

US008968247B2

(12) United States Patent Piskun et al.

(10) Patent No.: US 8,968,247 B2 (45) Date of Patent: Mar. 3, 2015

(54) LAPAROSCOPIC INSTRUMENT AND CANNULA ASSEMBLY AND RELATED SURGICAL METHOD

(71) Applicant: Covidien LP, Mansfield, MA (US)

(72) Inventors: **Gregory Piskun**, Morganville, NJ (US); **Anatoly Konik**, Haifa (IL); **Dan**

Rottenberg, Haifa (IL)

(73) Assignee: Covidien LP, Mansfield, MA (US)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35

U.S.C. 154(b) by 2 days.

(21) Appl. No.: 13/913,539

(22) Filed: Jun. 10, 2013

(65) **Prior Publication Data**

US 2013/0281783 A1 Oct. 24, 2013

Related U.S. Application Data

(60) Continuation of application No. 12/779,294, filed on May 13, 2010, now Pat. No. 8,460,271, which is a division of application No. 12/228,028, filed on Aug. 8, 2008, now abandoned, which is a division of application No. 10/895,546, filed on Jul. 21, 2004, now Pat. No. 7,753,901.

(51) **Int. Cl.**A61M 31/00 (2006.01)

A61M 5/32 (2006.01)

(Continued)

(52) U.S. Cl.

 (2013.01); A61B 2017/2929 (2013.01); A61B 2017/3445 (2013.01); A61B 2017/3447 (2013.01); A61B 2017/3449 (2013.01); (Continued)

(58) Field of Classification Search

See application file for complete search history.

(56) References Cited

U.S. PATENT DOCUMENTS

FOREIGN PATENT DOCUMENTS

EP 0567146 10/1993 EP 0592244 4/1994

(Continued)

OTHER PUBLICATIONS

International Search Report for PCT/US05/24636 dated Jun. 20, 2006. (2 pages).

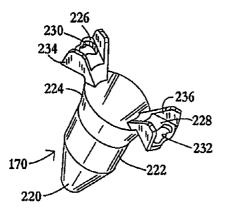
(Continued)

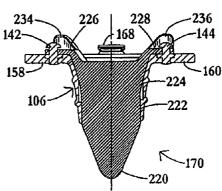
Primary Examiner — Andrew Gilbert

(57) ABSTRACT

A laparoscopic port assembly includes a cannula unit including three cannulas each extending at an acute angle relative to a base. The cannulas are flexible for receiving respective angulated laparoscopic instruments. The cannula unit is rotatingly received in a port holder for rotation about a longitudinal axis of the holder, the holder being disposable in an opening in a patient's skin.

8 Claims, 8 Drawing Sheets





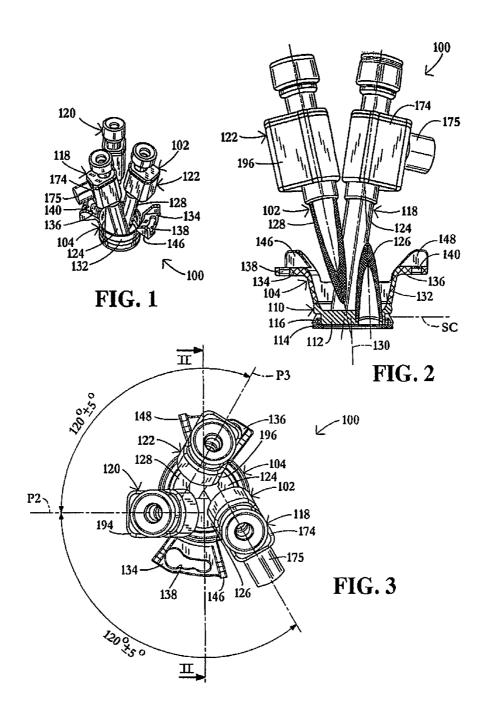
US **8,968,247 B2**Page 2

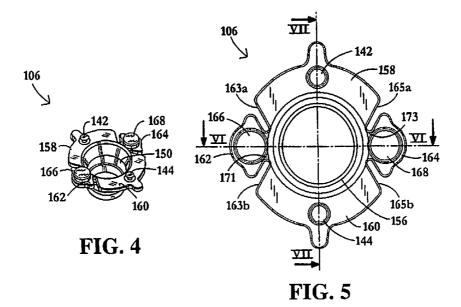
(51) Int. Cl.		6,468,292 B1		Mollenauer et al.
A61M 5/178	(2006.01)	6,488,620 B1		Segermark et al.
A61F 11/00	(2006.01)	6,488,692 B1 6,537,209 B1		Spence et al. Pinkhasik et al.
A61B 17/02	(2006.01)	6,551,270 B1		Bimbo et al.
A61B 17/34	(2006.01)	6,551,276 B1	4/2003	Mann et al.
A61B 17/29	(2006.01)	6,558,371 B2	5/2003	
(52) U.S. Cl.		6,578,577 B2		Bonadio et al. Butler et al.
CPC A61B 2017/3	466 (2013.01); A61B 2017/347	6,582,364 B2 6,623,426 B2		Bonadio et al.
(2013.0	01); <i>A61B 2017/3492</i> (2013.01)	6,669,674 B1		Macoviak et al.
USPC 604/164.	.01; 604/174; 604/513; 606/108	6,676,639 B1		Ternstrom
		6,706,050 B1		Giannadakis
(56) Referen	ices Cited	6,800,084 B2 6,811,532 B2		Davison et al. Ogura et al.
HC DATENT	C DOCK IN CENTER	6,846,287 B2		Bonadio et al.
U.S. PATENT	DOCUMENTS	6,878,110 B2		Yang et al.
5,490,843 A * 2/1996	Hildwein et al 604/164.03	6,890,295 B2		Michels et al.
	Pracas	6,899,673 B2 6,902,569 B2*		Ogura et al 606/108
	Leyva	6,916,331 B2		Mollenauer et al.
5,540,648 A 7/1996		6,945,932 B1		Caldwell et al.
	Williamson, IV Kronner	6,972,026 B1		Caldwell et al.
	Crainich	7,001,397 B2		Davison et al.
5,571,115 A 11/1996	Nicholas	7,008,377 B2 7,056,321 B2		Beane et al. Pagliuca et al.
	Marlow et al.	7,081,089 B2		Bonadio et al.
	Bolanos et al. Pasquallucci et al.	7,195,590 B2		Butler et al.
	Aranyi et al.	7,214,185 B1		Rosney et al.
5,634,911 A 6/1997	Hermann et al.	7,223,257 B2 7,223,278 B2		Shubayev et al. Davison et al.
	Mollenauer et al 606/213	7,225,278 B2 7,235,084 B2		Skakoon et al.
	Tangherlini et al.	7,238,154 B2	7/2007	Ewers et al.
	de la Torre et al. Bishop et al.	7,276,075 B1		Callas et al.
	de la Torre et al.	7,300,399 B2 7,316,699 B2*		Bonadio et al. McFarlane 606/168
	Christy	7,753,901 B2*		Piskun et al 604/539
	Oberlin et al.	7,850,600 B1	12/2010	
	Alexander, Jr. et al. Nicholas et al.	8,377,089 B2*		Lipchitz et al 606/185
· · · · · · · · · · · · · · · · · · ·	Bonadio	8,460,271 B2 *		Piskun et al 604/513
5,810,712 A 9/1998		2002/0038077 A1 2002/0049451 A1		de la Torre et al. Parmer et al.
· · · · · · · · · · · · · · · · · · ·	Flom et al 604/174	2002/0161378 A1*		Downing 606/108
	Hildwein et al 604/175 Wallace et al.	2002/0183594 A1	12/2002	Beane et al.
	Levitsky et al.	2003/0014076 A1		Mollenauer et al.
	Berkelaar	2003/0028179 A1 2003/0036748 A1	2/2003	Cooper et al.
-,,	Hermann et al.	2003/0030748 A1 2003/0040753 A1		Daum et al.
	Gilson	2003/0114832 A1		Kohler et al.
5,906,577 A 5/1999 5,916,147 A 6/1999	Beane et al.	2003/0135091 A1		Nakazawa
5,916,198 A 6/1999	Dillow	2003/0208122 A1 2003/0236549 A1		Melkent Bonadio et al.
5,931,832 A 8/1999	Jensen	2004/0024414 A1*		Downing 606/108
	Segermark et al.	2004/0049099 A1	3/2004	Ewers et al.
· · · · · · · · · · · · · · · · · · ·	de la Torre et al. Mollenauer et al.	2004/0167543 A1		Mazzocchi et al.
	de la Torre et al.	2004/0230100 A1 2005/0137609 A1	6/2005	Shluzas Guiraudon
6,018,094 A 1/2000		2005/0209607 A1*		Lipchitz et al 606/108
	de la Torre et al.	2005/0215863 A1	9/2005	Ravikumar et al.
6,033,378 A 3/2000 6,033,426 A 3/2000	Lundquist et al. Kaji	2005/0222582 A1		Wenchell
	Lucey	2005/0241647 A1 2005/0273133 A1		Nguyen et al. Shluzas et al.
, ,	Shimomura et al.	2005/02/3133 A1 2005/0288558 A1		Ewers et al.
	Termin et al.	2006/0020241 A1		Piskun et al.
	Macoviak et al. Beane et al.	2006/0041188 A1		Dirusso et al.
	Hart et al.	2006/0041232 A1		Stearns et al.
6,217,555 B1 4/2001	Hart et al.	2006/0149306 A1 2006/0192319 A1	8/2006	Hart et al. Solar
	Levinson	2006/0192319 A1 2006/0195119 A1		Mazzocchi et al.
	de la Torre et al.	2006/0241651 A1	10/2006	Wilk
	Kieturakis et al. de la Torre et al.	2006/0247498 A1		Bonadio et al.
	de la Torre et al.	2007/0203398 A1		Bonadio et al.
6,406,472 B1 6/2002	Jensen	2007/0208312 A1		Norton et al.
	Wenner et al.	2007/0270654 A1 2007/0276430 A1		Pignato et al. Lee et al.
, ,	Beane et al. Peterson	2007/0270430 A1 2008/0027476 A1	1/2007	
	Piskun	2008/0103366 A1		Banchieri et al.
6,458,077 B1 10/2002	Boebel et al.	2008/0200767 A1	8/2008	Ewers et al.
6,464,686 B1 10/2002	O'Hara et al.	2008/0255519 A1	10/2008	Piskun et al.

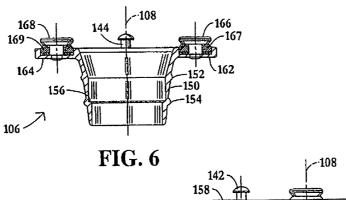
US 8,968,247 B2

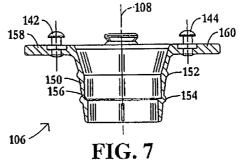
Page 3

(56)	Refere	nces Cited	WO WO	WO 02/17800 WO 03/001969	3/2002 1/2003	
2009/0012477 2010/0081880	7 A1 1/2009 0 A1 4/2010		WO WO WO WO	2006/100658 2006/113216 2008/015566 2008/121294	9/2006 10/2006 2/2008 10/2008	
2010/0113886 F0		Piskun et al 600/231 NT DOCUMENTS	Internat	OTHER Pl	UBLICATIONS PCT/US08/03991 dai	ted Jul. 30, 2008.
EP EP EP WO WO WO WO WO	1312318 1637086 1870043 2044889 93/14801 94/04067 97/42889 99/16368 O 00/18306	5/2003 3/2006 12/2007 4/2009 8/1993 3/1994 11/1997 4/1999 4/2000	pages). Europea pages). Europea	nn Search Report of E. nn Search Report for E nn Search Report for Ap (15 pages).	EP 09252168 dated M	Mar. 19, 2010 (3
WO	02/07611	1/2002	* cited	by examiner		









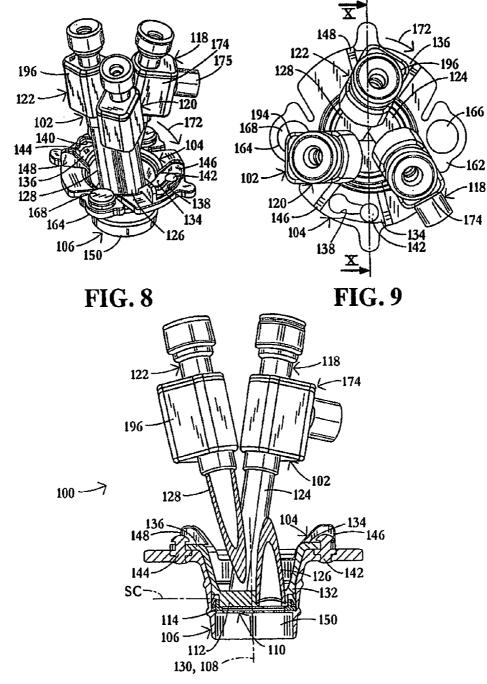


FIG. 10

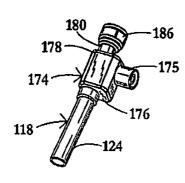


FIG. 11

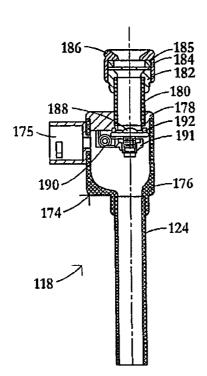


FIG. 13

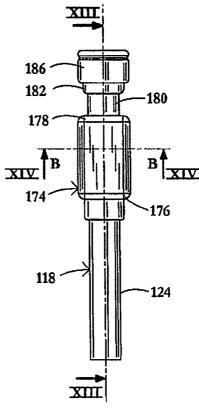


FIG. 12

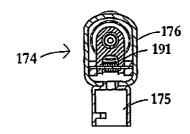


FIG. 14

FIG. 17

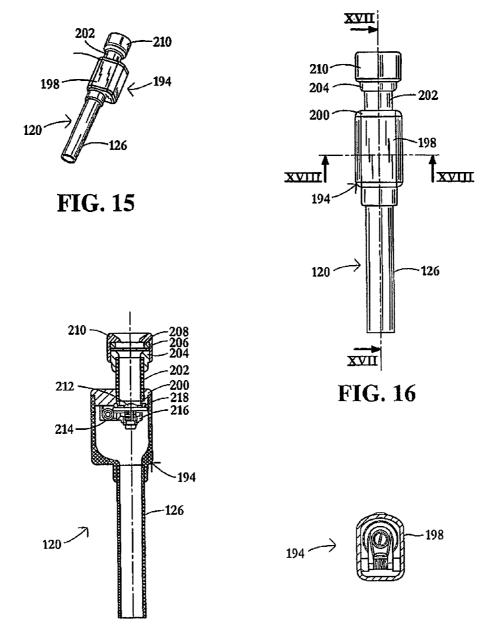
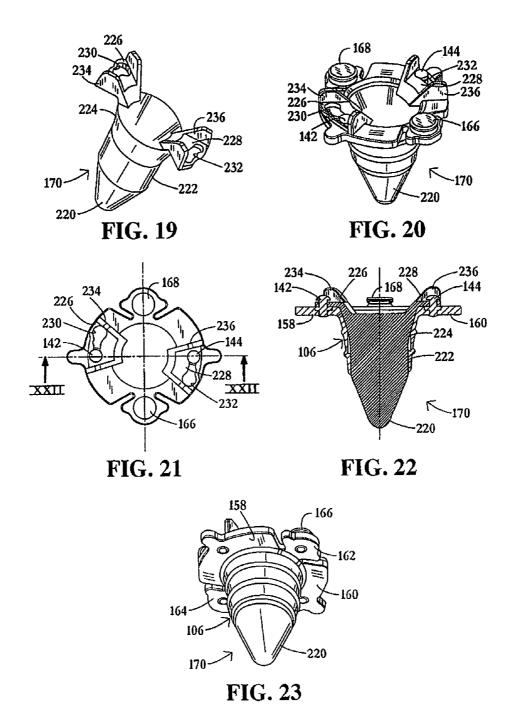
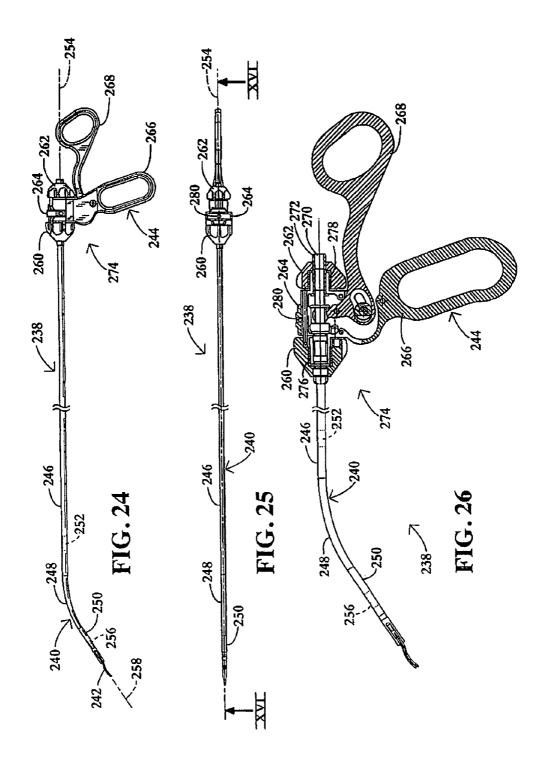
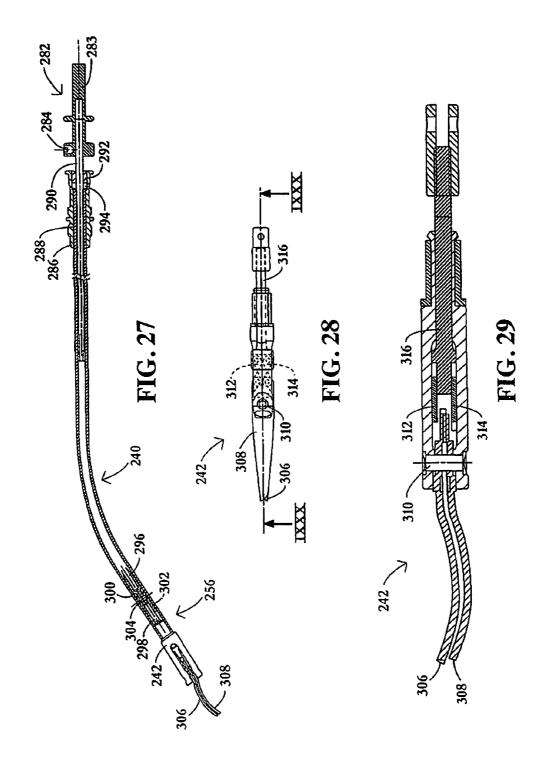


FIG. 18







LAPAROSCOPIC INSTRUMENT AND CANNULA ASSEMBLY AND RELATED SURGICAL METHOD

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation of, and claims priority to. U.S. patent application Ser. No. 12/779,294, filed on May 13, 2010 by Piskun et al, published as U.S. Patent Application 10 Publication No. US 2010/0222643 A1 and now U.S. Pat. No. 8,460,271, which is a divisional of, and claims priority to, U.S. patent application Ser. No. 10/895,546 by Piskun et al. filed on Jul. 21, 2004, now U.S. Pat. No. 7,753,901 issued on Jul. 13, 2010, entitled "LAPAROSCOPIC INSTRUMENT AND CANNULA ASSEMBLY AND RELATED SURGI-CAL METHOD", the entire contents of both of which are hereby incorporated by reference herein. This application incorporates by reference herein in its entirety PCT application PCT/US2005/024111 by Piskun et al. having an international filing date of Jul. 7, 2005 and published as WO 2006/ 019592 A2 "LAPAROSCOPIC INSTRUMENT AND CANNULA ASSEMBLY (sic) AND RELATED SURGI-CAL METHOD" on Feb. 23, 2006.

FIELD OF THE INVENTION

The present invention relates to surgical instruments, surgical port assemblies, and an associated method. The instruments, port assemblies and method are particularly useful in the performance of laparoscopic procedures entirely through the umbilicus.

BACKGROUND OF THE INVENTION

Abdominal laparoscopic surgery gained popularity in the late 1980's, when benefits of laparoscopic removal of the gallbladder over traditional (open) operation became evident. Reduced postoperative recovery time, markedly decreased post-operative pain and wound infection, and improved cosmetic outcome are well established benefits of laparoscopic surgery, derived mainly from the ability of laparoscopic surgeons to perform an operation utilizing smaller incisions of the body cavity wall.

Laparoscopic procedures generally involve insufflation of the abdominal cavity with CO2 gas to a pressure of around 15 mm Hg. The abdominal wall is pierced and a 5-10 mm in diameter straight tubular cannula or trocar sleeve is then inserted into the abdominal cavity. A laparoscopic telescope connected to an operating room monitor is used to visualize 50 the operative field, and is placed through (one of) the trocar sleeve(s). Laparoscopic instruments (graspers, dissectors, scissors, retractors, etc.) are placed through two or more additional trocar sleeves for the manipulations by the surgeon and surgical assistant(s).

Recently, so-called "mini-laparoscopy" has been introduced utilizing 2-3 mm diameter straight trocar sleeves and laparoscopic instruments. When successful, mini-laparoscopy allows further reduction of abdominal wall trauma and improved cosmesis. However, instruments used for mini-laparoscopic procedures are generally more expensive and fragile. Because of their performance limitations, due to their smaller diameter (weak suction-irrigation system, poor durability, decreased video quality), mini-laparoscopic instruments can generally be used only on selected patients with 65 favorable anatomy (thin cavity wall, few adhesions, minimal inflammation, etc.). These patients represent a small percent-

2

age of patients requiring laparoscopic procedure. In addition, smaller, 2-3 mm, incisions may still cause undesirable cosmetic outcomes and wound complications (bleeding, infection, pain, keloid formation, etc.).

Since the benefits of smaller and fewer body cavity incisions are proven, it would be attractive to perform an operation utilizing only a single incision in the navel. An umbilicus is the thinnest and least vascularized, and a well-hidden, area of the abdominal wall. The umbilicus is generally a preferred choice of abdominal cavity entry in laparoscopic procedures. An umbilical incision can be easily enlarged (in order to eviscerate a larger specimen) without significantly compromising cosmesis and without increasing the chances of wound complications. The placement of two or more standard (straight) cannulas and laparoscopic instruments in the umbilicus, next to each other, creates a so-called "chopstick" effect, which describes interference between the surgeon's hands, between the surgeon's hands and the instruments, and between the instruments. This interference greatly reduces the surgeon's ability to perform a described procedure.

Thus, there is a need for instruments and trocar systems, which allow laparoscopic procedures to be performed entirely through the umbilicus while at the same time reducing or eliminating the "chopstick effect". A laparoscopic procedure performed entirely through the umbilicus, using the laparoscopic instruments and trocar system according to an embodiment of the present invention, allows one to accomplish the necessary diagnostic and therapeutic tasks while further minimizing abdominal wall trauma and improving cosmesis.

OBJECTS OF THE INVENTION

The present invention provides instruments and cannula or port assemblies for the performance of surgical procedures, particularly including laparoscopic procedures, for instance, entirely through the umbilicus.

An object of the present invention is to provide an improved port assembly for facilitating access to internal organs of a patient during laproscopic procedures.

Another object of the present invention is to provide such a port assembly that provides enlarged workspace for the hands of the surgeon(s) when plural laparoscopic instruments are placed through the umbilicus.

An additional object of the present invention is to provide improved laparoscopic instruments for facilitating operations through the umbilicus.

These and other objects of the invention will be apparent from the drawings and descriptions herein. Although each object of the invention is believed to be attained by at least one embodiment of the invention, there is not necessarily any single embodiment that achieves all of the objects of the invention.

SUMMARY OF THE INVENTION

The present invention facilitates the performance of laparoscopic surgical procedures wherein several laparoscopic instruments are inserted into a patient through respective cannulas all extending through the same opening in the patient, for instance, through the umbilicus. The advantages of such an operation include minimizing trauma to the patient and accelerating the patient recovery.

A surgical port assembly that facilitates the performance of such a laparoscopic surgical procedure comprises, in accordance with the present invention, a cannula unit including at least one cannula member, and a holder disposable in an

opening in a patient's skin for receiving the cannula component so that the cannula component is movable relative to the holder during a surgical procedure. Preferably, the cannula unit is rotatable with respect to the holder about a longitudinal axis of the holder. It is contemplated that the holder is fastened to the patient during the surgical operation, so that the cannula unit is movable relative to the patient.

Pursuant to another feature of the present invention, the cannula unit comprises a base or frame that is removably attachable to the holder and that defines a closure surface extending, during the surgical procedure, substantially tangentially to the patient's skin at the opening. The cannula member is connected to the base and defines an access path through the closure surface. The cannula member extends at an acute angle relative to the closure surface so that the cannula is inclined relative to the patient's skin surface during the surgical procedure.

Pursuant to further features of the present invention, the cannula is flexible and has a relaxed configuration that is 20 linear. The base or frame is provided with a panel or wall forming the closure surface.

In a particular embodiment of the present invention, the cannula is one of a plurality of cannulas each extending at an acute angle relative to the closure surface so that the cannulas 25 are all inclined relative to the patient's skin surface during the surgical procedure.

Pursuant to additional features of the present invention, the cannula unit is partially insertable into the holder, while the port assembly further comprises a connector member for 30 removably attaching the cannula unit to the holder, the connector also being partially insertable into the holder.

The holder and the connector member may be provided with cooperating locking elements such as projections and slots for reversibly securing the cannula unit to the holder. 35 Thus, after placement of the holder in an opening in the patient (and after removal of an insert assist member from the holder), the base or frame of the cannula unit is inserted into the holder and secured thereto by an insertion and a rotation of the connector so that the projections and slots are interlocked. 40

The base or frame of the cannula unit may include a frustoconical portion insertable into the holder. Similarly, the connector may include a frustoconical portion insertable into the holder to secure the cannula unit to the holder. The holder is provided internally with a shoulder engaging a lower end of 45 the cannula unit.

As indicated above, in one embodiment of the present invention, a surgical port assembly comprises (a) a base or frame seatable in an opening in a patient's skin and defining a closure surface extending substantially tangentially to the 50 patient's skin at the opening during a surgical procedure and (b) a cannula connected to the base and defining an access path through the closure surface. The cannula extends at an acute angle relative to the closure surface so that the cannula is inclined relative to the patient's skin surface during the 55 surgical procedure. The cannula may be one of a plurality of cannulas each extending at an acute angle relative to the closure surface so that the cannulas are all inclined relative to the patient's skin surface during the laparoscopic procedure.

Where the closure surface is located in a main plane, the cannulas have linear configurations, and the base or frame has a longitudinal axis, each of the cannulas may define a respective secondary plane oriented perpendicularly to the main plane, each the secondary plane being spaced from the longitudinal axis. The cannulas may be three in number with the secondary planes disposed at angles of 120° relative to each other

4

A surgical method in accordance with the present invention comprises forming an opening in a patient, inserting a cannula holder through the opening, disposing a plurality of cannulas in the holder so that after inserting of the holder the cannulas traverse the holder and extend from outside the patient to inside the patient, thereafter inserting a plurality of elongate medical instruments through the respective cannulas, and rotating the cannulas and the instruments relative to the holder, about a longitudinal axis of the holder.

In accordance with another aspect of the present invention, where the cannulas are attached to a base member, the disposing of the cannulas in the holder includes inserting the base member into the holder, the base member being in rotatable engagement with the holder. The disposing of the cannulas in the holder may further include attaching a locking member to the holder to maintain the base member in rotatable engagement with the holder.

A laparoscopic medical instrument insertable through a laparoscopic trocar sleeve comprises, in accordance with the present invention, an elongate shaft, an operative tip disposed at one end of the shaft, and a first actuator disposed at an opposite end of the shaft. The actuator is operatively connected to the operative tip via the shaft for controlling the operation of the operative tip. The shaft has a straight proximal end portion, a curved middle portion and a straight distal end portion, the distal end portion extending at an angle with respect to the proximal end portion. The proximal end portion is provided with a first rotary joint so that the distal end portion and the operative tip are rotatable about a longitudinal axis of the proximal end portion. The distal end portion is provided with a second rotary joint so that the operative tip is rotatable about a longitudinal axis of the distal end portion. A second actuator disposed at the end of the shaft opposite the operative tip is operatively connected to the proximal end portion of the instrument shaft for rotating the distal end portion thereof and the operative tip about the longitudinal axis of the proximal end portion of the shaft. A third actuator disposed at the end of the shaft opposite the operative tip is operatively connected to the distal end portion of the shaft for rotating the operative tip about the longitudinal axis of the distal end portion.

The proximal end portion, the middle portion and the distal end portion of the instrument shaft are each substantially rigid throughout so that they cannot be bent. In one embodiment of the invention, the instrument shaft has a hockey-stick shape. Two laparoscopic surgical instruments each having a hockey stick shape are advantageously used in a crossed configuration, which markedly improves the degrees of freedom of the instruments, particularly during lateral (medial-lateral) movements.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a perspective view of a laparoscopic port having multiple cannulas, in accordance with the present invention.

FIG. 2 is a cross-sectional view of the laparoscopic port or cannula assembly of FIG. 1, taken along line II-II in FIG. 3.

FIG. 3 is a top view of the laparoscopic port or cannula assembly of FIG. 1.

FIG. 4 is a perspective view of an annular holder disposable in an opening in a patient for receiving the laparoscopic port or cannula assembly of FIGS. 1-3.

FIG. 5 is a top plan view of the port holder of FIG. 4.

FIG. 6 is a longitudinal cross-sectional view of the port holder of FIGS. 4 and 5, taken along line VI-VI in FIG. 5.

FIG. 7 is another longitudinal cross-sectional view of the port holder of FIGS. 4 and 5, taken along line VII-VII in FIG. 5

FIG. **8** is a perspective view of the laparoscopic port or cannula assembly of FIGS. **1-3**, together with the port holder of FIGS. **4-7**, showing the laparoscopic port or cannula assembly inserted into and attached to the port holder.

FIG. 9 is a top plan view of the laparoscopic port or cannula assembly of FIGS. 1-3 connected to the port holder of FIGS. 4-7, as shown in FIG. 8.

FIG. 10 is a longitudinal cross-sectional view taken along line X-X in FIG. 9.

FIG. 11 is a perspective view of a cannula with an insufflation valve, included in the laparoscopic port or cannula assembly of FIGS. 1-3 and 8-10.

FIG. 12 is a side elevational view of the cannula of FIG. 11, on a larger scale.

FIG. 13 is a longitudinal cross-sectional view of the cannula of FIGS. 11 and 12, taken along line XIII-XIII in FIG. 12.

FIG. 14 is a transverse cross-sectional view of the cannula of FIGS. 11 and 12, taken along line XIV-XIV in FIG. 12.

FIG. 15 is a perspective view of a cannula without an insufflation valve, included in the laparoscopic port or cannula assembly of FIGS. 1-3 and 8-10.

FIG. 16 is a side elevational view of the cannula of FIG. 16, on a larger scale.

FIG. 17 is a longitudinal cross-sectional view of the cannula of FIGS. 15 and 16, taken along line XVII-XVII in FIG. 16

FIG. **18** is a transverse cross-sectional view of the cannula of FIGS. **15** and **16**, taken along line XVIII-XVIII in FIG. **16**.

FIG. 19 is a perspective view of an insertion plug used to facilitate insertion of the port holder of FIGS. 4-7 in a patient at the beginning of a laparoscopic procedure.

FIG. **20** is a top perspective view of the insertion plug of FIG. **19** temporarily inserted in and attached to the port holder of FIGS. **4-7**.

FIG. 21 is a top plan view of the assembled insertion plug and port holder of FIG. 20.

FIG. 22 is a longitudinal cross-section taken along line XXII-XXII in FIG. 21.

FIG. 23 is a bottom perspective view of the assembled insertion plug and port holder of FIGS. 20-22.

FIG. **24** is a side elevational view of a laparoscopic instru- ⁴⁵ ment utilizable with the multiple-cannula port assembly of FIGS. **8-10**, in accordance with the present invention.

FIG. 25 is a top plan view of the laparoscopic instrument of FIG. 24.

FIG. ${\bf 26}$ is partially a side elevational view and partially a $\,$ 50 cross-sectional view

FIG. 27 is a partial longitudinal cross-sectional view also taken along line XXVI-XXVI in FIG. 25.

FIG. **28** is a longitudinal cross-sectional view, on a larger scale, of a distal end of the laparoscopic instrument of FIGS. 55 **24-27**.

FIG. 29 is a longitudinal cross-sectional view, on an even larger scale, taken along line IXXX-IXXX in FIG. 28.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

As depicted in FIGS. 1-3, a laparoscopic port or cannula assembly 100 comprises a cannula unit 102 and a connector 104 associated therewith for removably fastening the cannula 65 unit to an annular port holder 106 (FIGS. 4-8 and 10) that is disposed in an opening (e.g., formed in the umbilicus) in a

6

patient. Cannula unit 102 is coupled to port holder 106 by connector 104 so as to permit rotation of cannula unit 102 about a longitudinal axis 108 (FIGS. 7 and 10) of holder 106. Holder 106 has an inner side facing the patient and an outer side facing outside the patient during a laparoscopic procedure, and axis 108 traverses the holder from the inner side to the outer side thereof.

Cannula unit 102 comprises a base or frame 110 that is insertable into and removably attachable to port holder 106. Base or frame 110 includes a base member in the form of a planar panel or wall 112 that defines a closure surface or plane SC extending, during a laparoscopic surgical procedure, substantially tangentially and accordingly parallel to the patient's skin at the opening through with port holder 106 extends. Base or frame 110 further includes a seating ring 114 and a sealing ring 116.

Cannula unit 102 additionally comprises three cannula members 118, 120, 122 each connected to base or frame 110 and defining a respective access path through closure surface SC. Cannulas 118, 120, 122 are each connected at one end to base 110 and particularly panel or wall 112 at mutually spaced locations so as to enable simultaneous insertion of multiple instruments through closure surface SC at such locations, the cannulas each extending away from closure surface 25 SC so as to be located on only an outer side thereof opposite the patient during a surgical procedure. Cannula members 118, 120, 122 have outer surfaces (not separately designated) that are contiguous with, and stationary with respect to, planar panel or wall 112 at points of passage of the cannulas through the panel or wall. When cannula unit 102 and holder 106 are connected together (see FIG. 10), cannula members 118, 120, 122 extend into and are partially surrounded by said holder 106 and enable disposition of distal ends of three separate instruments in the patient at an opening or incision in the patient. Cannula members 118, 120, 122 each extend at an acute angle relative to closure surface SC so that the cannulas are inclined relative to the patient's skin surface during a laparoscopic surgical procedure. Cannula members 118, 120, 122 include flexible tubular portions 124, 126, 128 that have linear configurations in a relaxed or unstressed condition.

Each cannula member 118, 120, 122 defines a respective plane P1, P2, P3 (FIG. 3) oriented perpendicularly to closure surface or plane SC (the main plane) and spaced from a longitudinal axis 130 of base or frame 110. These secondary planes P1, P2, and P3 are disposed at angles of 120° relative to each other, as indicated in FIG. 3. Connector 104 is loosely coupled to cannula unit 102 so as to be freely movable along axis 130 of the cannula unit, between base 110 and valve components of cannulas 118, 120, 122.

Connector 104 includes a frustoconical portion 132 insertable into port holder 106 (see FIG. 10) and further includes a pair of flanges 134, 136 for temporarily locking cannula unit 102 to holder 106. To that end, flanges 134, 136 are provided with dual-lobed slots 138, 140 for receiving respective pins or projections 142, 144 on port holder 106 (see FIGS. 4, 5, 7, 8, 9). Flanges 134 and 136 are also provided with respective pairs of upturned ears 146 and 148 functioning in part as thumb and finger rests for swiveling connector 104 about axis 108 (and 130) to reversibly secure connector 106 and concomitantly cannula unit 102 to port holder 106.

As illustrated in FIGS. 4-7, port holder 106 includes an annular peripheral wall in the form of a tapered, slightly frustoconical sleeve 150 provided along an outer surface with a pair of annular beads or ribs 152 and 154 and along an inner surface with a shoulder 156 that serves as an abutment or rest for seating ring 114 of cannula unit 102. Peripheral wall or sleeve 150 is an outermost portion of holder 106 and is in

contact with tissues of said patient upon a disposition of the holder in an opening of the patient. At a wider end of sleeve 150, holder 106 includes a pair of diametrically opposed flanges 158, 160 and a pair of diametrically opposed ears 162, 164. Pins or projections 142, 144 are rigid with flanges 158, 5160, while ears 162, 164 carry respective flat-headed posts 166, 168 around which sutures are wound to fasten holder 106 to the skin of the patient. Rubber gaskets 167 and 169 may be provided for clamping suture threads to posts 166 and 168. Alternatively or additionally, sutures anchoring port holder 106 to the body wall of the patient may be inserted through slots 163a, 163b and 165a, 165b and partially wrapped around bases 171 and 173 of ears 162, 164 (see FIG. 5).

After placement of holder 106 in an opening in the patient (and after removal of an insert assist member 170, FIGS. 15 19-22, from the holder), base or frame 110 of cannula unit 102 is inserted into holder 106 until seating ring 114 engages shoulder 15 (see FIG. 10). Cannula unit 110 is secured to holder 106 by an insertion and a subsequent rotation of connector 104 relative to holder 106, as indicated by an arrow 172 oin FIGS. 8 and 9, so that projections 142, 144 and slots 138, 140 are interlocked (see FIGS. 8-10). After this locking of connector 104 to holder 106, cannula unit 102 is rotatable about axes 108 and 130 in opposition to a frictional drag force exerted by virtue of sealing ring 116.

As illustrated in FIGS. 11-14, cannula unit 118 includes valve component 174 connected to tubular portion 124. Valve component 124 includes an insufflation port 175 for receiving a tube (not shown) for guiding carbon dioxide gas from a pressurized source into the patient. As shown particularly in FIG. 13, valve component 124 of cannula member 118 includes a valve box or casing 176 with a cover or closure 178 to which an extension tube 180 is attached. At an end opposite valve casing 176, extension tube 180 is provided with a sleeve 182, a valve seal 184, a disc 185, and a cap 186. Valve 35 component 174 further includes a valve door 188 that is biased into a closure position shown in FIG. 13 by a helical or coil spring 190. Door 188 is supported by a mounting bracket and associated hardware 191. An O-ring seal 192 is provided for inhibiting the escape of insufflation gas from a patient 40 through extension tube 124 when a laparoscopic surgical instrument does not traverse cannula member 118.

As illustrated in FIGS. 1-3 and 8-10, cannula members 120 and 122 include respective valve components 194 and 196 connected to respective flexible tubular portions 126 and 128. 45 These valve components are structurally identical, a representative component 194 being depicted in FIGS. 15-18. Valve component 194 includes a valve box or casing 198 with a cover or closure 200 to which an extension tube 202 is attached. At an end opposite valve casing 198, extension tube 50 202 is provided with a sleeve 204, a valve seal 206, a disc 208, and a cap 210. Valve component 194 further includes a valve door 212 that is biased into a closure position shown in FIG. 17 by a helical or coil spring 214. Door 212 is supported by a mounting bracket and associated hardware 216. An O-ring 55 seal 218 is provided for inhibiting the escape of insufflation gas from a patient through extension tube 202 when a laparoscopic surgical instrument does not traverse the respective cannula member 120 (or 122).

As shown in FIG. 19, insert assist member 170 includes a 60 rounded conical tip 220, a cylindrical middle portion 222 and a slightly tapered or frustoconical outer portion 224. Outer portion 224 is provided with a pair of flanges 226, 228 for temporarily locking insert assist member 170 to port holder 106. To that end, flanges 226, 228 are provided with duallobed slots 230, 232 for receiving respective pins or projections 142, 144 on port holder 106, as depicted in FIGS. 20-22.

8

Flanges 226 and 228 are formed with respective pairs of upturned ears 234 and 236 that are manually engageable by a user to reversibly secure connector insert assist member 170 to port holder 106.

After a small incision or opening is made in a patient, port holder 106 with insert assist member 170 connected thereto is inserted through the incision. Sutures (not shown) are stitched to the patient and are wound around and tied to posts 166, 168 to firmly secure the port holder 106 to the patient. Insert assist member 170 is then removed, by a reverse rotation unlocking flanges 226, 228 from pins or projections 142, 144 and by separating the insert assist member from holder 106. Cannula unit 102 is then attached to holder 106 as described above.

FIGS. 24-29 depict a laparoscopic surgical instrument 238 insertable through a laparocopic trocar sleeve or cannula such as cannula member 118, 120, or 122 of the port assembly of FIGS. 1-3 and 8-10 for executing a laparoscopic surgical operation. Instrument 238 comprises an elongate shaft 240, an operative tip 242 disposed at one end of the shaft, and a hand-grip-type actuator 244 disposed at an opposite end of the shaft. Actuator 244 is operatively connected to operative tip 242 via shaft 240 for controlling the operation of the operative tip.

Shaft 240 has a straight proximal end portion 246, a curved 25 middle portion 248 and a straight distal end portion 250, the distal end portion extending at a non-zero angle with respect to the proximal end portion, as shown in FIGS. 24, 26 and 27. Proximal end portion 246 is provided with a first rotary joint 252 so that distal end portion 250 and operative tip 242 are rotatable about a longitudinal axis 254 of proximal end portion 246. Distal end portion 250 is provided with a second rotary joint 256 so that operative tip 242 is rotatable about a longitudinal axis 258 of the distal end portion. A rotary actuator or knob 260 disposed at the proximal end of instrument 238 is operatively connected to proximal end portion 246 of instrument 240 for rotating distal end portion 250 and operative tip 242 about longitudinal axis 254. Another rotary actuator or knob 262 disposed at the proximal end of instrument 238 is operatively connected to distal end portion 250 of shaft 240 for rotating operative tip 242 about longitudinal axis 258.

Proximal end portion 246, middle portion 248 and distal end portion 250 of instrument shaft 240 are each substantially rigid throughout and can only be rotated about joints 252 and 256 and not bent. The angle between axes 254 and 258 are such that shaft 240 has a shape reminiscent of a hockey stick. In an alternative embodiment of instrument 238, middle portion 248 of shaft 240 may be flexible to permit shaft 240 to alternately assume a linear configuration and the hockey-stick configuration of FIGS. 24 and 26. In that case, a handle assembly 274 is provided with an actuator (not shown) for enabling a bending of middle portion 248.

As shown in FIGS. 24-26, instrument 238 may be provided with further actuators, such as a slidable toggle switch 264, for example for performing a locking function or inducing a pivoting of operative tip 242 about an axis perpendicular to axis 258

Actuator 244 includes a hand grip member 266 fixed relative to shaft 240 and further includes a pivotable hand grip 268. A proximal end 270 of shaft 240 is journaled in a bearing 272 about which rotary knob 262. Actuator 244, rotary knobs 260 and 262, and toggle switch 264 are parts of a handle assembly 274 also incorporating yokes 276 and 278, a stopper pin 278, a set screw 280.

FIG. 27 illustrates further parts of a shaft assembly 282 including shaft 240, a slider member 283, a socket set screw 284, an outer bearing 286, a motion bar 288, a bend tube 290, a ring 292, and an O-ring seal 294.

Q

Rotary joint 256 is representative of joint 254 and comprises, as shown in FIG. 27, a proximal pin or inner shaft portion 296, a distal pin or inner shaft portion 298, a pair of coupling elements 300 and 302, and a transverse connector pin 304.

As illustrated in FIGS. 28 and 29, operative tip 242 exemplarily includes a pair of jaws 306 and 308 pivotably connected to a distal end of distal end portion 250 via a pivot pin 310. Jaws 306 and 308 are rotatable about pin 310 through the action of levers or arms 312, 314 that pivot in response to a 10 longitudinal motion of a tip rod 316.

After a deployment of cannula unit 102 in a patient as described above, operative tip 242 and shaft 240 of instrument 238 are insertable through a cannula member 118, 120, or 122, with the respective tubular portion 124, 126, or 128 15 bending to accommodate the bent shaft 240. The bent shape of shaft 240, as well as the rotary joints 252 and 256 facilitate the performance of laparoscopic surgical procedure using multiple laparoscopic instruments extending through a single opening in a patient, for instance, in the umbilicus. Such a 20 procedure involves the rotation of distal end portion 250 and operative tip 242 together about axis 254 and the rotation of operative tip about axis 258. In addition, the entire instrument assembly including cannula unit 102 and multiple instruments 238 can be rotated about collinear axes 108 and 130, to 25 optimize the simultaneous or successive access of multiple operative tips 242 to a surgical site inside a patient.

Although the invention has been described in terms of particular embodiments and applications, one of ordinary skill in the art, in light of this teaching, can generate additional 30 embodiments and modifications without departing from the spirit of or exceeding the scope of the claimed invention. Accordingly, it is to be understood that the drawings and descriptions herein are proffered by way of example to facilitate comprehension of the invention and should not be construed to limit the scope thereof.

What is claimed is:

- 1. An insert assist member for inserting a port holder in an opening at a surgical site comprising:
 - a proximal outer portion;
 - a cylindrical middle portion extending distally from the proximal outer portion; and
 - a distal tip portion extending distally from the cylindrical middle portion, the distal tip portion defining a rounded tip for insertion through an aperture defined in a port 45 holder configured for insertion in said opening at a surgical site; and,
 - wherein the proximal outer portion includes a pair of flange members each configured to define an aperture therethrough, the pair of flange members configured such that

10

the apertures receive and rotatably engage with corresponding projections extending proximally from the port holder.

- 2. The insert assist member according to claim 1, wherein the pair of flange members are configured such that the apertures disengage from the projections upon rotation of the pair of flange members in a direction opposite the direction in which the apertures engage with the projections.
- 3. The insert assist member according to claim 2, wherein the insert assist member is removable from the port holder upon rotation of the pair of flange members in the direction opposite the direction in which the apertures engage with the projections.
- **4**. The insert assist member according to claim **1**, wherein the apertures are each a dual-lobed slot configured to receive a corresponding projection extending proximally from the port holder.
- 5. The insert assist member according to claim 1, wherein the pair of flange members each are configured with respective pairs of upturned ears that are manually engageable to reversibly secure the insert assist member to the port holder.
- **6**. A method of inserting a port holder in an opening at a surgical site comprising:
 - providing an insert assist member configured to at least partially fit within the port holder to form a port holder and insert assist member assembly, the insert assist member having an atraumatic distal tip;
 - positioning the insert assist member within the port holder such that the atraumatic distal tip extends distally beyond a distal end of the port holder; and
 - inserting the port holder and insert assist member assembly in the opening; and,
 - wherein the port holder includes at least one proximally extending projection and said insert assist member defines an aperture for receiving the at least one projection, the step of positioning the insert assist member within the port holder further including aligning the port holder and insert assist member to place the projection into the aperture.
- 7. The method of inserting a port holder according to claim 6, further comprising securing the port holder to the surgical site via winding of sutures around posts configured and disposed in the port holder to enable one or more sutures to be wound around the posts to enable the port holder to be fastened to the surgical site.
- **8**. The method of inserting a port holder according to claim **6**, further comprising disengaging the insert assist member proximally from the port holder.

* * * * *



专利名称(译)	腹腔镜仪器和套管组件及相关手术	方法			
公开(公告)号	US8968247	公开(公告)日	2015-03-03		
申请号	US13/913539	申请日	2013-06-10		
[标]申请(专利权)人(译)	柯惠有限合伙公司				
申请(专利权)人(译)	COVIDIEN LP				
当前申请(专利权)人(译)	COVIDIEN LP				
[标]发明人	PISKUN GREGORY KONIK ANATOLY ROTTENBERG DAN				
发明人	PISKUN, GREGORY KONIK, ANATOLY ROTTENBERG, DAN				
IPC分类号	A61M31/00 A61M5/32 A61F11/00 A61M5/178 A61B17/02 A61B17/29 A61B17/34				
CPC分类号	A61B17/0218 A61B17/3403 A61B17/3421 A61B17/3423 A61B17/3462 A61B17/2909 A61B17/3498 A61B2017/2904 A61B2017/2905 A61B2017/292 A61B2017/2929 A61B2017/3445 A61B2017/3447 A61B2017/3449 A61B2017/3466 A61B2017/347 A61B2017/3449 A61B2017/3466 A61B2017/347 A61B2017/3492 A61B2017/3419				
审查员(译)	GILBERT , ANDREW				
优先权	10/895546 2010-07-13 US				
其他公开文献	US20130281783A1				
外部链接	Espacenet USPTO				

摘要(译)

腹腔镜端口组件包括套管单元,套管单元包括三个套管,每个套管相对 于基座以锐角延伸。套管是柔性的,用于接收相应的成角度腹腔镜器 械。套管单元可旋转地容纳在端口保持器中,以围绕保持器的纵向轴线 旋转,保持器可置于患者皮肤的开口中。

