



(11)

EP 2 476 384 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
18.07.2012 Bulletin 2012/29

(51) Int Cl.:
A61B 17/34 (2006.01)

(21) Application number: 12164164.1

(22) Date of filing: 21.11.2007

(84) Designated Contracting States:
AT BE BG CH CY CZ DE DK EE ES FI FR GB GR
HU IE IS IT LI LT LU LV MC MT NL PL PT RO SE
SI SK TR

(30) Priority: 22.11.2006 US 866939 P

(62) Document number(s) of the earlier application(s) in accordance with Art. 76 EPC:
07854750.2 / 2 086 430

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Remarks:

This application was filed on 13-04-2012 as a divisional application to the application mentioned under INID code 62.

(54) Trocar cannula with atraumatic tip

(57) A surgical access port is provided with a trocar cannula having a substantially rigid portion and an atraumatic distal tip. The atraumatic distal tip is substantially compliant relative to the rigid portion of the cannula. The trocar cannula provides unobstructed surgical access into a body cavity allowing the insertion and removal of surgical instruments through the trocar cannula and into the body cavity using a minimal sized incision or entry-way.

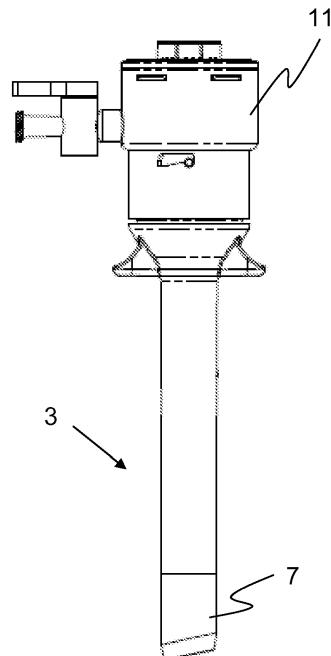


FIG. 1

DescriptionCross-reference to related applications

[0001] This application claims the benefit of U.S. Provisional Patent Application No. 60/866,939, filed on November 22, 2006, the entire disclosure of which is hereby incorporated by reference as if set forth in full herein.

Background

[0002] This invention generally relates to medical access devices and, more specifically, to a trocar cannula with an atraumatic tip.

[0003] During use of a trocar in a surgical procedure, such as a laparoscopic procedure, a trocar is placed across the abdominal wall and the trocar cannula is left disposed across the abdominal wall. The distal tip of the cannula is positioned on the anterior side of the abdominal wall. Laparoscopic procedures can utilize several trocars across the abdominal wall and therefore several cannulas may be disposed across the abdominal wall at the same time. With several cannulas disposed across the abdominal wall, it can be difficult to constantly observe and monitor the positioning of the distal tips of the cannulas while the laparoscopic procedure is being conducted. For example, because of the number of cannulas disposed across an abdominal wall in a laparoscopic procedure, it is possible for a cannula tip to unintentionally engage body tissue out of the surgeon's field of view provided by the laparoscope and camera. The trocar in accordance with aspects of the present invention minimizes the possibility for a trocar cannula to inadvertently engage and traumatize body tissue in a body cavity, due to accidental misuse, for example, by providing a cannula with a distal tip, which is elastomeric, compliant, soft, and generally atraumatic.

Summary

[0004] In one aspect, a trocar cannula comprises a elongate body having a proximal enlarged end and a distal tip end with a lumen extending from the proximal end to the distal end for inserting and removing surgical instruments through the lumen and an atraumatic tip integrated with and extending from the distal tip end of the elongate body, the tip being more compliant than the elongate body and formed from a material different from the elongate body.

[0005] In one aspect, a trocar cannula comprises a elongate body having a proximal enlarged end and a distal tip end with a lumen extending from the proximal end to the distal end for inserting and removing surgical instruments through the lumen and a flexible joint integrated with and extending from the distal tip end of the elongate body, the flexible joint being more compliant than the elongate body and formed from a material different from the elongate body. A tip is integrated with and ex-

tends from an opposing end of the flexible joint away from the distal tip end of the elongate body, the tip being less compliant than the flexible joint and being formed from a material different from the flexible joint.

[0006] In one aspect, a trocar cannula comprises a trocar seal housing having an instrument seal and a zero seal contained within the trocar seal housing, a elongate body having a proximal enlarged end having projections extending from an outer surface of the proximal enlarged end, a distal tip end having annular projections and a lumen extending from the proximal enlarged end to the distal tip end for inserting and removing surgical instruments through the lumen, the trocar seal housing releasably connected to plurality of projections of the proximal enlarged end with the zero seal situated within the proximal enlarged end and the instrument seal situated within the zero seal and an atraumatic tip fixedly interlocked with the annular projections of the distal tip end of the elongate body, the tip being deformable, formed from a material different from the elongate body and having a flexural modulus of about 1,500 psi and an ultimate elongation of about 400%, and the elongate body having a flexural modulus substantially greater than the flexural modulus of the tip and an ultimate elongation substantially less than the ultimate elongation of the tip.

[0007] Many of the attendant features of the present invention will be more readily appreciated as the same becomes better understood by reference to the foregoing and following description and considered in connection with the accompanying drawings in which like reference symbols designate like parts throughout.

Brief Description of the Drawings

35 **[0008]**

FIGs. 1-2 are side views of a trocar system in accordance with various aspects of the present invention;

40 FIGs. 3-4 are side views of an operational use of a trocar system in accordance with various aspects of the present invention;

FIG. 5 is a cross-sectional side view of a trocar cannula in accordance with various aspects of the present invention;

45 FIG. 6A is an enlarged cross-sectional side view of a portion of a trocar cannula in accordance with various aspects of the present invention;

FIG. 6B is an enlarged side view of a portion of a trocar cannula without a tip in accordance with various aspects of the present invention;

FIG. 7 is a side view of a trocar system in accordance with various aspects of the present invention;

FIG. 8 is a side view of a trocar system in accordance with various aspects of the present invention;

50 FIG. 9A is an enlarged cross-sectional side view of a portion of a trocar cannula in accordance with various aspects of the present invention;

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FIG. 9B is an enlarged side view of a portion of a trocar cannula without a tip in accordance with various aspects of the present invention; FIG. 10 is a side view of a trocar system in accordance with various aspects of the present invention; FIG. 11 is a side view of a trocar cannula in accordance with various aspects of the present invention; FIG. 12 is a side view of a trocar cannula in accordance with various aspects of the present invention; FIG. 13 is a side view of a trocar cannula in accordance with various aspects of the present invention; FIG. 14 is an enlarged cross-sectional side view of a portion of a trocar cannula in accordance with various aspects of the present invention; and FIG. 15 is an enlarged cross-sectional side view of a portion of a trocar cannula in accordance with various aspects of the present invention.

Detailed Description

[0009] Generally, an access port or trocar is provided with a cannula having an atraumatic distal tip. The trocar is used during minimally invasive surgery to provide an access channel into the body through which a surgeon may insert medical instruments. The cannula in one aspect has a proximal portion, which is formed of a rigid polymer and a distal portion formed of an elastomeric polymer. The elastomeric cannula tip has sufficient column strength to enable passage through a body wall when used in combination with an obturator yet has a lower durometer hardness and is more compliant as compared to the proximal portion of the cannula. The low durometer and compliance of the atraumatic cannula tip prevents the cannula tip from potentially traumatizing, due to accidental misuse, for example, or engaging adjacent body tissue during a surgical procedure such as a laparoscopic procedure. Some trocar cannulas are typically formed of a single material such as stainless steel or polycarbonate. These trocar cannulas can be rigid and include tapered distal tips, which can engage body tissue and may cause trauma to the engaged body tissue, due to, for example, accidental misuse.

[0010] Referring to FIGS. 1-14, a surgical access port, e.g., a trocar, which comprises a trocar seal housing 11, a trocar cannula 3, and/or an obturator is provided. In this description, "proximal" or "proximally" refers to that portion of the instrument, component, or element that extends toward the user. "Distal" or "distally" refers to that portion of the instrument, component, or element that extends away from the user. The trocar is configured to access a body cavity and to maintain positive pressure at its distal end to prevent loss of surgical insufflation gas such as carbon dioxide used, for example, in laparoscopic procedures to insufflate the body cavity. The trocar seal and trocar cannula is also configured to sealingly engage surgical instruments of various diameters, which would typically be inserted through the trocar, to prevent loss of surgical gas during use of such instruments and/or

when no instrument is inserted. In one aspect, the trocar seal housing is releasably attachable to the trocar cannula to allow the seal to be removed during surgery to enable the extraction of tissue specimens through the trocar. The trocar in one aspect has or is included with an optical obturator having a tip, which includes a smooth outer surface and has a high degree of optical clarity.

[0011] The trocar seal housing 11 in one aspect can be easily detached or removed from a proximal enlarged end of the trocar cannula 3 and easily attached or reattached to the trocar cannula 3 for example during a surgical procedure. During surgery, small tissue specimens may be extracted from a body cavity through a trocar to enable pathological analysis of the tissue specimen. The integrity of the tissue specimen can be maintained or the maintenance facilitated by avoiding or minimizing withdrawal of delicate tissue specimens through a trocar seal. As such, in one aspect, the trocar seal housing 11 is arranged to be removed from the trocar cannula 3 to enable extraction of tissue specimens from a body cavity while maintaining the integrity of the tissue specimen. The trocar seal housing 11 also easily reattaches to the trocar cannula 3 after its initial removal during a surgical procedure.

[0012] Laparoscopic surgery of the abdominal area typically requires the introduction of an insufflation gas into the peritoneal cavity of the patient. The insufflation gas is usually pressurized to about 10 mm Hg above atmospheric pressure. This in turn lifts the abdominal wall away from the organs underlying it. Pressurized insufflation gas in one aspect is introduced through the stopcock of the trocar seal housing 11 into the trocar cannula 3. Trocar seal housing 11, accommodating instrument and/or zero seals, prevents the gas from escaping proximally out from the cannula 3 and allows the insertion and removal of a laparoscope and surgical instruments into and out of the surgical site.

[0013] Also, in one aspect, the trocar seal housing 11 is easily removable from the trocar cannula 3 to enable rapid desufflation of an insufflated body cavity. In one aspect, a trocar lock releasably attaches the trocar cannula to the trocar seal housing. For example, towards the end of a laparoscopic surgical procedure, release of the insufflation gas such as carbon dioxide from the peritoneal cavity of the patient is performed. By opening one or more stopcock valves on the trocar seal, desufflation can be achieved. The flow rate through the stopcock valves, however, can be slow with regard to evacuation of the carbon dioxide from the peritoneal cavity and therefore the time expended to evacuate the insufflation gas can be excessive. By removing the seal housing 11 from the cannula 11, the cannula provides an unobstructed outlet for the insufflation gas to escape thereby decreasing desufflation time.

[0014] During an operational exemplary use, a laparoscope 21 is inserted into and through trocar seal housing 11 and into cannula 3. An endoscopic video camera is attached to the proximal end of the laparoscope. As the

surgeon advances the trocar through the body wall, the surgeon can visually observe the tissue through the laparoscope via a video monitor, which is connected to the endoscopic video camera.

[0015] Referring to FIGS. 1-6, the trocar system, i.e., a surgical access port for entry into a body cavity, in one aspect is provided in that an atraumatic distal tip 7 of a trocar cannula 3 prevents or minimizes potentially destructive engagement with laparoscopic instrumentation such as an articulating laparoscope 21. For example, if an articulating laparoscope is placed through a typical cannula, the distal tip of the laparoscope then articulated, and the laparoscope then withdrawn from the cannula, the flexible joint of the laparoscope can contact the distal tip of the cannula resulting in damage to the flexible joint of the laparoscope. The damage to the laparoscope may be such as to require immediate replacement of the laparoscope resulting in an immediate delay in the surgical procedure. Articulating laparoscopes are also typically very expensive in comparison with non-articulating laparoscopes and therefore, it can be very costly for a hospital or a manufacturer to replace or repair a damaged articulating laparoscope. The trocar in accordance with one aspect of the present invention addresses this by providing a cannula 3 with an elastomeric distal tip 7, which is compliant, and has a low durometer. The atraumatic elastomeric distal tip 7 on the cannula 3 in one aspect is sufficiently compliant and soft so as to prevent or minimize potential damage to instrumentation, which engages the cannula tip 7 during withdrawal of the instrumentation.

[0016] In one aspect, the cannula 3 with the atraumatic tip 7 is a high durometer rigid polymer cannula formed of a material such as polycarbonate. The atraumatic tip comprises a low durometer elastomeric material such as polyurethane. The cannula in one aspect is injection molded and the atraumatic elastomeric tip is then over-molded onto a distal tip end of an elongate body of the cannula. The atraumatic tip is held in place via a mechanical interlock fit. The mechanical interlock fit includes a plurality of annular ribs 9 on the cannula tip 4, which are over-molded with the atraumatic elastomeric tip 7. The annular ribs 9 prevent or resist axial movement of the atraumatic elastomeric tip 7 relative to the rigid polycarbonate cannula. The length of the atraumatic elastomeric tip 7 can range from approximately .25" to approximately 1" long. This length enables the cannula 3 to be disposed across the abdominal wall such that the rigid portion 5 of the cannula 3 is positioned within the abdominal wall to hold the abdominal wall in a retracted position. The retracted abdominal wall also serves to aid with anchoring the cannula 3 in place and prevents axial movement of the cannula during the surgical procedure. The atraumatic elastomeric tip 7 is positioned within the peritoneal cavity and in one aspect is the only portion of the cannula to be disposed within the peritoneal cavity.

[0017] In one aspect, the cannula tip or distal tip end 4 of the cannula has a wall thickness smaller than re-

maining portions of the cannula 3. A proximal portion of the atraumatic tip 7 has a wall thickness substantially similar to cannula tip 4 and in one aspect, a distal portion of the atraumatic tip 7 has a wall thickness substantially similar to a wall thickness of a proximal portion of the cannula 3. In one aspect, the atraumatic elastomeric tip 7 is formed from a material, e.g., a transparent polyurethane material, or otherwise configured/arranged to ensure that visibility through the tip is maintained. The transparent elastomeric tip assists in positioning the cannula 3 within the abdominal wall. By placing the laparoscope lens just proximal to the tip of the cannula 3, the abdominal wall can be viewed through the cannula tip 6 while positioning the cannula such that only the atraumatic elastomeric tip 7 of the cannula is disposed within the peritoneal cavity. The atraumatic elastomeric tip 7 in one aspect is formed with a contrasting tint as compared to the rigid portion 5 of the cannula to further aid with positioning of the cannula within the abdominal wall and the peritoneal cavity.

[0018] In one aspect, the rigid portion 5 of the cannula 3 is formed from polyethylene, polysulfone, polyethersulfone, polyetherimide, polycarbonate, polyurethane, liquid crystal polymer, nylon, polyester, polypropylene, or ABS (Acrylonitrile Butadiene Styrene). In one aspect, the atraumatic elastomeric tip portion of the cannula is formed from silicone, polyurethane, polyester, polystyrene, nylon, polyvinyl chloride, mylar, polyethylene, Kraton® thermoplastic elastomers, C-Flex® thermoplastic elastomers, Versaflex® thermoplastic elastomers, Santoprene® thermoplastic elastomers, Carbothane® thermoplastic polyurethanes, copolymer/mineral oil gels, polyisoprene, or natural rubber.

[0019] In one aspect, the cannula 3 has an ultimate elongation less than the ultimate elongation of the atraumatic tip 7. For example, the ultimate elongation of the cannula 3 is about 120% versus the ultimate elongation of the atraumatic tip 7 being about 410%. In one aspect, the ultimate elongation of the cannula 3 is within the range of about 2% to about 150% versus the ultimate elongation of the atraumatic tip 7 being in the range of about 300% to about 1,000%. In one aspect, the ultimate elongation of the tip is at least three times greater than the ultimate elongation of the cannula. The large or greater ultimate elongation of the tip ensures that the tip does not tear off when a surgical instrument is inserted through the cannula, moved off-axis and/or portions of the instrument articulated off-axis relative to the longitudinal axis of the cannula which may interfere and/or damage the instrument and/or tip portions of which may fall into the surgical site.

[0020] In one aspect, the atraumatic elastomeric tip 7 of the cannula 3 is formed from an elastomeric material which further softens upon extended exposure to body temperatures. The elastomeric material can therefore be less compliant during the initial insertion of the trocar when lower compliance is used providing a lower insertion force, and upon extended exposure to body temper-

atures, the atraumatic cannula tip would then soften resulting in a more compliant and therefore a more atraumatic cannula tip. Examples of elastomeric materials with these properties are the Carbothane® thermoplastic polyurethanes available from Noveon.

[0021] In one aspect, the cannula 3 with the atraumatic elastomeric tip 7 is provided with inside diameters ranging from 1mm to 30mm. A typical wall thickness for the cannula is about .25mm to 1 mm. The cannula with the atraumatic elastomeric tip in one aspect is formed using a dual-shot molding process whereby the rigid portion is first molded and then the elastomeric portion is then over-molded onto the rigid portion using a dual-shot injection mold and a dual-shot injection molding press. In one aspect, the cannula 3 with the atraumatic elastomeric tip 7 can be used in conjunction with non-bladed dilating obturators, non-shielded bladed obturators, shielded bladed obturators, electrosurgical obturators, and blunt tip obturators. In one aspect, the cannula 3 with the atraumatic elastomeric tip 7 can have a semi-rigid or flexible proximal portion to enable placement of the cannula through a body conduit such as a urethra or ureter. The proximal portion of the cannula or catheter would be configured to be less flexible than the distal tip portion of the cannula. In one aspect, the proximal portion 8 of the cannula 3 is enlarged accommodating instrument and/or zero seals, surgical instruments with different diameters and orientation of such instruments, providing finger holds or grips, suture tie slots, and/or a releasable connection to the seal housing 11.

[0022] In one aspect, the cannula 3 with the atraumatic elastomeric tip 7 is formed of a single material. The cannula has thinned wall sections or axial slots 25 at its distal tip to provide the distal tip of the cannula with greater flexibility as compared to the proximal portions of the cannula (FIG. 7). In FIG. 8, the cannula in one aspect is formed of a single material with a flexible joint 27 between the proximal portion and the distal portion of the cannula. The flexible joint, e.g., bellows, would enable the distal tip or portion of the cannula to pivot in response to contact with body tissue or inserted instrumentation. In one aspect, the flexible joint 27 integrated with and extending from the proximal portion of the cannula 3. The flexible joint 27 is more compliant than the proximal or rigid portion 5 of the cannula 3 and is formed from a material that is different from the material of the other portions of the cannula 3. The tip or distal portion of the cannula is integrated with and extending from an opposing end of the flexible joint away from the distal tip end of the elongate body. In one aspect, the tip is less compliant than the flexible joint and is formed from a material different from the material of the flexible joint. The tip or distal portion in one aspect is formed from a material corresponding to the material of the proximal portion of cannula 3. In one aspect, the flexible joint is formed of thermoplastic polyurethane and has a flexural modulus substantially smaller than the flexural modulus of the cannula 3.

[0023] In one aspect, the atraumatic tip 7 has a flexural

modulus that is about 1,500 psi and the flexural modulus of the cannula is about 330,000 psi. In one aspect, the flexural modulus of the cannula is greater than at least 100,000 psi and the tip 7 is less than at least 10,000 psi.

5 In one aspect, the flexural modulus of the atraumatic tip is within the range of about 500 psi to about 15,000 psi versus the cannula is in the range of about 100,000 psi to about 500,000 psi. The flexural modulus of the cannula is about ten times greater than the flexural modulus of 10 the tip. The greater flexural modulus of the tip relative to the cannula allows the tip 7 to deflect, bend and/or deform to avoid unintended contact and complications with tissue/organs and to avoid potential interference and/or damage to flexible or deflectable instruments articulated 15 during a surgical procedure or instruments being withdrawn with portions of the instrument deflected or articulated off-center or alignment with the longitudinal axis of the cannula. The pronounced difference ensures the tip to deflect relative to the cannula allowing the remaining 20 portion to remain still and thus not pull, tear or enlarge the minimally sized surgical incision or entryway. Also, the difference ensures that tip deflects or does not otherwise interfere with a surgical instrument, e.g., a laparoscope, obturator and others, having a tubular portion or 25 shaft with properties similar to the remaining portions of the cannula inserted through the cannula and moved off-axis or portions of the instrument articulated off-axis.

[0024] The atraumatic elastomeric tip 7 in one aspect is bonded with an adhesive to the rigid portion of the 30 cannula. The atraumatic elastomeric tip in one aspect is ultrasonically welded or thermally welded to the rigid portion of the cannula. In one aspect, the atraumatic elastomeric tip is formed of a material arranged to chemically bond with the rigid portion 5 of the cannula 3 during injection molding. For example, the cannula 3 may be 35 formed of polypropylene and the elastomeric tip 7 formed of a C-Flex® thermoplastic elastomer specifically formulated to chemically bond to polypropylene during injection molding.

40 **[0025]** In one aspect, the atraumatic elastomeric tip 7 is coated or treated to reduce the friction associated with the movement of instrumentation when the instrumentation contacts the elastomeric tip. The coating or treatment can also reduce the force used to place the trocar through 45 the abdominal wall. Examples of coatings and treatments include parylene coatings, hydrophilic coatings, plasma surface treatments, and chlorination treatments. In one aspect, the atraumatic elastomeric tip is formed from a radiopaque material. A typical radiopaque material includes barium sulfate as an additive.

50 **[0026]** Referring now to FIG. 9, in one aspect, the tip of the rigid portion 5 of the cannula 3 is formed with annular barbs 15 to provide a mechanical lock with the over-molded elastomeric tip 7. The tip 4 of the rigid portion 5 of the cannula 3 in one aspect is formed with a series of 55 directionally alternating annular barb configurations, such that the barbs create a mechanical lock increasing axial tension strength of the tip to the rest of the cannula

and preventing the elastomeric tip 7 from moving in either a proximal or a distal direction relative to the rigid portion 5 of the cannula 3. The tip 4 of the rigid portion 5 of the cannula 3 in one aspect is formed with annular grooves to provide a mechanical lock with the over-molded elastomeric tip 7. The tip 4 of the rigid portion 5 of the cannula 3 in one aspect is formed with holes and/or axial grooves to provide a mechanical lock with the over-molded elastomeric tip 7. The tip 4 of the rigid portion 5 of the cannula 3, in one aspect, is formed with a thread to provide a mechanical lock with the over-molded elastomeric tip. [0027] The inside diameter of a distal tapered portion 17 of the atraumatic elastomeric tip 7 in one aspect is configured to slightly interfere with the outside diameter of an obturator 31 such that when the obturator is inserted into the cannula 3, the obturator expands the elastomeric distal tip of the cannula to create a smooth transition between the distal tip of the cannula and the obturator (FIG. 10). The smooth transition between the distal tip of the cannula and the obturator can decrease the possibility for body tissue to wedge between the cannula and the obturator during insertion through the abdominal wall and can therefore reduce the insertion force required to place the trocar through the abdominal wall. The interference fit can also serve to create a seal between the distal tip of the obturator 31 and the cannula 3. The seal can prevent or minimize insufflation gasses from flowing between the interface of the distal tip of the cannula and the obturator. In one aspect, to facilitate the interference fit, the atraumatic tip 7 has an inner diameter that is less than an inner diameter of the elongate body or distal tip end of the cannula 3. The atraumatic tip 7, in one aspect, is chamfered or tapered narrowing and transitioning from a larger inner diameter corresponding to the inner diameter of the rigid portion and/or the distal tip end 4 of the elongate body to the smaller inner diameter of the atraumatic tip 7 (FIG. 15). As such, the taper also provides a smooth transition or lead-in (reducing "catch" points) for the obturator or other surgical instruments being inserted into the cannula and through the tip.

[0028] The greater or larger ultimate elongation of the tip relative to the rigid portion 5 of the cannula 3, as noted above, also assists in the interference fit between the tip and the inserted obturator. The tip 7 is allowed to stretch to accommodate an obturator having an outer diameter larger than the inner diameter of the tip, thereby providing a tight instrument seal and a smooth transition between instrument and tip and tip to cannula. The difference ensures that the tip stretches or expands relative to the cannula allowing the tip to allow passage of the inserted surgical instrument, e.g., a laparoscope, obturator, and others, having an outer diameter larger than the inner diameter of the tip. An interference fit and/or instrument seal is thereby provided with the tip and the inserted surgical instrument.

[0029] The cannula 3 with the atraumatic elastomeric tip 7 in one aspect is formed of reusable materials to enable the product to be autoclave sterilized and re-used.

Reusable materials for the cannula and elastomeric tip in one aspect can be polysulfone, polyetherimide, polyethersulfone, silicone, and polyisoprene. The cannula 3 with the atraumatic elastomeric tip 7 in one aspect is formed of stainless steel with a bonded elastomeric tip formed of silicone or polyisoprene to enable the product to be autoclave sterilized and re-used.

[0030] The cannula 3 with the atraumatic tip 7 in one aspect has a short length over-molded section at its distal tip. The short length over-molded section can provide for an atraumatic elastomeric cannula tip, yet utilize no or minimal column strength or less axial column strength as compared to a longer length over-molded section. The length of the shortened section in one aspect varies from .025" to .250".

[0031] In FIG. 11, the cannula 3 in one aspect has a flexible elastomeric joint 33 between the rigid portion 5 of the cannula and the distal tip 7' of the cannula. The distal tip of the cannula in one aspect is either flexible or rigid. The flexible joint 33 allows the distal tip of the cannula to deflect in response to contact between, for example, body tissue or an instrument thereby preventing potential trauma to the body tissue or potential damage to the instrument. The flexible joint 33 in one aspect has a bellows configuration with a minimal axial length. The minimal axial length of the flexible joint 33 in combination with a rigid or semi-rigid distal tip 7 in one aspect can provide the cannula 3 with greater column strength as compared to a cannula with an over-molded elastomeric tip. The flexible joint in one aspect is formed from silicone, polyurethane, Kraton® thermoplastic elastomers, C-Flex® thermoplastic elastomers, Versaflex® thermoplastic elastomers, polyisoprene, Santoprene® thermoplastic elastomers, Carbothane® thermoplastic polyurethanes, copolymer/mineral oil gels, or natural rubber.

[0032] In FIG. 12, in one aspect, the cannula 3 with the flexible joint 33 has a seal 35 formed of a gel material located at the distal tip 7' of the cannula. The seal 35 in one aspect is configured to maintain a seal in the absence of inserted instrumentation and the seal, in one aspect, is configured to maintain a seal in the presence of inserted instrumentation. The flexible joint 33 allows the seal 35 and the distal tip 7' to pivot in response to the lateral movement of inserted instrumentation to ensure that a seal 35 is maintained during off-axis movement of the instrumentation. The seal 35 could be formed of a single piece component with the gel for example shaped as a disc with a slit in the center of the disc. In one aspect, as shown in FIG. 13, seal 35' is formed of two opposed gel rollers, such that instrumentation would be inserted between the rollers. The cannula 3 with the seal 35 can also be formed without the flexible joint 33. The elongation and sealing properties of the gel material can enable a seal to be maintained during off-axis movement of inserted instrumentation. The gel material in one aspect is formulated of an SEBS (Styrene Ethylene Butylene Stryrene) copolymer and a mineral oil.

[0033] Referring now to FIG. 14, the atraumatic elas-

tomeric cannula tip 7 in one aspect is formed with an internal wire form 19 to provide greater rigidity and column strength as compared to a cannula tip without an internal wire form. The wire form 19 in one aspect is configured in the shape of a coil spring and a polymer is fused or over-molded over the wire form resulting in a cannula tip 7 with an embedded wire form. The wire form in one aspect also comprises a series of wires radially spaced and embedded within the atraumatic elastomeric cannula tip.

[0034] Accordingly, a trocar cannula with an atraumatic tip is provided. Although the present invention has been described in certain specific aspects, many additional modifications and variations would be apparent to those skilled in the art. It is therefore to be understood that the present invention may be practiced otherwise than specifically described, including various changes in the size, shape and materials, without departing from the scope and spirit of the present invention. Thus, embodiments of the present invention should be considered in all respects as illustrative and not restrictive.

Claims

1. A trocar cannula (3) for receiving an articulating laparoscope (21) comprising:

an elongate body (5) having a proximal enlarged end and a distal tip end (4) with a lumen extending from the proximal end to the distal end for inserting and removing surgical instruments through the lumen; and
 an atraumatic tip (7) extending from the distal tip end (4) of the elongate body (5);
characterized in that the elongate body (5) is formed of a rigid polymer and the tip (7) is deflectable relative to the elongate body (5) to avoid interference with the articulating laparoscope.

2. The cannula of claim 1, wherein the distal tip end (4) of the elongate body (5) comprises annular ribs (9) and wherein the atraumatic tip (7) is fixedly interlocked with the annular ribs (9).

3. The cannula of claim 2, wherein the atraumatic tip (7) is formed of a material different than that of the elongate body (5) and the atraumatic tip (7) has a flexural modulus of about 1,500 psi and an ultimate elongation of about 400% and the elongate body (5) has a flexural modulus substantially greater than the flexural modulus of the atraumatic tip (7) and an ultimate elongation substantially less than the ultimate elongation of the atraumatic tip (7).

4. The cannula of claim 3, wherein the flexural modulus of the elongate body (5) is approximately 300,000

psi and the ultimate elongation of the elongate body is approximately 100%.

5. The cannula of any of the preceding claims, wherein the atraumatic tip (7) has an inner diameter less than an interior diameter of the elongate body (5).

6. The cannula of claim 5, wherein the inner diameter of the atraumatic tip (7) is tapered.

7. The cannula of any of the preceding claims, wherein the inner diameter of the atraumatic tip (7) is chamfered.2.

15 8. The cannula of claim 1, further comprising a flexible elastomeric joint (33) between the elongate body (5) and the atraumatic tip (7).

20 9. The cannula of claims 1 or 8, wherein the atraumatic tip (7) is flexible.

10. The cannula of claims 1 or 8, wherein the atraumatic tip (7) is rigid.

25 11. The cannula of any of claims 8-10, further comprising a seal (35) formed of a gel material located at the atraumatic tip (7).

30 12. The cannula of claim 11, wherein the seal (35) comprises two opposed gel rollers.

13. The cannula of any of claims 11-12, wherein the gel material comprises a styrene ethylene butylene styrene copolymer and a mineral oil.

35 14. The cannula of claim 1, wherein the elongate body (5) and the atraumatic tip (7) are formed of a single material.

40 15. The cannula of claim 14, wherein the atraumatic tip (7) further comprises axial slots (25) to provide the atraumatic tip (7) with flexibility.

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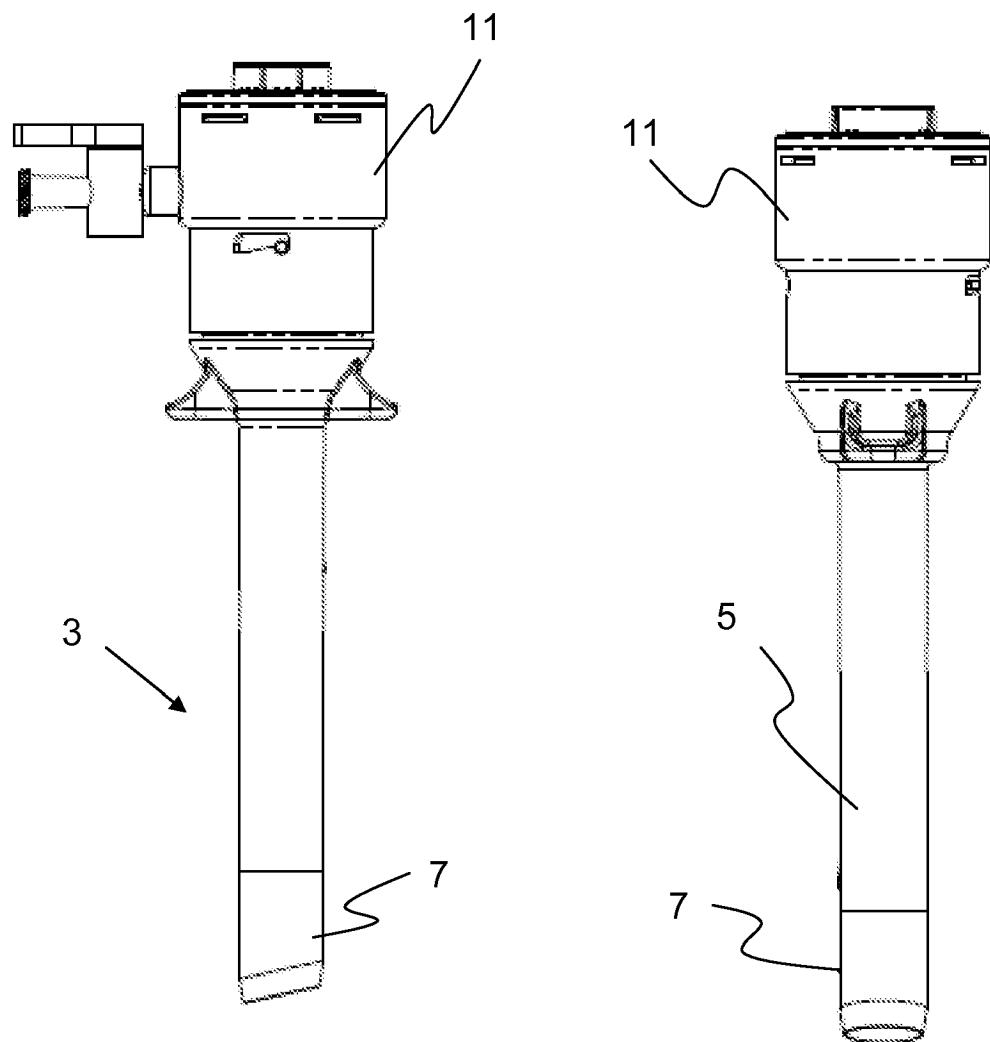
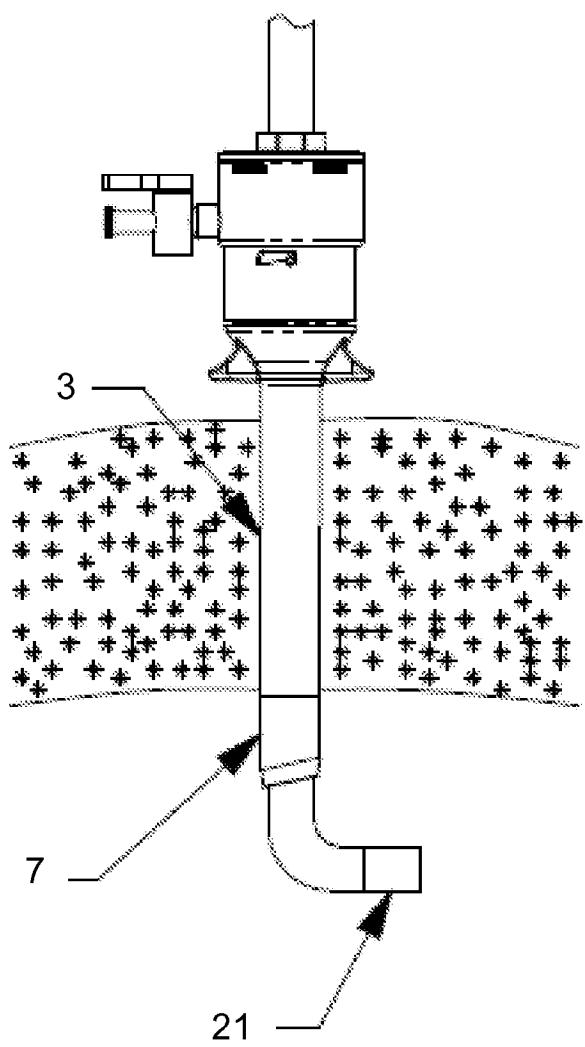
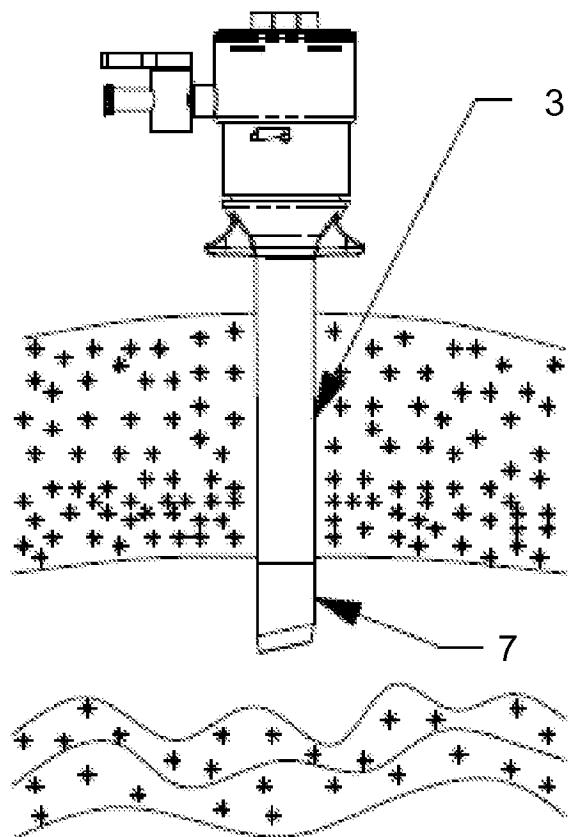


FIG. 1

FIG. 2



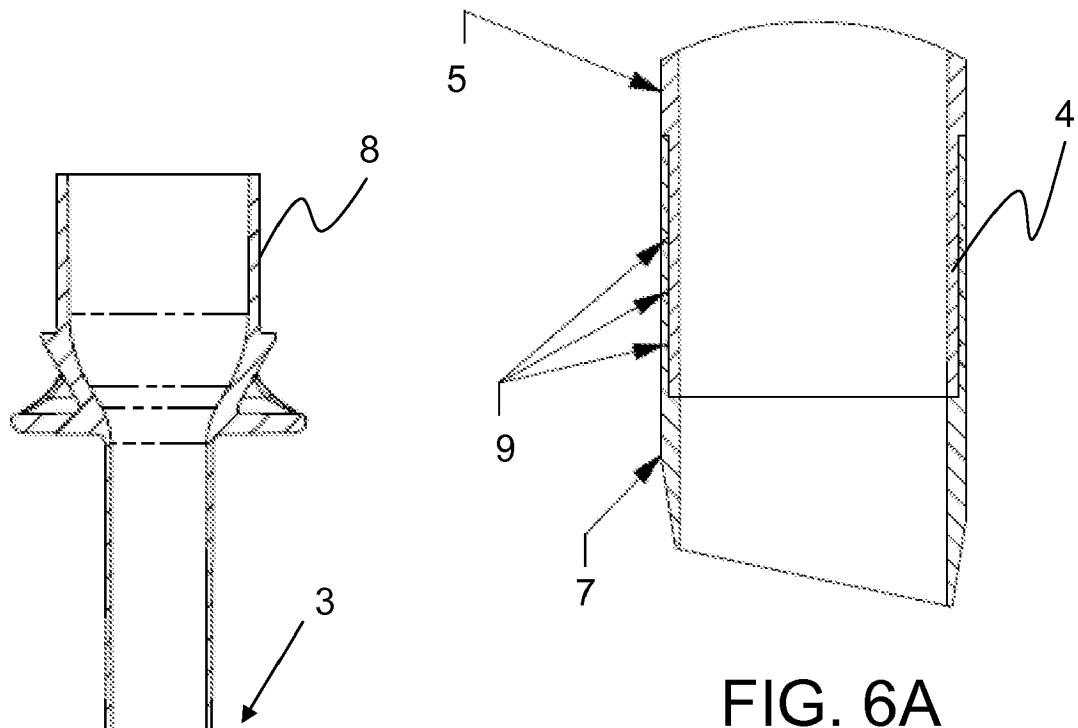


FIG. 6A

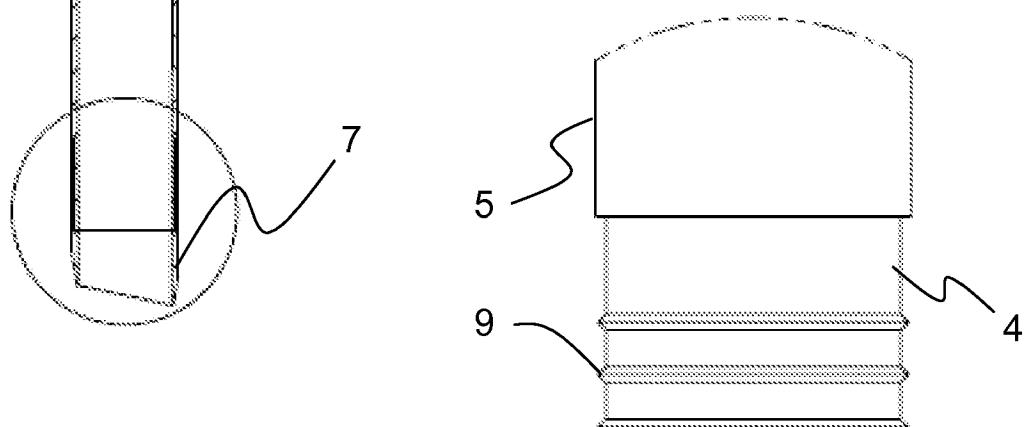


FIG. 5

FIG. 6B

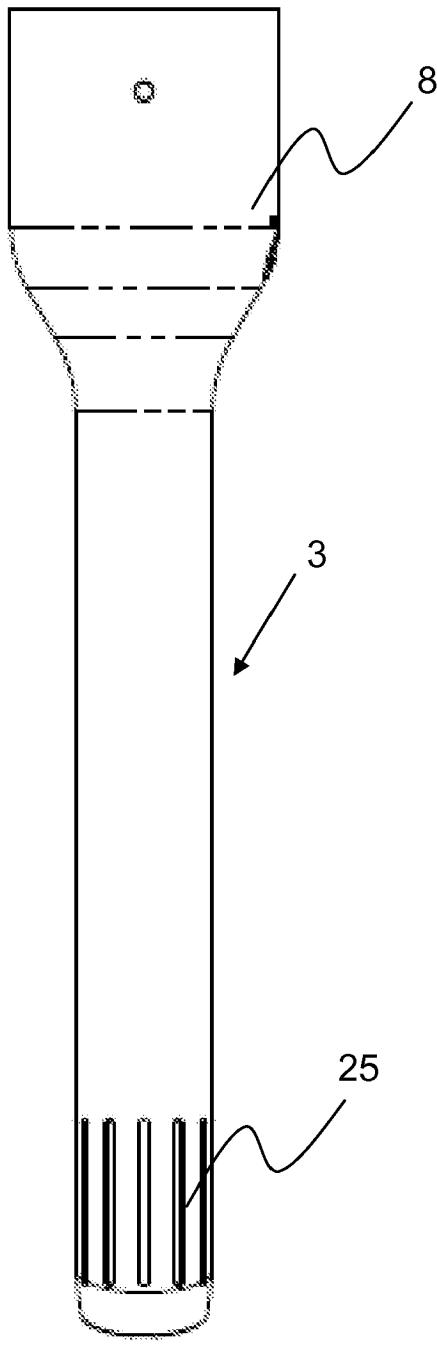


FIG. 7

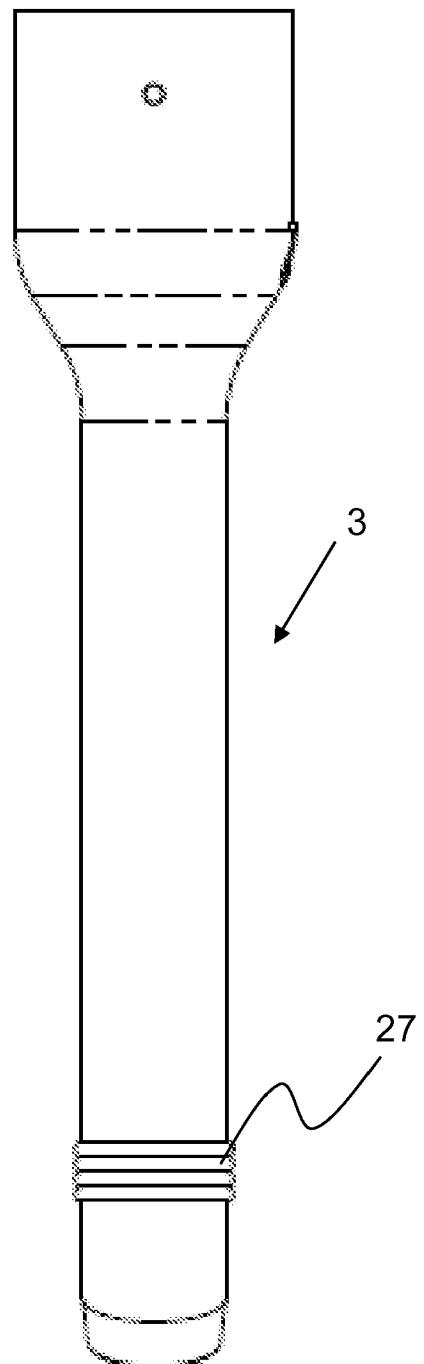


FIG. 8

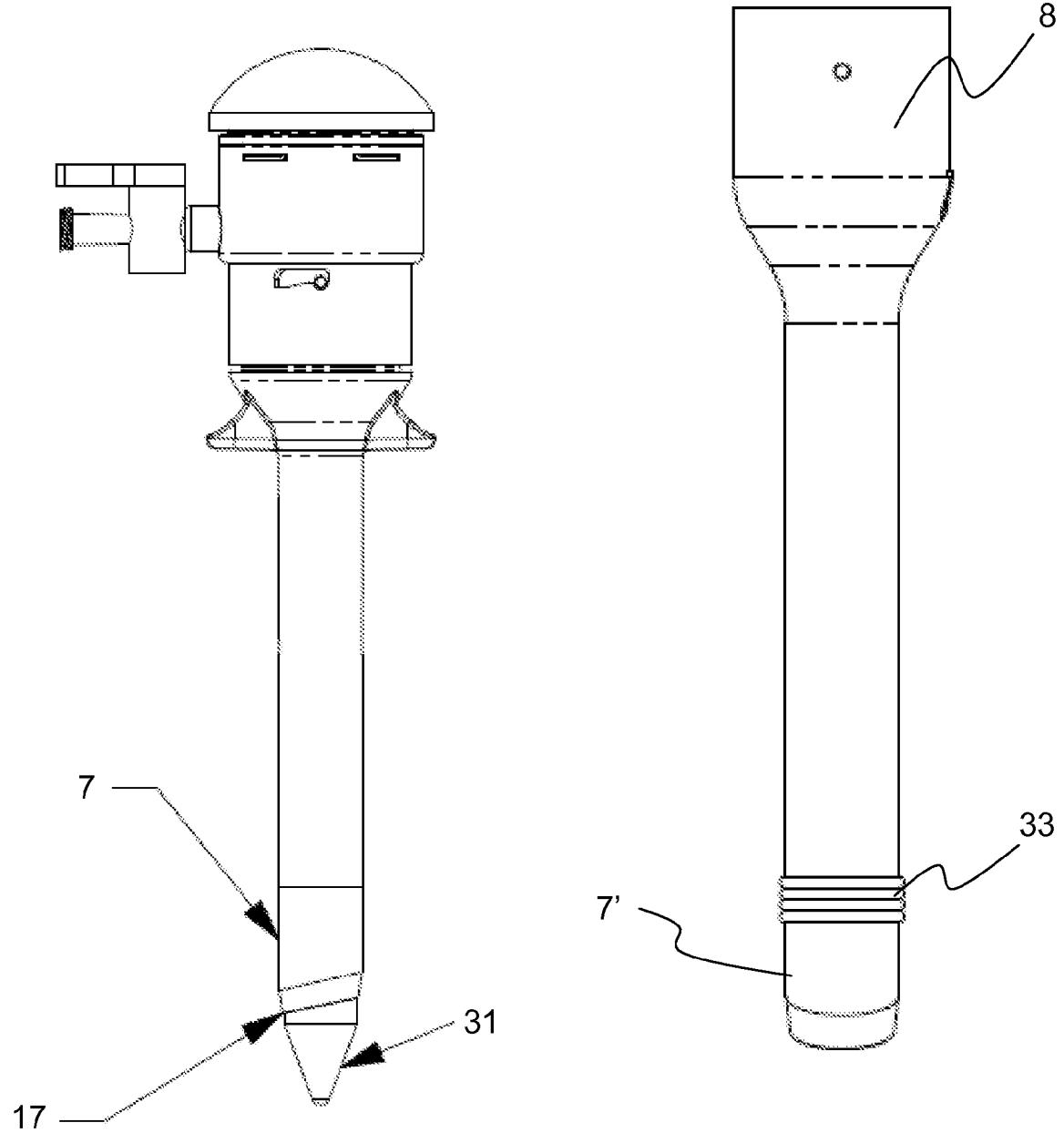


FIG. 10

FIG. 11

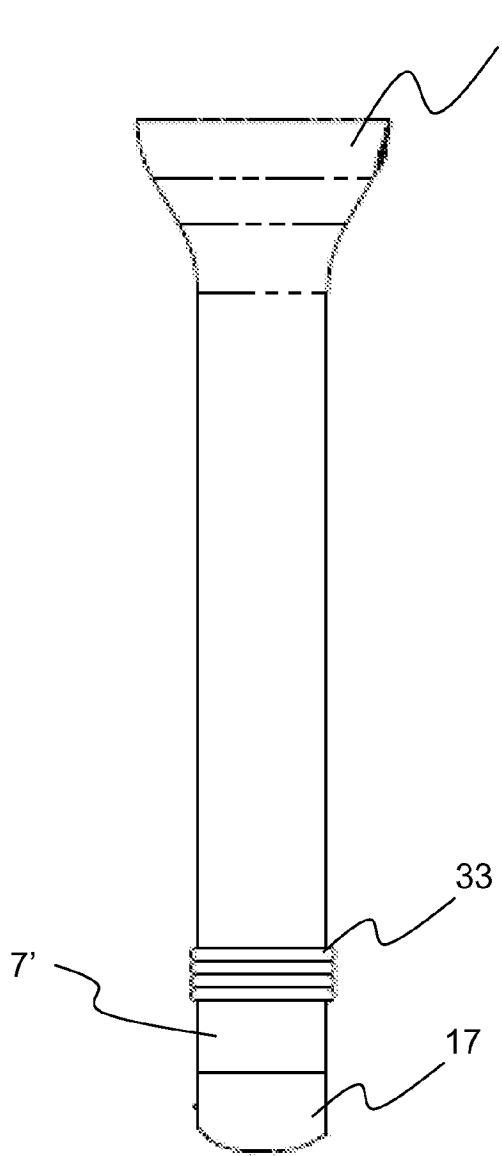


FIG. 12

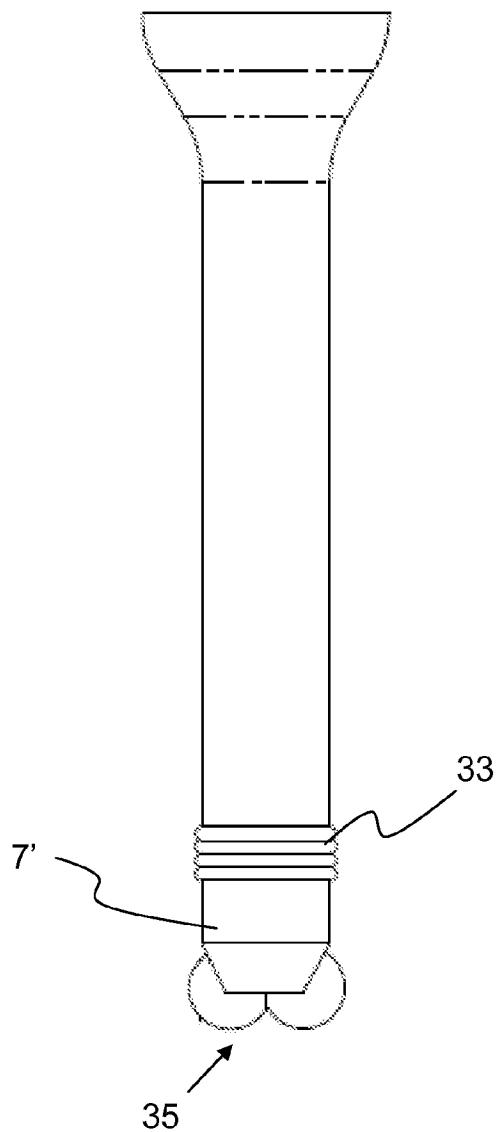


FIG. 13

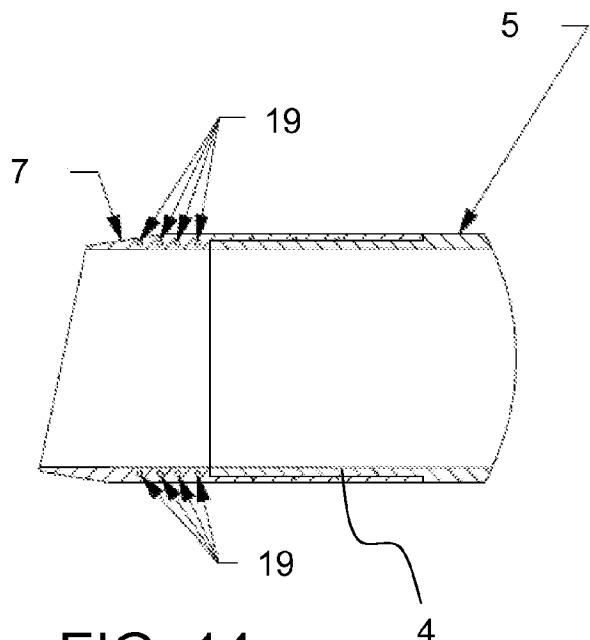


FIG. 14

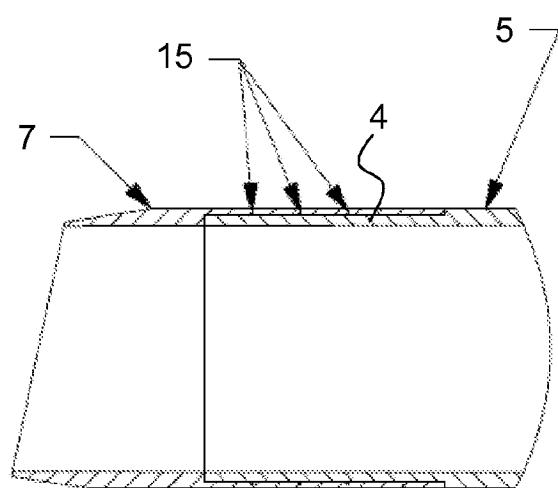


FIG. 9A

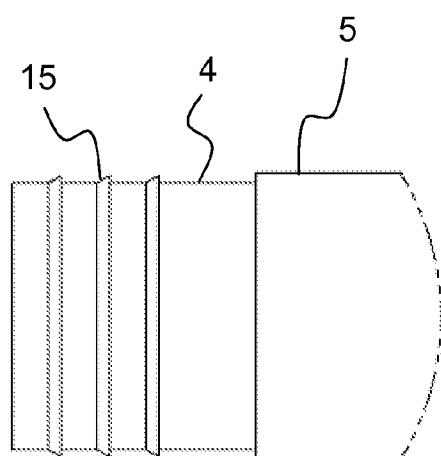


FIG. 9B

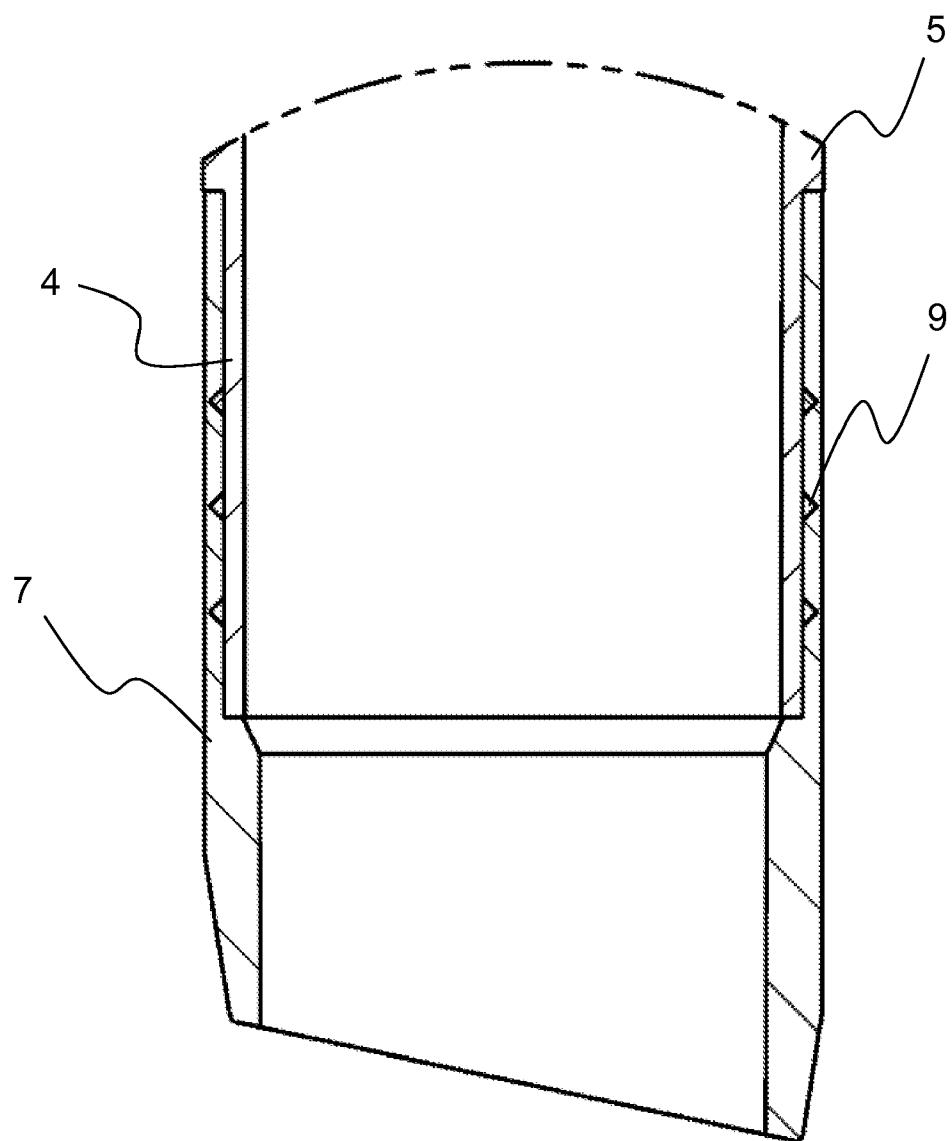


FIG. 15



EUROPEAN SEARCH REPORT

Application Number

EP 12 16 4164

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (IPC)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
X	US 4 795 426 A (JONES JEFFREY S [US]) 3 January 1989 (1989-01-03) * column 2, lines 36-66; claim 1; figure 3 * ----- X US 5 257 973 A (VILLASUSO RAUL [US]) 2 November 1993 (1993-11-02) * columns 3-4; figures 2-3 *----- X US 2005/059934 A1 (WENCHELL THOMAS [US] ET AL) 17 March 2005 (2005-03-17) * paragraphs [0049], [0051], [0077]; figures 2,3 *----- X US 2005/137524 A1 (SAKAL ROBERT [US] ET AL) 23 June 2005 (2005-06-23) * paragraph [0039] *----- X WO 03/088854 A (TYCO HEALTHCARE [US]; JANKOWSKI BRUCE [US]) 30 October 2003 (2003-10-30) * page 7, line 10 - page 9, line 32; claim 1 *-----	1,3-6,9, 10,14 1-7,9,10 1,3,4,9, 10 1,3,4,6, 14,15 1,3-6,9, 10,15	INV. A61B17/34
			TECHNICAL FIELDS SEARCHED (IPC)
			A61B
The present search report has been drawn up for all claims			
3	Place of search Munich	Date of completion of the search 29 May 2012	Examiner Chopinaud, Marjorie
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			
T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

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ON EUROPEAN PATENT APPLICATION NO.**

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This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

29-05-2012

Patent document cited in search report		Publication date	Patent family member(s)			Publication date
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US 5257973	A	02-11-1993	NONE			
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- US 86693906 P [0001]

专利名称(译)	Trocar套管与无创伤尖端		
公开(公告)号	EP2476384A1	公开(公告)日	2012-07-18
申请号	EP2012164164	申请日	2007-11-21
[标]申请(专利权)人(译)	应用医疗资源		
申请(专利权)人(译)	应用医疗资源CORPORATION		
当前申请(专利权)人(译)	应用医疗资源CORPORATION		
[标]发明人	PRAVONGVIENGKHAM KENNII KAHLE HENRY ALBRECHT JEREMY J TAYLOR SCOTT V		
发明人	PRAVONGVIENGKHAM, KENNII KAHLE, HENRY ALBRECHT, JEREMY J TAYLOR, SCOTT V		
IPC分类号	A61B17/34		
CPC分类号	A61B17/3421 A61B17/3494 A61B2017/00473 A61B2017/3441 A61B2090/08021 A61B2017/3456		
优先权	60/866939 2006-11-22 US		
其他公开文献	EP2476384B1		
外部链接	Espacenet		

摘要(译)

外科进入端口设置有套管针套管，该套管针套管具有基本上刚性的部分和无创伤的远侧尖端。无创伤远端尖端相对于套管的刚性部分基本上是柔顺的。套管针套管提供无阻碍的外科手术进入体腔，允许使用最小尺寸的切口或入口通过套管针插管插入和移除手术器械并进入体腔。

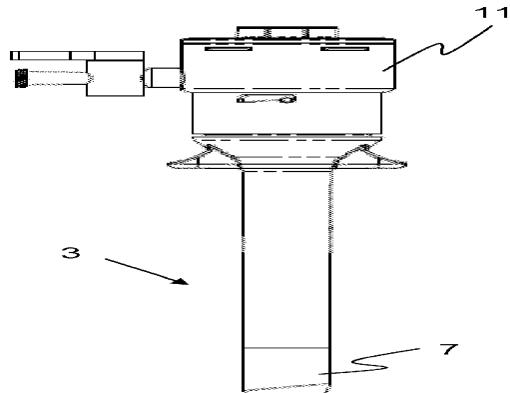


FIG. 1