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(54) **ENDOSCOPIC STAPLING DEVICE**

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Description

FIELD OF THE INVENTION

[0001] The present invention relates generally to the field of systems and methods for performing endoscopic surgery, and specifically to systems and methods for endoscopic stapling of tissue within body cavities.

BACKGROUND OF THE INVENTION

[0002] An anatomical view of a human stomach S and associated features is shown in Fig. 1A. The esophagus E delivers food from the mouth to the proximal portion of the stomach S. The z-line or gastro-esophageal junction Z is the irregularly-shaped border between the thin tissue of the esophagus and the thicker tissue of the stomach wall. The gastro-esophageal junction region G is the region encompassing the distal portion of the esophagus E, the z-line, and the proximal portion of the stomach S.

[0003] Stomach S includes a fundus F at its proximal end and an antrum A at its distal end. Antrum A feeds into the pylorus P which attaches to the duodenum D, the proximal region of the small intestine. Within the pylorus P is a sphincter that prevents backflow of food from the duodenum D into the stomach. The middle region of the small intestine, positioned distally of the duodenum D, is the jejunum J.

[0004] Fig. 1B illustrates the tissue layers forming the stomach wall. The outermost layer is the serosal layer or "serosa" S and the innermost layer, lining the stomach interior, is the mucosal layer or "mucosa" MUC. The sub-mucosa SM and the multi-layer muscularis M lie between the mucosa and the serosa.

[0005] There are a number of applications for endoscopic application of fasteners such as staples to tissue within a body cavity. Some of those applications involve forming tissue structures such as plications or folds in tissue of the body cavity.

[0006] Several prior applications, including International Application No. WO 2005/037152 having an international filing date of October 8, 2004 and U.S. Application US 2008065122 describe methods according to which medical implants are coupled to tissue structures formed within the stomach. According to these applications, devices for inducing weight loss (e.g. by restricting and/or obstructing flow of food into the stomach, and/or by occupying a portion of the stomach volume) may be coupled to tissue tunnels or plications formed from stomach tissue.

[0007] For example, US 2008065122 describes a restrictive and/or obstructive implant system for inducing weight loss. In one embodiment, flexible loops are coupled to tissue plications formed in the gastroesophageal junction region of the stomach. An implant, such as a flow restrictive and/or obstructive implant, is passed through the loops 2 and thus retained in the stomach.

[0008] In other instances, tissue plications may them-

selves be sufficient to provide the necessary treatment. For example, the plications may be used to reduce stomach volume or form a flow restriction within the stomach as disclosed in WO 2005/037152, and in US2007/0219571, upon which the two part form of appended claim 1 is based.

[0009] Other types of implants may be coupled to such plications or other tissue structures for a variety of purposes. These implants include, but are not limited to prosthetic valves for the treatment of gastro-esophageal reflux disease, gastric stimulators, pH monitors and drug eluting devices that release drugs, biologics or cells into the stomach or elsewhere in the GI tract. Such drug eluting devices might include those which release leptin (a hormone which creates feelings of satiety), Ghrelin (a hormone which creates feelings of hunger), octreotide (which reduces Ghrelin levels and thus reduces hunger), Insulin, chemotherapeutic agents, natural biologics (e.g. growth factor, cytokines) which aid in post surgery trauma, ulcers, lacerations etc. Still other implants might be of a type which might provide a platform to which specific cell types can adhere, grow and provide biologically-active gene products to the GI tract, and/or a platform for radiation sources that can provide a local source of radiation for therapeutic purposes, or provide a platform whereby diagnostic ligands are immobilized and used to sample the GI tract for evidence of specific normal or pathological conditions, or provide an anchor point for imaging the GI tract via cameras and other image collecting devices.

[0010] The prior applications listed above, address the desirability of forming tissue plications, pockets or tunnels in a way that regions of serosal tissue (i.e. the tissue on the exterior surface of the stomach) are retained in contact with one another. Over time, adhesions formed between the opposed serosal layers create strong bonds that can facilitate retention of the plication/pocket/tissue over extended durations, despite the forces imparted on them by stomach movement and implanted devices.

[0011] Regardless of the application for which a plication is being formed, it is highly desirable to form that plication using steps carried out from within the stomach using instruments passed down the esophagus, rather than using more invasive surgical or laparoscopic methods. The present application describes endoscopic staplers which may be passed transorally into the stomach and used to form serosal-to-serosal plications in a stomach wall.

SUMMARY OF THE INVENTION

[0012] The invention is a stapler device (12) for applying a staple to a tissue as set out in appended claim 1. Preferred embodiments are defined by the dependent claims.

[0013] Also disclosed is a method for capturing a tissue fold by the steps of:

(a) drawing tissue (17) into a vacuum chamber (21) defined by first and second relatively movable members (25, 27) and a membrane, such as an elastomeric or pleated membrane (24), extending between the two,
 (b) advancing a first tissue-contact plate (78) and a second tissue-contact plate (96) contained within the first and second members, respectively, toward one another by independent movement of the first and second tissue-contact plates (78, 96) staple within their associated members (25, 27, respectively, and by moving the second member toward the first member;
 (c) by continued application of vacuum during said advancing, continuing to draw tissue into said chamber until a tissue fold (17a) is captured between the first and second tissue-contact plates.

[0014] The features of the invention will become more fully apparent when the following detailed description of the invention is read in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

[0015] Fig. 1A is a schematic illustration of a human stomach and a portion of the small intestine, as known in the prior art.

[0016] Fig. 1B is a cross-sectional perspective view of a portion of a stomach wall, illustrating the layers of tissue forming the wall, also as known in the prior art.

[0017] Fig. 2 illustrates an endoscopic stapling system or instrument constructed in accordance with an embodiment of the invention.

[0018] Figs. 3A - 3C are perspective views showing the stapler head or device of the stapling system of Fig. 2 in three different positions.

[0019] Fig. 4 is a perspective view of the stapler head or device, with the membrane removed, showing first and second members in the device.

[0020] Fig. 5 is a perspective view of the proximal end of the staple housing of the stapler device of Fig. 4.

[0021] Fig. 6 is a perspective view of the distal end of the staple housing of the stapler device of Fig. 4.

[0022] Fig. 7 is an exploded perspective view showing elements advanceable within the staple housing during compression and stapling operations.

[0023] Fig. 8 is a plan view of a staple reinforcement device.

[0024] Fig. 9 is a side elevation view of a staple cartridge.

[0025] Fig. 10 is a perspective view of the staple housing similar to Fig. 6, but showing some of the elements of Fig. 7 within the housing.

[0026] Figs. 11A - 11D are a series of schematic representations of the hydraulic chamber and pistons, illustrating operation of an exemplary hydraulic system during tissue compression and stapling.

[0027] Fig. 11E is similar to Fig. 11D and shows an alternative piston configuration.

[0028] Fig. 12 is a perspective view of the anvil housing of the stapler head of Fig. 4.

5 **[0029]** Fig. 13 is a perspective view of the anvil support.

[0030] Fig. 14 is a plan view of the anvil.

[0031] Fig. 15A is a cross-sectional side view of the cutting device and a first embodiment of a cutting board.

10 **[0032]** Fig. 15B is a cross-sectional side view of the cutting device and a second embodiment of a cutting board.

[0033] Fig. 16 is a perspective view of the hinged arm assemblies of the stapler head of Fig. 4.

15 **[0034]** Fig. 17 is a top plan view of the stapler head of Fig. 4 in the streamlined position for introduction into the body. Both the membrane and the membrane raiser are not shown for purposes of clarity.

[0035] Fig. 18 is similar to Fig. 17 and illustrates hidden features of Fig. 17.

20 **[0036]** Fig. 19 is a perspective view of the stapler head in an intermediate, partially expanded, position.

[0037] Fig. 20 is a plan view similar to Fig. 17 but showing the stapler head in the intermediate position.

25 **[0038]** Fig. 21 is similar to Fig. 20 and illustrates hidden features of Fig. 20.

[0039] Fig. 22 is a perspective view of the stapler head in a fully expanded, full compression position.

[0040] Fig. 23 is a plan view similar to Fig. 20 but showing the stapler head in the full compression position.

30 **[0041]** Fig. 24 is similar to Fig. 23 and illustrates hidden features of Fig. 24.

[0042] Figs. 25A - 25B are perspective views showing the staple housing, cartridge and a portion of the membrane raiser. These figures illustrate the steps of detaching a staple cartridge from the staple housing.

35 **[0043]** Fig. 26 is a perspective view of the stapler instrument of Fig. 2, with the staple head removed.

[0044] Fig. 27A is a plan view of the articulating section of the stapler of Fig. 2, showing interleaved drive fluid lines.

40 **[0045]** Fig. 27B shows a drive fluid line having an alternate longitudinally expandable shape.

[0046] Fig. 28 is a cross-sectional side view of the handle of the stapler instrument of Fig. 2.

45 **[0047]** Fig. 29 is a perspective view of the handle of the stapler of Fig. 2.

[0048] Figs. 30A and 30B are plan views of the proximal face of the staple housing, showing a method for attaching the end plate of the stapler handle to the staple housing.

50 **[0049]** Figs. 31A - 31E are a series of drawings schematically illustrating use of the system of Fig. 2 to form a plication in a stomach.

55 **[0050]** Figs. 32A-32C are a series of perspective views illustrating use of the stapler of Fig. 2 to acquire, compress, and then staple stomach wall tissue to form a plication in the stomach. The membrane is not shown in these drawings.

[0051] Fig. 33 is a top plan view of a plication formed in body tissue.

[0052] Figs. 34 and 35 are perspective views of an alternative stapler head equipped to carry additional tools.

[0053] Figs. 36 and 37 show alternative embodiments of stapler-alignment structure in the stapler device of the invention.

DETAILED DESCRIPTION OF THE DRAWINGS

[0054] The present application describes endoscopic fastener-applying devices which in preferred embodiments may be passed transorally into the stomach and used to plicate stomach tissue.

[0055] In the disclosed embodiments, tissue is drawn inwardly into a vacuum chamber, although tissue may be drawn inwardly using other components (e.g. graspers) that do not involve the use of a vacuum. When a portion the interior stomach wall is drawn inwardly, sections of serosal tissue on the exterior of the stomach are positioned facing one another. The disclosed fastener applying device allows the opposed sections of tissue to be moved into contact with one another, and delivers fasteners that will hold the tissue sections together until at least such time as serosal bonds form between them. Each of these steps may be performed wholly from the inside of the stomach and thus can eliminate the need for any surgical or laparoscopic intervention. After one or more plications is formed, medical devices (including, but not limited to any of the types listed above) may be coupled to the plication(s) for retention within the stomach.

[0056] The disclosed embodiments include an optional feature that forms a hole or cut in a plication using the fastener-applying device. This hole or cut might be formed so that a portion of a medical implant may be passed through or linked to the hole/cut, or it may be formed so as to provoke a healing response that will contribute to the strength of the resulting tissue bond.

[0057] In the description of the embodiments given below, the fastener-applying devices are described as being staplers, and exemplary methods are given with respect to the formation of plications in stomach tissue. It should be understood, however, that the embodiments described herein include features having equal applicability for applying other types of fasteners, and for applying staples or other fasteners for purposes other than formation of plications. More specifically, the term "staple" is used herein to designate any type of fastener that (i) can be pushed through tissue, and (ii) has one or more leg members that when forced against an anvil are crimped to secure the fastener to the tissue and hold tissue fastened tissue fold together. The disclosed embodiments and methods will also find use in parts of the body outside the GI system. Additionally, although the disclosed embodiment features circular stapling and cutting of a concentric hole, modifications are conceivable

in which linear stapling can be accomplished, as well as circular or linear stapling without cutting.

[0058] Fig. 2 illustrates one embodiment of a system or instrument 10 for tissue stapling that is suitable for endoscopic use, as well as surgical or laparoscopic use if desired.

[0059] Generally speaking, system 12 includes a stapler or stapler instrument 12 having a stapler head or device 14 positioned on a distal portion of a shaft 16. A handle 18 on the shaft 16 controls articulation of the stapler head 14 and actuation of the tissue acquisition, tissue compression, and stapling functions of the stapler head 14. Vacuum and fluid sources 20, 31 in the system are fluidly coupled to the handle 18 for use in tissue acquisition, compression and stapling as discussed below. The vacuum source 20 may be the "house vacuum" accessible through a coupling on the wall of the operating room, or an auxiliary suction pump. The stapler may include a switch 21 allowing the user to control airflow between the vacuum source and stapler.

[0060] The stapler device also serves to capture a tissue fold for stapling, and is thus also referred to herein as a tissue capture device for immobilizing a tissue fold, e.g., for fastening the sides of the fold. The tissue capture device may operate independently for capturing tissue, e.g., absent a separate stapling mechanism, or may be combined with the stapling elements, as illustrated.

[0061] The fluid source 31 may be a single source of drive fluid (e.g. water, saline, oil, gas) or multiple sources, but in each case the fluid source preferably includes two actuators separately used to control flow into each of two hydraulic lines (one for tissue compression and one for stapling). An endoscope 22 in the system is insertable through a lumen in the shaft 16 permits visualization of the plication procedure. The system may optionally include an overtube, such an endoscopic guide tube 23, having a lumen for receiving the stapler 12.

[0062] Referring to Fig. 3A, a covering or membrane 24 in the stapler encloses the stapler mechanism to form a vacuum chamber 21 (Figs. 17-23) within the stapler head 14. The side exposed to the tissue to be plicated remains uncovered by the membrane 24 to allow tissue to be drawn into the chamber during use. For example, the membrane 24 may include a side opening 26 as shown in Fig. 3B. Membrane 24 is preferably formed of silicone, elastomeric material, or any other inelastic or elastic flexible or deformable biocompatible material, such as a pleated mylar film, capable of forming a vacuum chamber 21 that will expand in volume to accommodate tissue drawn into chamber 21. Also shown in Figs. 3A-3C is an articulating section 128 connecting the instrument shaft to the staple head, and described below with reference to Figs. 26 and 27.

[0063] At least a portion of the membrane is at least partially transparent. In being at least partially transparent, the membrane is formed of a material, or includes sections of material, that will allow the user to see through the membrane well enough to confirm (via endoscopic

observation) that an appropriate volume of tissue has been acquired into the stapler head prior to staple application. The opening 26 may be surrounded by a reinforced section 27 formed of material that will strengthen the area around the opening 26. Reinforced section 27 may be formed of a thicker section of the membrane material, and/or a higher durometer material. Alternatively, reinforcing ribs or other structures or element may be formed into or onto the membrane material, or embedded in the membrane material.

Stapler Head or Device

[0064] The stapler head 14 is designed to have a minimum profile during insertion to the plication site, and to then transform into a much larger profile device having to large internal volume. For example, in one embodiment the vacuum chamber might have an initial internal volume of 0.2 cubic inches, and an expanded volume of 0.6 cubic inches (i.e. the internal chamber volume after subtracting the volume occupied by the stapler head components positioned within the vacuum chamber). This large internal volume allows a large volume of tissue to be drawn into the vacuum chamber and stapled. In this way, the stapler head creates a large plication without requiring invasive techniques for insertion. The unique features of the stapler head allow in situ volumetric expansion of the stapler head using a minimum of motion and force input. In particular, as will be seen below with respect to Figs. 32A-C and Fig. 33, the plication can be sized such that the staples applied to the tissue, such that the two annular staple arrays seen in Fig. 33 are well spaced from the edges of the stapled tissue, minimizing the risk of tissue tearing around the staples.

[0065] Features of the stapler head are shown in Figs. 4-10. For clarity, the membrane is not shown in these figures. Referring to Fig. 4, stapler head 14 generally includes a first, staple member 25 comprising a proximal staple housing 28, a second, anvil member 27 comprising a distal anvil housing 30, and at least one elongate member but preferably a pair of hinged arm assemblies 32 that operatively connect the two housing as described below.

[0066] The staple housing and anvil housing are arranged to allow tissue to be compressed between contact surfaces on each of the staple housing and the anvil housing. In the disclosed embodiment, the contact surfaces are on a staple holding portion of the staple housing, i.e., the outer face of the staple holder, and an anvil on the anvil housing. Considering only the tissue-capture operation of the device, staple holder 78 (shown in Fig. 7) functions as a tissue-capture plate having a front, tissue-contacting surface 83, and anvil 96 ((shown in Figs. 13 and 14) functions as a second tissue-capture plate having a tissue contacting surface 103 confronting surface 83, where these two surfaces serve to capture the tissue fold during operation of the device, as will be described more detail with respect to Figs. 32A - 32C.

[0067] The arm assemblies 32 extend between the staple housing 28 and anvil housing 30 on opposite sides of the stapler head 14. Proximal and distal pins 34, 36 pivotally couple each arm assembly 32 to the staple housing 28 and the anvil housing 30. An expansion member comprising a membrane raiser 37 also extends between the staple housing 28 and the anvil housing 30. Although the membrane 24 is not shown in Fig. 4, it should be understood that the membrane raiser 37 is positioned opposite the opening 26 (Fig. 3B) in the membrane. In the illustrated embodiment, membrane raiser 37 includes a link 38 pivotally mounted to the staple housing by a pin 42, a corresponding link 40 pivotally mounted to the anvil housing by pin 44, and spring wires 46 coupling the links 38, 40 to one another.

Staple Housing

[0068] Turning to a more detailed discussion of the stapler head components, the staple housing 28 can be seen separated from other components in Figs. 5 and 6. As shown in Fig. 5, proximal face 48 of the staple housing includes input ports 50a, 50b through which fluid is directed for hydraulic actuation of the tissue compression, stapling, and optional cutting operations of the stapler head. Seals 51 surround the ports 50a, 50b to minimize fluid leakage.

[0069] Vacuum ports 52 are fluidly coupled to a vacuum source 20 (Fig. 2) that is selectively activated to create negative pressure in the vacuum chamber for tissue acquisition. The vacuum ports 52 are connected to the vacuum source 20 by flexible tubing (not shown) in the stapler shaft 16 (Fig. 2). Mounting holes 54 are used to mount the stapler head 14 to the shaft 16 through the articulating section 128.

[0070] The staple housing 28 includes upper and lower sections 58a, 58b above and below open side sections 56. The upper section 58a includes a recess 60 within which the pivot pin 42 for link 38 (Fig. 4) is mounted. As best shown in Fig. 6, bores 62 are positioned in the upper and lower sections 58a, 58b to receive pins 34 (Fig. 4) that serve as the proximal pivot points for arm assemblies 32. Guide slots 64 extend longitudinally through the upper and lower sections 58a, 58b.

[0071] Referring to Fig. 6, a hydraulic chamber 66 is disposed within the staple housing 28. Within the hydraulic chamber 66 (Fig. 6) is a dedicated hydraulic circuit for driving the tissue compression and stapling functions of the stapler. Chamber 66 is fluidly coupled to the fluid input ports 50a, 50b (Fig. 5). As will be discussed in detail in connection with Figs. 11A-11D, fluid driven into the hydraulic chamber 66 via input ports 50a, 50b sequentially advances a system of hydraulic pistons (not shown) that act on other components to compress the tissue, and that drive the staples and cutting element through the compressed tissue.

[0072] Fig. 7 illustrates components of the stapler head that are driven by the hydraulic system for compression,

stapling, and cutting. For clarity, these components are shown separated from the staple housing and from each other. In this discussion, the components that are driven by the hydraulic system will be described. The hydraulic system itself is described in a later section in connection with Figs. 11A - 11D.

[0073] In particular, Fig. 7 illustrates a drive member which takes the form of a disk 68 in the staple housing. In the assembled housing, disk 68 is positioned such that it will be pushed distally by a hydraulic compression piston (piston 106 in Figs. 11A-11E). As will be seen in Figs. 11A-11E, the drive member is movable between a first, retracted position shown in Fig. 11A to a second, extended position shown in Figs. 11C and 11D. Although the drive member illustrated here is driven by, but separate from piston 106, it will be appreciated that these two components can be formed of a single-piece member, i.e., as a single-piece drive member including both piston and disc. As will be seen below, the drive member is coupled to the arm assemblies 32, the anvil housing, and the staple housing so that advancing the drive member distally (toward its extended position) effects tissue compression by bringing the contact surfaces of the staple housing and anvil housing relatively towards one another. The combination of disk 68, its driving piston 106, and assembly arm 32 coupling the two housings is also referred to herein collectively as a drive assembly, indicated at 29 in Fig. 10. The drive assembly may further include drive links 114 in the anvil member, which are operatively linked to assemblies 32 as described below.

[0074] As seen best in Fig. 7, disk 68 includes mounting bores 70, a central opening 72, and alignment posts 74. Referring briefly to Fig. 10, in the assembled stapler head, disk 68 is coupled to the stapler housing 28, and its axial movement therein constrained, by pins 84 that extend through the housing's guide slots 64 and through mounting bores 70 in the disk 68.

[0075] A portion of the staple housing 28 contains, i.e., is loaded to contain, staples to be fired into the tissue. The staples are contained within a staple holder, such as staple cartridge 78, on the staple housing. The staple holder may have a number of different configurations. For example, it may be an integral portion of the staple housing, or a separate part mounted or attached to the staple housing, and/or it may be moveable relative to the body of the staple housing to effect tissue compression prior to stapling. In any of these examples, the staple holder may be a removeable/replaceable cartridge, and/or it may be refillable by inserting additional staples into it. In other embodiments, the staple holder may be neither replaceable nor refillable, i.e., i.e., in a dive intended for one-time use.

[0076] In the disclosed embodiment, the staple holder is a removeable staple cartridge 78 that can be replaced with another cartridge after staple firing. In this embodiment, the staple cartridge is moveable relative to the body of the staple housing to compress the tissue prior to staple firing.

[0077] Referring again to Fig. 7, staple cartridge 78 is positionable within the staple housing, distal to the disk 68, such that distal advancement of the disk by the compression piston pushes the cartridge from a first, retracted position distally to a second, extended position to compress tissue disposed between the cartridge and anvil. Grooves 79 on the exterior of the cartridge slide over corresponding ones of the alignment posts 74 during insertion of the cartridge into the stapler head. Fig. 10 shows the alignment posts prior to loading of a cartridge into the staple housing. As shown, the alignment posts 74 may have tapered ends to facilitate loading of the cartridge over the posts. It will be appreciated that the alignment posts hold the cartridge against angular movement within housing 28 during stapler operation.

[0078] Again referring to Fig. 7, cartridge 78 includes a number of staple locations 80, each housing a staple, such as staples 158 seen in Fig. 33. The staple cartridge is equipped with bosses 81 to retain a staple line reinforcement device 83 of the type shown in Fig. 8 and disclosed in detail in commonly-owned U.S. Application No. 11/542,457, entitled ENDOSCOPIC PLICATION DEVICES AND METHODS, filed October 3, 2006, and published September 20, 2007 as US 20070219571. To summarize briefly, this type of reinforcement device 83 may be a ring or other element positionable against the distal face of the staple cartridge. When the ring is placed on the cartridge, openings 85 in the ring align with prongs of some of the staples in the cartridge. When staples are driven from the cartridge, these prongs pass through associated ones of the openings 85 and capture the ring 83 against the adjacent body tissue.

[0079] Referring to Figs. 7 and 9, a number of undercut bosses 81 on the anvil-facing side of the cartridge may be used to lock the reinforcement device 83 in place on the face of the staple cartridge. Other positive shapes, such as mushrooms, hooks, and tilted bosses could be used to accomplish the same end. Negative shapes, such as pockets or grooves formed into the surface of the cartridge, may also be employed to engage corresponding features on the reinforcement device 83. As another alternative, the reinforcement device may be held in place on the cartridge using adhesives.

[0080] In the embodiment shown, a cutter element 86 extends through the central opening 72 (Fig. 7) of the disk 68. The cutter element is shown as a tubular punch having a sharpened wall and a lumen 87, but may be provided in alternative forms. A staple pusher 76 is mounted to the cutter element, distally of the disk as can be seen in the assembled view of Fig. 10. Staple pusher 76 includes pusher elements 82 proportioned to slide into the cartridge's staple locations 80 as the staple pusher 76 is advanced into the staple cartridge 78, thus driving the staples from the cartridge. A hydraulically-driven staple piston (shown at 116 in Figs. 11A-11E) in the hydraulic chamber 66 (carried within a hydraulic chamber formed by piston 106) is coupled to the cutter element 86 such that advancement of the stapler piston advances

the staple pusher 76 and cutter element 86 in a distal direction.

Fluid Drive System

[0081] The fluid drive system used to actuate compression, stapling and cutting may be configured in various ways. The following paragraphs describe one exemplary configuration for the fluid drive system, which in this embodiment is a hydraulic system. Figs. 11A and 11B schematically show the fluid flow in the hydraulic chamber 66 of the stapler housing 28 during both compression and stapling stages of actuation. Referring to Fig. 11A, compression piston 106 is disposed within hydraulic chamber 66. Disk 68 (also shown in Figs. 7 and 10) is positioned in contact with or slightly distal to piston 106. Compression piston 106 is generally cup-shaped, having a rear wall 108 and a side wall 110 enclosing an interior 111. O-ring seals 112 are spaced-apart on a proximal portion of the side wall 110. Channels 115 are formed through the side wall 110, between the o-ring seals 112.

[0082] A second piston, referred to as the staple piston 116, is positioned in the interior 111 of compression piston 106, against the rear wall 108. Although not shown in Figs. 11A-11D, cutting element 86 (Fig. 7), with the staple pusher 76 thereon, is positioned in contact with or slightly distal to the staple piston 116. An o-ring seal 118 surrounds a portion of the staple piston 116 that is distal to the channels 115 in the compression piston.

[0083] A first fluid channel 120 extends from fluid port 50a in the stapler housing 28 to a proximal section of the hydraulic chamber 66. A second fluid channel 122 extends from fluid port 50b in the stapler housing to a more distal section of the hydraulic chamber 66. Fluid flow from port 50a and fluid channel 120 against the compression piston cylinder is shown in Fig. 11A. Fluid pressure within the hydraulic chamber 66 advances the compression piston 106, with the stapler piston 116 within it, in a distal direction, from a first, retracted position, shown in Fig. 11A to a second, extended position shown in Figs. 11C and 11D. Fig. 11B shows the compression piston 106 approaching the end of its travel, i.e., fully extended position. Once the compression piston reaches the end of its travel as shown in Fig. 11C, channel 115 in the compression piston 106 aligns with channel 122 in the housing, allowing fluid introduced through fluid port 50b to enter the interior of the compression piston 106 via channel 122. The fluid entering the interior of the compression piston drives the staple piston distally as shown in Fig. 11D, from a first, retracted position shown in Figs. 11A-11C, to a second, extended position shown in Fig. 11D. In an alternative embodiment shown in Fig. 11E, a third piston 117 is provided for separately driving the cutting element 86. In this embodiment, fluid introduced into a third drive fluid port 50c causes advancement of the third piston 117 from a first, retracted position to a second extended position (not shown). The pistons 106, 116 and 117 and associated fluid paths may be arranged so that

fluid cannot enter the interior of the stapler piston to advance the cutting piston 117 until compression piston 106 has traveled to the tissue-compression position and stapler piston 116 has in turn traveled to the stapling position.

[0084] The anvil housing (identified by numeral 30 in Fig. 4) in anvil member 27 will next be described with reference to Fig. 12. The anvil housing 30 includes mounting bores 88 for receiving pivot pins 36 at the distal end of the hinged arm assemblies 32. The upper section of the anvil housing 30 includes a section 94 through which the pivot pin 44 for link 40 (Fig. 4) is mounted.

[0085] A central bore 90 extends longitudinally through the anvil housing 30. An anvil support 92 (Fig. 13) is longitudinally slidable within the bore. Both the bore 90 and the anvil support 92 are preferably formed to have non-circular cross-sections (such as the illustrated rectangular cross-section) with flat bearing surfaces to prevent rotation of the piston within the bore.

[0086] Fig. 13 shows the anvil support 92 separated from the anvil housing 30. The distal portion of the anvil support 92 is split into upper and lower plates 95a, b. Plate 95a has a bore 93 axially aligned with a similar bore in plate 95b. The proximal portion of the anvil support 92 carries the anvil 96. As shown in Fig. 14, anvil 96 includes a plurality of indentations 98 positioned such when staples are driven from the staple cartridge, each staple leg engages one of the indentations, which causes the staple leg to fold or crimp. In the embodiment shown, the anvil is designed for a staple array having two annular rings of offset staples, five staples per ring. A central opening 97 extends through the anvil 96 and is contiguous with a lumen in the anvil support 92.

[0087] The anvil 96 and the staple cartridge 78 (Fig. 7) are the two parts of the stapler head which exert force on the tissue to be stapled. As shown in Figs. 9 and 14, the preferred anvil and cartridge are designed to use a minimal amount of material surrounding the indentations 98 of the anvil 96 and the staple locations 80 of the cartridge 78 - so that the amount of anvil/cartridge surface area contacting the tissue is as small as possible. When subjected to a constant force, a smaller footprint will damage less tissue than would a larger footprint, since a smaller area of tissue is squeezed between the anvil and cartridge. However, the tissue that does get squeezed experiences more pressure from the given force because the force is distributed over a smaller area. In other words, the minimized footprint creates more pressure on the tissue with less force. This is advantageous from a mechanical standpoint because the stapler head need not supply or withstand as much force as would be needed with a larger-footprint cartridge and anvil.

[0088] Referring to Fig. 7, in the illustrated embodiments, the staple cartridge 78 has an outer wall that tracks the contours of the staples housed within it, thus forming a number of pedals 73 surrounding the outer staple positions or slots 80a, with the grooves 79 disposed between the pedals, adjacent to the inner staple

positions 80b. Rather than providing each staple position to be fully surrounded by cartridge material, the staple positions 80a, 80b preferably each include a back wall 71a and a retaining element attached to the wall and positioned to retain a staple between the retaining element and the back wall. In Fig. 7, the retaining element comprises a pair of wings 71b that curve inwardly from the back wall 71 to define a slot that is sufficiently bounded to retain a staple within the staple position, but that is preferably not bounded around its full circumference. The anvil has a similar pedal arrangement, as shown in Fig. 13.

[0089] Referring again to Fig. 13, a plate 99 is positioned on the anvil 96 such that the distally-advancing cutting element 86 will advance into contact with the plate 99 during tissue cutting. In one embodiment, the plate 99 may be seated within the opening 97 in the anvil. The plate 99, which will also be referred to as the "cutting board", has a hole 101 in it which relieves the pressure of the captured tissue and prevents hydraulic locking, a condition in which the punch and plate create a closed volume. If it is desired to move the cutting element 86 after contact is made, pressure will increase inside this closed volume and it will resist further motion. This may prevent or adversely affect tissue cutting.

[0090] The cutting board is preferably designed so as to not serve as a hard stop against advancement of the cutting element 86. If the cutting element 86 is stopped by the cutting board, the stapling piston will also be stopped and incomplete staple formation may result. Therefore, it is preferred that the cutting element 86 is allowed to penetrate or displace the cutting board during and after the tissue is cut; that is the cutter can advance slightly once initial contact with the board is made.

[0091] Figs. 15A and 15B illustrate the cutting element 86 advanced into contact with different embodiments of cutting boards. In the Fig. 15A embodiment, the material of cutting board 99a is a relatively soft material, such as an elastomeric silicone, which is cut or penetrated by the advancing cutting element as shown. This material allows the sharp distal end of the cutting element to move into the cutting board during the final stage of staple formation. In the Fig. 15B embodiment, the cutting board 99b can be made of a harder material positioned with a compressible object such as an elastomeric spring 99c behind it. In the figure, this spring is an o-ring. Advancement of the cutting element 86 against the cutting board 99b causes the cutting board to be displaced distally against the spring 99c. The advancing cutting element 86 experiences increasing resistance as the o-ring is compressed. Other spring shapes and materials, such as coiled wire, spring washers and leaf springs can be used to achieve the same result. The chamfer 99d on the surface of the cutting board 99b may help to align the cutting element 86 as it is forced into contact with the cutting board.

Arm Assemblies

[0092] Following is a discussion of the features of the arm assemblies 32. Fig. 16 shows the arm assemblies 32 separated from the other elements of the stapler head. In general, each arm assembly has a first arm section 100 pivotally coupled to the staple housing and a second arm section 102 pivotally coupled between the first arm section and the anvil housing. While not present in the illustrated embodiment, additional arm sections may be positioned between the first and second arm sections.

[0093] That is, each arm assembly includes a proximal arm 100 and a distal arm 102 joined to one another to form a hinge 104. Each of the proximal arms 100 has a longitudinal cutout 108 and an arm spreader 113 pivotally mounted within the cutout 108. The distal end of each arm spreader 113 includes a bore 112. Pin 84 is positioned within the bore 112. As disclosed in connection with Fig. 10, this pin 84 extends through the disk 68 and has ends that ride within the slots 64 (Fig. 6) on the lower and upper sections of the stapler housing. Longitudinal movement of the disk 68 within the stapler housing will thus advance the pins 84 within their corresponding slots 64, causing the arm spreaders 113 to pivot relative to the pins 84 and to thus drive the arm assemblies 32 outwardly. Additional specifics concerning movement of the arm assemblies 32 is set forth in the section entitled Stapler Head Operation.

[0094] Distal arms 102 of the arm assemblies include pins 36 which, as discussed, are pivotally mounted to the anvil housing 30 (Fig. 4). A pair of drive links 114 are provided, each of which has a first end pivotally attached to a corresponding one of the distal arms 102 and a second end pivotally coupled to a common pin 116. In the assembled stapler head, pin 116 is positioned in the bores 93 of the upper and lower plates 95a, 95b of the anvil support (see plates 95a, b in Fig. 12). As detailed in the Stapler Head Operation section below, when the arm spreaders 113 drive the arm assemblies 32 outwardly, drive links 114 act on the pin 116 to push the anvil support in a proximal direction, causing the anvil to advance proximally towards the staple cartridge.

[0095] Alternatively, the anvil may be driven within its housing independently of the arm assemblies, by a direct hydraulic drive mechanism carried in the anvil housing. Preferably, the drive mechanisms in the two housings are coordinated so that the staple holder and anvil are moved toward one another in their respective housings at the same time.

Stapler Head Operation

[0096] The following discussion centers on the manner in which the arm assemblies function to expand the vacuum chamber and to compress tissue that has been drawn into the chamber using suction. As an initial step preceding chamber expansion, the stapler head is positioned with the opening 26 in the membrane 24 in contact

with tissue at the location at which plication creation is desired. Vacuum source 20 (Fig. 2) is activated to apply vacuum to the inside of the vacuum chamber defined by the membrane. Tissue in contact with the opening 26 (Fig. 3B) will be drawn into the vacuum chamber between the staple housing 28 and the anvil housing 30. After the tissue is drawn in, the stapler profile is changed, expanding the volume of the chamber within the membrane.

[0097] The streamlined position of the stapler head 28 prior to expansion is shown in Figs. 4, 17 and 18. In particular, the hinged arm assemblies 32 and membrane raisers 37 are in generally straight orientations. The proximal arms 100 serve as the drive arms for chamber expansion and tissue compression. Motion of these arms is initiated when water under pressure is forced into the hydraulic circuit of the staple housing. Referring to Fig. 19, the fluid pressure advances disk 68 (by action of the compression piston 106, not shown in Fig. 19). Disk 68 in turn pushes the staple cartridge 78 toward the anvil 96 as shown in Figs. 19-21, causing the staple cartridge 78 to extend further from the staple housing 28.

[0098] Both the disk 68 and the arm spreaders 113 are coupled to the pins 84. For this reason, the longitudinal movement of the disk 68 within the stapler housing 28 will carry the pins 84 distally within their corresponding slots 64. The arm spreaders 113 will consequently pivot relative to the pins 84, driving the proximal arms 100 outwardly. Outward movement of proximal arms 100 at hinge 104 causes the distal arms 102 to also pivot outwardly at hinge 104, forming an angle between the proximal and distal arms 100, 102. Naturally, formation of the angle between the arms 100, 102 shortens the effective length between the remote ends of the arms, causing the distal pins 36 of the distal arms 102 to carry the anvil housing 30 towards the staple cartridge. The pivoting movement of the distal arms 102 further causes drive links 114 to act on pin 116 to push the anvil support in a proximal direction. This moves the anvil support relative to the anvil housing in a proximal direction at the same time the anvil housing is also moving proximally.

[0099] In essence, one motion, that of the hydraulically driven compression piston, creates at least three motions, illustrated by arrows A1, A2 and A3 in Figs. 19-21. These three motions include: the staple cartridge 78 moving relative to the staple housing in a direction towards the anvil 96 (arrow A1), the anvil housing 30 moving toward the staple housing 28 (arrow A2) and the anvil 96 itself moving relative to the anvil housing 30 in a direction towards the cartridge (arrow A3). This compound motion of the anvil toward the staple cartridge enables a small displacement of the compression piston to quickly compress tissue in the grip of stapler. The multiplication of motion also enhances force transmission between the two housings by keeping the angle at hinge 104, between the proximal (driven) arm and the distal (drive) arm, as large as possible.

[0100] The relative motion of the two housings 28, 30 toward each other also drives upward links 38, 40 and

their interconnecting spring wires 46 on the top of the stapler head 14. Together, the links and spring wires raise the top of the membrane, creating more volume to accommodate expansion of the tissue during compression.

[0101] Compression of the tissue is halted when the pins 84 traveling in slots 64 in the staple housing 28 reach the limit of travel, as shown in Figs. 22 - 24. Thus, the slots and associated components are dimensioned to set the desired separation distance between the tissue contact surfaces on the stapler side and the anvil side of the stapler head. Exemplary separation distances for use in stomach wall plications might include approximately 0.06-0.07 inches (e.g. for use with staples having legs of 5.5mm length) or 0.109 inches for 6.5mm leg length staples. Application of additional pressure into the hydraulic circuit will not compress the tissue any further.

[0102] Moreover, because of the piston arrangement, the stapling function is effectively locked out until tissue compression is complete. With this arrangement, fluid introduced via the fluid port 50b (Fig. 11A) into the staple fluid channel 122 prior to completion of tissue compression will leak until the two o-rings 112 of the compression piston 106 are straddling the inlet 114. This design prevents premature staple firing.

[0103] At the fully compressed position, the arm spreaders 113 are nearly perpendicular to the longitudinal centerline of the stapler head. Once tissue is compressed between cartridge 78 and anvil 96, the tissue is ready for stapling.

[0104] Stapling is initiated by introducing hydraulic fluid through port 50b (Fig. 5). The staple piston advances, pushing cutting element 86 (Figs. 7 and 10) towards the anvil 96. Because the staple pusher 76 is mounted to the cutter 86, this action carries the staple pusher 76 through the cartridge 78 where it simultaneously pushes all staples through the tissue. Staple piston travel is limited by internal stops, and is preset to yield optimal staple formation.

[0105] During compression, as the angle at the hinge 104 of arm assemblies 32 reaches its minimum, the force required to resist separation of the staple and anvil housings increases. These forces increase further when the forces of staple crushing are exerted on the anvil by the staple piston. To compensate, the arm spreaders 113 serve as displacement struts to channel at least a portion of these forces into the disk 68. These forces, if not reacted by the pusher disk, would pull in the arms 100, 102 and potentially release the compression on the tissue, causing incomplete staple formation or tissue cutting. In this way, a truss-like structure is created for force displacement.

[0106] When staples have been formed, staple pressure is released and a spring (not shown) returns the staple pusher 72 to its base position. Releasing fluid pressure will allow the deflected spring wires 46 on membrane raiser 37 to return the staple head to its minimum profile configuration and release the plication from the stapler. Once outside the patient, the used staple cartridge can

be ejected and a new one installed.

[0107] Figs. 25A - 25C illustrate one method for retaining a removable staple cartridge 78 within the staple housing. The cartridge is spring-loaded into the staple housing and retained by two latches 170 (one visible), each pivotable relative to a fulcrum 172. As shown, the fulcrum 172 may be coupled to the disk 68 by pin 84. Each latch 170 includes a catch 174 which engages a corresponding catch 176 on the cartridge. The latch 170 is preferably spring biased to urge the catch 174 inwardly towards the cartridge.

[0108] Depressing the proximal end 175 of each latch 170 as shown by arrow P in Fig. 25B pivots the latch against this bias, causing ejection of the staple cartridge. A new staple cartridge may then be positioned with its grooves 79 aligned with alignment posts 74 as shown in Fig. 25C and then pushed towards the staple housing. As the new cartridge slides into position, catch 174 rides over the tapered proximal portion 178 of the catch 176. Once catch 174 passes over the distal end 180 of the catch 176, it drops inwardly towards the cartridge due to its spring bias, thus engaging the cartridge. When the cartridge is properly seated, a click will be felt or heard as the latches engage the new cartridge.

Stapler alignment during stapling

[0109] in operation, as the two stapler members are brought together, a tissue fold is captured between the outer face of the staple cartridge and the confronting face of the anvil. As this tissue capture is occurring, as just prior to and during the stapling of the tissue fold, variations in the thickness and or compressibility of regions of the captured tissue may bias the stapling operation off-center, i.e., off axis, causing the staple legs to be received out of position within their associated anvil indentations, with the result that one or more of the staples may fastened defectively, e.g., without complete folding of the staple legs, and/or the array of staples may be offset with respect to the center axis of the device, causing some staples, for example, to be too close to the hole cut in the center of the stapled fold.

[0110] The insure that the stapling operations occurs successfully for each staple array in the cartridge (for example, two concentric arrays of five staples each), the device of the invention may include alignment structure to hold to proximal and distal members of the device in axial alignment just prior to and during the stapling operation. The alignment structure is shown in side sectional views in Figs. 36 and 37. Shown in Fig. 36 is the portion of the staple device 12 that includes anvil 96 and cutting board 99a on distal member 27, and cutter 86 on proximal member 25. In operation, the two members, and the staple cartridge and anvil carried thereon, are first moved together, to capture a tissue fold between the confronting faces of the cartridge and anvil, as illustrated in Figs. 22 and 23. Staple piston 116 is then moved from its retracted to its extended position, as seen in Fig. 24, to move the

cutter against cutting board 99a, as shown, and drive the staples in the staple cartridge against the anvil.

[0111] The alignment structure includes a pin held axially within cutter 86 for movement therewith, and a pin-receiving slot formed in a bushing 164 supported within a portion of housing 27 attached to anvil 96 for axial movement therewith. In the embodiment shown in Fig. 36, bushing 164 is spring loaded, through a spring 166, within housing 27 in a direction that resists movement of the pin toward the bushing, allowing the cutter and attached pin, which has now penetrated through the captured tissue, to seat in the beveled end of the bushing. As the cutter, pin, and staple pusher continue to move toward the second member, ultimately forming a hole in the tissue and ejecting staples from the staple cartridge through the tissue and against the anvil, the two device members are maintained in axial alignment, so that the stapling operation occurs with all staples positioned in alignment with the respective to the associated anvil indentations. When this operation is complete, the staple piston and drive member are retracted to release the stapled tissue and return the device to its linear condition.

[0112] The embodiment of device 12 shown in Fig. 37 is similar to that just described, but where the alignment pin 168 carried in the housing of cutter is spring biased, by a spring 174, in the direction of the second member, and a pin-receiving slot 170 is in a fixed-position bushing 172 carried below the anvil (not shown) in member 27. In this embodiment, motion of the two members toward one another, once the pin has been initially seated in the bushing slot, is accommodated by movement of the pin in a direction away from the second member, as the staple piston continues to move toward its fully extended position, to cut and staple the tissue captured between the two members, as described above.

Stapler Shaft and Handle

[0113] Referring again to Fig. 2, the stapler shaft 16 connecting the handle 18 and the stapler head 14 is flexible enough to conform to the curvature of the upper digestive tract, yet maintains the ability to transmit enough torque to rotate the stapler head. The shaft is formed with sufficient stiffness to allow it to be pushed down esophageal guide tube 23. Suitable materials include

[0114] Fig. 26 shows a distal portion of the shaft 16, with the stapler head removed from the shaft. As shown, shaft 16 includes an endoscope lumen 124 through which an endoscope is advanced to allow visualization of a stapling operation. Side lumens 126 may also be provided for receiving other instruments useful during the procedure.

[0115] An articulating section 128 is positioned at the distal end of the shaft 16, between the shaft 16 and the stapler head 14 so as to allow the stapler head to be articulated relative to the shaft. Tubing coupled to the vacuum source and the source of hydraulic fluid extends from the handle and through the shaft 16 and the artic-

ulating section 128.

[0116] Fig. 27A shows one configuration that may be used for the hydraulic fluid lines 130. During use, the hydraulic fluid lines are subjected to significant deflection and elongation in the articulating section of the stapler. They are also subjected at times to fluid pressure which may be in excess of 1,500 psi. Typically, hydraulic lines in industrial applications are flexible and have a working loop of extra tubing that accommodates length changes during use. The illustrated configuration for the hydraulic lines is a lower profile solution particularly suitable for an endoscopic device having space constraints. A preferred hydraulic line is a tube 130 having a portion that is shaped into a longitudinally expandable shape so that it can accommodate effective length changes during bending. The longitudinally expandable portion of the tube is preferably disposed within the articulating section 128 of the stapler 12. In a preferred design, the longitudinally expandable shape is a coil shape as shown in Fig. 27A. In alternate embodiments, the tube 130 may be formed into other longitudinally expandable shapes, such as regular or irregular undulating shapes (Fig. 27B).

[0117] The preferred material for the tubes 130 is stainless steel hypotube, although other materials may instead be used. In the preferred stapler configuration, two drive fluid lines are provided, one for actuating tissue compression, and the other for staple application (and cutting when used). In the present embodiment, the tubes are coiled together as shown in Fig. 27A. In alternate embodiments, two or more coiled tubes may be nested one inside the other. As the articulating section bends, it forces the coiled tubes 130 to bend and to change length in response to bending. The coiled tubes behave just as coiled wires would during these motions and are thus able to change length, deflect, and follow the contour of the articulating section without compromising flow through the lumens of the tubes or imparting undue stress to the connections at either end of the hydraulic system.

[0118] The longitudinally expandable shapes for the fluid lines may be suitable for use in allowing delivery of fluid to the operative ends of other types of articulating medical devices, such as catheters or endoscopic devices for delivering therapeutic agents or irrigation fluids past an articulating or bendable section of the device.

[0119] Referring again to Fig. 26, articulating section 128 is comprised of a spine formed of a plurality of links 132 strung over a pair of pull cables 134 (only one shown in Fig. 26). In one embodiment, engagement of the pull cables allows the stapler head 14 to be articulated in two directions through a range of motion of approximately 90 degrees in one direction (see Fig. 3B) to 175 degrees in the opposite direction (see Fig. 3C). Each pull cable is anchored at or near the stapler head, such as at the distal most link 132 of the stapler housing 28.

[0120] The more proximal portions of the pull cables 134 extend the length of the shaft 16 and terminate in the handle 18. Referring to Fig. 28, the handle 18 includes a rotating knob 136 that may be selectively rotated in a

clockwise or counterclockwise to articulate the stapler head up or down. Rotation in one direction applies tension to one of the pull cables to cause the stapler head to bend downwardly, whereas rotation in the opposite direction puts tension on the other cable, causing the head to bend upwardly.

[0121] In a preferred handle configuration, the knob 136 includes an internal threaded bore 138. Knob 136 is partially restrained within the handle 18 so that it remains fixed within the handle but can rotate freely. A carriage 140 having a threaded exterior surface is positioned within the threaded bore 128 of the knob. The threads within the bore 138 are engaged with the threads on the carriage 140 so that rotation of the knob causes the carriage 140 to translate, but not rotate, within the handle.

[0122] Each of the two pull cables, identified in Fig. 28 as cables 134a and 134b, is terminated on a different member in the handle. Cable 134a is mounted on the sliding carriage and cable 134b is mounted to a stationary part of the handle 18. Each cable extends through a corresponding sheath. Cable 134a extends through a sheath 135a having a proximal end fixed to a stationary part of the handle 18. Cable 134b extends through a sheath 135b having a proximal end mounted to the sliding carriage.

[0123] The cables 134a,b and sheaths 135a,b are arranged such that translation of the carriage in one direction will cause deflection of the stapler head in one direction, and translation of the carriage on the other direction will deflect the stapler head in another direction.

[0124] Referring to Fig. 28, if knob 136 is rotated to causes the carriage 140 to translate to the left of the page, cable 134a will be tensioned and cable 134b will slacken, causing the stapler head to articulate in a first direction (e.g. upwardly). Rotation of the knob 136 in the opposite direction will advance the carriage to the right of the page, releasing tension on cable 134a and pushing sheath 135b over the cable 134b towards the distal end of the staple head, causing articulation in the second direction (e.g. downwardly) as the sheath 135b is advanced against a distal portion of the shaft 16. The proximal portion of sheath 135b is provided with sufficient working length prevent it from being placed under tension when the carriage moves distally. The positioning of the knob is advantageous in that the hand movement required for stapler articulation is always the same, regardless of the rotational orientation of the stapler. Also, the use of the threaded knob can prevent unintentional relaxation of the deflection angle, even if the knob is provided without a lock to retain its rotational position.

[0125] Referring to Figs. 28 and 29, the endoscope lumen 124 extends along the center axis of the stapler. The positioning of the lumen and the coaxial relationship of the articulation knob in relative to the endoscope 124 allows the endoscope and stapler to be rotated independently without one interfering with one another. Thus, if the user chooses to change the rotational orientation of the stapler head 14 within the body, s/he may rotate the

handle 18 and shaft 16 while maintaining the rotational position of the endoscope.

[0126] For cost efficiency, the stapler 12 may be designed to permit the stapler head 14 to be discarded while allowing the shaft 16 and handle 18 to be sterilized and re-used. One mechanism for removably coupling the stapler head to the shaft 16 is illustrated, although others are readily conceivable (e.g. a slip coupling type arrangement). Referring to Fig. 26, an end plate 142 is mounted to the distalmost one of the links 132. Each of the end plate 142 and the corresponding rear surface of the stapler head are provided with latch features that allow the end plate and stapler head to be engaged to one another.

[0127] End plate 142 includes a cantilevered pin 144 having a peg 145 (which may be a spring pin), a central opening 146, and a pair of u-shaped catches or engagement edges 148 along its edges. Hydraulic feed holes or openings 156a, b are formed through the end plate 142. The hydraulic tubes that deliver hydraulic fluid to the stapler head (see tubes 130 of Fig. 27) are preferably welded to the end plate to allow fluid from the tubes to be directed through the feed holes 156a, b.

[0128] Figs. 30A and 30B show the rear surface 48a of the staple housing, which has been somewhat modified relative to Fig. 5. In this variation of the rear surface 48a, the hydraulic input ports or openings 50a, 50b are repositioned as shown. Additionally, the rear surface 48a has been modified to include a pair of catches or engagement edges in the form of undercut bosses 150, plus an aligning pin 152, and a hole 154. The snap-on connection of shaft to stapler device;

[0129] Figs. 30A and 30B show the end plate 142 positioned against the rear surface 48a of the staple housing. The other features of the articulating section 128 are not shown in Figs. 30A and 30B for clarity. To attach the stapler head to the shaft 16, the plate 142, attached to the handle assembly, is pressed against the rear surface 48a of the staple housing as shown in Fig. 30A. As the plate is pushed, it is rotated in a clockwise direction, causing the peg 145 (Fig. 26) of the cantilevered pin 144 to engage hole 154 in the rear surface of the staple housing. When this latch is engaged, hydraulic feed holes 156a, b of the end plate 142 are lined up with the hydraulic inlets 50a, 50b on the staple head as shown in Fig. 30B. At the same time, portions of the end plate surrounding u-shaped catches 148 slide beneath the undercut bosses 152. Pressing the plate compresses the face-sealing o-rings surrounding the hydraulic input ports 50a, 50b. Compression on the o-rings is maintained by engagement of the catches and the undercut bosses overhanging the end plate. To remove the stapler head from the housing, the stapler housing is twisted in a counterclockwise direction to disengage the end plate 142 from the rear surface 48a. The stapler shaft and handle may then be sterilized in preparation for mounting of a fresh stapler head.

Exemplary Procedure

[0130] One example of a method for using the system 10 will next be described in the context of formation of plications in stomach wall tissue, with particular reference to Figs. 31-33.

[0131] As an initial step (Fig. 2), endoscopic guide tube 23 is advanced into the stomach via the mouth and esophagus. The endoscope 22 is inserted into the endoscope channel in the stapler handle (not shown) and advanced down the lumen of the stapler handle. The stapler/endoscope are simultaneously passed through the endoscopic guide tube towards the stomach. Once the stapler and endoscope reach the gastroesophageal junction region of the stomach, the position of the stapler is maintained while the endoscope is advanced further into the stomach.

[0132] The stapler head 14 is advanced to the desired depth and location in the stomach. Using the articulation controls on the stapler handle, the angular orientation of the stapler head is adjusted to allow positioning of the stapler head 12 at the pre-identified target tissue as shown in Fig. 31 A. The opening 26 in the membrane 24 is positioned against the target tissue. The endoscope 22 is placed in a retroflexed position as shown.

[0133] The vacuum source 20 (Fig. 2) is coupled to the vacuum port on the handle external to the body, and vacuum pressure is applied to draw tissue 17 through the opening 26 and into the vacuum chamber defined by membrane 24 as shown in Figs. 31B and 32A. Acquisition of the target tissue will be readily identified endoscopically through the wall of transparent membrane 24 on the stapler head.

[0134] The fluid source (is shown) is coupled to the handle. Once it has been visually confirmed that a sufficient amount of tissue has been acquired, fluid is introduced to cause compression of the tissue and expansion of the arm assemblies 32 and membrane raiser 37 as shown in Figs. 32B and 31C. As can be seen, the expansion of the arm assemblies and the membrane allows a large volume of tissue to be acquired into the vacuum chamber and displaced further into the chamber during tissue compression. As noted earlier, the drawing in of tissue into the expanded chamber during operation, well "above" the staple holder and anvil in Figs. 32B and 32C, provides a relatively large margin of tissue around the stapled portion of the tissue, reducing the risk of tissue tearing or tissue-fold weakness near the stapled portion of the tissue. The captured tissue fold is indicated at 17a.

[0135] Once the tissue has been compressed, additional hydraulic fluid is introduced to cause stapling and cutting of the tissue as shown in Figs. 31D and 32C, forming a plication P, indicated at 17b in Fig. 33. The compression and stapling hydraulic sources are then deactivated to release fluid pressure within the hydraulic circuit. With the hydraulic pressure relieved, the spring wires of the membrane raiser 37 help to restore the stapler head 14 to its original streamlined configuration, al-

lowing the stapler head to be withdrawn from the tissue as shown in Fig. 31E. The stapler head may be articulated relative to the shaft to assist in moving the stapler head away from the plication P.

[0136] In a preferred plication configuration shown in Fig. 33 the staples 158 are arranged in two concentric rings of five staples, with the staple reinforcement device 83 retained by the staples and distributing forces around the staple pattern as shown. The plication P includes a hole H formed by the cutting element, through which various implants or anchors for various implants can be placed.

[0137] If multiple plications are needed, the stapler 12 is briefly withdrawn from the endoscopic guide tube and the staple cartridge is replaced in the manner described in connection with Figs. 25A - 25C. The procedure is repeated until all desired plications have been formed.

[0138] The system may be packaged with instructions for use instructing the user to use the various disclosed features to perform a stapling procedure using methods disclosed herein.

Alternate Embodiments

[0139] The basic architecture of the stapler disclosed above can be used as a foundation for other stapling tools. Figs. 34-35 show a modified stapler in which the membrane and membrane raiser have been removed, and in which the staple housing 28 has been modified for the attachment of tools. As shown in Fig. 34, the staple housing 28 includes a pair of grooves 160 proportioned to receive tools 162. Tools 162 may be seated in these grooves 160 and mounted to the staple housing as shown in Fig. 35. This attachment will provide for a stable base from which to actuate the tools. The tools may be self-articulating, or the staple housing 28 may be equipped with devices 164 for moving the tools between streamlined positions for insertion of the assembly into a body cavity, and a deployed position such as that shown in Fig. 35. Tools similar to those in Fig. 35 might be used for tissue acquisition, by reaching between the cartridge and anvil and used to engage tissue and pull the tissue into position between the cartridge and anvil so that it may be stapled, or otherwise affected by various features added to or in place of the anvil and cartridge. Procedures which may benefit from adaptation of the stapler include, but are not limited to gastropasty, stoma adjustment, polypectomy, lead placement, bleeding control, perforation or hole closure, biopsy and tumor removal.

[0140] The disclosed systems provide convenient embodiments for carrying out the disclosed compression and stapling functions. However, there are many other widely varying instruments or systems may alternatively be used within the scope of the present invention, as defined by the appended claims.

Claims

1. A surgical stapler device (14) for applying a staple to a tissue, comprising,
a staple member (25) having a staple housing (28) and a staple holder (78) ;
an anvil member (27) having an anvil housing (30) and an anvil (96) carried on the anvil housing (30), and
a drive assembly (29) including a drive member (68) operatively connected to the staple holder (78) for movement within the staple housing (28) from a retracted position to an extended position, and an arm assembly (32) operatively coupled to the staple and anvil members (25,27), such that movement of the drive member (68) from its retracted to its extended position is effective to (i) move the anvil member (27) toward the staple member (A2);
characterised in that the staple holder (78) is independently movable with respect to the staple housing; and **in that** the said movement of the drive member (68) from its retracted to its extended position is effective to (ii) move the staple holder with respect to the staple housing (78) toward the anvil (96) (A1).
2. The device of claim 1, wherein the anvil (96) is independently movable in the anvil housing (30), and the arm assembly (32) is operatively coupled to the anvil (96), such that movement of the drive member (68) from its retracted to its extended position is effective to (i) move the staple holder with respect to the staple housing toward the anvil (A1), (ii) move the anvil member toward the staple member (A2), and (iii) move the anvil with respect to the anvil housing toward the staple holder (A3).
3. The device of claim 2, wherein the anvil member (27) includes drive links (114) operatively connected to the arm assembly (32) for moving the anvil (96) toward the staple holder (78) (A3) as the drive assembly (106) is moved from its retracted to its extended position.
4. The device of claim 1, wherein the staple holder (78) and anvil (96) have confronting surfaces that define, in combination with arm assembly (32), a chamber (21), and the effect of movement of the drive member (68) from its retracted to its extended position is to squeeze tissue captured within the chamber to form a captured tissue fold.
5. The device of claim 4, wherein the chamber (21) is covered by a membrane (24), allowing tissue to be drawn into the chamber, with application of a vacuum to the membrane.
6. The device of claim 5, wherein the membrane (24)

has an opening (26) on one side thereof, for slicking tissue into the chamber, and movement of the drive member (68) from its retracted to extended positions, with application or vacuum to the chamber, is effective to draw tissue into the side of the chamber opposite the opening.

7. The device of claim 4, wherein the drive member includes a disc (68) that travels within the staple housing (25), and carries at least one pin (84) that moves within a slot (64) in the staple housing (25), limiting the extent of travel of the drive member toward its extended position to the extent of travel allowed for the pin (84) within the slot (64). 10
8. The device of claim 7, which further includes at least one arm spreader (113) pivotally connecting said disc to the arm assembly (32), for spreading the arm assembly outwardly as the disc travels from its retracted to its extended position. 20
9. The device of claim 1, wherein the drive member includes a drive piston (106) connected to a disc (68), and which further includes a staple piston (116) carried for movement within the drive piston (106) between retracted and extended positions and a staple pusher (76) adapted to engage one or more staples in the staple holder (78) and eject the one or more staples from the holder against the anvil (96), when the staple pusher is moved with the staple piston from a retracted to extended position. 25 30
10. The device of claim 9, for use with a staple holder (78) designed to hold an annular array of staples, and the staple pusher (78) is designed to engage and eject the array of staples in the holder simultaneously. 35
11. The device of claim 10, wherein the staple holder (78) is a replaceable staple cartridge adapted to be inserted into the staple housing to travel therein with the drive member (68) between retracted and extended positions. 40
12. The device of claim 7, wherein the disc (68) includes at least one axially extending post (84) adapted to engage the staple cartridge, with such received in the device, and prevent angular movement of the cartridge within the staple housing. 45
13. The device of claim 12, which further includes a cartridge-side reinforcing ring (83) disposed against the cartridge (78), and having openings (85) for receiving staples therethrough, for attaching the reinforcing ring (83) to the side of stapled tissue facing the cartridge. 50 55
14. The device of claim 9, which further includes a tissue

cutter (86) mounted on the staple piston (116), adapted to cut a hole in a tissue fold held between the staple holder (78) and anvil (96), as the tissue fold is being stapled by movement of the staple piston (116) from its retracted to its extended position.

15. The device of claim 13, wherein the anvil member includes a compressible cutting board (99a, 99d), allowing the cutter (86) to be advanced by movement of the staple piston (116) beyond its point of initial contact with the cutting board.
16. The device of claim 15, wherein the cutting board (99a) is formed of a material, such as silicone, that can be penetrated by the cutter. 15
17. The device of claim 15, wherein the cutting board (99d) is spring biased in the direction opposing movement of the cutter. 20
18. The device of claim 9, which further includes an alignment pin (160, 168) carried on one of the staple and anvil members, and a pin-receiving bushing (164, 170) carried on the other of the members, said pin and bushing being positioned such that movement of the staple piston toward its extended position causes the pin to mate with the bushing, thus to hold the two members in axial alignment as staples are ejected with further movement of the staple piston (116) toward its extended position. 30
19. The device of claim 18, wherein one of the pin and pin-receiving bushing is retractable under a spring bias. 35
20. The device of claim 1 which is carried at the distal end of a shaft (16) carrying hydraulic fluid to the device, wherein the staple and anvil members of the device are proximal and distal members, respectively, with the shaft being operatively connected to the proximal member of the device.
21. A medical instrument (10) for stapling stomach tissue comprising
the stapler device (14) of claim 1,
a shaft (16) having a proximal-end handle and a distal end,
an articulating section (128) connecting the distal end of the shaft to the first, proximal member of the stapling device, and
a hydraulic fluid line (130) contained within the shaft effective for carrying hydraulic fluid under a pressure of at least about 69 Bar (1,000 psi) to the drive mechanism in the staple member, wherein the hydraulic fluid line within the articulating section has a coiled or sinusoidal configuration, to accommodate off-axis movement of the device with respect to the shaft.

22. The instrument of claim 21, wherein the articulating section (128) has a spine (128) formed of a plurality of links (132) formed over the coiled or sinusoidal portion of the hydraulic line (130).

23. The instrument of claim 21, wherein the drive member (68) is hydraulically driven and the stapler device includes a hydraulically driven staple piston (116), said shaft (16) and articulating section (128) carry separate hydraulic lines (130) for the drive member and staple piston, and the lines (130) have an interleaved coil configuration within the articulating section.

24. A tissue capture device for capturing and immobilizing a tissue fold, comprising the stapler device of claim 1;
the staple member (25) having a first tissue-contact plate (78) independently movable within the staple member housing,
the anvil member (27) having a second tissue-contact plate (96) independently movable within the anvil member housing,
wherein movement of the drive member (68) from its retracted to its extended position is effective to (i) move the first tissue-contact plate (78) within the staple member housing toward the second tissue contact plate (96) (A1), and (ii) move the anvil member (27) toward the staple member (A2),
an elastomeric membrane extending between the two members and defining a tissue capture chamber between the two tissue-contact plates, and having an opening therein for drawing tissue into the chamber, upon application of a vacuum to the chamber, and
an expansion member (37) operatively coupled to the two members for expanding the chamber membrane outwardly, on the side of the membrane opposite said opening, upon movement of the drive member from its retracted to its expanded condition, causing additional tissue to be drawn into the chamber between the two tissue-contact plates, until the tissue is clamped between two tissue-contact plates.

25. A medical instrument (10) for stapling stomach tissue comprising
the stapler device (14) of claim 1, a shaft (16) having a proximal-end handle and a distal end, an articulating section (128) connecting the distal end of the shaft to the first, proximal member of the stapling device, and a hydraulic fluid line (130) contained within the shaft capable of carrying hydraulic fluid under a pressure of up to 103 Bar (1,500 psi) or greater, to the drive mechanism (106) in the staple member (25), wherein the hydraulic fluid line within the articulating section has a coiled or sinusoidal configuration, to accommodate off-axis movement of the device with respect to the shaft.

26. The instrument of claim 25, wherein the articulating section (128) has a spine (128) formed of a plurality of links (132) formed over the coiled or sinusoidal portion of the hydraulic line (130).

27. The instrument of claim 25, wherein the drive member (68) is hydraulically driven and the stapler device includes a hydraulically driven staple piston (116), and the shaft (16) and articulating section (128) carry separate hydraulic lines (130) for the drive member and staple piston, and the lines (130) have an interleaved coil configuration within the articulating section.

28. The medical instrument (12) of claim 21 having a hydraulically driven device (14) for capturing and/or stapling tissue and an elongate shaft (16) for delivering hydraulic fluid to the device, a snap-on connection between the distal end of a shaft and the proximal face of a staple member in a hydraulically activated staple device, comprising
at the distal shaft end, a plate having one or more openings for supplying hydraulic fluid under pressure from hydraulic lines attached to the openings at the input side of the plate, and an engagement edge adjacent each opening,
a plate-receiving face on the staple device including correspondingly positioned openings for receiving fluid into the device, where the openings communicate at the output side of the face with hydraulic supply lines within the staple device, and an undercut boss adjacent each opening, such that when the shaft plate is placed against the staple-device face, and rotated slightly to wedge the engagement edge (s) on the shaft plate within the undercut regions of the bosses of the device plate, an O-ring located between the plate and face at each opening is compressed, thus sealing the connection between the aligned openings, and
a lock mechanism on the shaft plate that engages the device face to lock the plate against rotational movement once the fluid-supply opening are received.

29. The device of claim 8, wherein movement of the pin within its slot in a retracted to extended position is effective to move the arm assembly outwardly, wherein the arm spreader provides a truss-like support between the staple body and the associated assembly arm.

Patentansprüche

1. Chirurgische Klammervorrichtung (14) zum Anbringen einer Klammer an ein Gewebe, die Folgendes umfasst,
ein Klammerelement (25) mit einem Klammergehäu-

- se (28) und einem Klammerhalter (78);
 ein Ambosselement (27) mit einem Ambossgehäuse (30) und einem auf dem Ambossgehäuse (30) getragenen Amboss (96), und
 eine Antriebsbaugruppe (29), umfassend ein Antriebselement (68), das zur Bewegung in dem Klammergehäuse (28) von einer eingefahrenen Stellung zu einer ausgefahrenen Stellung wirksam mit dem Klammerhalter (78) verbunden ist, und eine Armbaugruppe (32), die wirksam an das Klammerelement und das Ambosselement (25, 27) gekoppelt ist, so dass Bewegung des Antriebselements (68) von seiner eingefahrenen zu seiner ausgefahrenen Stellung dazu wirksam ist: (i) das Ambosselement (27) auf das Klammerelement zuzubewegen (A2);
dadurch gekennzeichnet, dass der Klammerhalter (78) unabhängig gegenüber dem Klammergehäuse bewegbar ist; und dass die genannte Bewegung des Antriebselements (68) von seiner eingefahrenen zu seiner ausgefahrenen Stellung dazu wirksam ist: (ii) den Klammerhalter gegenüber dem Klammergehäuse (78) auf den Amboss (96) zuzubewegen (A1).
2. Vorrichtung nach Anspruch 1, wobei der Amboss (96) unabhängig im Ambossgehäuse (30) bewegbar ist und die Armbaugruppe (32) wirksam an den Amboss (96) gekoppelt ist, so dass Bewegung des Antriebselements (68) von seiner eingefahrenen zu seiner ausgefahrenen Stellung dazu wirksam ist: (i) den Klammerhalter gegenüber dem Klammergehäuse auf den Amboss zuzubewegen (A1), (ii) das Ambosselement auf das Klammerelement zuzubewegen (A2), und (iii) den Amboss gegenüber dem Ambossgehäuse auf den Klammerhalter zuzubewegen (A3).
 3. Vorrichtung nach Anspruch 2, wobei das Ambosselement (27) Antriebsgestänge (114) umfasst, die wirksam mit der Armbaugruppe (32) verbunden sind, um den Amboss (96) auf den Klammerhalter (78) zuzubewegen (A3), wenn die Antriebsbaugruppe (106) von ihrer eingefahrenen zu ihrer ausgefahrenen Stellung bewegt wird.
 4. Vorrichtung nach Anspruch 1, wobei der Klammerhalter (78) und der Amboss (96) einander zugewandte Oberflächen haben, die in Kombination mit der Armbaugruppe (32) eine Kammer (21) definieren, und die Wirkung von Bewegung des Antriebselements (68) von seiner eingefahrenen zu seiner ausgefahrenen Stellung darin besteht, in der Kammer gefangenes Gewebe zu quetschen, um eine gefangene Gewebefalte zu bilden.
 5. Vorrichtung nach Anspruch 4, wobei die Kammer (21) von einer Membran (24) bedeckt ist, so dass zugelassen wird, dass mit Anlegen eines Unterdrucks an die Membran Gewebe in die Kammer gezogen wird.
 6. Vorrichtung nach Anspruch 5, wobei die Membran (24) eine Öffnung (26) an einer Seite derselben hat, um Gewebe in die Kammer zu saugen, und Bewegung des Antriebselements (68) von seiner eingefahrenen zu seiner ausgefahrenen Stellung, mit Anlegen von Unterdruck an die Kammer, wirksam ist, Gewebe in die der Öffnung gegenüberliegende Seite der Kammer zu ziehen.
 7. Vorrichtung nach Anspruch 4, wobei das Antriebselement eine Scheibe (68) umfasst, die in dem Klammergehäuse (25) verfahren wird und mindestens einen Stift (84) trägt, der sich in einem Schlitz (64) in dem Klammergehäuse (25) bewegt und das Ausmaß des Verfahrens des Antriebselements auf seine ausgefahrne Stellung zu auf das für den Stift (84) in dem Schlitz (64) zugelassene Ausmaß des Verfahrens begrenzt.
 8. Vorrichtung nach Anspruch 7, die weiter mindestens einen Armspreizer (113) umfasst, der die genannte Scheibe drehbar mit der Armbaugruppe (32) verbindet, um die Armbaugruppe nach außen zu spreizen, wenn die Scheibe von ihrer eingefahrenen zu ihrer ausgefahrenen Stellung verfahren wird.
 9. Vorrichtung nach Anspruch 1, wobei das Antriebselement einen Antriebskolben (106) umfasst, der mit einer Scheibe (68) verbunden ist und das weiter einen Klammerkolben (116) umfasst, der zur Bewegung mit dem Antriebskolben (106) zwischen eingefahrenen und ausgefahrenen Stellungen getragen wird und einen Klammerdrücker (76) umfasst, der dazu angepasst ist, mit einer oder mehreren Klammern in dem Klammerhalter (78) in Eingriff zu treten und die eine oder die mehreren Klammern aus dem Halter gegen den Amboss (96) auszustoßen, wenn der Klammerdrücker mit dem Klammerkolben von einer eingefahrenen zu einer ausgefahrenen Stellung bewegt wird.
 10. Vorrichtung nach Anspruch 9 zur Verwendung mit einem Klammerhalter (78), der dazu ausgeführt ist, eine ringförmige Gruppierung von Klammern zu halten, und wobei der Klammerdrücker (78) dazu ausgeführt ist, gleichzeitig mit der Gruppierung von Klammern in dem Halter in Eingriff zu treten und sie auszustoßen.
 11. Vorrichtung nach Anspruch 10, wobei es sich bei dem Klammerhalter (78) um eine austauschbare Klammerpatrone handelt, die dazu angepasst ist, in das Klammergehäuse eingesetzt zu werden, um darin mit dem Antriebselement (68) zwischen einer eingefahrenen und einer ausgefahrenen Stellung verfahren zu werden.

12. Vorrichtung nach Anspruch 7, wobei die Scheibe (68) mindestens einen sich axial erstreckenden Pfosten (84) umfasst, der dazu angepasst ist, mit der Klammerpatrone in Eingriff zu treten, wenn diese in der Vorrichtung aufgenommen ist, und die Winkelbewegung der Patrone in dem Klammergehäuse zu verhindern.
13. Vorrichtung nach Anspruch 12, die weiter einen patronenseitigen Verstärkungsring (83) umfasst, der an der Patrone (78) anliegend angeordnet ist und Öffnungen (85) zum Aufnehmen von Klammern dadurch hindurch hat, um den Verstärkungsring (83) an der der Patrone zugewandten Seite von geklammertem Gewebe zu befestigen.
14. Vorrichtung nach Anspruch 9, die weiter eine Gewebeschneidvorrichtung (86) umfasst, die an dem Klammerkolben (116) angebracht ist und dazu angepasst ist, ein Loch in eine Gewebefalte zu schneiden, die zwischen dem Klammerhalter (78) und dem Amboss (96) gehalten wird, wenn die Gewebefalte durch Bewegung des Klammerkolbens (116) von seiner eingefahrenen in seine ausgefahrene Stellung geklammert wird.
15. Vorrichtung nach Anspruch 13, wobei das Ambosselement eine zusammendrückbare Schneidplatte (99a, 99d) umfasst, die zulässt, dass die Schneidvorrichtung (86) durch Bewegung des Klammerkolbens (116) über den Punkt ihres ersten Kontakts mit der Schneidplatte hinaus voranbewegt wird.
16. Vorrichtung nach Anspruch 15, wobei die Schneidplatte (99a) aus einem Material, wie etwa Silikon, gebildet ist, in das die Schneidvorrichtung eindringen kann.
17. Vorrichtung nach Anspruch 15, wobei die Schneidplatte (99d) in der der Bewegung der Schneidvorrichtung entgegengesetzten Richtung eine Feder-
vorspannung aufweist.
18. Vorrichtung nach Anspruch 9, die weiter einen Führungszapfen (160, 168), der auf dem Klammerelement oder dem Ambosselement getragen wird, und eine Zapfenaufnahmebuchse (164, 170), die auf dem jeweils anderen der Elemente getragen wird, umfasst, wobei der genannte Zapfen und die genannte Buchse derart positioniert sind, dass Bewegung des Klammerkolbens auf seine ausgefahrene Stellung zu dazu führt, dass der Zapfen mit der Buchse zusammenpasst, um so die zwei Elemente in axialer Ausrichtung zu halten, wenn Klammern mit weiterer Bewegung des Klammerkolbens (116) auf seine ausgefahrene Stellung zu ausgestoßen werden.
19. Vorrichtung nach Anspruch 18, wobei der Zapfen oder die Zapfenaufnahmebuchse unter einer Feder-
vorspannung einfahrbar ist.
20. Vorrichtung nach Anspruch 1, die am distalen Ende eines Schafts (16) getragen wird, der Hydraulikfluid zu der Vorrichtung leitet, wobei es sich bei dem Klammerelement und dem Ambosselement der Vorrichtung um ein proximales bzw. ein distales Element handelt, wobei der Schaft wirksam mit dem proximalen Element der Vorrichtung verbunden ist.
21. Medizinisches Instrument (10) zum Klammern von Magengewebe das Folgendes umfasst die Klammervorrichtung (14) nach Anspruch 1, einen Schaft (16) mit einem Griff am proximalen Ende und einem distalen Ende einen Gelenkabschnitt (128), der das distale Ende des Schafts mit dem ersten proximalen Element der Klammervorrichtung verbindet, und eine in dem Schaft enthaltene Hydraulikfluidleitung (130), die dazu wirksam ist, unter einem Druck von mindestens ungefähr 69 bar (1000 psi) stehendes Hydraulikfluid zu dem Antriebsmechanismus in dem Klammerelement zu leiten, wobei die Hydraulikfluidleitung in dem Gelenkabschnitt eine gewickelte oder sinusförmige Ausbildung hat, um die nicht in Achsenrichtung erfolgende Bewegung der Vorrichtung gegenüber dem Schaft zu berücksichtigen.
22. Instrument nach Anspruch 21, wobei der Gelenkabschnitt (128) ein Rückgrat (128) hat, das aus mehreren über dem gewickelten oder sinusförmigen Abschnitt der Hydraulikleitung (130) gebildeten Gliedern (132) gebildet ist.
23. Instrument nach Anspruch 21, wobei das Antriebs-
element (68) hydraulisch angetrieben ist und die Klammervorrichtung einen hydraulisch angetriebenen Klammerkolben (116) umfasst, wobei der genannte Schaft (16) und der Gelenkabschnitt (128) getrennte Hydraulikleitungen (130) für das Antriebs-
element und den Klammerkolben tragen und die Leitungen (130) in dem Gelenkabschnitt eine Ausbildung mit überlappenden Wicklungen aufweisen.
24. Gewebefangvorrichtung zum Einfangen und Immobilisieren einer Gewebefalte, umfassend die Klammervorrichtung nach Anspruch 1; wobei das Klammerelement (25) eine erste Gewebekontaktplatte (78) aufweist, die unabhängig in dem Klammerelementgehäuse bewegbar ist, wobei das Ambosselement (27) eine zweite Gewebekontaktplatte (96) aufweist, die unabhängig in dem Ambosselementgehäuse bewegbar ist, wobei Bewegung des Antriebs-
elements (68) von seiner eingefahrenen zu seiner ausgefahrenen Stellung dazu wirksam ist (i) die erste Gewebekontakt-

platte (78) in dem Klammerelementgehäuse auf die zweite Gewebekontaktplatte (96) zuzubewegen (A1), und (ii) das Ambosselement (27) auf das Klammerelement (A2) zuzubewegen, wobei sich eine Elastomermembran zwischen den zwei Elementen erstreckt und eine Gewebefangkammer zwischen den zwei Gewebekontaktplatten definiert und die eine Öffnung darin hat, um bei Anlegen eines Unterdrucks an die Kammer, Gewebe in die Kammer zu ziehen, und wobei ein Ausdehnungselement (37) wirksam an die zwei Elemente gekoppelt ist, um die Kammermembran auf der der genannten Öffnung gegenüberliegenden Seite der Membran bei Bewegung des Antriebselements von seinem eingefahrenen zu seinem ausgefahrenen Zustand nach außen auszuweichen und zu bewirken, dass zusätzliches Gewebe in die Kammer zwischen den zwei Gewebekontaktplatten gezogen wird, bis das Gewebe zwischen den zwei Gewebekontaktplatten eingeklemmt ist.

25. Medizinisches Instrument (10) zum Klammern von Magengewebe das Folgendes umfasst:

die Klammervorrichtung (14) nach Anspruch 1, einen Schaft (16) mit einem Griff an einem proximalen Ende und einem distalen Ende, einen Gelenkabschnitt (128), der das distale Ende des Schafts mit dem ersten, proximalen Element der Klammervorrichtung verbindet, und eine Hydraulikfluidleitung (130), die in dem Schaft enthalten ist und fähig ist, unter einem Druck von bis zu 103 bar (1500 psi) oder mehr stehendes Hydraulikfluid zu dem Antriebsmechanismus (106) in dem Klammerelement (25) zu leiten, wobei die Hydraulikfluidleitung in dem Gelenkabschnitt eine gewickelte oder sinusförmige Ausbildung aufweist, um die nicht in Achsenrichtung erfolgende Bewegung der Vorrichtung gegenüber dem Schaft zu berücksichtigen.

26. Instrument nach Anspruch 25, wobei der Gelenkabschnitt (128) ein Rückgrat (128) hat, das aus mehreren über dem gewickelten oder sinusförmigen Abschnitt der Hydraulikleitung (130) gebildeten Gliedern (132) gebildet ist.

27. Instrument nach Anspruch 25, wobei das Antriebselement (68) hydraulisch angetrieben ist und die Klammervorrichtung einen hydraulisch angetriebenen Klammerkolben (116) umfasst, und der Schaft (16) und der Gelenkabschnitt (128) getrennte Hydraulikleitungen (130) für das Antriebselement und den Klammerkolben tragen und die Leitungen (130) in dem Gelenkabschnitt eine Ausbildung mit überlappenden Wicklungen aufweisen.

28. Medizinisches Instrument (12) nach Anspruch 21 mit

einer hydraulisch angetriebenen Vorrichtung (14) zum Einfangen und/oder Klammern von Gewebe und einem langgestreckten Schaft (16) zum Zuführen von Hydraulikfluid an die Vorrichtung, einer einrastenden Verbindung zwischen dem distalen Ende eines Schafts und der proximalen Stirnfläche eines Klammerelements einer hydraulisch betätigten Klammervorrichtung, umfassend eine Platte am distalen Schaftende, mit einer oder mehreren Öffnungen zum Zuführen von unter Druck stehendem Hydraulikfluid aus Hydraulikleitungen, die an den Öffnungen an der Eingangsseite der Platte befestigt sind, und mit einer Eingriffskante angrenzend an jede Öffnung, eine Plattenaufnahme-Stirnfläche an der Klammervorrichtung, umfassend entsprechend positionierte Öffnungen zum Empfangen von Fluid in der Vorrichtung, wobei die Öffnungen an der Ausgangsseite der Stirnfläche mit Hydraulik-Versorgungsleitungen in der Klammervorrichtung in Verbindung stehen, und einen hinterschnittenen Ansatz angrenzend an jede Öffnung, so dass wenn die Schaftplatte gegen die Stirnfläche der Klammervorrichtung platziert wird und geringfügig gedreht wird, um die Eingriffskante (n) an der Schaftplatte in den hinterschnittenen Regionen der Ansätze der Vorrichtungsplatte zu verkeilen, ein zwischen der Platte und der Stirnfläche an jeder Öffnung befindlicher O-Ring zusammengedrückt wird, wodurch die Verbindung zwischen den aufeinander ausgerichteten Öffnungen abgedichtet wird, und einen Sperrmechanismus an der Schaftplatte, der mit der Vorrichtungsfläche in Eingriff tritt, um die Platte gegen Drehbewegung zu sperren, nachdem die Fluidversorgungsöffnung empfangen wurde.

29. Vorrichtung nach Anspruch 8, wobei Bewegung des Stifts in seinem Schlitz in einer eingefahrenen zu einer ausgefahrenen Stellung wirksam ist, um die Armspreizer eine fachwerkartige Abstützung zwischen dem Klammerkörper und dem zugehörigen Baugruppenarm vorsieht.

Revendications

- Dispositif d'agrafage chirurgical (14) pour l'application d'une agrafe sur un tissu, comprenant, un élément d'agrafe (25) comprenant un boîtier d'agrafe (28) et un porte-agrafe (78); un élément d'enclume (27) comprenant un boîtier d'enclume (30) et une enclume (96) supportée par le boîtier d'enclume (30) et; un mécanisme d'entraînement (29) comprenant un ensemble d'entraînement (68) opérationnellement relié au porte-agrafe (78) pour un déplacement au sein du boîtier d'agrafe (28) depuis une position ré-

- tractée à une position déployée, et un ensemble de bras (32) opérationnellement relié à l'agrafe et aux éléments d'enclumes (25, 27), de manière à ce que le déplacement de l'élément d'entraînement (68) de sa position rétractée à sa position déployée permette (i) de déplacer l'élément d'enclume (27) en direction de l'élément d'agrafe (A2);
- caractérisé en ce que** le porte-agrafe (78) se déplace indépendamment par rapport au boîtier d'agrafe et **en ce que** ledit déplacement de l'élément d'entraînement (68) de sa position rétractée à sa position déployée permet (ii) de déplacer le porte-agrafe par rapport au boîtier d'agrafe (78) en direction de l'enclume (96) (A1).
2. Dispositif selon la revendication 1, dans lequel l'enclume (96) se déplace indépendamment dans le boîtier d'enclume (30) et l'ensemble de bras (32) est opérationnellement relié à l'enclume (96) de manière à ce que le déplacement de l'élément d'entraînement (68) de sa position rétractée à sa position déployée permette de (i) déplacer le porte-agrafe par rapport au boîtier d'agrafe en direction de l'enclume (A1), (ii) de déplacer l'élément d'enclume en direction de l'élément d'agrafe (A2) et (iii) de déplacer l'enclume par rapport au boîtier d'enclume en direction du porte-agrafe (A3).
 3. Dispositif selon la revendication 2, dans lequel l'élément d'enclume (27) comprend des maillons d'entraînement (14) opérationnellement reliés à l'ensemble de bras (32) pour déplacer l'enclume (96) en direction du porte-agrafe (78) (A3) à mesure que l'ensemble d'entraînement (106) est déplacé de sa position rétractée à sa position déployée.
 4. Dispositif selon la revendication 1, dans lequel le porte-agrafe (78) et l'enclume (96) ont des surfaces se faisant face qui définissent, en association avec l'ensemble de bras (32), une chambre (21) et l'effet du déplacement de l'élément d'entraînement (68) de sa position rétractée à sa position déployée est de pincer le tissu capturé au sein de la chambre pour former un pli de tissu capturé.
 5. Dispositif selon la revendication 4, dans lequel la chambre (21) est couverte d'une membrane (24) permettant au tissu d'être tiré dans la chambre par l'application d'un vide à la membrane.
 6. Dispositif selon la revendication 5, dans lequel la membrane (24) comprend une ouverture (26) sur un côté pour aspirer le tissu dans la chambre et le déplacement de l'élément d'entraînement (68) de sa position rétractée à sa position déployée, par l'application d'un vide dans la chambre, permet de tirer le tissu dans le côté de la chambre face à l'ouverture.
 7. Dispositif selon la revendication 4, dans lequel l'élément d'entraînement comprend un disque (68) qui se déplace au sein du boîtier d'agrafe (25) et transporte au moins une broche (84) qui se déplace au sein d'une fente (64) dans le boîtier d'agrafe (25) limitant la course de déplacement de l'élément d'entraînement en direction de sa position déployée à la course de déplacement permise par la broche (84) au sein de la fente (64).
 8. Dispositif selon la revendication 7, qui comprend en outre au moins un écarteur de bras (113) reliant par pivot ledit disque à l'ensemble de bras (32) pour écarter l'ensemble de bras vers l'extérieur à mesure que le disque se déplace de sa position rétractée à sa position déployée.
 9. Dispositif selon la revendication 1, dans lequel l'élément d'entraînement comprend un piston d'entraînement (106) relié à un disque (68) et qui comprend en outre un piston d'agrafage (116) supporté en vue d'un déplacement au sein du piston d'entraînement (106) entre les positions rétractées et déployées et un poussoir d'agrafes (76) adapté pour engager une ou plusieurs agrafes dans le porte-agrafe (78) et pour éjecter la une ou plusieurs agrafes du porte-agrafe contre l'enclume (96) lorsque le poussoir d'agrafes est déplacé avec le piston d'agrafage d'une position rétractée à une position déployée.
 10. Dispositif selon la revendication 9, destiné à être utilisé avec un porte-agrafe (78) conçu pour contenir une matrice annulaire d'agrafes, et le poussoir d'agrafes (78) est conçu pour engager et éjecter simultanément la matrice d'agrafes contenue dans le porte-agrafe.
 11. Dispositif selon la revendication 10, dans lequel le porte-agrafe (78) est une cartouche d'agrafes remplaçable adaptée pour être insérée dans le boîtier d'agrafe pour se déplacer en son sein avec l'élément d'entraînement (68) se trouvant entre les positions rétractées et déployées.
 12. Dispositif selon la revendication 7, dans lequel le disque (68) comprend au moins un montant s'étendant axialement (84) adapté pour placer en prise la cartouche d'agrafes avec les agrafes reçues dans le dispositif et pour empêcher le mouvement angulaire de la cartouche à l'intérieur du boîtier d'agrafe.
 13. Dispositif selon la revendication 12, qui comprend en outre un anneau de renforcement du côté de la cartouche (83) disposé contre la cartouche (78) et possédant des ouvertures (85) à travers lesquelles les agrafes sont reçues de manière à attacher l'anneau de renforcement (83) au côté du tissu agrafé faisant face à la cartouche.

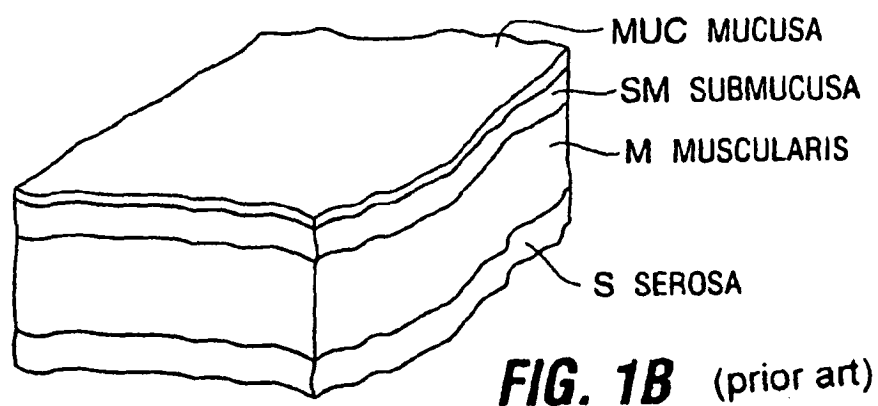
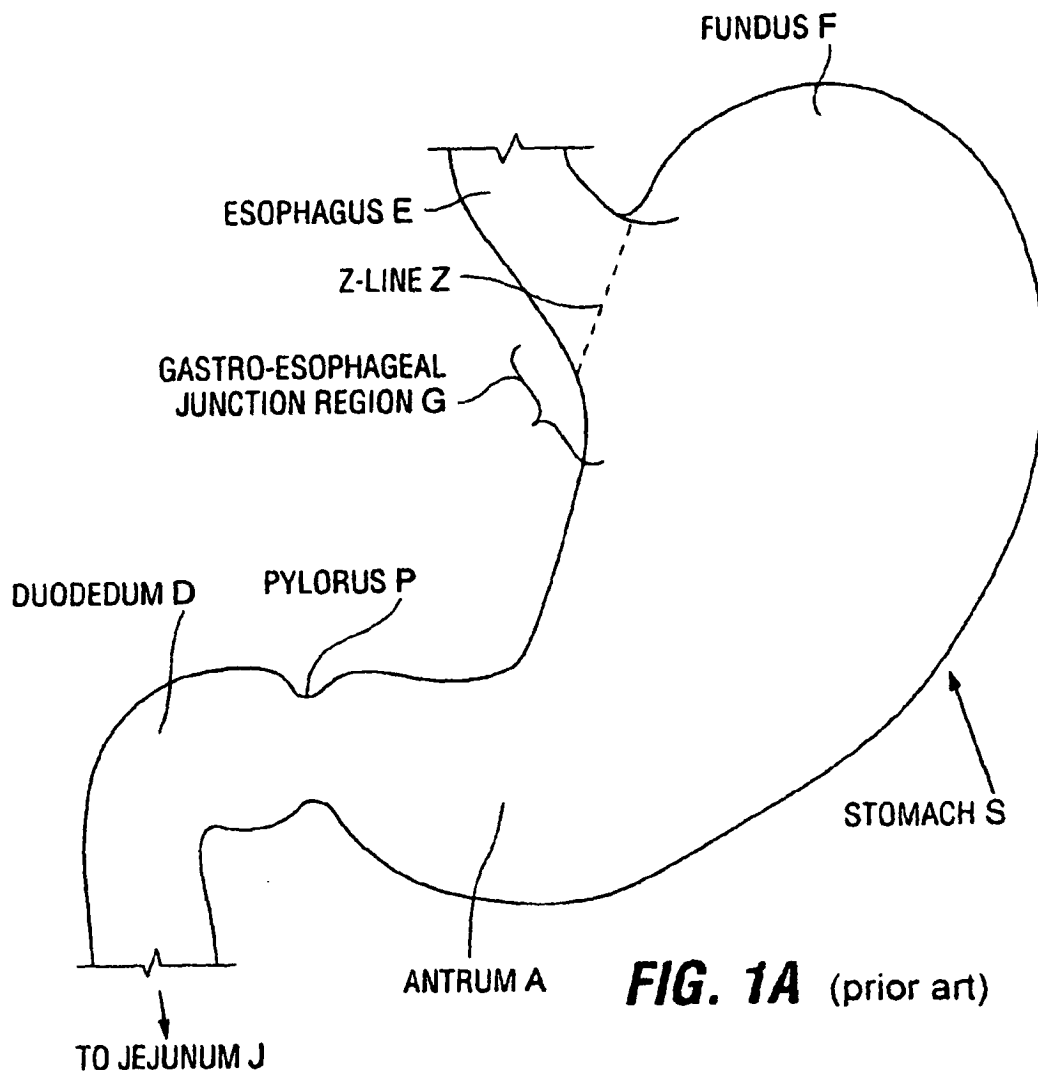
14. Dispositif selon la revendication 9, comprenant en outre un coupeur de tissu (86) monté sur le piston d'agrafage (116), adapté pour couper un évidement dans un pli de tissu maintenu entre le porte-agrafes (78) et l'enclume (96) à mesure que le pli de tissu est agrafé par le déplacement du piston d'agrafage (116) de sa position rétractée à sa position déployée. 5
15. Dispositif selon la revendication 13, dans lequel l'élément d'enclume comprend une planche à découper compressible (99a, 99d), permettant au coupeur (86) d'être avancé par le déplacement du piston d'agrafage (116) au-delà de son point de contact initial avec la planche à découper. 10
16. Dispositif selon la revendication 15, dans lequel la planche à découper (99a) est formée d'un matériau tel que la silicone qui peut être pénétré par le coupeur. 15
17. Dispositif selon la revendication 15, dans lequel la planche à découper (99d) est sollicitée par un ressort dans un sens de déplacement opposé à celui du coupeur. 20
18. Dispositif selon la revendication 9, qui comprend en outre une broche d'alignement (160, 168) supportée sur l'un des éléments d'agrafes et d'enclume, et un coussinet de réception de la broche (164, 170) supporté sur l'autre des éléments, lesdits broche et coussinet étant positionnés de telle sorte que le déplacement du piston d'agrafage en direction de sa position déployée cause l'accouplement de la broche avec le coussinet de manière à maintenir les deux éléments en alignement axial à mesure que les agrafes sont éjectées avec un déplacement additionnel du piston d'agrafage (116) en direction de sa position déployée. 25
19. Dispositif selon la revendication 18, dans lequel un de la broche et du coussinet de réception de la broche est rétractable sous une sollicitation par ressort. 30
20. Dispositif selon la revendication 1, qui est supporté à l'extrémité distale d'un arbre (16) transportant du fluide hydraulique au dispositif, dans lequel les éléments d'agrafe et d'enclume du dispositif sont des éléments proximaux et distaux, respectivement, avec l'arbre étant opérationnellement relié à l'élément proximal du dispositif. 35
21. Instrument médical (10) destiné à l'agrafage de tissus de l'estomac comprenant le dispositif d'agrafage (14) selon la revendication 1, un arbre (16) possédant une poignée d'extrémité proximale et une partie d'extrémité distale, une section de jonction (128) reliant l'extrémité distale de l'arbre au premier élément proximal du dispositif d'agrafage, et 40
- une conduite de fluide hydraulique (130) contenue à l'intérieur de l'arbre permettant de transporter le fluide hydraulique sous une pression d'au moins environ 69 bar (1000 psi) jusqu'au mécanisme d'entraînement contenu dans l'élément d'agrafe, dans lequel la conduite de fluide hydraulique au sein de la section de jonction a une configuration à serpents ou sinusoïdale pour prendre en charge le déplacement hors de l'axe du dispositif par rapport à l'arbre. 45
22. Instrument selon la revendication 21, dans lequel la section de jonction (128) possède une artère dorsale (128) comportant une pluralité de liaisons (132) formée pardessus la partie à serpents ou sinusoïdale de la conduite hydraulique (130). 50
23. Instrument selon la revendication 21, dans lequel l'élément d'entraînement (68) est à entraînement hydraulique et le dispositif d'agrafage comprend un piston d'agrafage à entraînement hydraulique (116), ledit arbre (16) et la section de jonction (128) supportant des conduites hydrauliques distinctes (130) de l'élément d'entraînement et du piston d'agrafage, et les conduites (130) ayant une configuration à serpents imbriqués au sein de la section de jonction. 55
24. Dispositif de capture de tissus destiné à la capture et à l'immobilisation d'un pli de tissu, comprenant le dispositif d'agrafage selon la revendication 1 ; l'élément d'agrafe (25) possédant une première plaque de contact du tissu (78) se déplaçant indépendamment au sein du boîtier de l'élément d'agrafe, un élément d'enclume (27) possédant une deuxième plaque de contact du tissu (96) se déplaçant indépendamment au sein du boîtier d'enclume, dans lequel le déplacement de l'élément d'entraînement (68) de sa position rétractée à sa position déployée permet (i) de déplacer la première plaque de contact du tissu (78) à l'intérieur du boîtier de l'élément d'agrafe en direction de la deuxième plaque de contact du tissu (96) (A1) et (ii) de déplacer l'élément d'enclume (27) en direction de l'élément d'agrafe (A2), 50
- une membrane élastomérique s'étendant entre les deux éléments et définissant une chambre de capture du tissu entre les deux plaques de contact du tissu et possédant une ouverture en son sein pour tirer le tissu dans la chambre lors de l'application d'un vide à la chambre, et 55
- un élément d'expansion (37) opérationnellement relié aux deux éléments pour l'expansion de la membrane de la chambre vers l'extérieur, du côté de la membrane se trouvant en face de ladite ouverture, lors du déplacement de l'élément d'entraînement de sa condition rétractée à sa condition déployée, causant le tirage de tissu additionnel dans la chambre

entre les deux plaques de contact du tissu, jusqu'à ce que le tissu soit pincé entre les deux plaques de contact du tissu.

25. Instrument médical (10) destiné à l'agrafage de tissus de l'estomac comprenant le dispositif d'agrafage (14) selon la revendication 1, un arbre (16) possédant une poignée d'extrémité proximale et une partie d'extrémité distale, une section de jonction (128) reliant l'extrémité distale de l'arbre au premier élément proximal du dispositif d'agrafage et une conduite de fluide hydraulique (130) contenue à l'intérieur de l'arbre apte à transporter le liquide hydraulique sous une pression de jusqu'à 103 bar (1500 psi) ou plus, jusqu'au mécanisme d'entraînement (106) contenu dans l'élément d'agrafe (25), dans lequel la conduite de fluide hydraulique au sein de la section de jonction possède une configuration à serpentins ou sinusoïdale pour prendre en charge un déplacement hors de l'axe du dispositif par rapport à l'arbre. 5 10 15 20
26. Instrument selon la revendication 25, dans lequel la section de jonction (128) possède une arête dorsale (128) comportant une pluralité de liaisons (132) formées pardessus la partie à serpentins ou sinusoïdale de la conduite hydraulique (130). 25
27. Instrument selon la revendication 25, dans lequel l'élément d'entraînement (68) est à entraînement hydraulique et le dispositif d'agrafage comprend un piston d'agrafage (116) à entraînement hydraulique et l'arbre (16) et la section de jonction (128) supporte des conduites hydrauliques distinctes (130) de l'élément d'entraînement et du piston d'agrafage (130) et les conduites (130) possèdent une configuration à serpentins imbriqués à l'intérieur de la section de jonction. 30 35
28. Instrument médical (12) selon la revendication 21 comportant un dispositif à entraînement hydraulique (14) destiné à la capture et/ou à l'agrafage de tissus et un arbre allongé (16) pour la délivrance de fluide hydraulique au dispositif, un raccord encliquetable entre l'extrémité distale d'un arbre et la face proximale d'un élément d'agrafe compris dans un dispositif d'agrafage à activation hydraulique, comprenant à l'extrémité distale de l'arbre, une plaque comportant une ou plusieurs ouvertures destinées à l'alimentation de fluide hydraulique sous pression depuis des conduites hydrauliques fixées aux ouvertures sur le côté d'entrée de la plaque et un bord d'engagement adjacent à chaque ouverture, une face de réception de la plaque sur le dispositif d'agrafage comportant des ouvertures positionnées de façon correspondante pour la réception du fluide dans le dispositif, dans laquelle les ouvertures communiquent sur le côté de sortie de la face avec les 40 45 50 55

conduites d'alimentation hydrauliques contenues à l'intérieur du dispositif d'agrafage et avec une protubérance en contre-dépouille adjacente à chaque ouverture, de sorte que lorsque la plaque de l'arbre est placée contre la face du dispositif d'agrafage et qu'elle est mise légèrement en rotation pour caler le/les bord(s) d'engagement sur la plaque de l'arbre au sein des zones en contre-dépouille des protubérances de la plaque du dispositif, un joint torique localisé entre la plaque et la face à chaque ouverture est comprimé, scellant ainsi la connexion entre les ouvertures alignées et un mécanisme de verrouillage se trouvant sur la plaque de l'arbre s'engage dans la face du dispositif pour verrouiller la plaque contre l'action rotative une fois que l'alimentation en fluide est reçue des orifices.

29. Dispositif selon la revendication 8, dans lequel le déplacement de la broche à l'intérieur de sa fente en position rétractée à déployée permet de déplacer l'assemblage du bras vers l'extérieur, dans lequel l'écarteur de bras fournit un support en treillis entre le corps de l'agrafe et l'assemblage de bras associé.



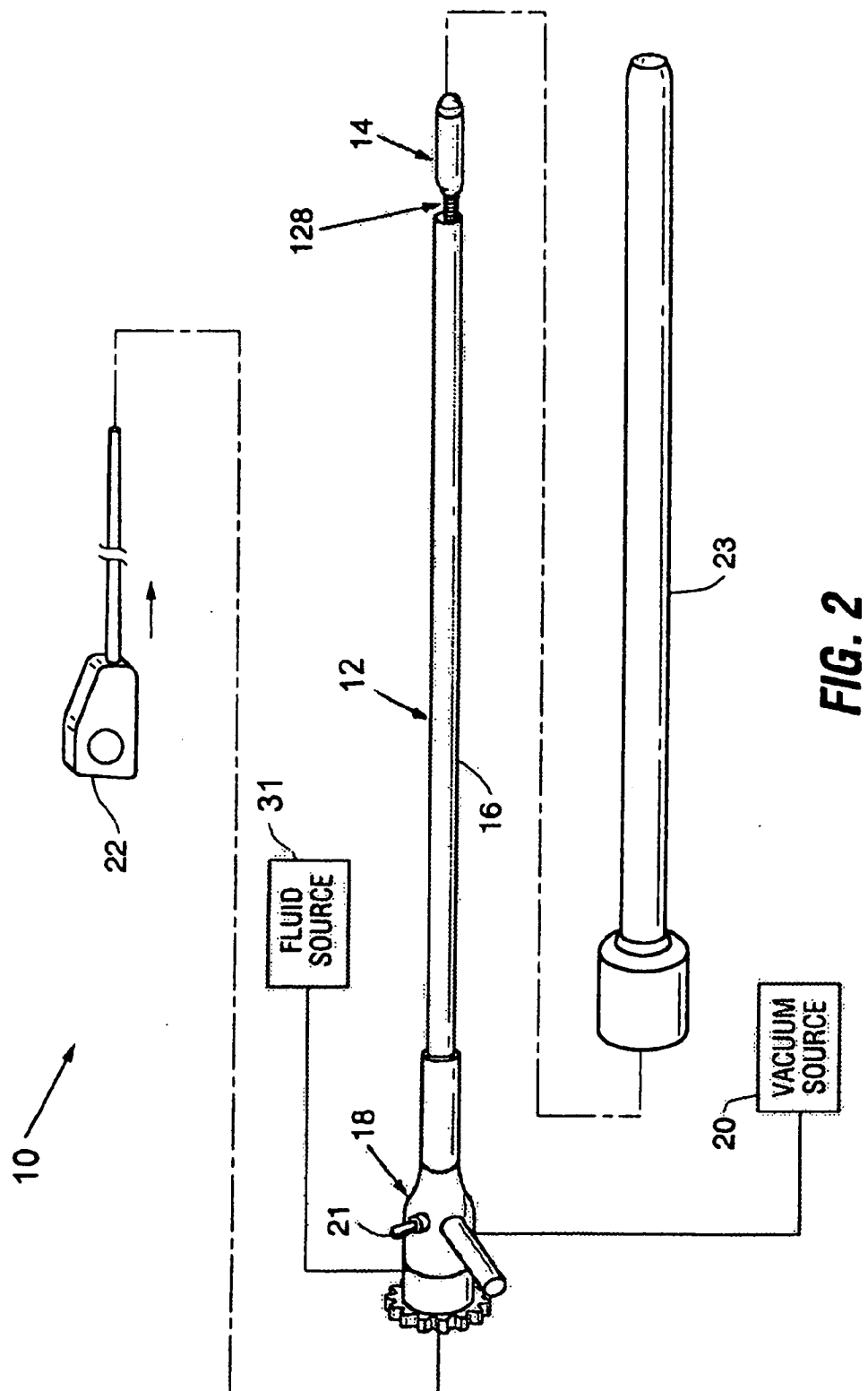
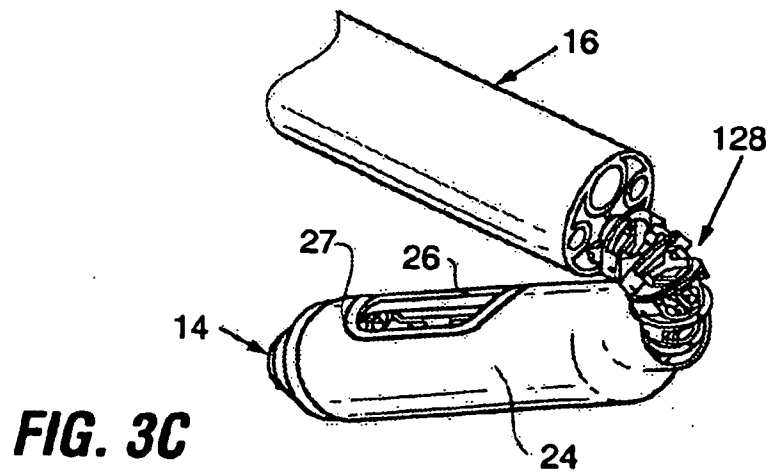
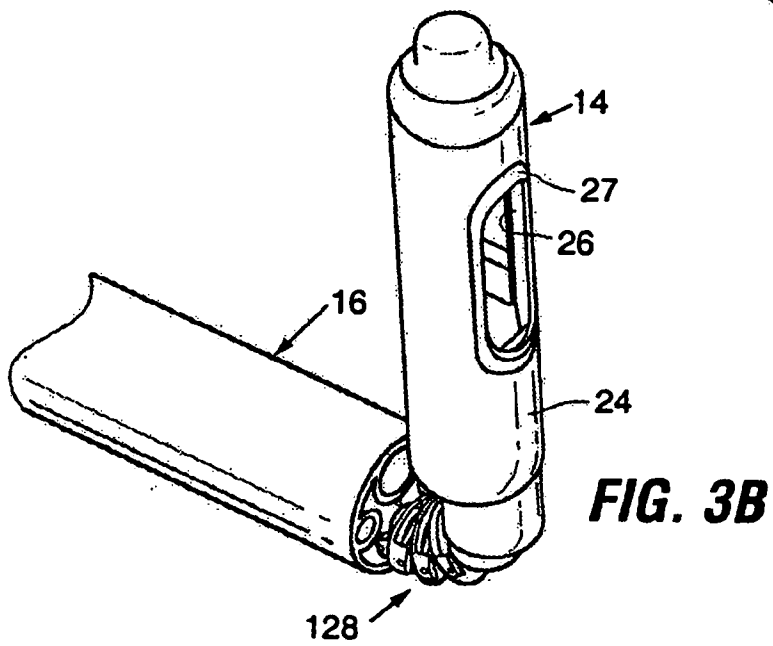
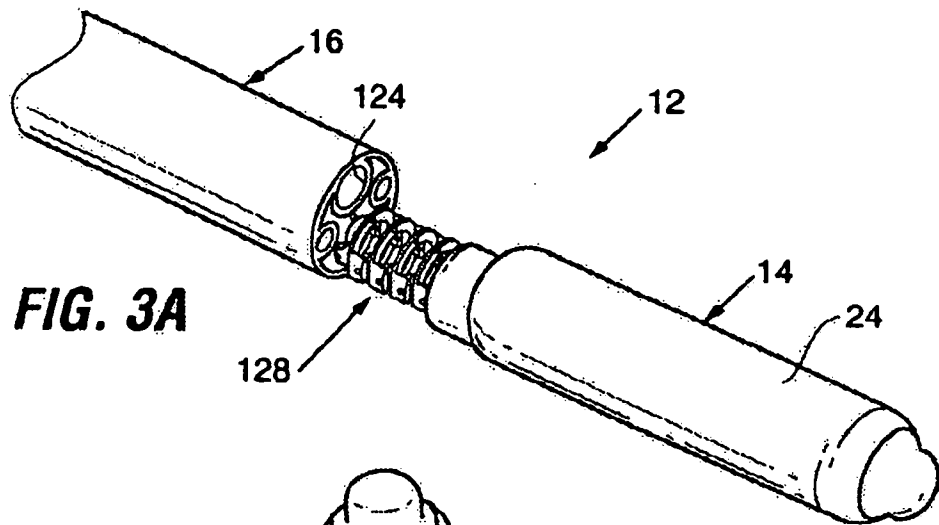


FIG. 2



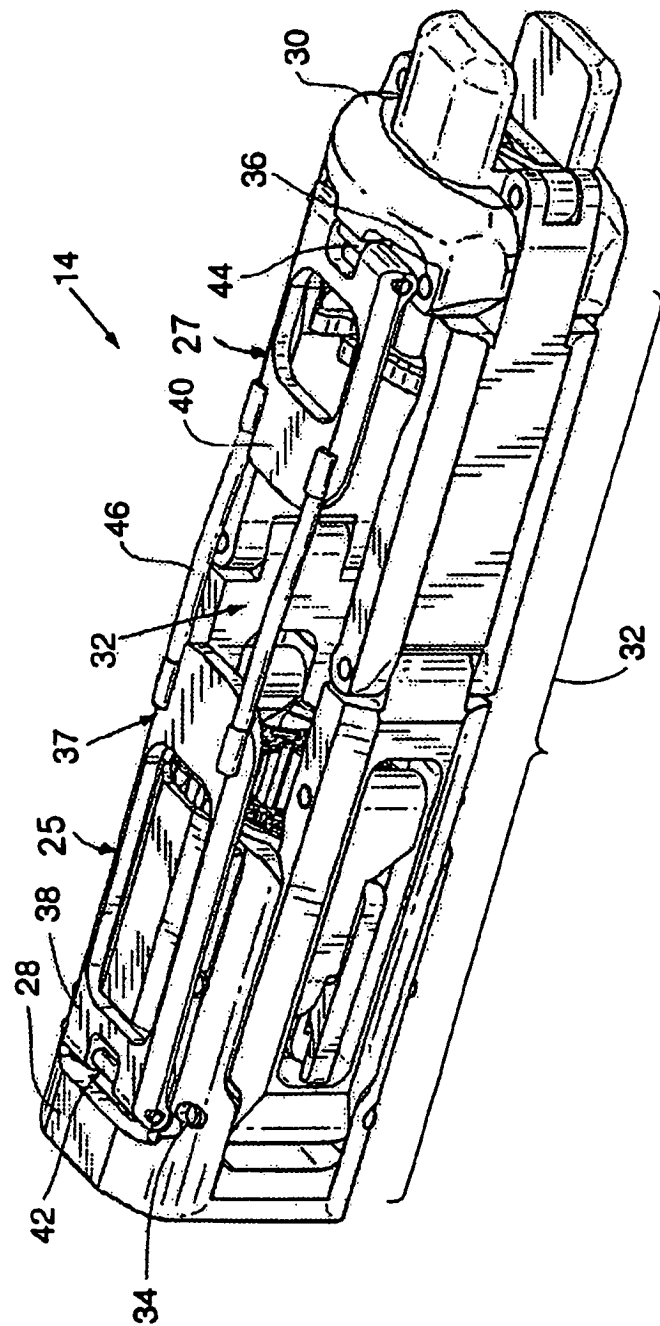
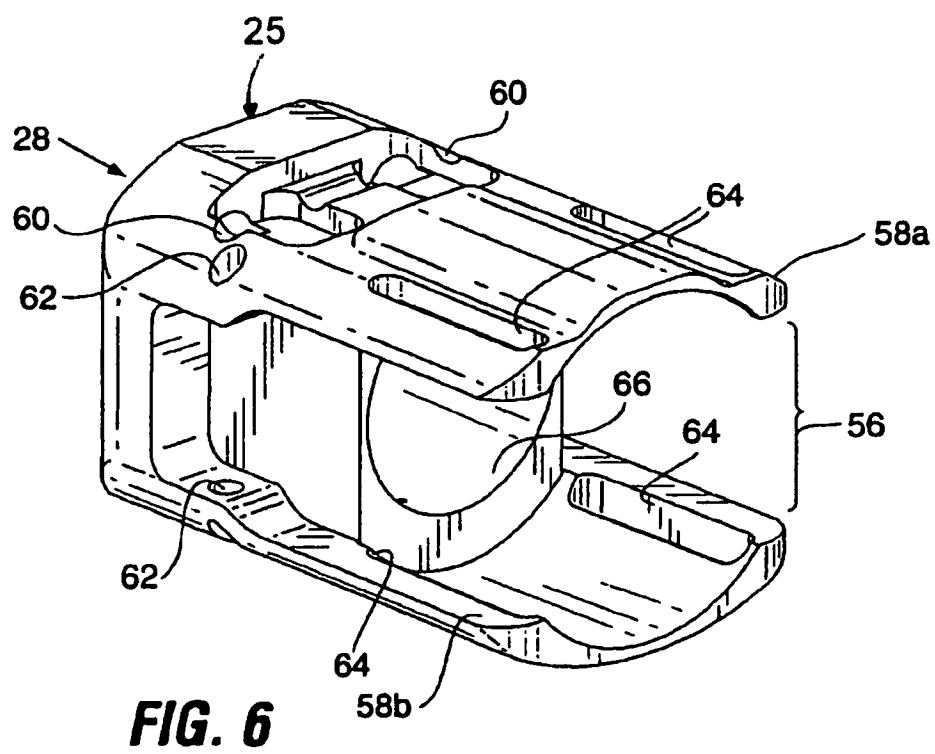
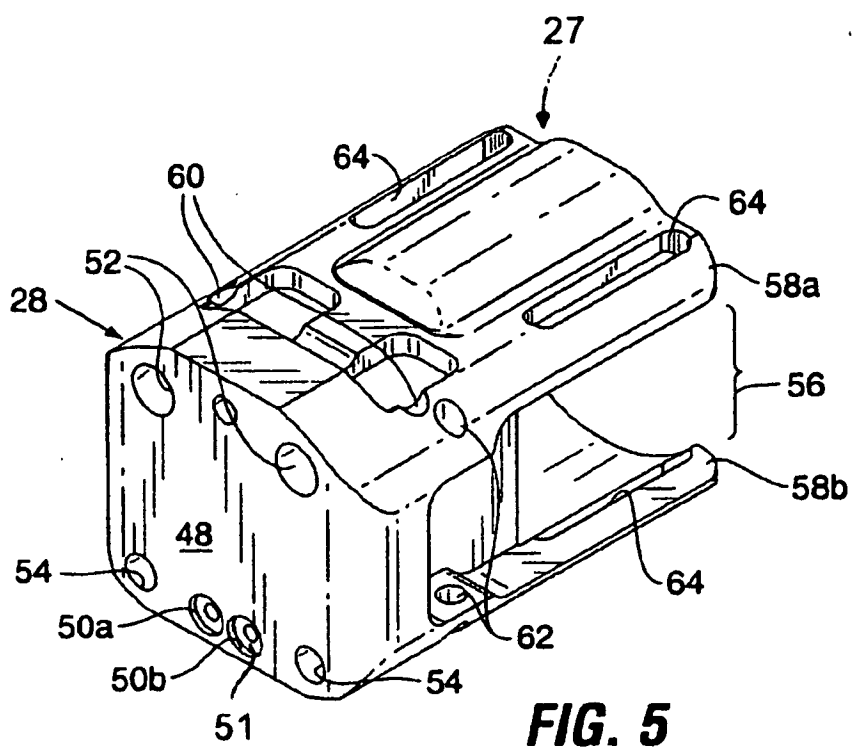
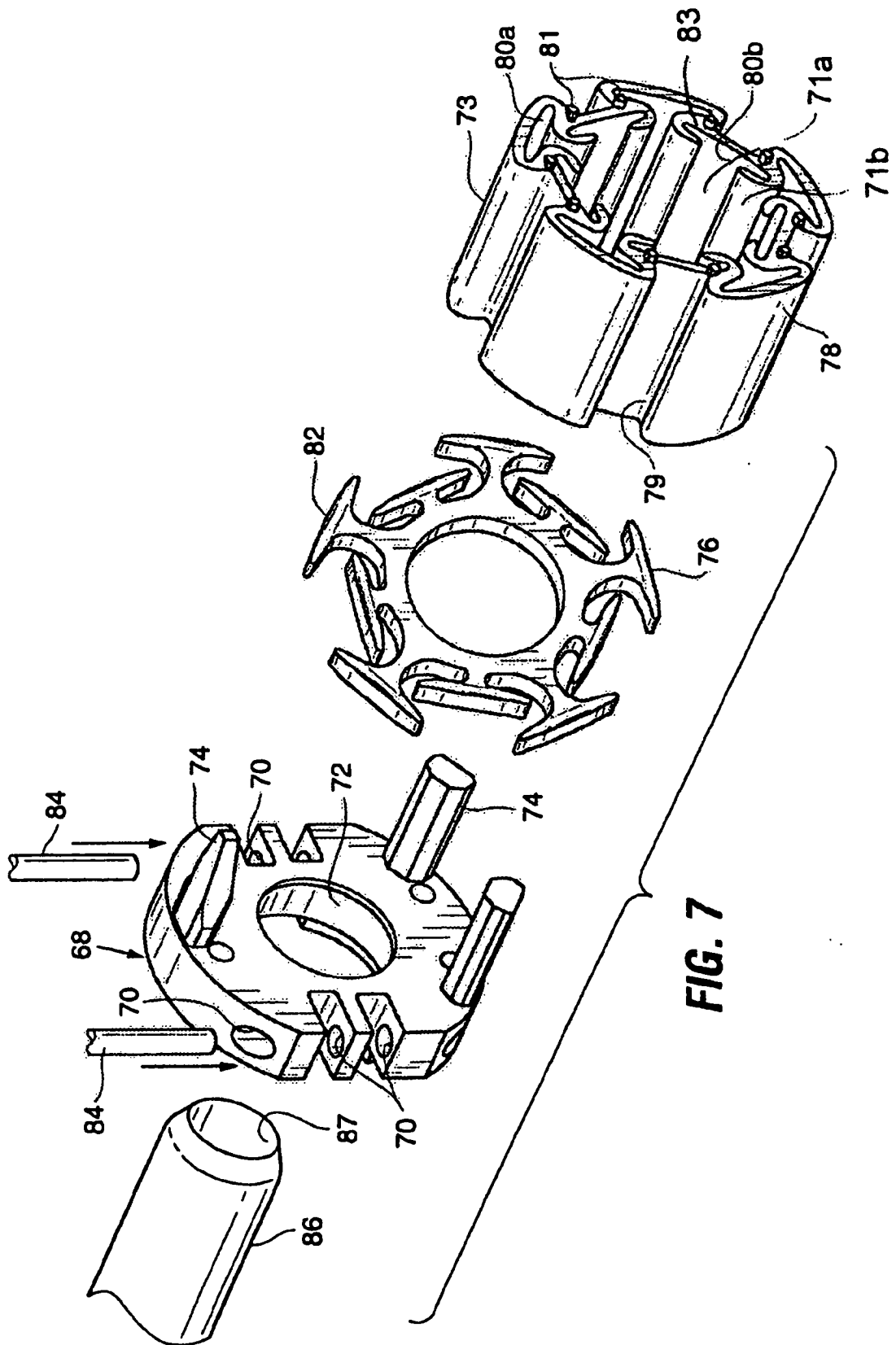


FIG. 4





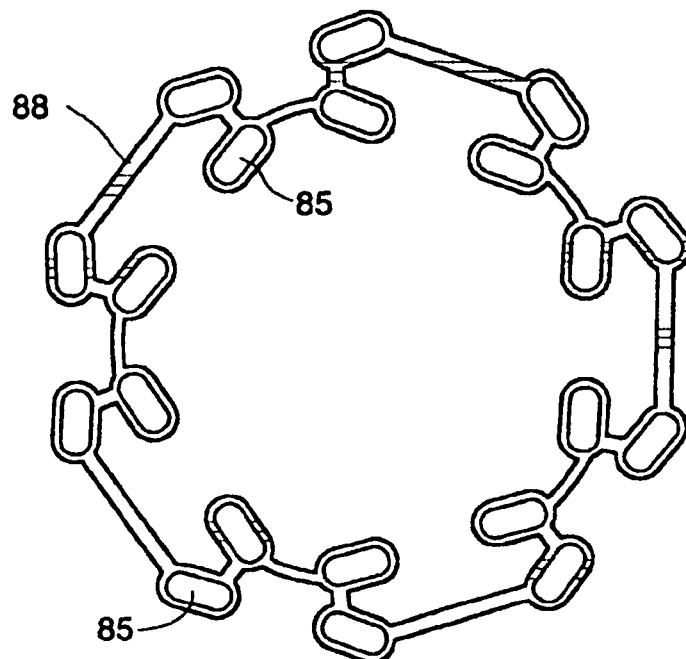


FIG. 8

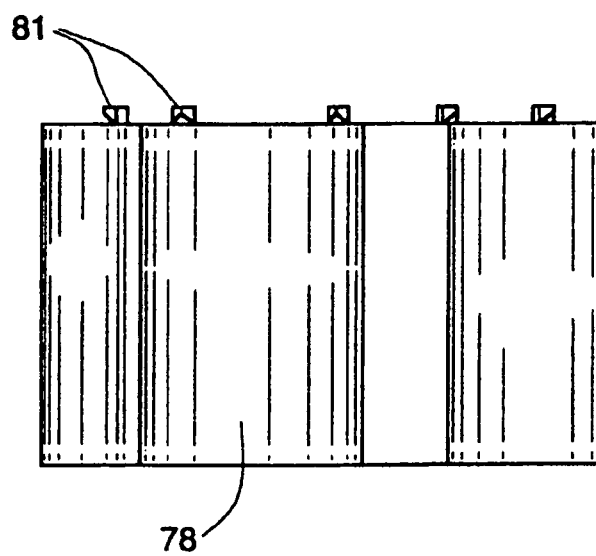


FIG. 9

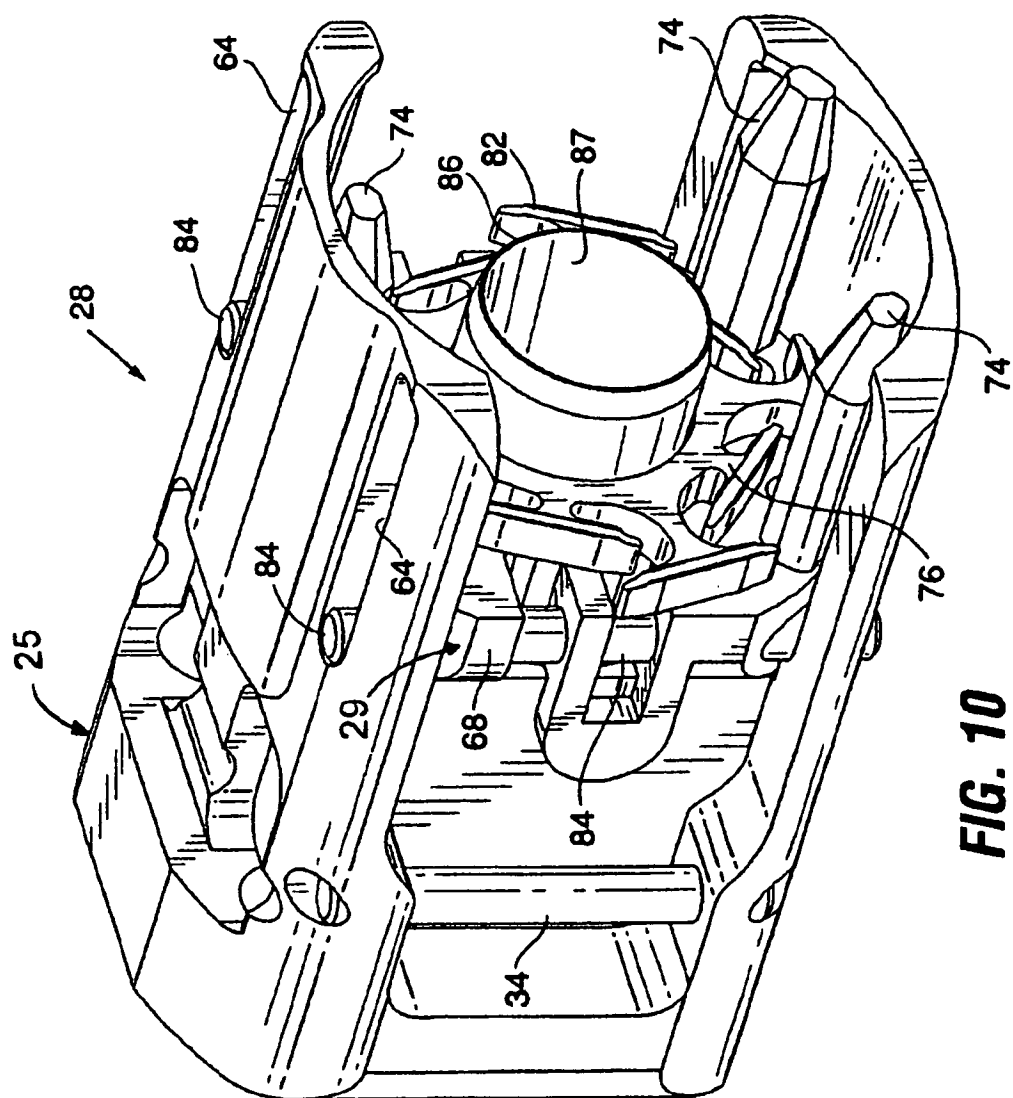


FIG. 10

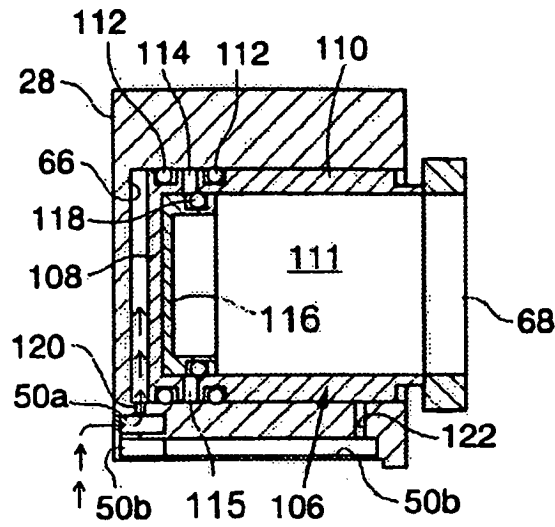


FIG. 11A

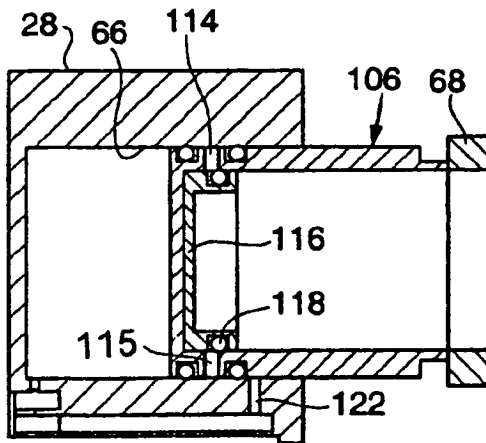


FIG. 11B

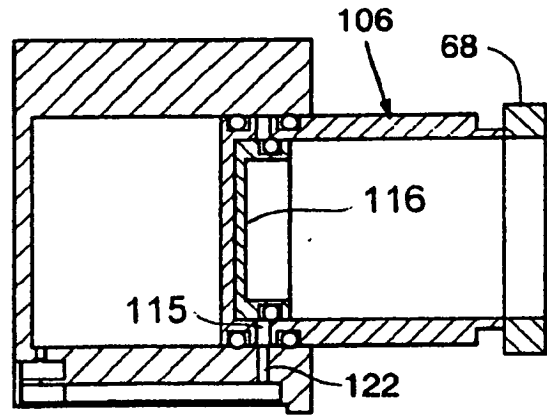


FIG. 11C

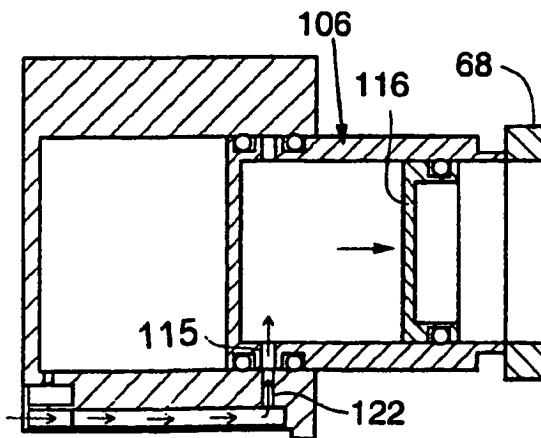


FIG. 11D

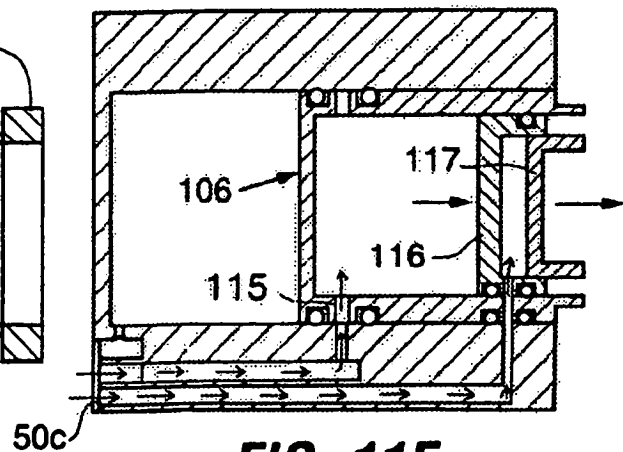
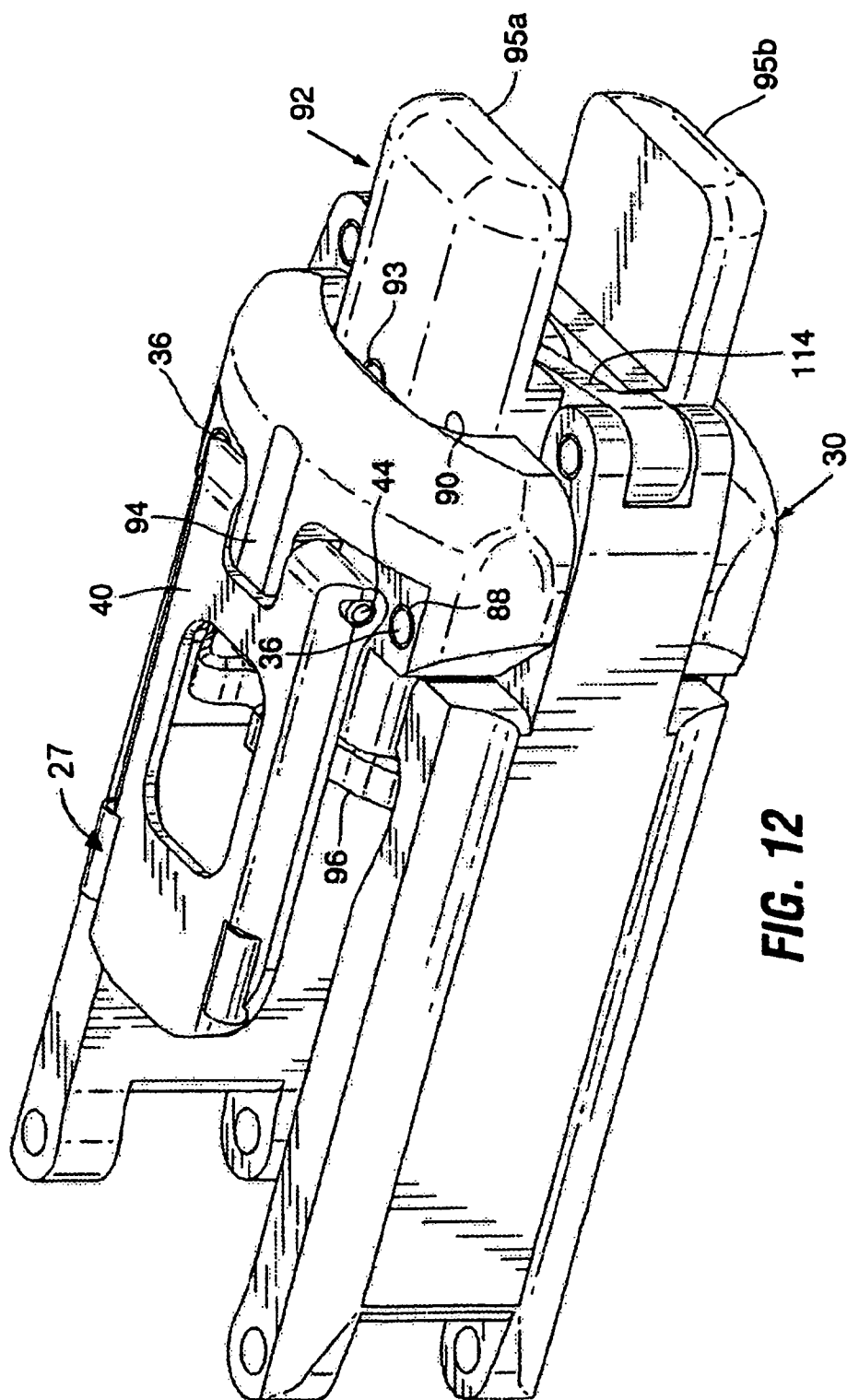
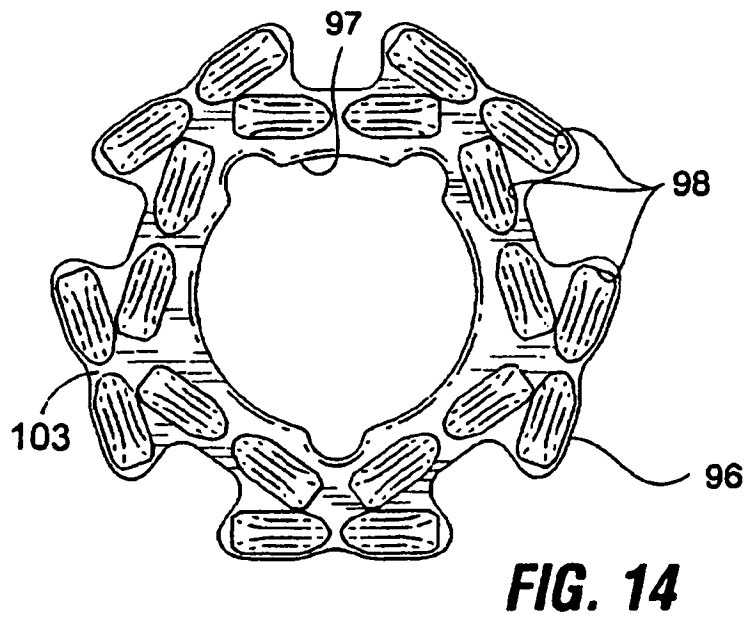
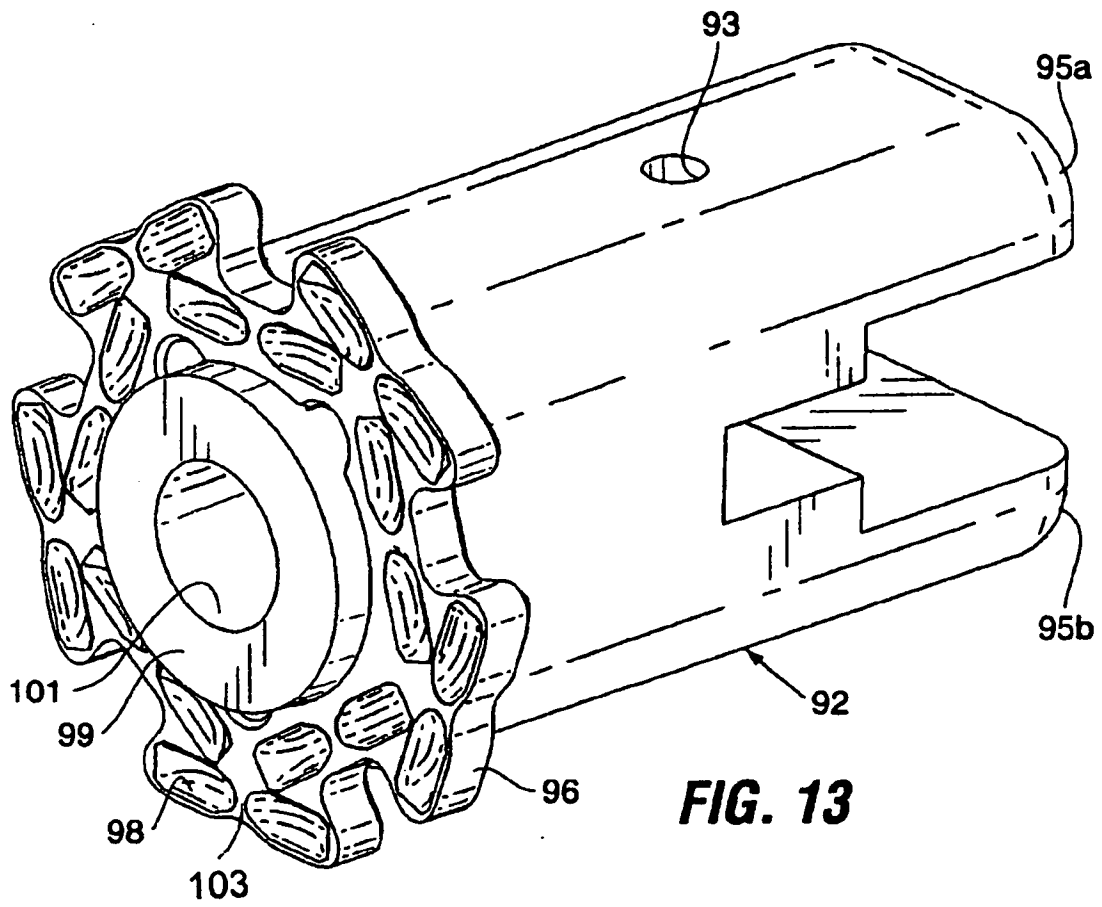


FIG. 11E





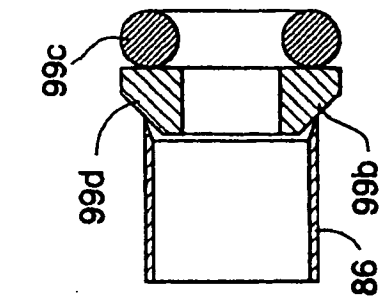


FIG. 15B

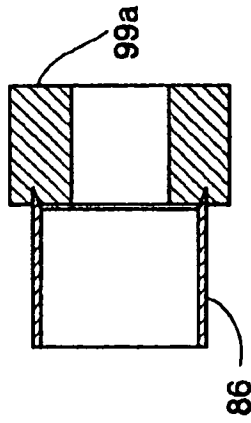


FIG. 15A

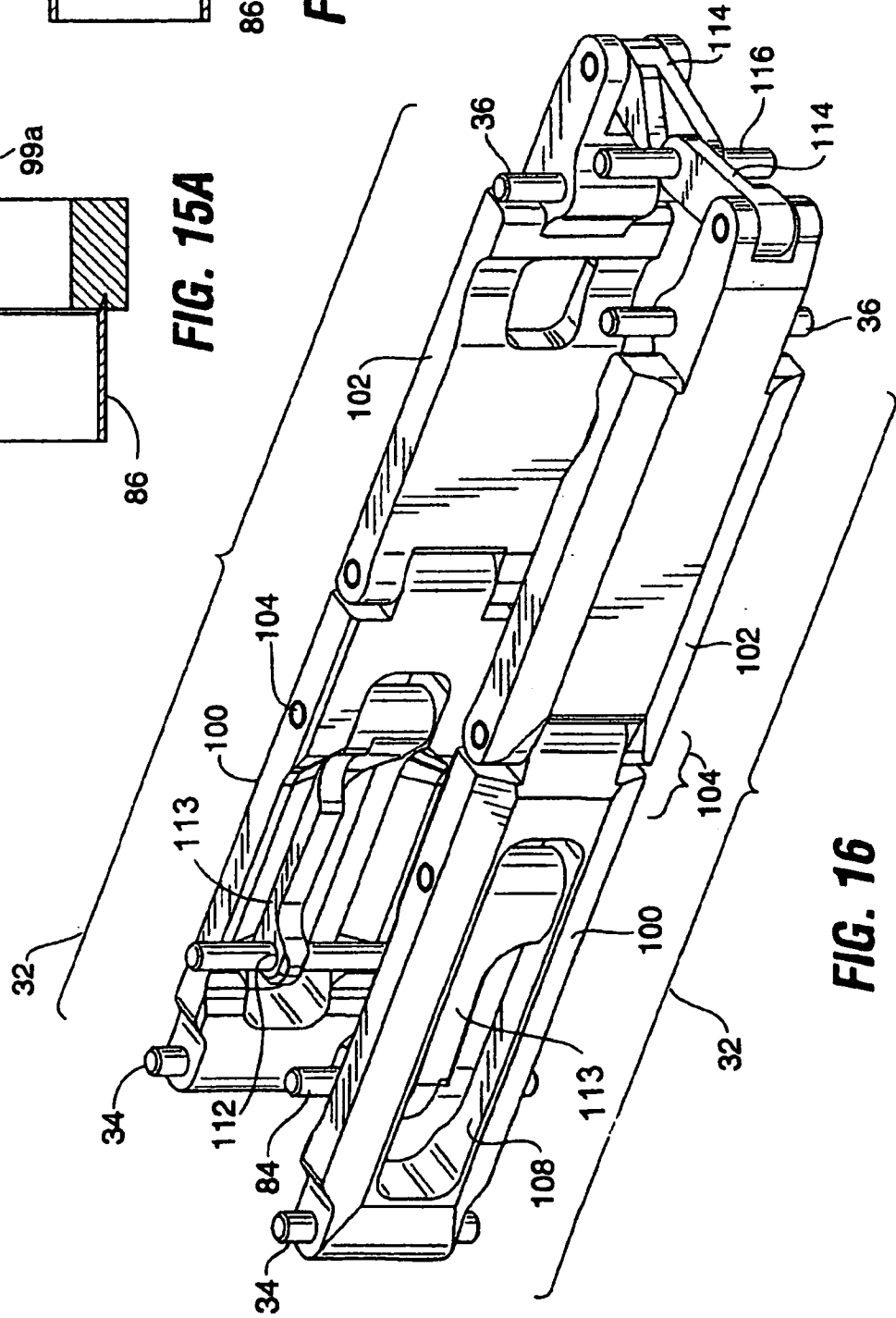


FIG. 16

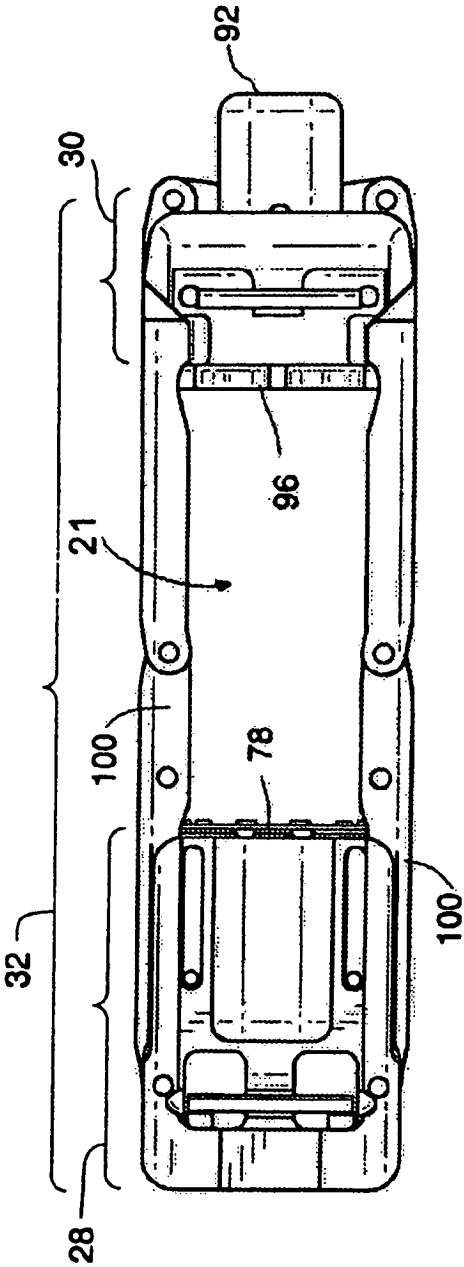


FIG. 17

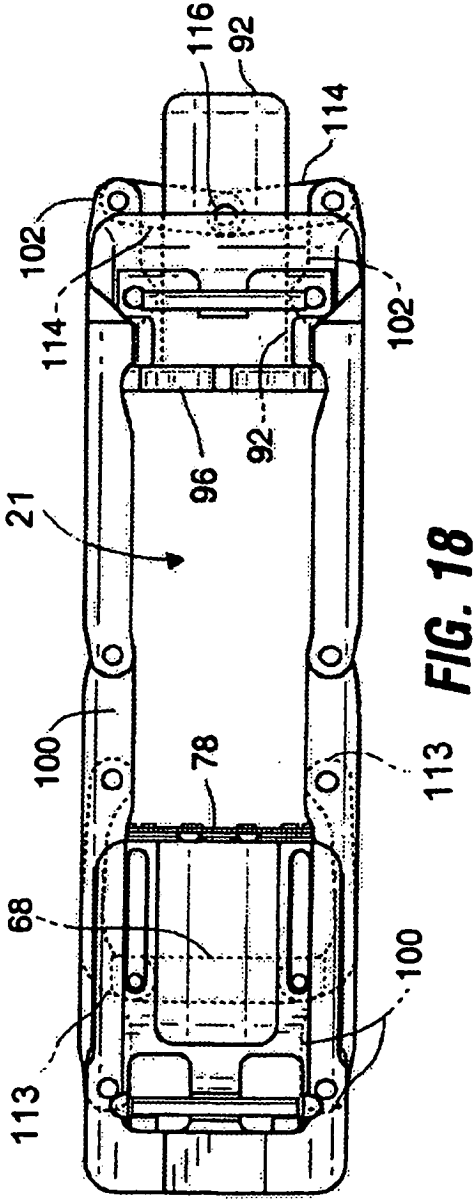


FIG. 18

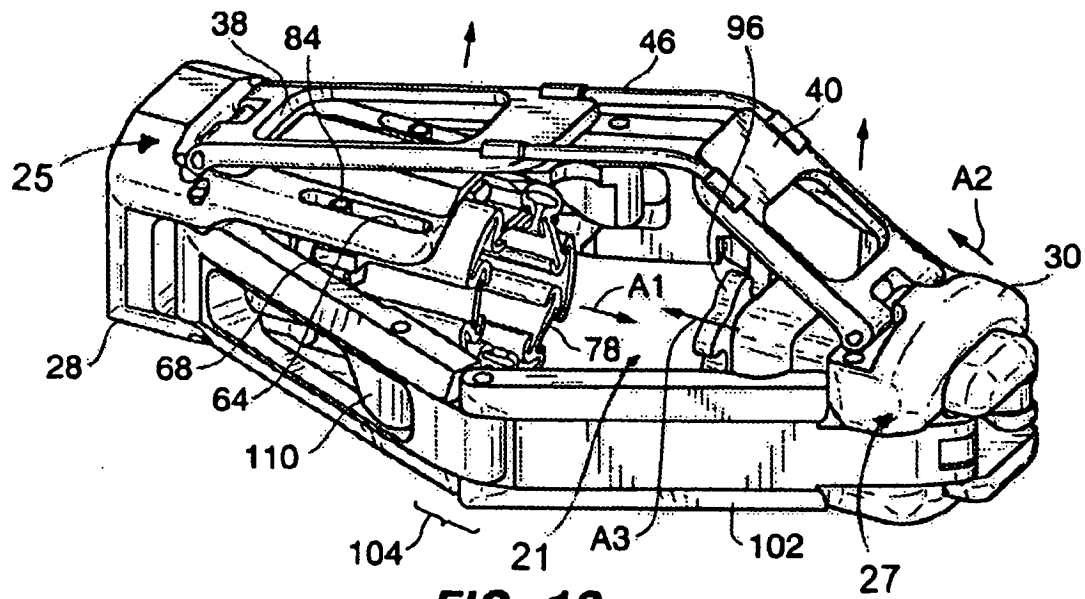


FIG. 19

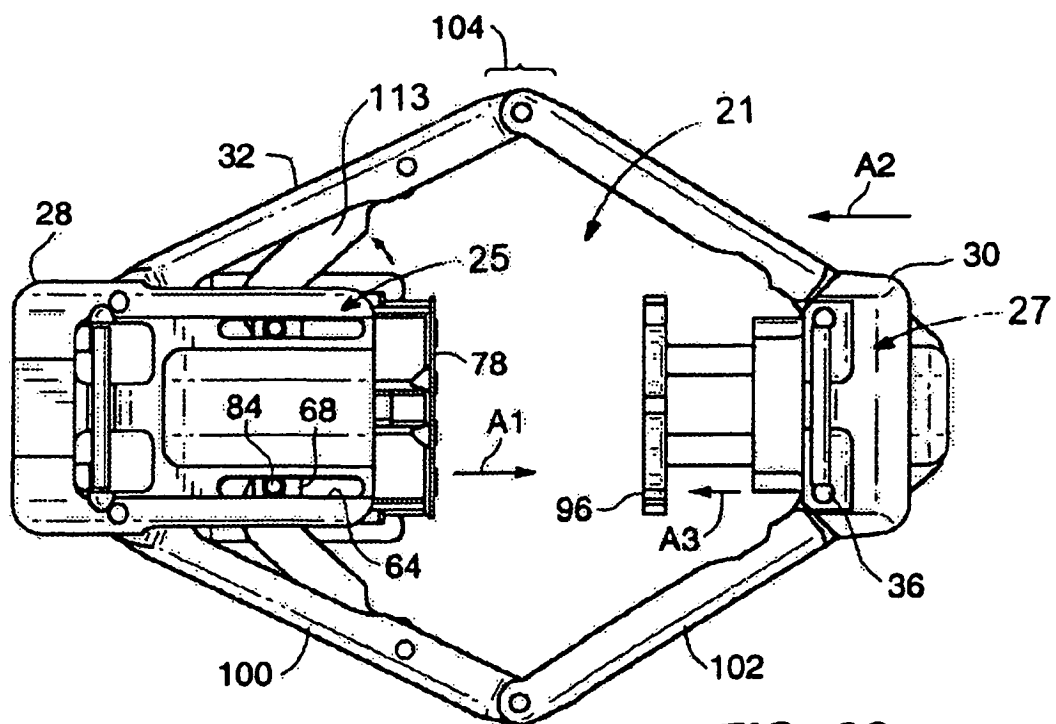


FIG. 20

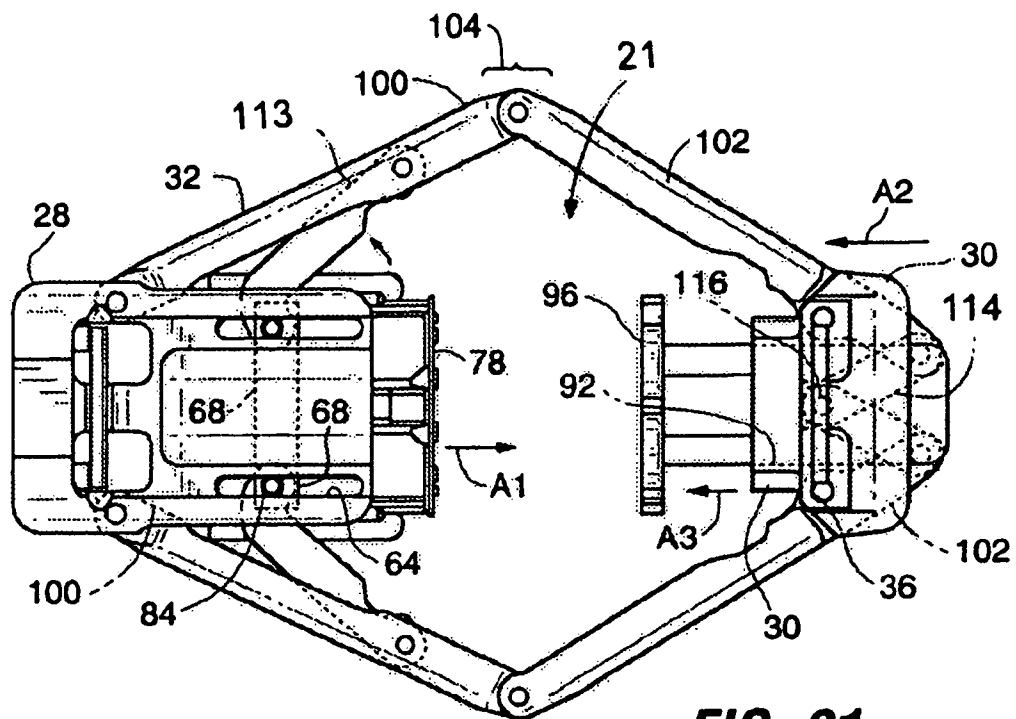


FIG. 21

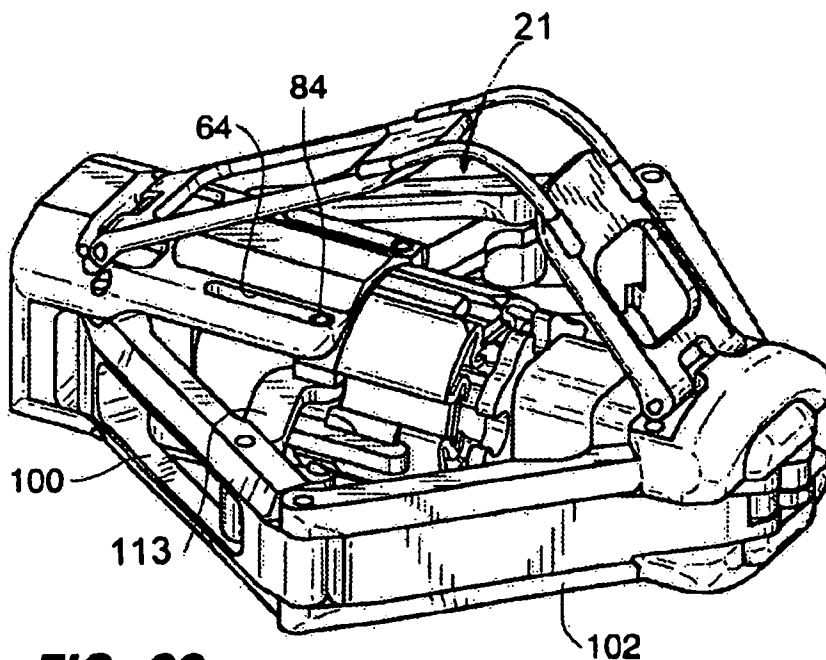
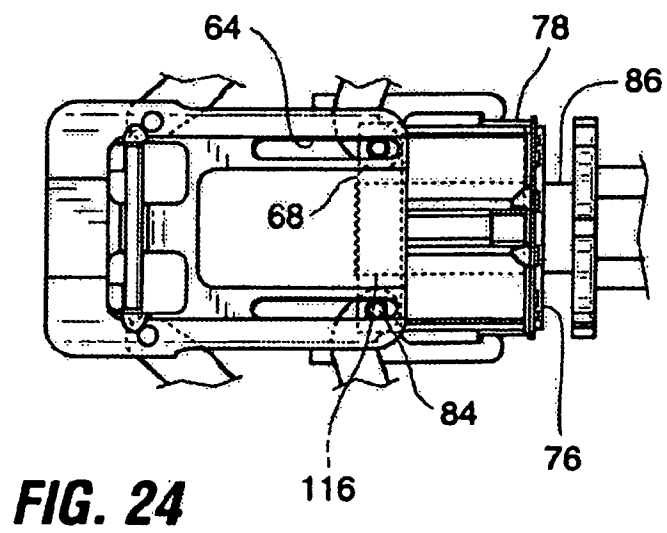
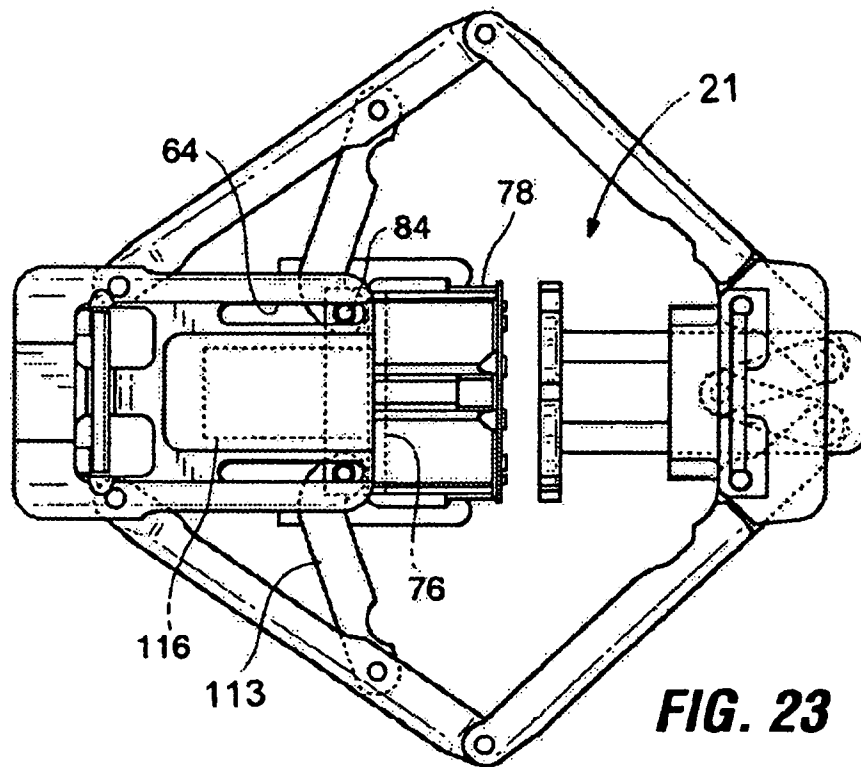
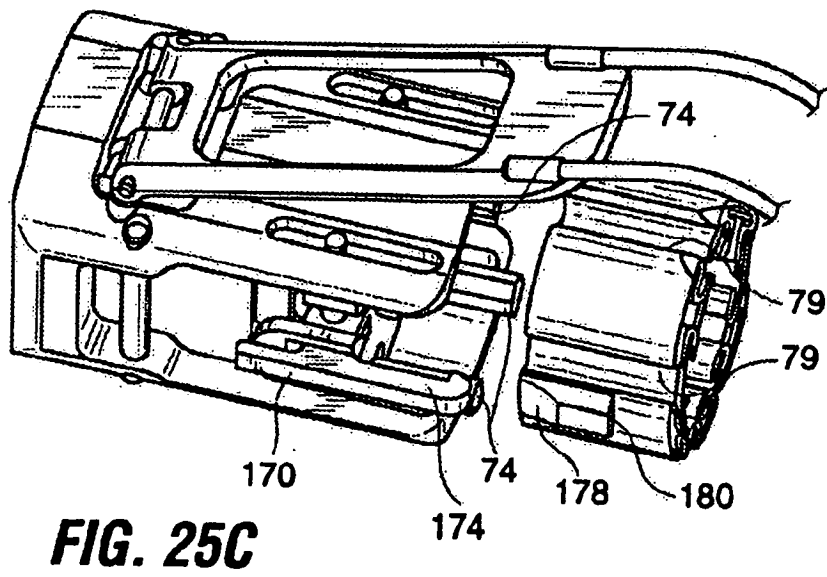
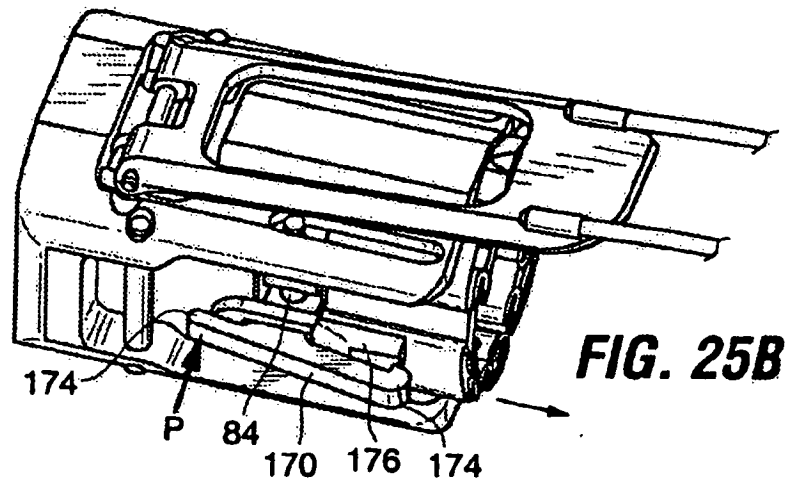
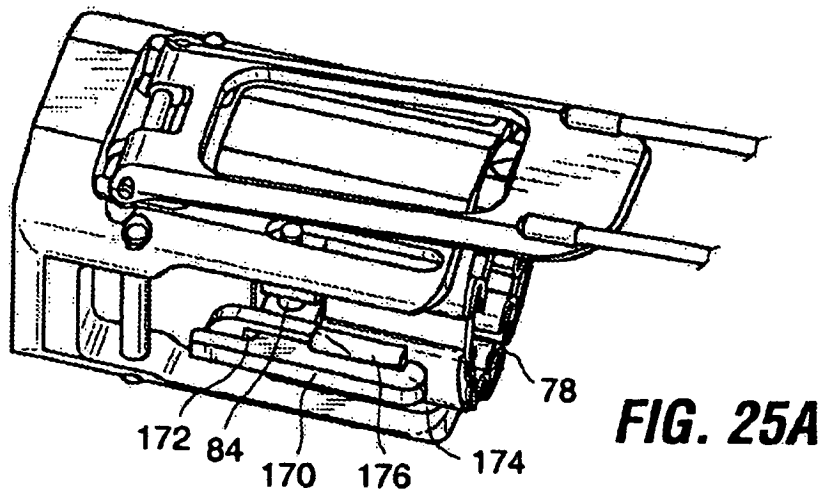
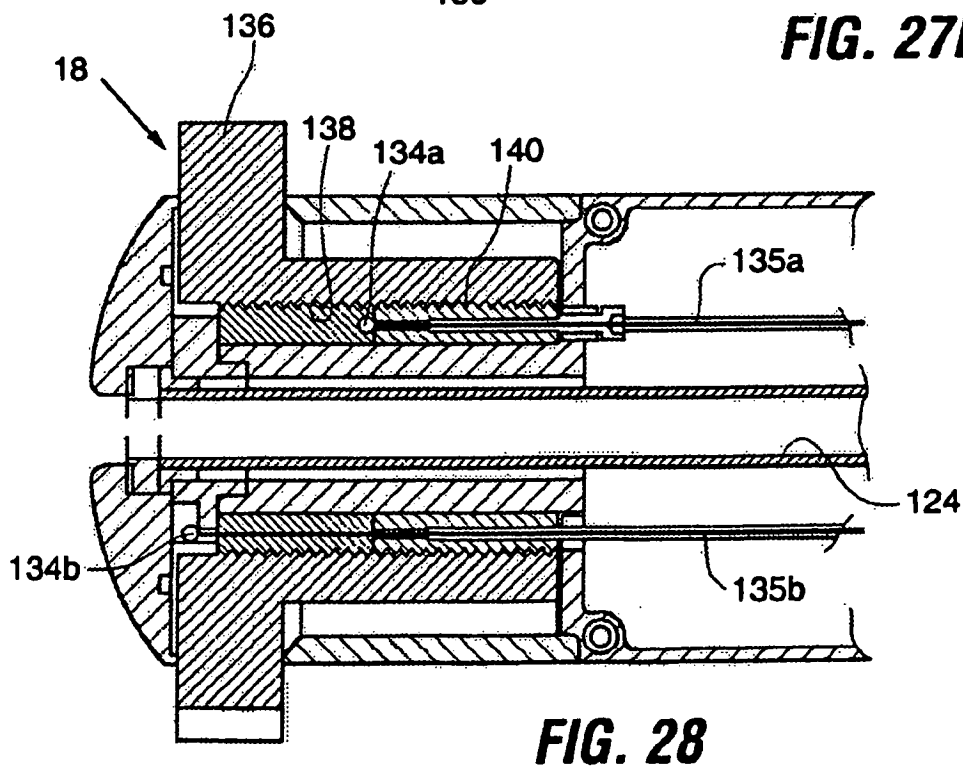
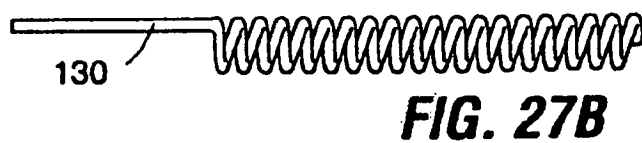
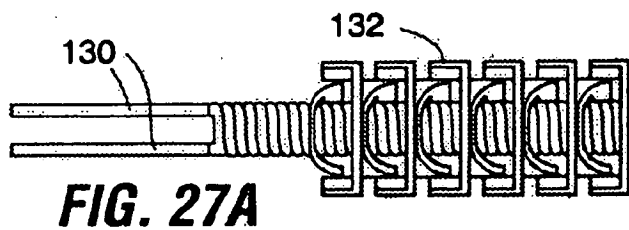
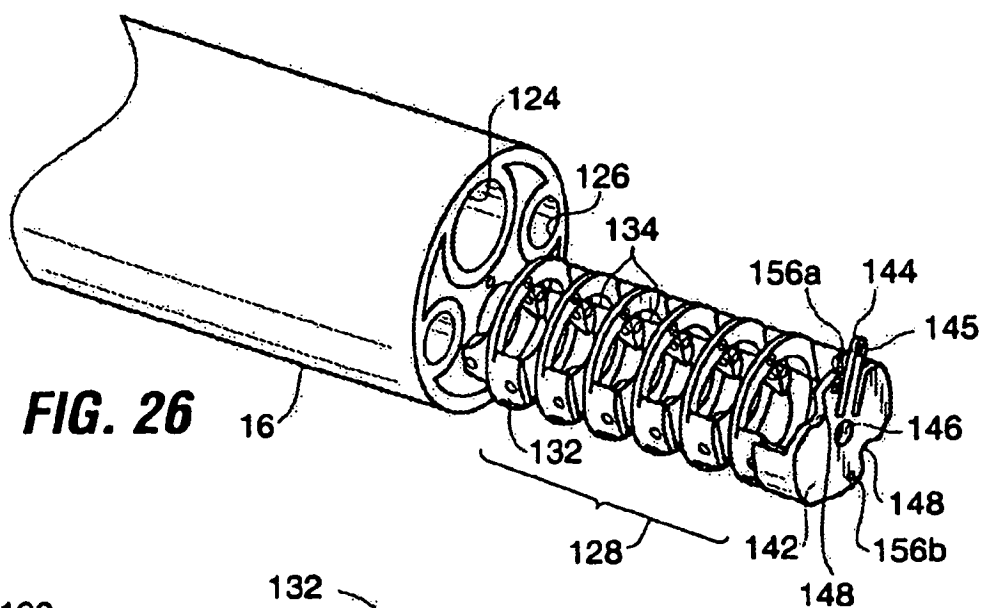
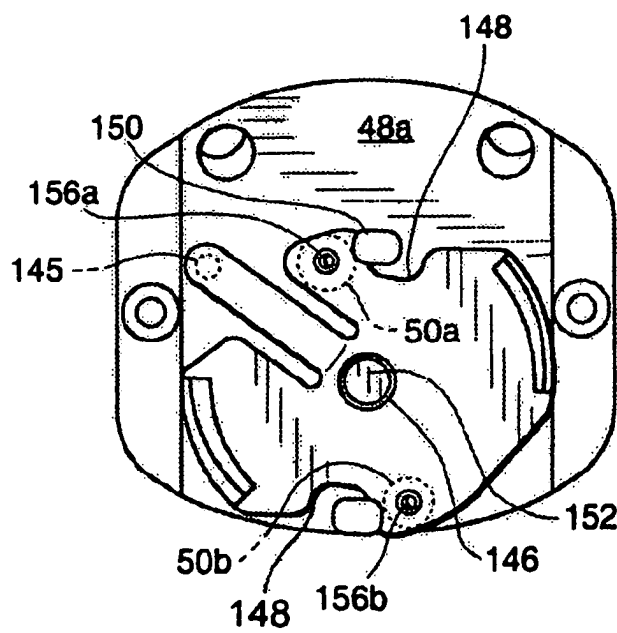
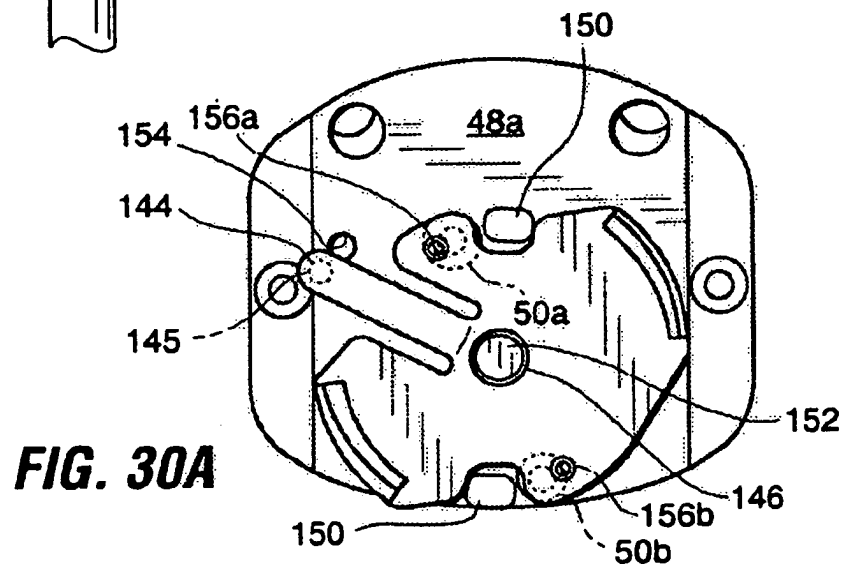
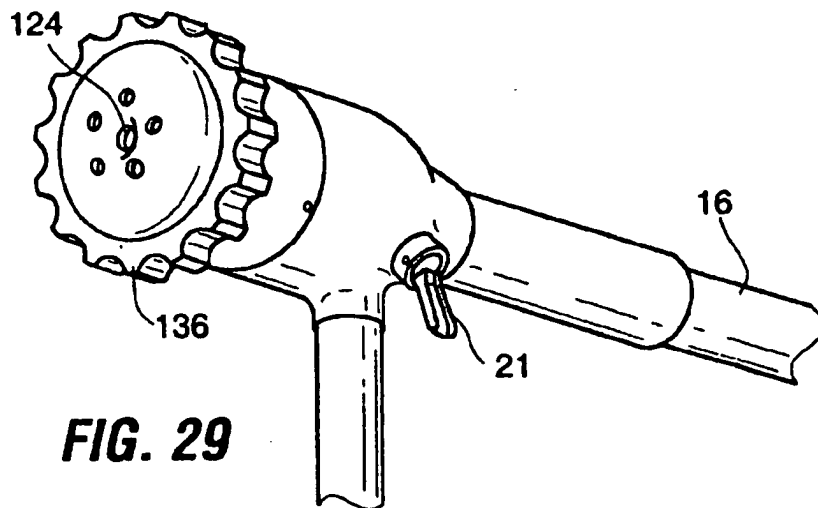


FIG. 22









snap rings

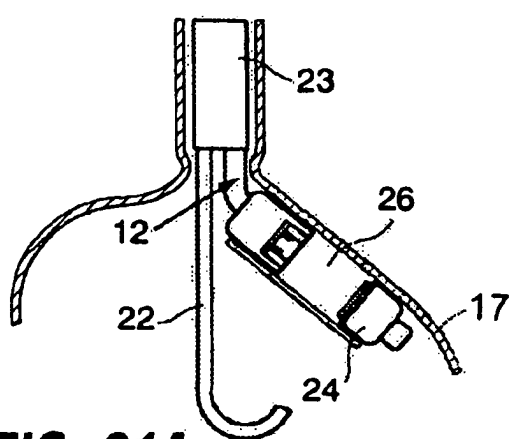


FIG. 31A

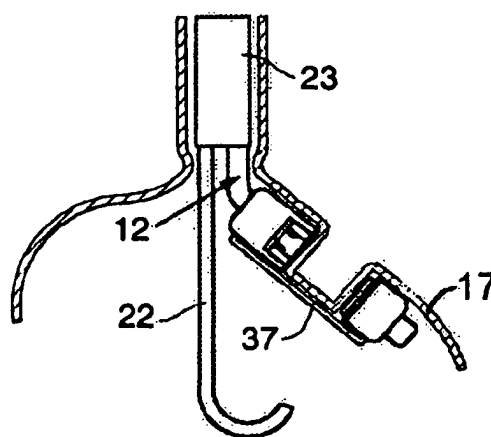


FIG. 31B

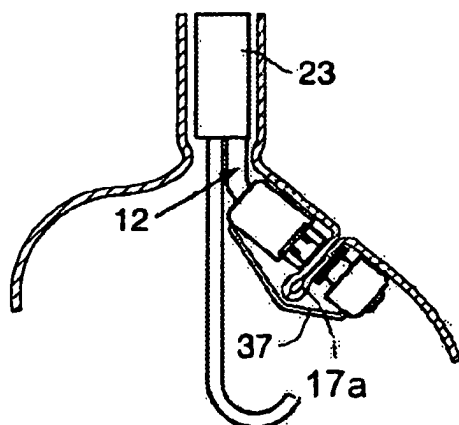


FIG. 31C

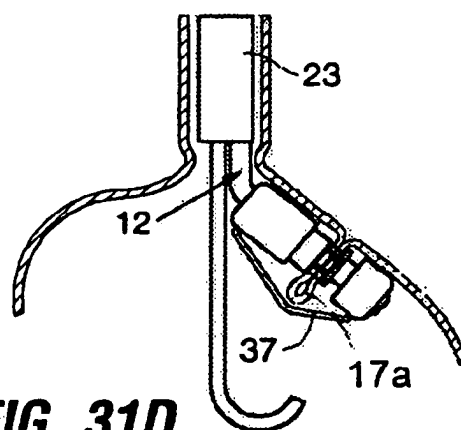


FIG. 31D

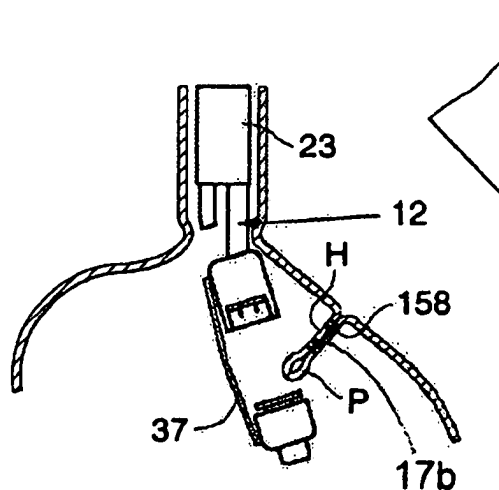


FIG. 31E

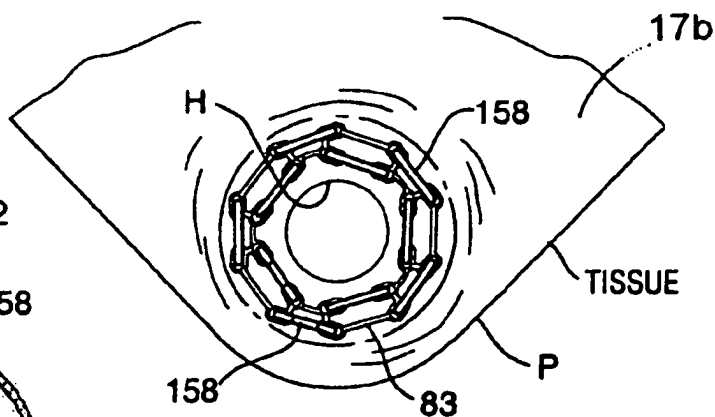


FIG. 33

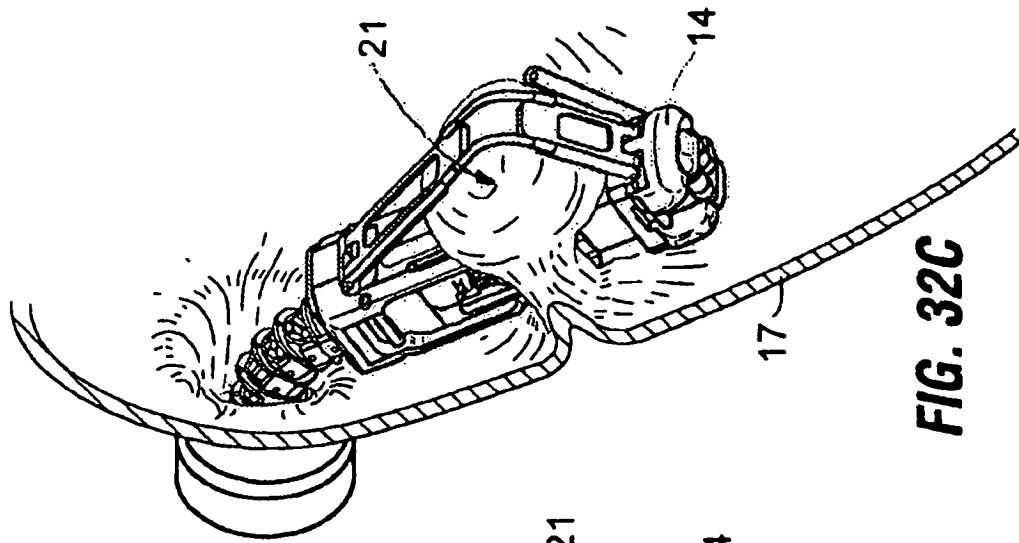


FIG. 32C

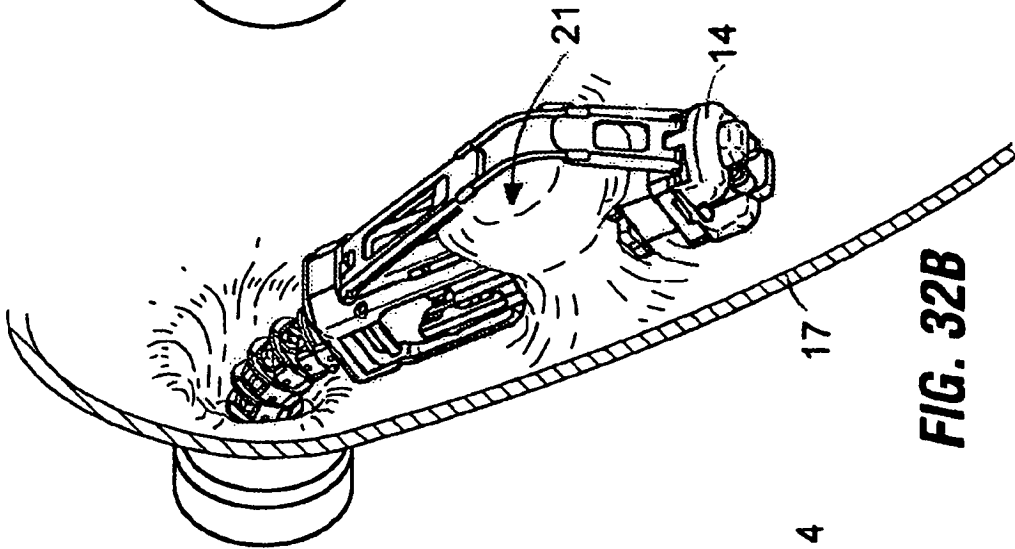


FIG. 32B

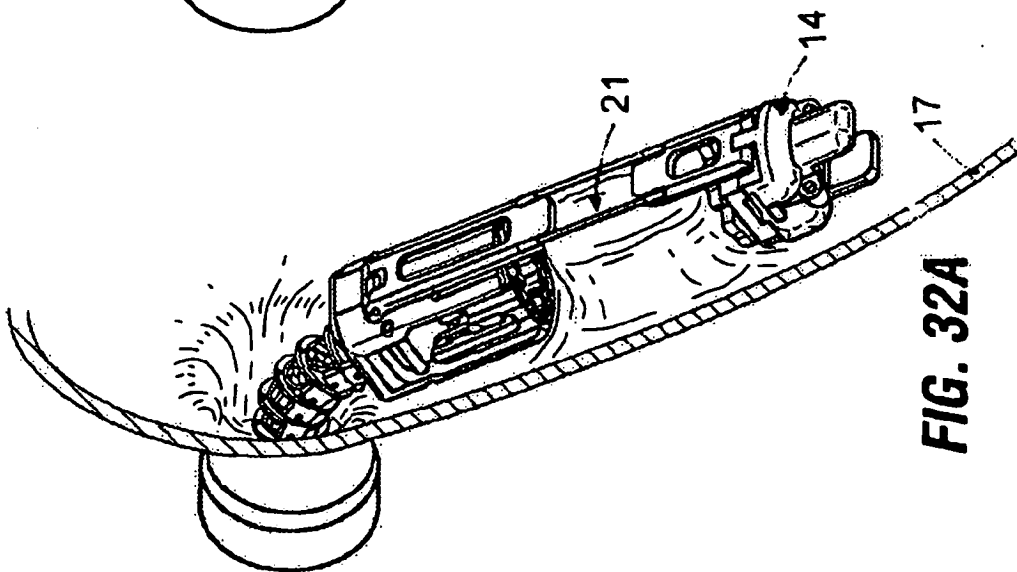


FIG. 32A

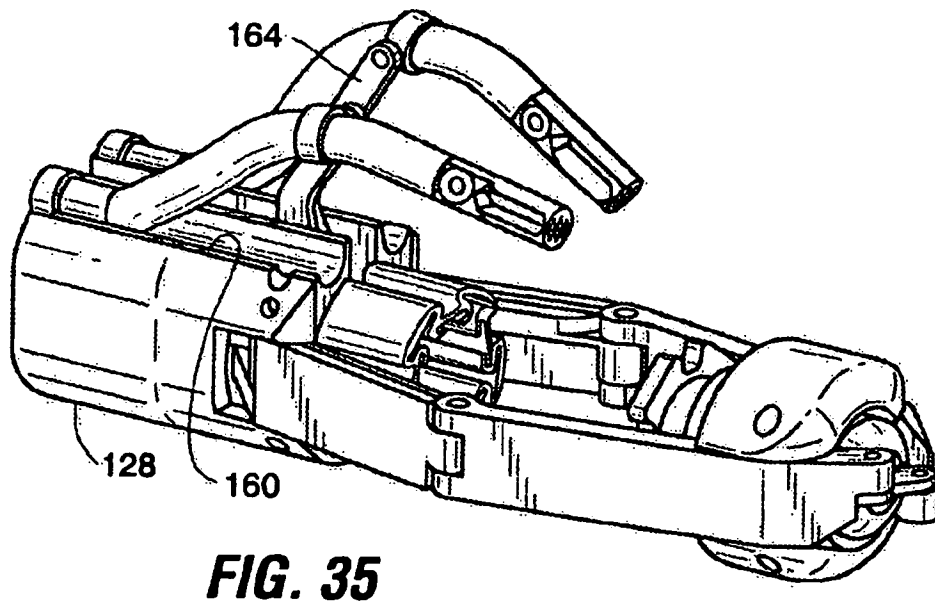
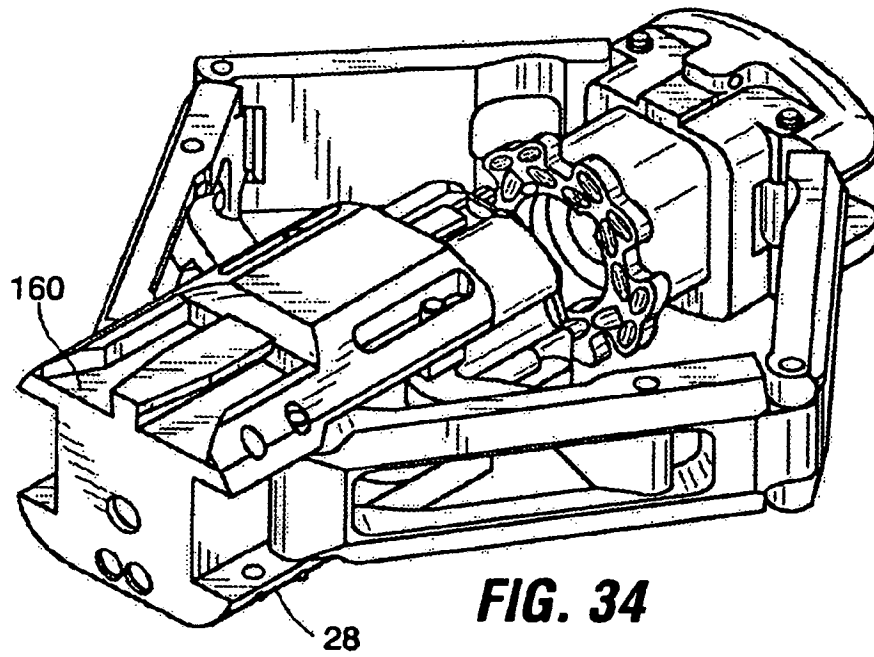


FIG. 37

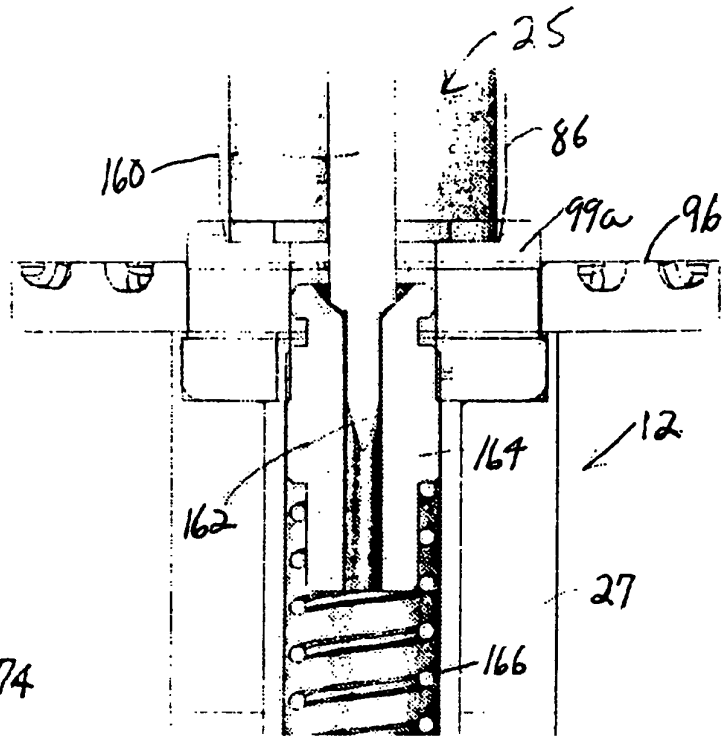
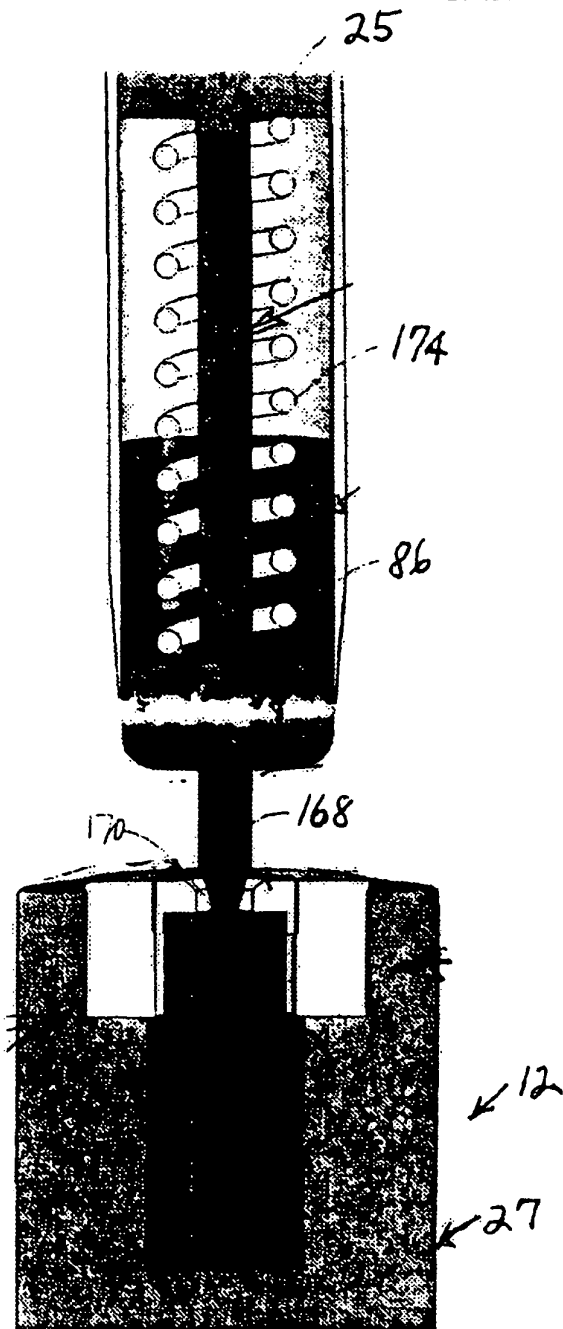


FIG. 36

REFERENCES CITED IN THE DESCRIPTION

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[标]申请(专利权)人(译)	BAROSENSE		
申请(专利权)人(译)	BAROSENSE INC.		
当前申请(专利权)人(译)	BAROSENSE INC.		
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摘要(译)

本文描述了一种用于将一个或多个紧固件施加到身体组织的吻合装置和方法。在一个实施例中，紧固件施加装置（优选地是订书机）经口部通入胃中并用于通过从胃内部接合组织并将其向内拉入而使胃组织复位。在所公开的实施例中，组织被向内抽入真空室，使得胃外部的浆膜组织部分彼此面对地定位。所公开的订书机允许组织的相对部分移动成彼此接触，并且优选地递送钉以维持组织部分之间的接触，至少直到在它们之间形成浆膜结合。这些步骤中的每一个可以完全从胃内部进行，因此可以消除对任何手术或腹腔镜介入的需要。在形成一个或多个褶皱之后，医疗装置可以可选地连接到褶皱以保持在内胃。

