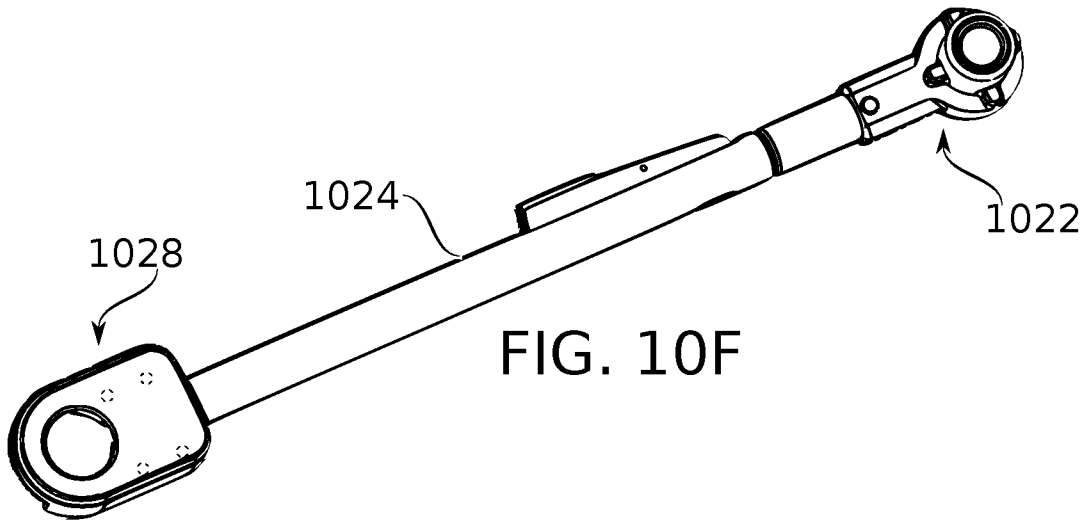


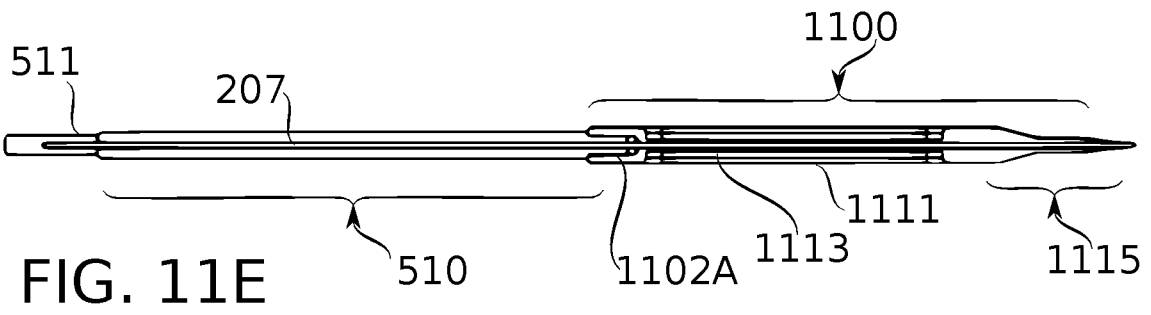
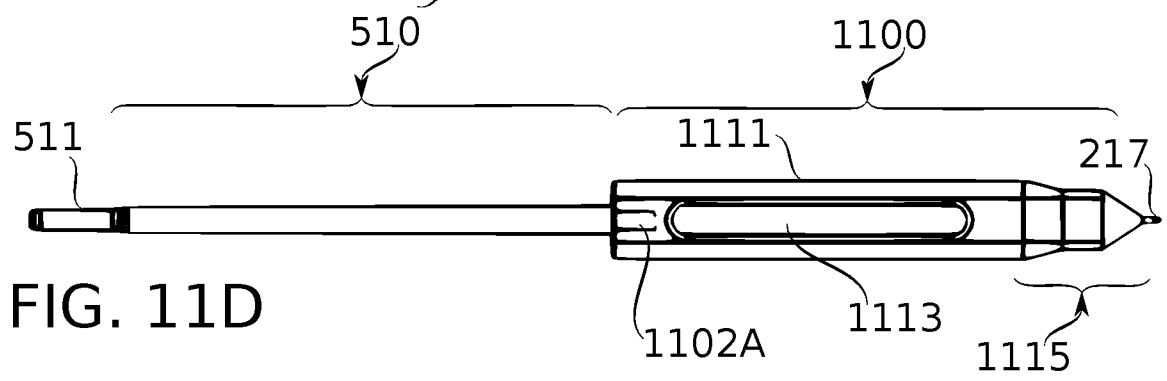
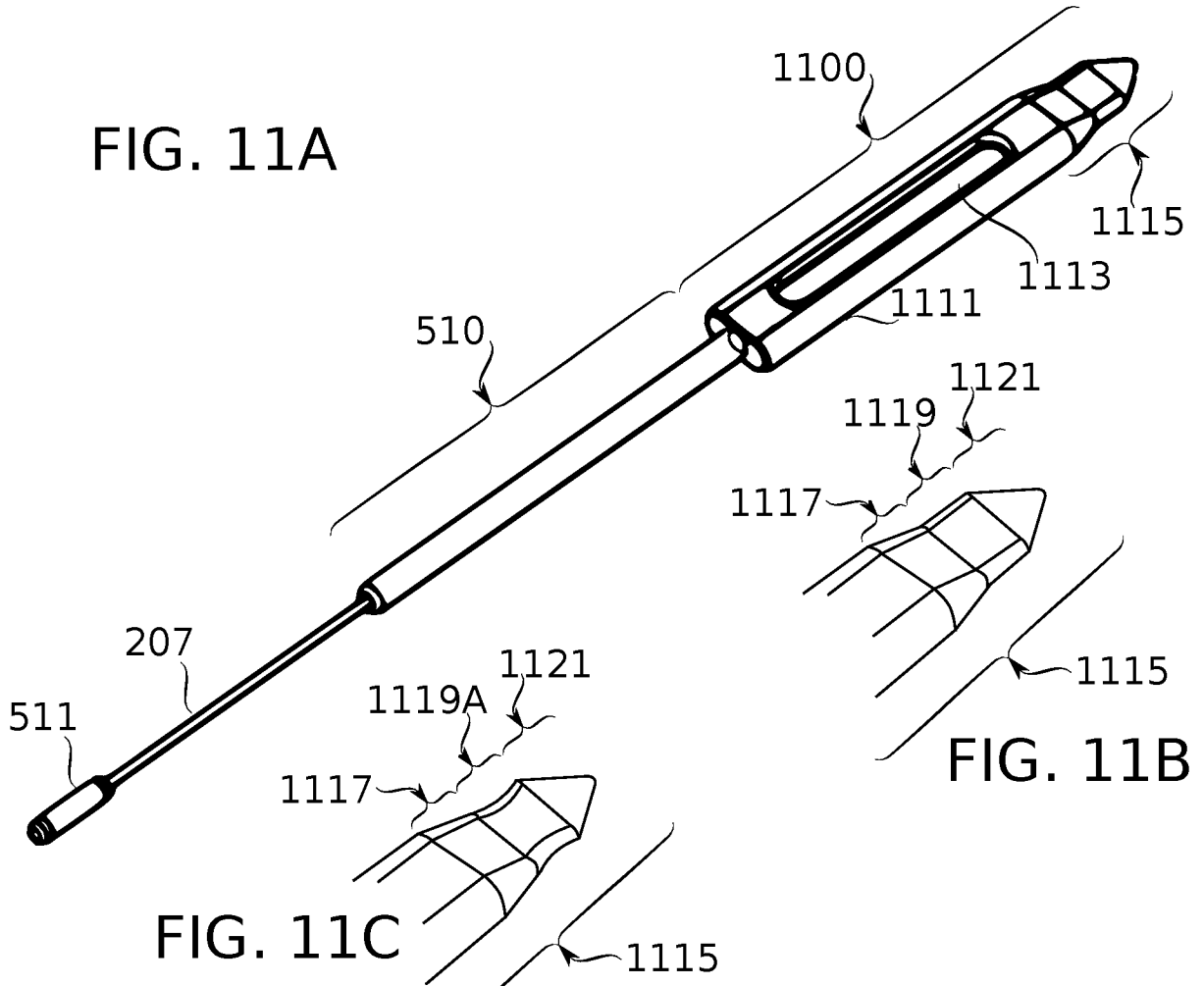
FIG. 8B











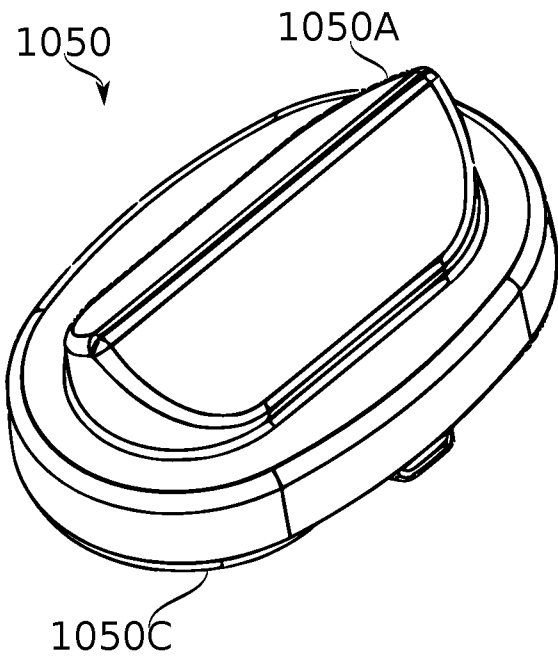


FIG. 12A

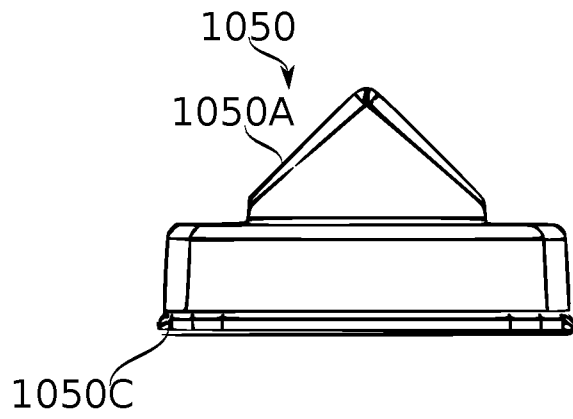


FIG. 12B

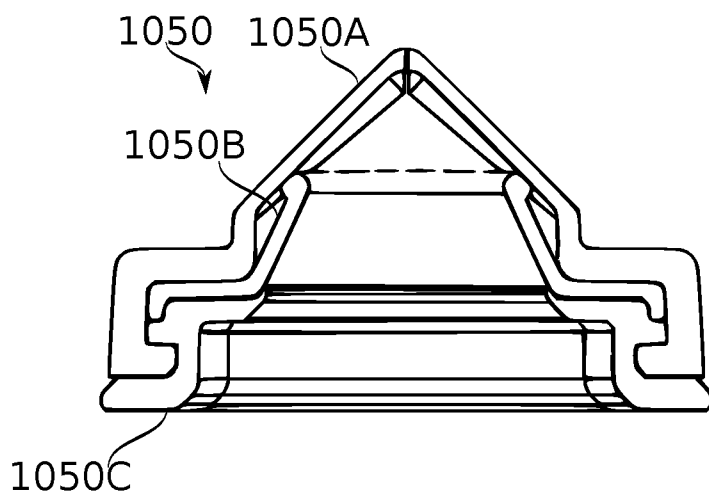


FIG. 12C

INTERNATIONAL SEARCH REPORT

International application No.

PCT/IL 18/50934

A. CLASSIFICATION OF SUBJECT MATTER
 IPC(8) - A61B 1/32, A61B 17/3209, A61B 17/42 (2018.01)
 CPC - A61B 17/3423, A61B 17/3421, A61B 17/34, A61B 17/00234, A61B 2017/3445, A61B 2017/00278, A61B 2017/348

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

See Search History Document

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

See Search History Document

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

See Search History Document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2015/0011978 A1 (TERUMO KABUSHIKI KAISHA) 8 January 2015 (08.01.2015), entire document	1, 10-11
X ----- Y	US 8,608,652 B2 (Voegele et al.) 17 December 2013 (17.12.2013), entire document	1-9, 12, 17, 41 ----- 13-16, 37-40, 42-44
X ----- Y	US 5,762,629 A (Kambin) 9 June 1998 (09.06.1998), entire document	18-27 ----- 39
Y	US 6,156,006 A (Brosens et al.) 5 December 2000 (05.12.2000), entire document	13-16, 37-40, 42-44

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

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier application or patent but published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search
 3 December 2018

Date of mailing of the international search report

04 FEB 2019

Name and mailing address of the ISA/US
 Mail Stop PCT, Attn: ISA/US, Commissioner for Patents
 P.O. Box 1450, Alexandria, Virginia 22313-1450
 Facsimile No. 571-273-8300

Authorized officer:

Lee W. Young

PCT Helpdesk: 571-272-4300
 PCT OSP: 571-272-7774

INTERNATIONAL SEARCH REPORT

International application No.

PCT/IL 18/50934

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

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

1. Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

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

This International Searching Authority found multiple inventions in this international application, as follows:
This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be searched, the appropriate additional search fees must be paid.

Group I: Claims 1-27 and 37-44 directed to a stepped dilator; a cannula comprising a cross section with a long and short axis; a kit for providing intraperitoneal access comprising a stepped dilator, cannula, and trocar needle; and a method of gaining intraperitoneal access comprising a stepped dilator.

Group II: Claim 28-36 directed to a cannula comprising an edge defining and aperture.

Group III: Claim 45 directed to a method for gaining intraperitoneal access comprising a camera and illumination device.

Group IV: Claims 46-50 directed to a kit for setting a position of a robotic arm system.

-*Continued on extra sheet*-

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-27 and 37-44

Remark on Protest

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

-*--continuation of Box III-*--

SPECIAL TECHNICAL FEATURES

The invention of Group I includes the special technical features of a stepped dilator including a dilator body, a first tapered region, a second tapered region, and an isolating region; a cannula including a cross section of an inner lumen, a long axis, a short axis; a kit for providing intraperitoneal access including a stepped dilator, a cannula, at least two cylindrical members, a trocar needle, a handle region, and an inner dilator, not required by the claims of Group II, III, and IV.

The invention of Group II includes the special technical feature of a cannula comprising an edge defining aperture with a first edge portion and a second edge portion on one side of aperture and an opposite side of aperture, respectively, not required by the claims of Group I, III, and IV.

The invention of Group III includes the special technical features of a camera, a field of view of a camera, an intraperitoneally positioned illumination device, and one or more images, not required by the claims of Group I, II and IV.

The invention of Group IV includes the special technical features of a robotic arm system, a motor unit, at least one robotic arm, a cannula body, a mounting block, a spacing arm, an aligning arm, a stowed position, a deployed position, and a predetermined position, not required by the claims of Group I, II, and III.

COMMON TECHNICAL FEATURES

Groups I, II, and IV share the common technical feature of a cannula (generally) that can provide intraperitoneal access for use with a trocar needle.

However, this shared technical feature does not represent a contribution over prior art as being anticipated by US 8,608,652 B2 to Voegelé et al. (hereinafter Voegelé), which discloses a cannula that provides intraperitoneal access for use with a trocar (column 6, ln 60 -column 7, ln 11; fig. 8 see cannula 300 providing intraperitoneal access in the system of the trocar 400, dilator 200, and cannula 300).

Groups I, II, and III share the common technical feature of a method for gaining intraperitoneal access (generally).

However, this shared technical feature does not represent a contribution over prior art as being anticipated by Voegelé, which discloses a method for gaining intraperitoneal access (see fig. 8, 9A, and 9B; column 20, ln 51-65)

Groups I and IV share the common technical feature of a kit for treating a body orifice (generally).

However, this shared technical feature does not represent a contribution over prior art as being anticipated by Voegelé, which discloses a kit for treating a body orifice (column 6, ln 50-54 kit includes speculum, surgical delivery device, and transorifice device).

As the common technical features were known in the art at the time of the invention, these cannot be considered special technical feature that would otherwise unify the groups.

Therefore, Groups I-III lack unity under PCT Rule 13 because they do not share a same or corresponding special technical feature.