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(54) **ENDOSCOPIC STAPLING DEVICES**

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(56) **References Cited**

U.S. PATENT DOCUMENTS

1,408,865 A 3/1922 Codwetl
3,663,965 A 5/1972 Lee et al.
(Continued)

FOREIGN PATENT DOCUMENTS

AU 629664 2/1991
CH 680263 A5 7/1992
(Continued)

OTHER PUBLICATIONS

International Search Report from PCT Patent Application No. PCT/US2008/008726 mailed Oct. 16, 2008.

(Continued)

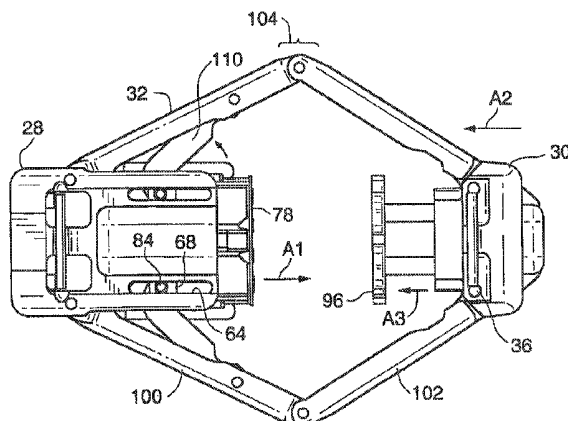
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(57) **ABSTRACT**

Described herein are endoscopic staplers and methods used to apply one or more fasteners to body tissue. In one embodiment, a fastener-applying device, which is preferably a stapler, is passed transorally into the stomach to plicate stomach tissue by engaging tissue from inside the stomach and drawing it inwardly. In the disclosed embodiments, the tissue is drawn into a tissue chamber, causing sections of serosal tissue to be positioned facing one another. The disclosed staplers allow opposed sections of tissue to be moved into contact with another, and preferably deliver staples for maintaining contact between tissue sections at least until serosal bonds form. 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 are formed, medical devices may optionally be coupled to the plication(s) for retention within the stomach.

18 Claims, 22 Drawing Sheets



continuation of application No. 12/050,169, filed on Mar. 18, 2008, now Pat. No. 8,020,741, and a division of application No. 12/052,997, filed on Mar. 21, 2008, now Pat. No. 7,909,222, which is a division of application No. 12/050,169, filed on Mar. 18, 2008, now Pat. No. 8,020,741, said application No. 13/053,133 is a division of application No. 12/053,010, filed on Mar. 21, 2008, now Pat. No. 7,922,062, which is a division of application No. 12/050,169, filed on Mar. 18, 2008, now Pat. No. 8,020,741, said application No. 13/053,133 is a division of application No. 12/053,027, filed on Mar. 21, 2008, now Pat. No. 7,909,219, which is a division of application No. 12/050,169, filed on Mar. 18, 2008, now Pat. No. 8,020,741, said application No. 13/053,133 is a division of application No. 12/053,066, filed on Mar. 21, 2008, now Pat. No. 7,909,223, which is a division of application No. 12/050,169, filed on Mar. 18, 2008, now Pat. No. 8,020,741, said application No. 13/053,133 is a division of application No. 12/053,182, filed on Mar. 21, 2008, now Pat. No. 7,913,892, which is a division of application No. 12/050,169, filed on Mar. 18, 2008, now Pat. No. 8,020,741.

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- U.S. PATENT DOCUMENTS

4,134,405	A	1/1979	Smit	
4,207,890	A	6/1980	Mamajek et al.	
4,246,893	A	1/1981	Berson	
4,315,509	A	2/1982	Smit	
4,331,277	A	5/1982	Green	
4,383,634	A *	5/1983	Green	A61B 17/072 227/135
4,403,604	A	9/1983	Wilkinson et al.	
4,416,267	A	11/1983	Garren et al.	
4,417,360	A	11/1983	Moasser	
4,441,215	A	4/1984	Kaster	
4,467,804	A	8/1984	Hardy et al.	
4,485,805	A	12/1984	Foster, Jr.	
4,501,264	A	2/1985	Rockey	
4,607,618	A	8/1986	Angelchik	
4,612,933	A	9/1986	Brinkerhoff et al.	
4,617,932	A	10/1986	Kornberg	
4,641,653	A	2/1987	Rockey	
4,648,383	A	3/1987	Angelchik	
4,694,827	A	9/1987	Weiner et al.	
4,723,547	A	2/1988	Kullas et al.	
4,747,849	A	5/1988	Galtier	
4,846,836	A	7/1989	Reich	
4,848,367	A	7/1989	Avant et al.	
4,899,747	A	2/1990	Garren et al.	

4,925,446	A	5/1990	Garay et al.
4,946,440	A	8/1990	Hall
4,969,896	A	11/1990	Shors
4,997,084	A	3/1991	Opie et al.
5,006,106	A	4/1991	Angelchik
5,037,021	A	8/1991	Mills et al.
5,061,275	A	10/1991	Wallsten et al.
5,084,061	A	1/1992	Gau et al.
5,088,979	A	2/1992	Filipi et al.
5,163,952	A	11/1992	Froix
5,211,658	A	5/1993	Clouse
5,234,454	A	8/1993	Bangs
5,246,456	A	9/1993	Wilkinson
5,259,399	A	11/1993	Brown
5,263,629	A	11/1993	Trumbull et al.
5,290,217	A	3/1994	Campos
5,306,300	A	4/1994	Berry
5,314,473	A	5/1994	Godin
5,327,914	A	7/1994	Shlain
5,345,949	A	9/1994	Shlain
5,355,897	A	10/1994	Pietrafitta et al.
5,401,241	A	3/1995	Delany
5,403,326	A	4/1995	Harrison et al.
5,405,377	A	4/1995	Cragg
5,431,673	A	7/1995	Summers et al.
5,486,187	A	1/1996	Schenck
5,514,176	A	5/1996	Bosley, Jr.
5,535,935	A	7/1996	Vidal et al.
5,542,949	A	8/1996	Yoon
5,562,239	A	10/1996	Boiarski et al.
5,571,116	A	11/1996	Bolanos et al.
5,577,654	A	11/1996	Bishop
5,593,434	A	1/1997	Williams
5,597,107	A	1/1997	Knodel et al.
5,609,624	A	3/1997	Kalis
5,628,786	A	5/1997	Banas et al.
5,630,539	A	5/1997	Plyle et al.
5,647,526	A	7/1997	Green et al.
5,653,743	A	8/1997	Martin
5,662,713	A	9/1997	Andersen et al.
5,673,841	A	10/1997	Schulze et al.
5,674,241	A	10/1997	Bley et al.
5,706,998	A	1/1998	Plyle et al.
5,709,657	A	1/1998	Zimmon
5,720,776	A	2/1998	Chuter et al.
5,749,918	A	5/1998	Hogendijk et al.
5,762,255	A	6/1998	Chrisman et al.
5,771,903	A	6/1998	Jakobsson
5,785,684	A	7/1998	Zimmon
5,792,119	A	8/1998	Marx
5,820,584	A	10/1998	Crabb
5,839,639	A	11/1998	Sauer et al.
5,848,964	A	12/1998	Samuels
5,855,311	A	1/1999	Hamblin et al.
5,855,601	A	1/1999	Bessler et al.
5,856,445	A	1/1999	Korsmeyer
5,861,036	A	1/1999	Godin
5,868,141	A	2/1999	Ellias
5,887,594	A	3/1999	LoCicero, III
5,897,562	A	4/1999	Bolanos et al.
5,910,144	A	6/1999	Hayashi et al.
5,922,019	A	7/1999	Hank et al.
5,947,983	A	9/1999	Solar et al.
5,993,473	A	11/1999	Chan et al.
5,993,483	A	11/1999	Gianotti
6,016,848	A	1/2000	Egrees
6,051,015	A	4/2000	Maahs
6,086,600	A	7/2000	Kortenbach
6,098,629	A	8/2000	Johnson et al.
6,102,922	A	8/2000	Jakobsson et al.
6,113,609	A	9/2000	Adams
6,120,534	A	9/2000	Ruiz
6,146,416	A	11/2000	Andersen et al.
6,159,146	A	12/2000	El Gazayerli
6,159,238	A	12/2000	Killion et al.
6,197,022	B1	3/2001	Baker
6,206,930	B1	3/2001	Burg et al.
6,245,088	B1	6/2001	Lowery
6,251,132	B1	6/2001	Ravenscroft et al.

(56)

References Cited

U.S. PATENT DOCUMENTS

6,254,642	B1	7/2001	Taylor	7,214,233	B2	5/2007	Gannoe et al.
6,258,120	B1	7/2001	McKenzie et al.	7,220,237	B2	5/2007	Gannoe et al.
6,264,700	B1	7/2001	Kilcoyne et al.	7,220,284	B2	5/2007	Kagan et al.
6,287,334	B1	9/2001	Moll et al.	7,223,277	B2	5/2007	DeLegge
6,302,917	B1	10/2001	Dua et al.	7,229,428	B2	6/2007	Gannoe et al.
6,358,197	B1	3/2002	Silverman et al.	7,229,453	B2	6/2007	Anderson et al.
6,416,522	B1	7/2002	Strecker	7,255,675	B2	8/2007	Gertner et al.
6,425,916	B1	7/2002	Garrison et al.	7,261,722	B2	8/2007	McGuckin, Jr. et al.
6,454,785	B2	9/2002	De Hoyos Garza	7,288,101	B2	10/2007	Deem et al.
6,460,543	B1	10/2002	Forsell	7,306,614	B2	12/2007	Weller et al.
6,461,366	B1	10/2002	Seguin	7,315,509	B2	1/2008	Jeong et al.
6,494,888	B1	12/2002	Laufer et al.	7,316,716	B2	1/2008	Egan
6,494,895	B2	12/2002	Addis	7,320,696	B2	1/2008	Gazi et al.
6,503,264	B1	1/2003	Birk	7,326,207	B2	2/2008	Edwards
6,506,196	B1	1/2003	Laufer et al.	7,335,210	B2	2/2008	Smit
6,527,784	B2	3/2003	Adams et al.	7,347,863	B2	3/2008	Rothe et al.
6,540,789	B1	4/2003	Silverman et al.	7,347,875	B2	3/2008	Levine et al.
6,544,291	B2	4/2003	Taylor	7,354,454	B2	4/2008	Stack et al.
6,547,801	B1	4/2003	Dargent et al.	7,399,304	B2	7/2008	Gambale et al.
6,558,400	B2	5/2003	Deem et al.	7,431,725	B2	10/2008	Stack et al.
6,558,429	B2	5/2003	Taylor	7,461,767	B2	12/2008	Viola et al.
6,572,627	B2	6/2003	Gabbay	7,470,251	B2	12/2008	Shah
6,572,629	B2	6/2003	Kaloo	7,485,142	B2	2/2009	Milo
6,575,896	B2	6/2003	Silverman et al.	7,503,922	B2	3/2009	Deem et al.
6,592,596	B1	7/2003	Geitz et al.	7,520,884	B2	4/2009	Swanstrom et al.
6,596,023	B1	7/2003	Nunez et al.	7,575,586	B2	8/2009	Berg et al.
6,607,555	B2	8/2003	Patterson et al.	7,608,114	B2	10/2009	Levine et al.
6,627,206	B2	9/2003	Lloyd	7,615,064	B2	11/2009	Bjerken
6,632,227	B2	10/2003	Adams	7,628,821	B2	12/2009	Stack et al.
6,663,639	B1	12/2003	Laufer et al.	7,662,161	B2	2/2010	Briganti et al.
6,675,809	B2	1/2004	Stack et al.	7,670,279	B2	3/2010	Gertner
6,733,512	B2	5/2004	McGhan	7,674,271	B2	3/2010	Bjerken
6,740,098	B2	5/2004	Abrams et al.	7,695,446	B2	4/2010	Levine et al.
6,740,121	B2	5/2004	Geitz et al.	7,699,863	B2	4/2010	Marco et al.
6,746,460	B2	6/2004	Gannoe et al.	7,708,181	B2	5/2010	Cole et al.
6,755,869	B2	6/2004	Geitz	7,717,843	B2	5/2010	Balbierz et al.
6,764,518	B2	7/2004	Godin	7,721,932	B2	5/2010	Cole et al.
6,773,440	B2	8/2004	Gannoe et al.	7,731,757	B2	6/2010	Taylor et al.
6,773,441	B1	8/2004	Laufer et al.	7,744,613	B2	6/2010	Ewers et al.
6,790,214	B2	9/2004	Kraemer et al.	7,744,627	B2	6/2010	Orban et al.
6,790,237	B2	9/2004	Stinson	7,753,870	B2	7/2010	Demarais et al.
6,821,285	B2	11/2004	Laufer et al.	7,766,861	B2	8/2010	Levine et al.
6,835,200	B2	12/2004	Laufer et al.	7,819,836	B2	10/2010	Levine et al.
6,845,776	B2	1/2005	Stack et al.	7,846,138	B2	12/2010	Dann et al.
6,916,332	B2	7/2005	Adams	7,846,174	B2	12/2010	Baker et al.
6,932,838	B2	8/2005	Schwartz et al.	7,881,797	B2	2/2011	Griffin et al.
6,960,233	B1	11/2005	Berg et al.	7,892,214	B2	2/2011	Kagan et al.
6,966,875	B1	11/2005	Longobardi	7,892,292	B2	2/2011	Stack et al.
6,981,978	B2	1/2006	Gannoe	2001/0011543	A1	8/2001	Forsell
6,981,980	B2	1/2006	Sampson et al.	2001/0020189	A1	9/2001	Taylor
6,994,715	B2	2/2006	Gannoe et al.	2001/0020190	A1	9/2001	Taylor
7,011,094	B2	3/2006	Rapackie et al.	2001/0021796	A1	9/2001	Silverman et al.
7,020,531	B1	3/2006	Colliu et al.	2001/0044595	A1	11/2001	Reydel et al.
7,025,791	B2	4/2006	Levine et al.	2002/0022851	A1	2/2002	Kaloo et al.
7,033,373	B2	4/2006	de la Torre et al.	2002/0055757	A1	5/2002	Torre et al.
7,033,384	B2	4/2006	Gannoe et al.	2002/0072761	A1	6/2002	Abrams et al.
7,037,344	B2	5/2006	Kagan et al.	2002/0082621	A1	6/2002	Shurr et al.
7,056,305	B2	6/2006	Garza	2002/0099439	A1	7/2002	Schwartz et al.
7,066,945	B2	6/2006	Hashiba et al.	2002/0183767	A1	12/2002	Adams et al.
7,083,629	B2	8/2006	Weller et al.	2002/0183768	A1	12/2002	Deem et al.
7,090,699	B2	8/2006	Geitz	2002/0188354	A1	12/2002	Peghini
7,097,650	B2	8/2006	Weller et al.	2003/0009236	A1	1/2003	Godin
7,097,665	B2	8/2006	Stack et al.	2003/0040804	A1	2/2003	Stack et al.
7,111,627	B2	9/2006	Stack et al.	2003/0040808	A1	2/2003	Stack et al.
7,112,186	B2	9/2006	Shah	2003/0065359	A1	4/2003	Weller et al.
7,120,498	B2	10/2006	Imran et al.	2003/0093117	A1	5/2003	Saadat et al.
7,121,283	B2	10/2006	Stack et al.	2003/0109892	A1	6/2003	Deem et al.
7,146,984	B2	12/2006	Stack et al.	2003/0109931	A1	6/2003	Geitz
7,147,140	B2	12/2006	Wukusick et al.	2003/0120289	A1	6/2003	McGuckin, Jr. et al.
7,152,607	B2	12/2006	Stack et al.	2003/0158569	A1	8/2003	Wazne
7,160,312	B2	1/2007	Saadat et al.	2003/0191476	A1	10/2003	Smit
7,172,613	B2	2/2007	Wazne	2003/0191525	A1	10/2003	Thornton
7,175,638	B2	2/2007	Gannoe et al.	2003/0199989	A1	10/2003	Stack et al.
7,175,660	B2	2/2007	Cartledge et al.	2003/0199990	A1	10/2003	Stack et al.
7,211,114	B2	5/2007	Bessler et al.	2003/0199991	A1	10/2003	Stack et al.
				2003/0208209	A1	11/2003	Gambale et al.
				2003/0220660	A1	11/2003	Kortenbach et al.
				2004/0006351	A1	1/2004	Gannoe et al.
				2004/0024386	A1	2/2004	Deem et al.

(56)

References Cited

U.S. PATENT DOCUMENTS

2004/0030347 A1 2/2004 Gannoe et al.
 2004/0044353 A1 3/2004 Gannoe
 2004/0044354 A1 3/2004 Gannoe et al.
 2004/0044357 A1 3/2004 Gannoe et al.
 2004/0044364 A1 3/2004 DeVries et al.
 2004/0059289 A1 3/2004 Garza et al.
 2004/0068276 A1 4/2004 Golden et al.
 2004/0082963 A1 4/2004 Gannoe et al.
 2004/0088023 A1 5/2004 Imran et al.
 2004/0092892 A1 5/2004 Kagan et al.
 2004/0092974 A1 5/2004 Gannoe et al.
 2004/0093091 A1 5/2004 Gannoe et al.
 2004/0098043 A1 5/2004 Trout
 2004/0107004 A1 6/2004 Levine et al.
 2004/0117031 A1 6/2004 Stack et al.
 2004/0133219 A1 7/2004 Forsell
 2004/0138761 A1 7/2004 Stack et al.
 2004/0143342 A1 7/2004 Stack et al.
 2004/0148034 A1 7/2004 Kagan et al.
 2004/0153167 A1 8/2004 Stack et al.
 2004/0158331 A1 8/2004 Stack et al.
 2004/0162568 A1 8/2004 Saadat et al.
 2004/0172141 A1 9/2004 Stack et al.
 2004/0172142 A1 9/2004 Stack et al.
 2004/0186502 A1 9/2004 Sampson et al.
 2004/0210243 A1 10/2004 Gannoe et al.
 2004/0215216 A1 10/2004 Gannoe et al.
 2004/0220682 A1 11/2004 Levine et al.
 2004/0225183 A1 11/2004 Michlitsch et al.
 2004/0225305 A1 11/2004 Ewers et al.
 2004/0236419 A1 11/2004 Milo
 2004/0243152 A1 12/2004 Taylor et al.
 2004/0243223 A1 12/2004 Kraemer et al.
 2004/0267378 A1 12/2004 Gazi et al.
 2005/0004430 A1 1/2005 Lee et al.
 2005/0004681 A1 1/2005 Stack et al.
 2005/0033326 A1 2/2005 Briganti et al.
 2005/0033345 A1 2/2005 DeLegge
 2005/0049718 A1 3/2005 Dann et al.
 2005/0075654 A1 4/2005 Kelleher
 2005/0080444 A1 4/2005 Kraemer et al.
 2005/0085787 A1 4/2005 Laufer et al.
 2005/0096673 A1 5/2005 Stack et al.
 2005/0096750 A1 5/2005 Kagan et al.
 2005/0149114 A1 7/2005 Cartledge et al.
 2005/0159769 A1 7/2005 Alverdy
 2005/0177181 A1 8/2005 Kagan et al.
 2005/0183732 A1 8/2005 Edwards
 2005/0192599 A1 9/2005 Demarais
 2005/0192615 A1 9/2005 Torre et al.
 2005/0216040 A1 9/2005 Gertner et al.
 2005/0216042 A1 9/2005 Gertner
 2005/0228504 A1 10/2005 Demarais et al.
 2005/0240279 A1 10/2005 Kagan et al.
 2005/0245965 A1 11/2005 Orban et al.
 2005/0247320 A1 11/2005 Stack et al.
 2005/0250980 A1 11/2005 Swanstrom et al.
 2005/0251158 A1 11/2005 Sadat et al.
 2005/0251162 A1 11/2005 Rothe et al.
 2005/0256533 A1 11/2005 Roth et al.
 2005/0256587 A1 11/2005 Egan
 2005/0261712 A1 11/2005 Balbierz et al.
 2005/0267405 A1 12/2005 Shah
 2005/0267499 A1 12/2005 Stack et al.
 2005/0267595 A1 12/2005 Chen et al.
 2005/0267596 A1 12/2005 Chen et al.
 2005/0273060 A1 12/2005 Levy et al.
 2006/0015006 A1 1/2006 Laurence et al.
 2006/0020278 A1 1/2006 Burnette et al.
 2006/0058829 A1 3/2006 Sampson et al.
 2006/0129094 A1 6/2006 Shah
 2006/0151568 A1 7/2006 Weller et al.
 2006/0155259 A1 7/2006 MacLay
 2006/0155311 A1 7/2006 Hashiba et al.
 2006/0178560 A1 8/2006 Saadat et al.

2006/0178691 A1 8/2006 Binmoeller
 2006/0195139 A1 8/2006 Gertner
 2006/0253142 A1 11/2006 Bjerken
 2006/0271076 A1 11/2006 Weller et al.
 2006/0282095 A1 12/2006 Stokes et al.
 2006/0287734 A1 12/2006 Stack et al.
 2007/0010864 A1 1/2007 Dann et al.
 2007/0027548 A1 2/2007 Levine et al.
 2007/0032800 A1 2/2007 Ortiz et al.
 2007/0043384 A1 2/2007 Ortiz et al.
 2007/0055292 A1 3/2007 Ortiz et al.
 2007/0060932 A1 3/2007 Stack et al.
 2007/0149994 A1 6/2007 Sosnowski et al.
 2007/0175488 A1 8/2007 Cox et al.
 2007/0191870 A1 8/2007 Baker et al.
 2007/0191871 A1 8/2007 Baker et al.
 2007/0198074 A1 8/2007 Dann et al.
 2007/0219571 A1* 9/2007 Balbierz A61F 5/0036
 606/153
 2007/0239284 A1 10/2007 Skervén et al.
 2007/0260327 A1 11/2007 Case et al.
 2007/0276428 A1 11/2007 Haller et al.
 2007/0276432 A1 11/2007 Stack et al.
 2008/0033574 A1 2/2008 Bessler et al.
 2008/0065122 A1 3/2008 Stack et al.
 2008/0097510 A1 4/2008 Albrecht et al.
 2008/0116244 A1 5/2008 Rethy et al.
 2008/0190989 A1* 8/2008 Crews A61B 17/068
 227/176.1
 2008/0195226 A1 8/2008 Williams et al.
 2008/0208355 A1 8/2008 Stack et al.
 2008/0208356 A1 8/2008 Stack et al.
 2008/0269797 A1 10/2008 Stack et al.
 2008/0294179 A1 11/2008 Balbierz et al.
 2008/0319471 A1 12/2008 Sosnowski et al.
 2009/0018558 A1 1/2009 Laufer et al.
 2009/0024143 A1 1/2009 Crews et al.
 2009/0030284 A1 1/2009 Cole et al.
 2009/0125040 A1 5/2009 Hambley et al.
 2009/0171383 A1 7/2009 Cole et al.
 2009/0177215 A1 7/2009 Stack et al.
 2009/0182424 A1 7/2009 Marco et al.
 2009/0236388 A1 9/2009 Cole et al.
 2009/0236389 A1 9/2009 Cole et al.
 2009/0236390 A1 9/2009 Cole et al.
 2009/0236391 A1 9/2009 Cole et al.
 2009/0236392 A1 9/2009 Cole et al.
 2009/0236394 A1 9/2009 Cole et al.
 2009/0236396 A1 9/2009 Cole et al.
 2009/0236397 A1 9/2009 Cole et al.
 2009/0236398 A1 9/2009 Cole et al.
 2009/0236400 A1 9/2009 Cole et al.
 2009/0236401 A1 9/2009 Cole et al.
 2009/0299487 A1 12/2009 Stack et al.
 2010/0016988 A1 1/2010 Stack et al.
 2010/0100109 A1 4/2010 Stack et al.
 2010/0116867 A1 5/2010 Balbierz et al.
 2010/0204719 A1 8/2010 Balbierz et al.

FOREIGN PATENT DOCUMENTS

DE 08708978 U1 11/1987
 EP 1256318 A1 11/2002
 EP 0775471 1/2005
 EP 1492478 1/2005
 EP 1602336 12/2005
 FR 2768324 3/1999
 JP 09-168597 6/1997
 WO WO 91/01117 2/1991
 WO WO 95/25468 A1 9/1995
 WO WO 97/47231 12/1997
 WO WO 00/12027 3/2000
 WO WO 00/32137 6/2000
 WO WO 00/78227 12/2000
 WO WO 01/41671 6/2001
 WO WO 01/45485 6/2001
 WO WO 01/49359 7/2001
 WO WO 01/66018 9/2001
 WO WO 01/85034 11/2001

(56)

References Cited

FOREIGN PATENT DOCUMENTS

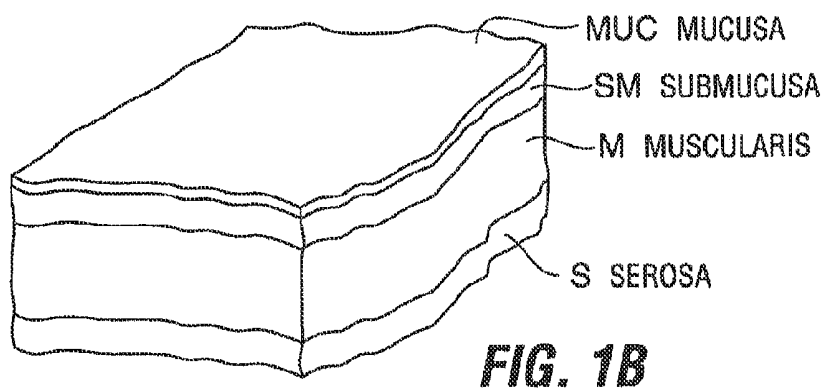
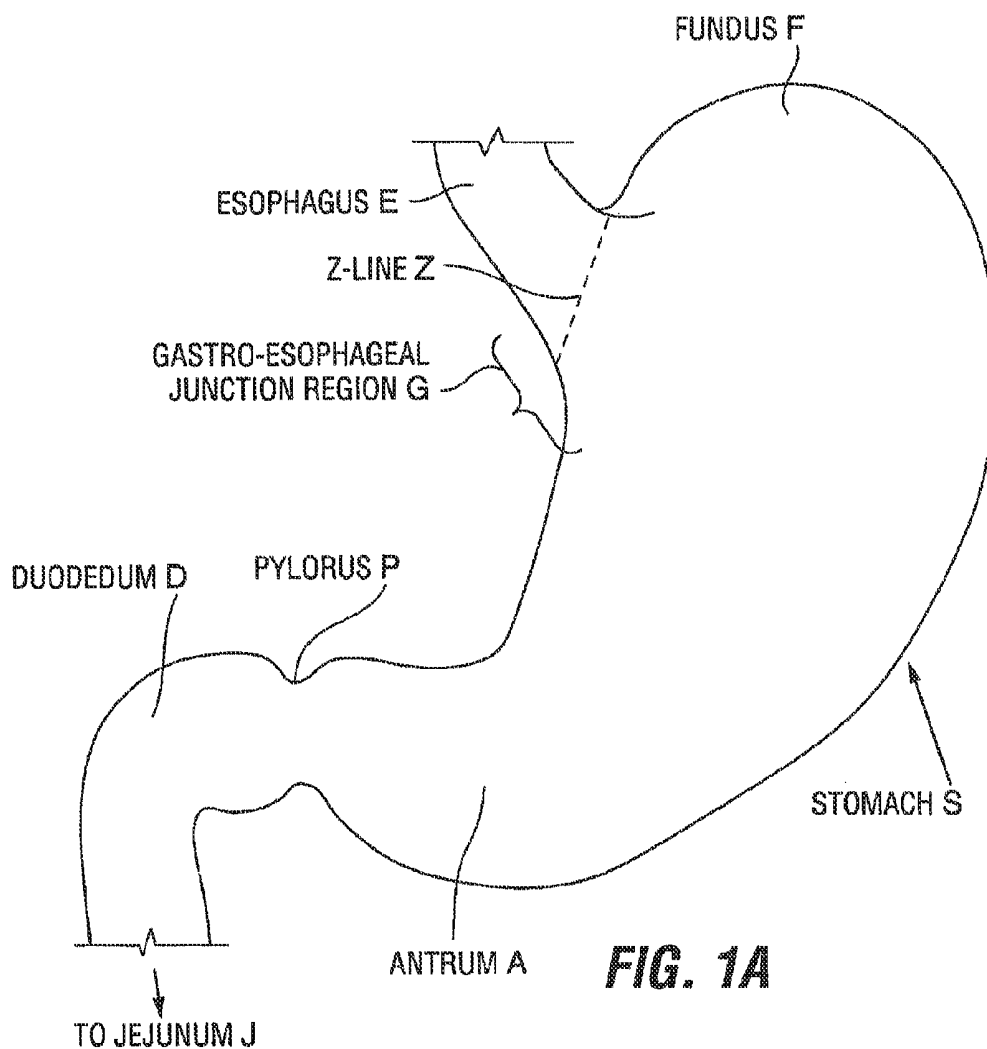
WO	WO 01/89393	11/2001
WO	WO 02/060328	8/2002
WO	WO 03/017882	3/2003
WO	WO 03/086246	10/2003
WO	WO 03/088247	10/2003
WO	WO 03/090633	11/2003
WO	WO 03/094784	11/2003
WO	WO 03/094785	11/2003
WO	WO 03/099137	12/2003
WO	WO 03/105698 A2	12/2003
WO	WO 2004/019765	3/2004
WO	WO 2004/019787	3/2004
WO	WO 2004/032760	4/2004
WO	WO 2004/037064	5/2004
WO	WO 2004/041133	5/2004
WO	WO 2004/064680	8/2004
WO	WO 2004/064685	8/2004
WO	WO 2004/080336	9/2004
WO	WO 2004/110285	12/2004
WO	WO 2005/037152	4/2005
WO	WO 2005/079673	9/2005
WO	WO 2005/096991	10/2005
WO	WO 2005/105003	11/2005
WO	WO 2006/016894	2/2006
WO	WO 2006/055365	5/2006
WO	WO 2006/127593	11/2006
WO	WO 2007/041598	4/2007
WO	WO 2008/030403	3/2008
WO	WO 2008/033409	3/2008
WO	WO 2008/033474	3/2008
WO	WO 2008/141288	11/2008
WO	WO 2009/011881	1/2009
WO	WO 2009/011882	1/2009
WO	WO 2009/086549	7/2009
WO	WO 2009/117533	9/2009
WO	WO 2010/054399	5/2010
WO	WO 2010/054404	5/2010

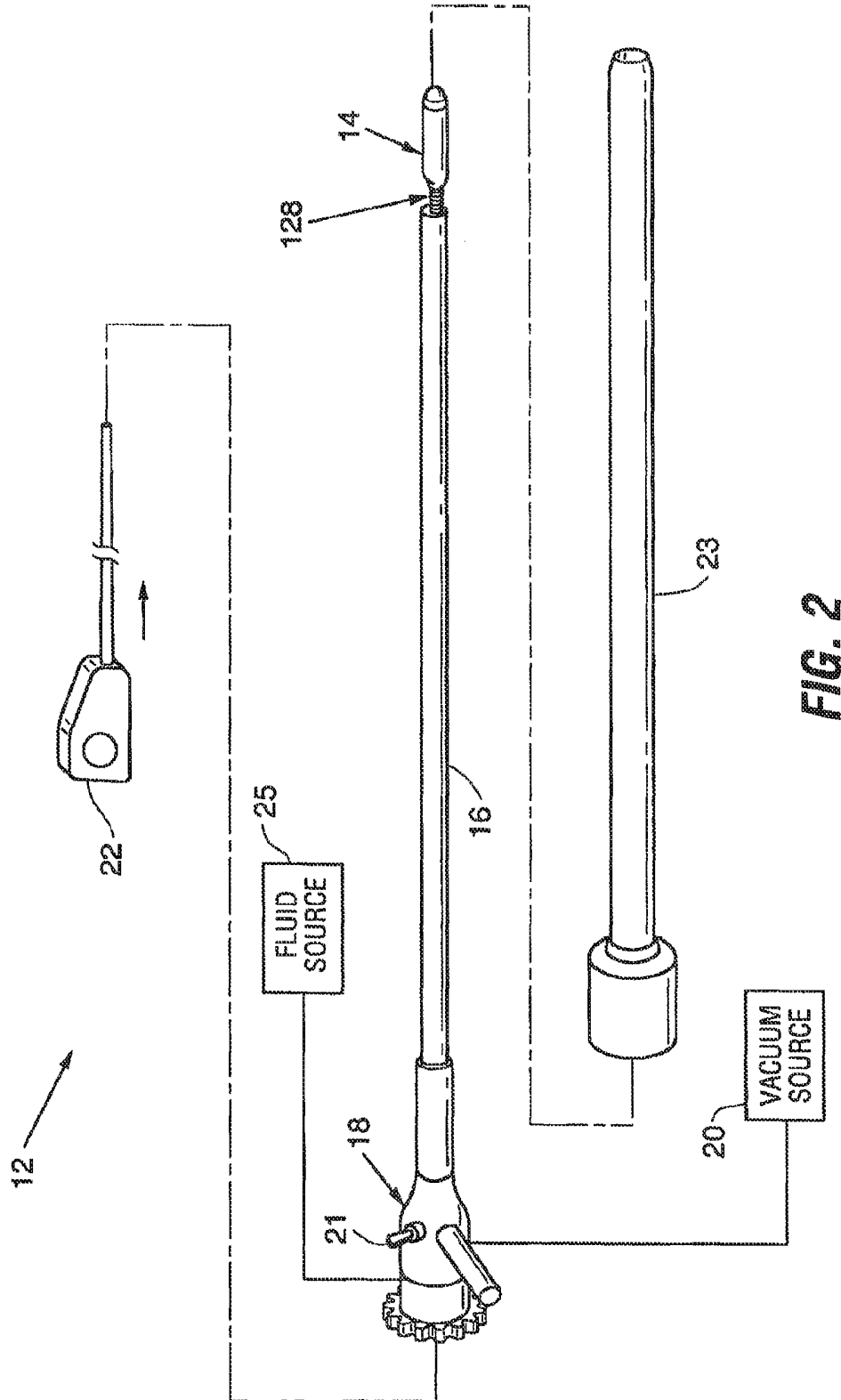
OTHER PUBLICATIONS

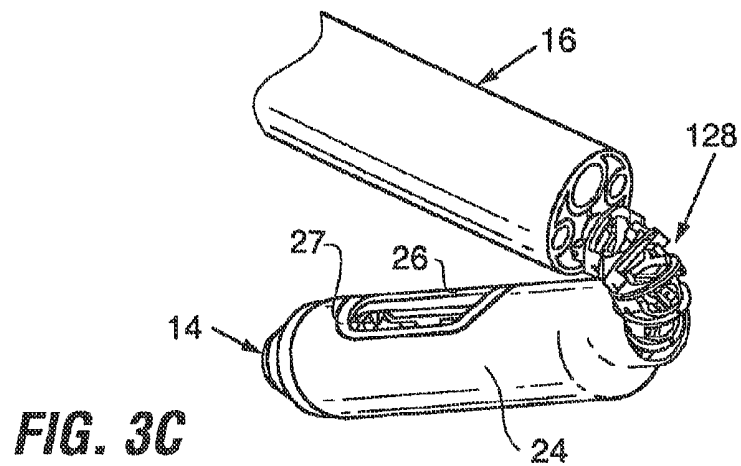
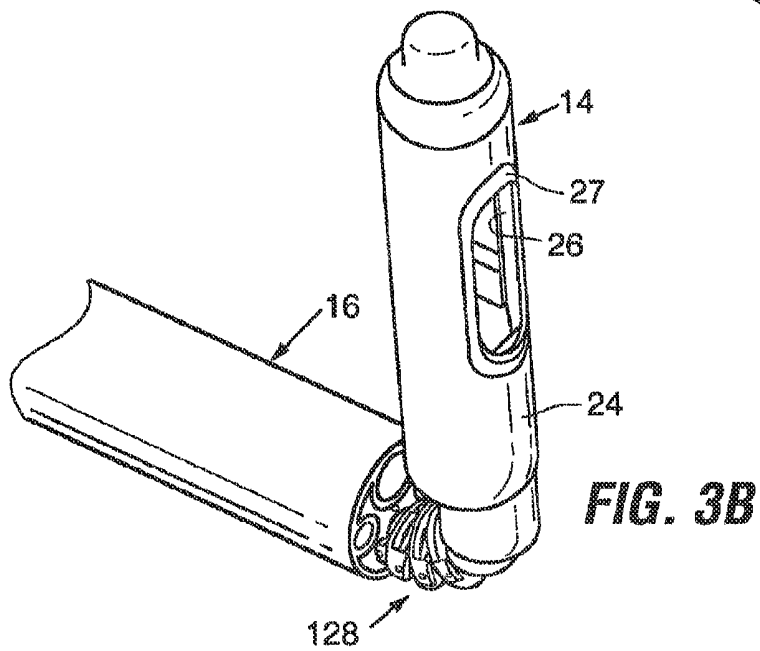
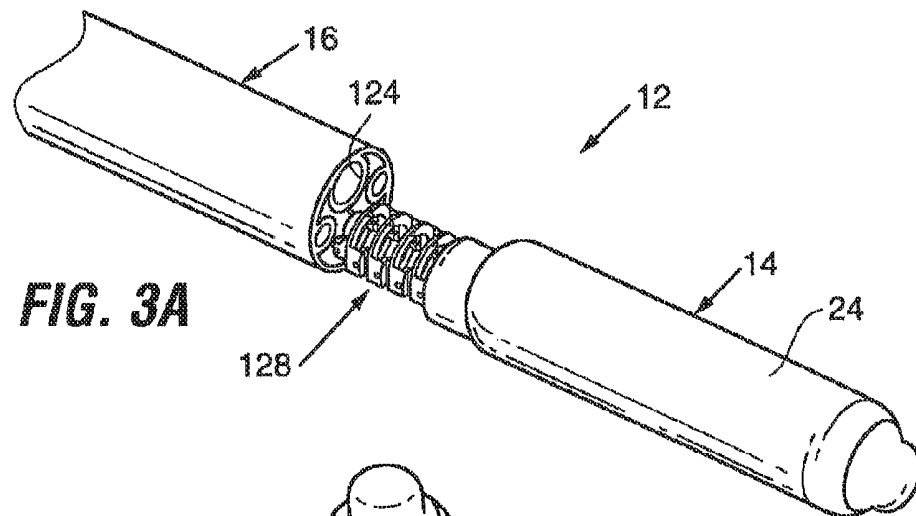
International Search Report from PCT Patent Application No. PCT/US2008/008729 mailed Aug. 18, 2009.
 International Search Report from PCT Patent Application No. PCT/US2008/063440 mailed Aug. 1, 2008.

International Search Report from PCT Patent Application No. PCT/US2008/088581 mailed Feb. 26, 2009.
 International Search Report from PCT Patent Application No. PCT/US2009/037586 mailed Sep. 28, 2009.
 International Search Report from PCT Patent Application No. PCT/US2009/063925 mailed Jan. 12, 2010.
 International Search Report from PCT Patent Application No. PCT/US2009/063930 mailed Jan. 12, 2010.
 International Search Report from PCT Patent Application No. PCT/US2002/027177 mailed Feb. 14, 2003.
 International Search Report from PCT Patent Application No. PCT/US2003/004378 mailed Aug. 13, 2003.
 International Search Report from PCT Patent Application No. PCT/US2003/033605 mailed Mar. 29, 2004.
 International Search Report from PCT Patent Application No. PCT/US2003/033606 mailed Mar. 29, 2004.
 International Search Report from PCT Patent Application No. PCT/US2003/004449 mailed Aug. 13, 2003.
 International Search Report from PCT Patent Application No. PCT/US2004/006695 mailed Sep. 8, 2004.
 International Search Report from PCT Patent Application No. PCT/US2004/033007 mailed Feb. 9, 2005.
 International Search Report from PCT Patent Application No. PCT/US2005/014372 mailed Jul. 28, 2005.
 International Search Report from PCT Patent Application No. PCT/US2006/019727 mailed Apr. 19, 2007.
 International Search Report from PCT Patent Application No. PCT/US2006/038684 mailed Feb. 14, 2007.
 International Search Report from PCT Patent Application No. PCT/US2007/019227 mailed Feb. 20, 2008.
 International Search Report from PCT Patent Application No. PCT/US2007/019833 mailed Feb. 20, 2008.
 International Search Report from PCT Patent Application No. PCT/US2007/019940 mailed Mar. 14, 2008.
 Felsher, et al., "Mucosal apposition in endoscopic suturing", Gastrointestinal Endoscopy, vol. 58, No. 6, pp. 867-870, (2003).
 Stecco, et al., "Trans-oral plication formation and gastric implant placement in a canine model", Stecco Group, San Jose and Barosense Inc., Redwood City, CA (2004).
 Stecco, et al. "Safety of a gastric restrictive implant in a canine model", Stecco group, San Jose and Barosense, Inc., Redwood CA (2004).

* cited by examiner







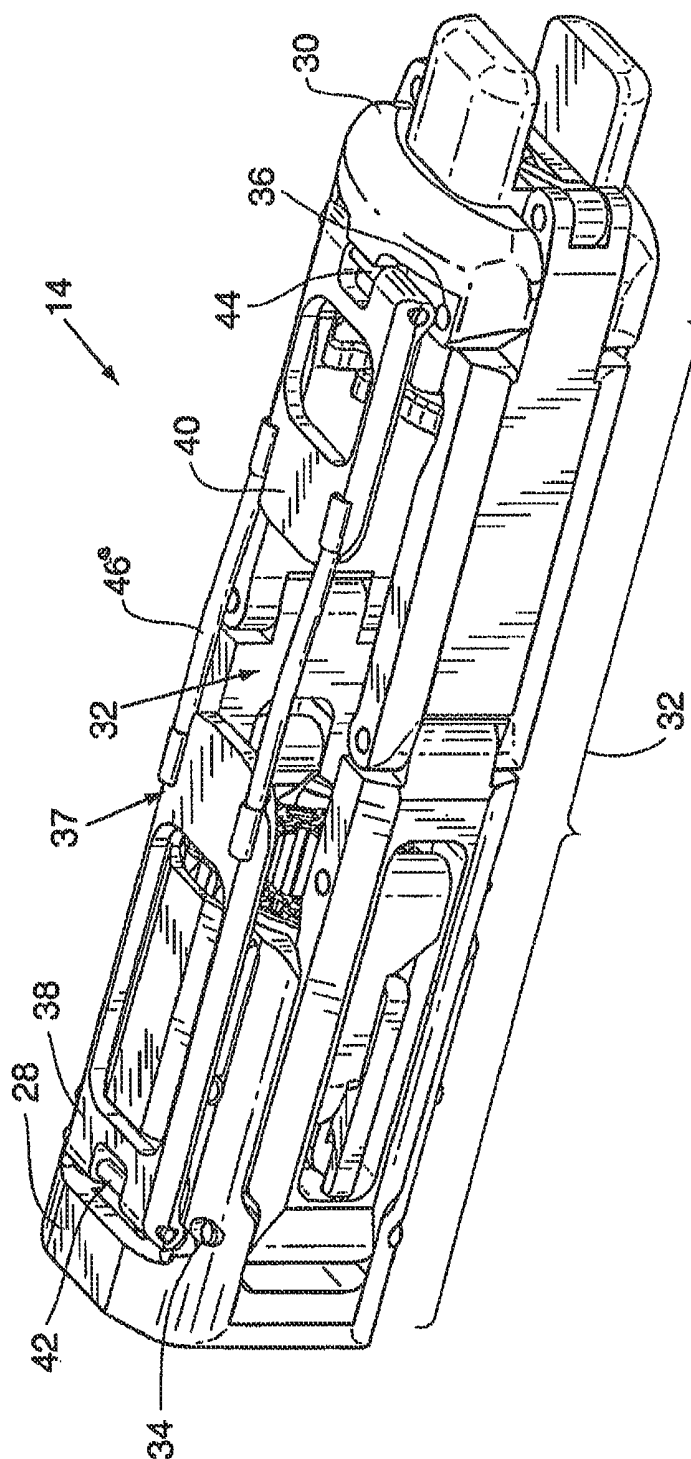


FIG. 4

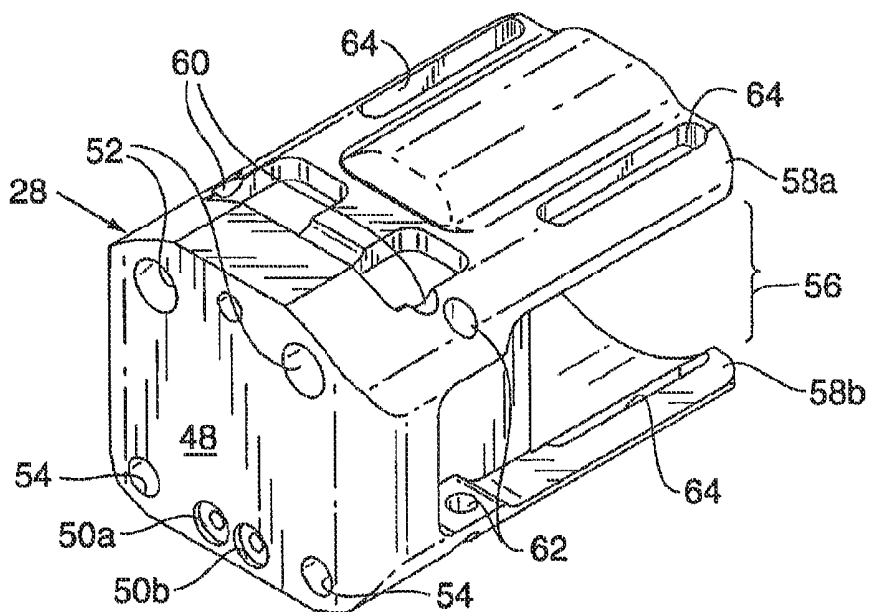


FIG. 5

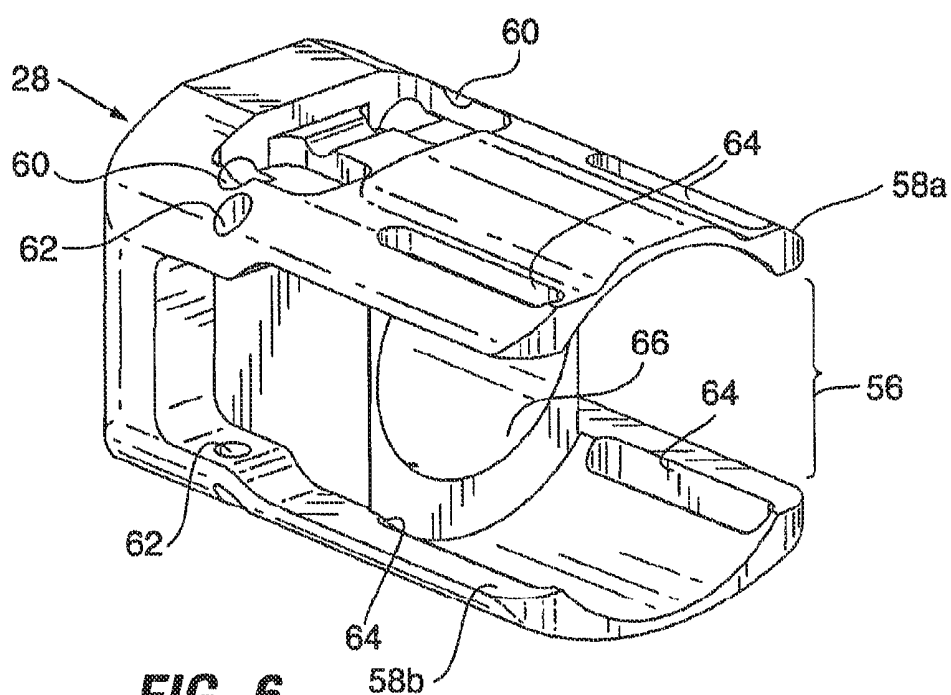
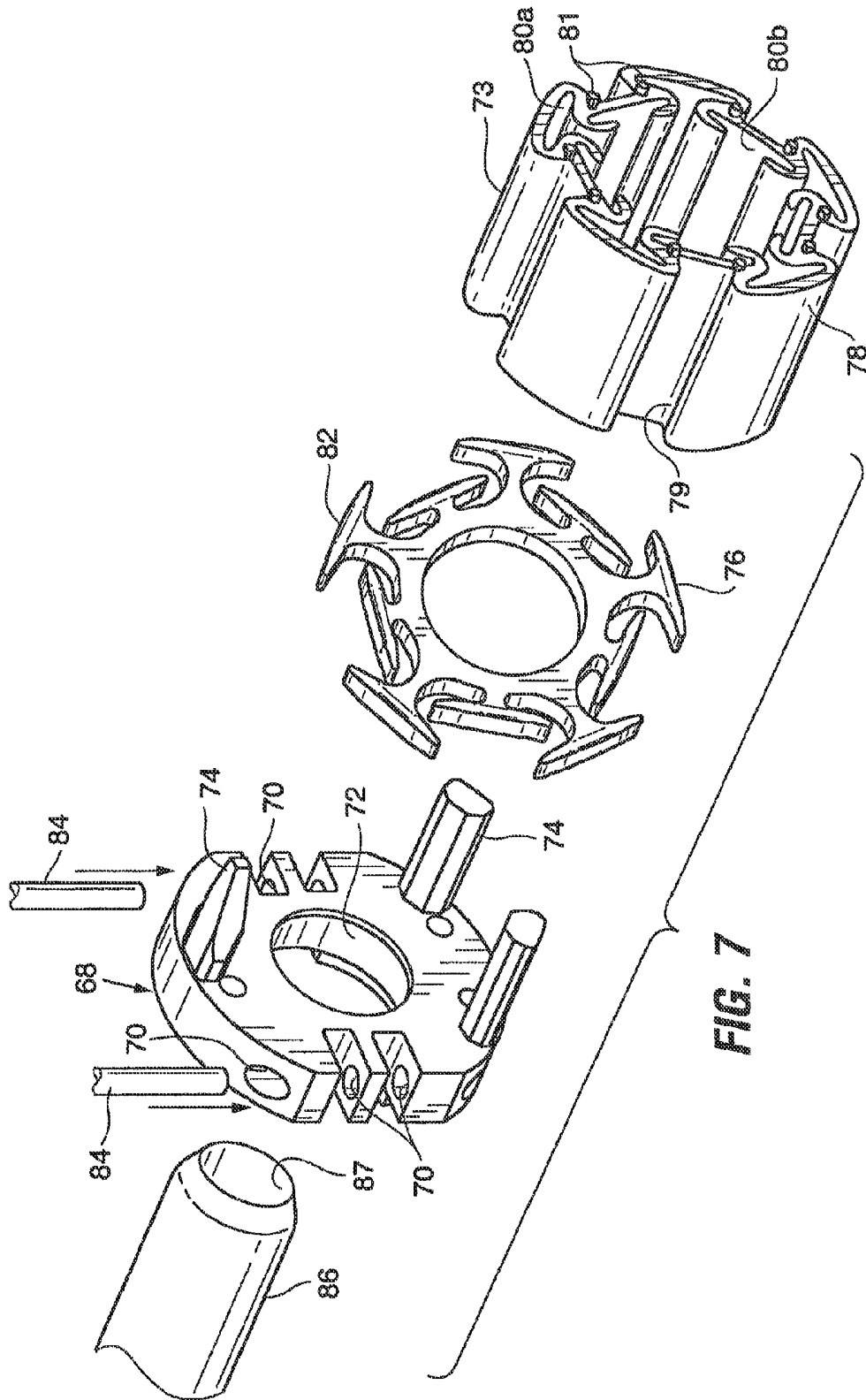
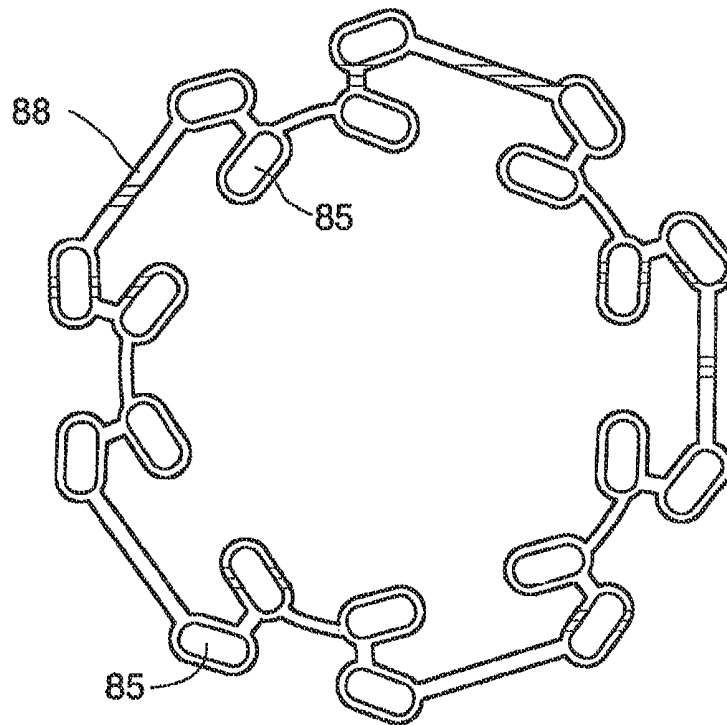
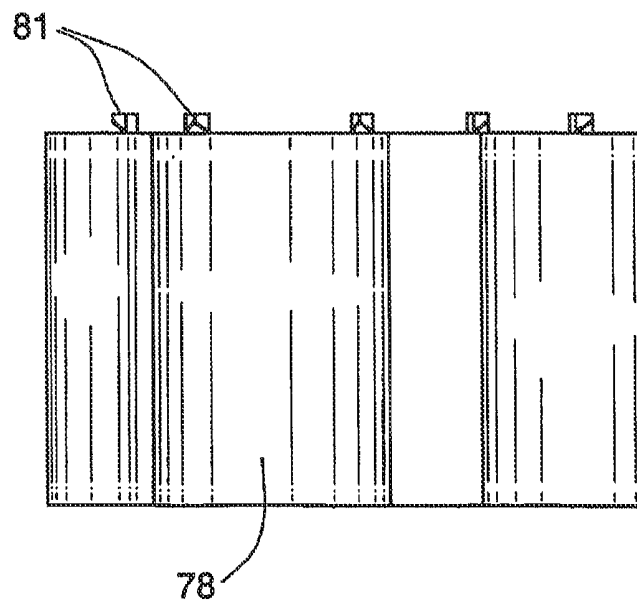
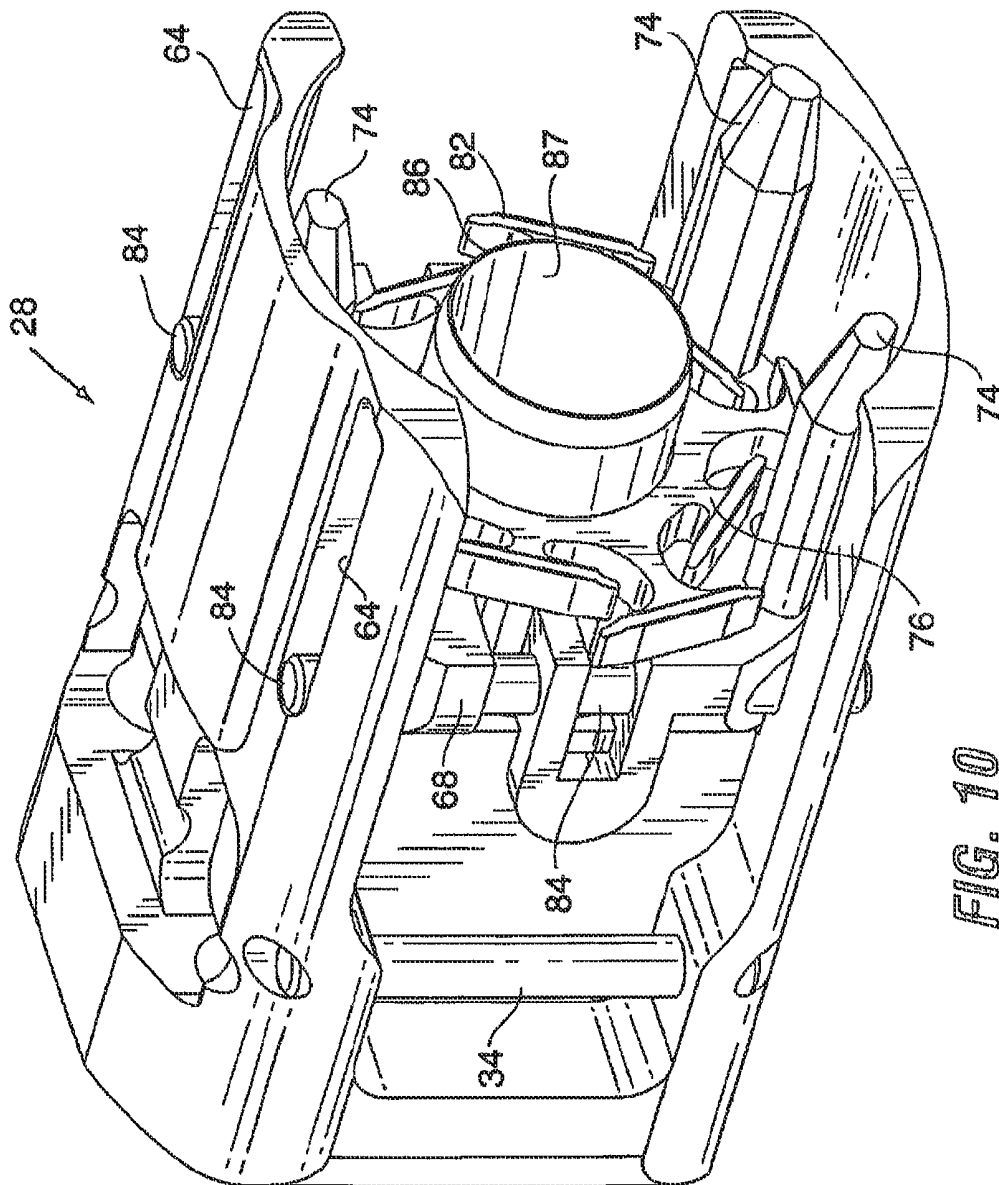


FIG. 6



**FIG. 8****FIG. 9**



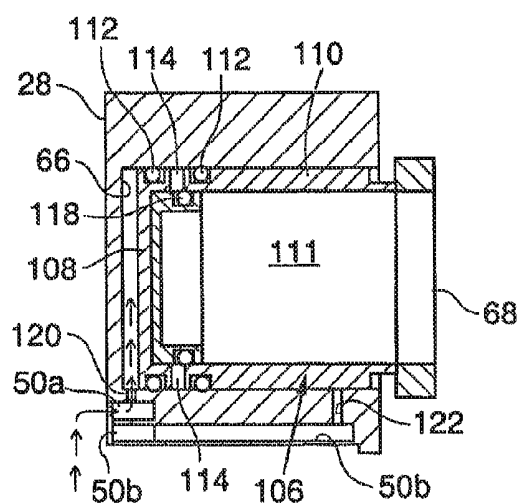


FIG. 11A

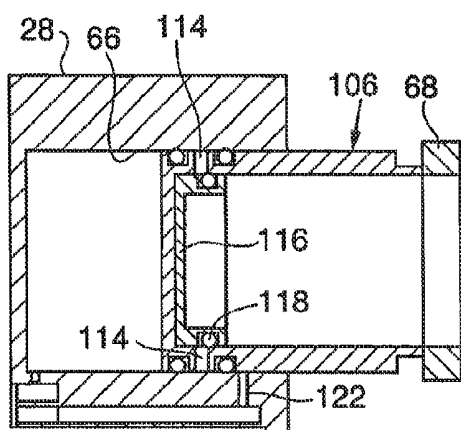


FIG. 11B

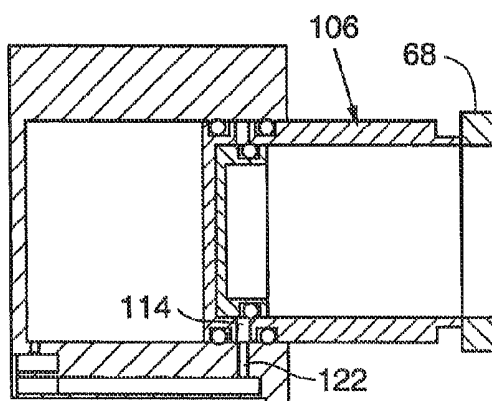


FIG. 11C

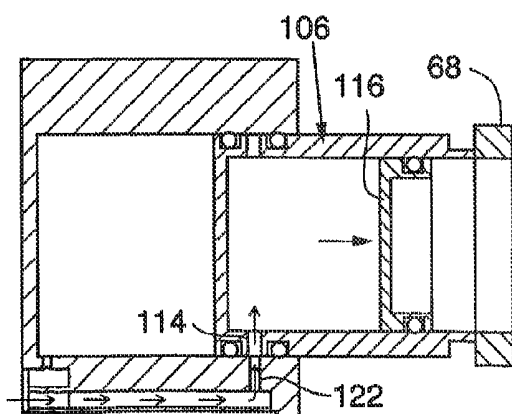


FIG. 11D

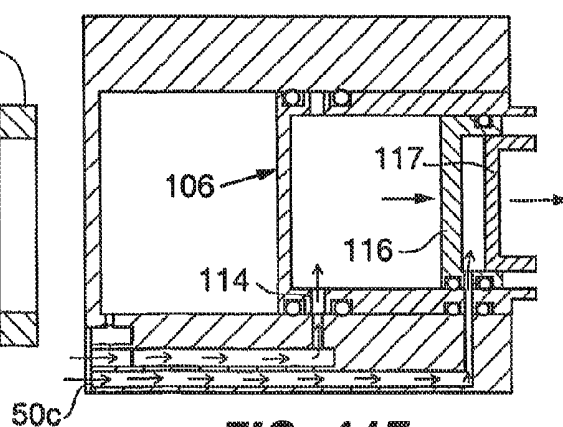
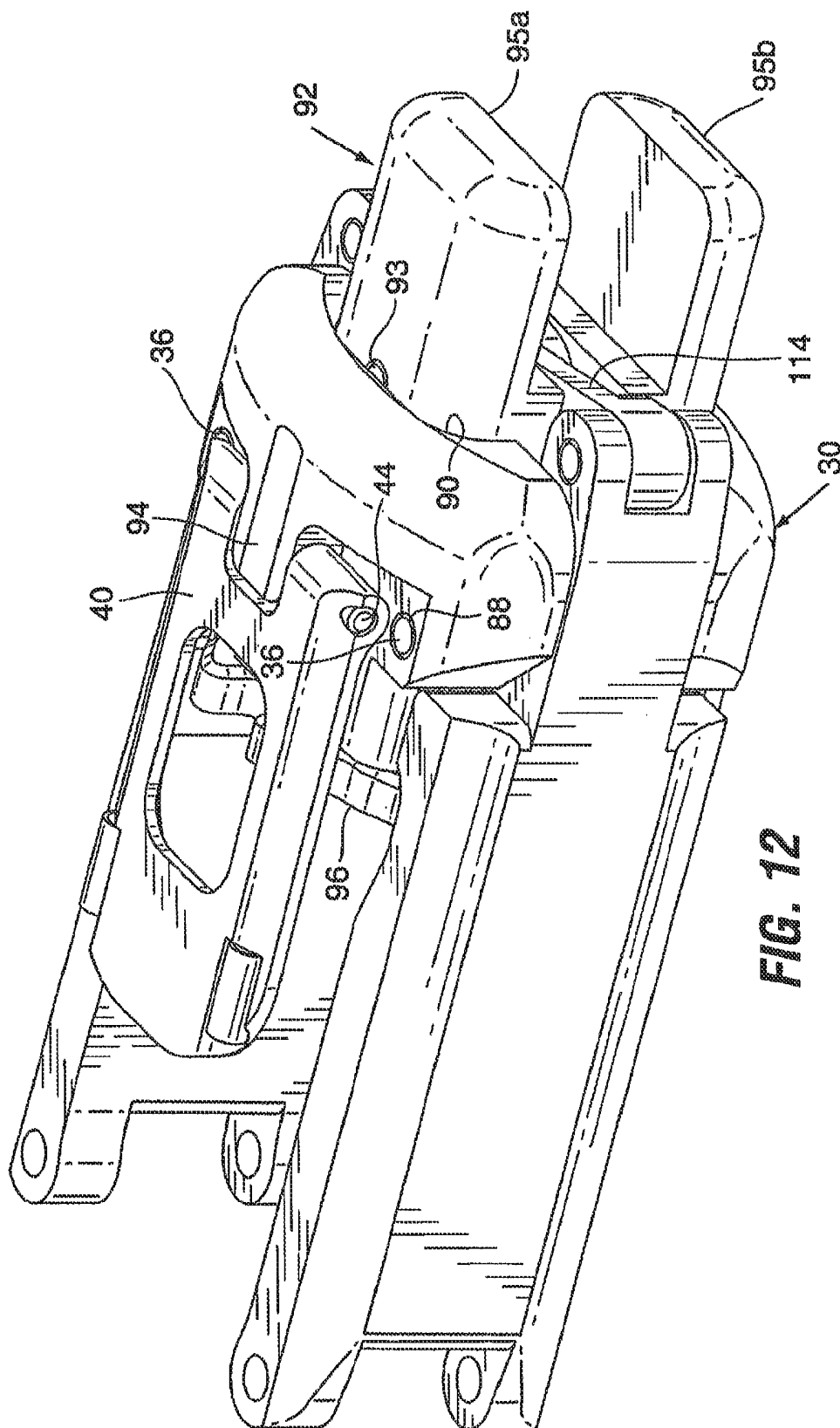
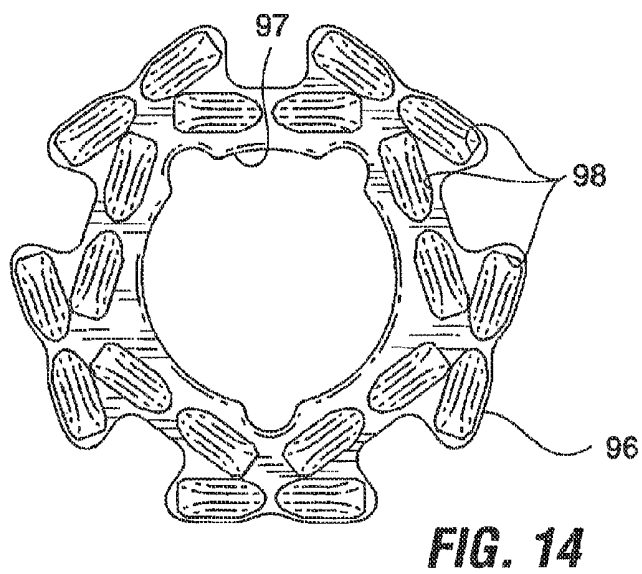
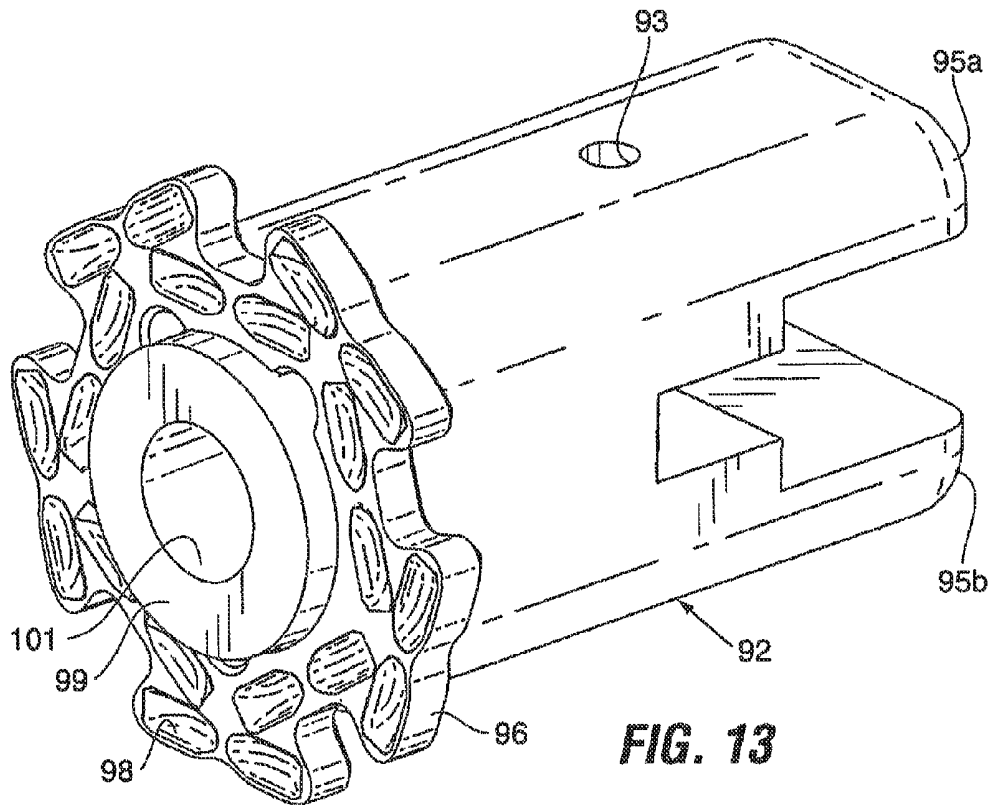


FIG. 11E





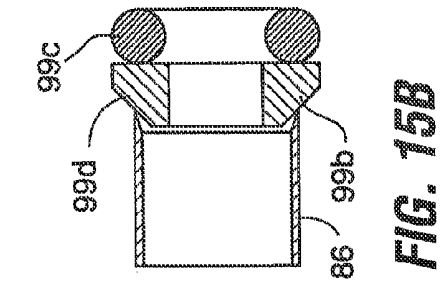


FIG. 15A

FIG. 15B

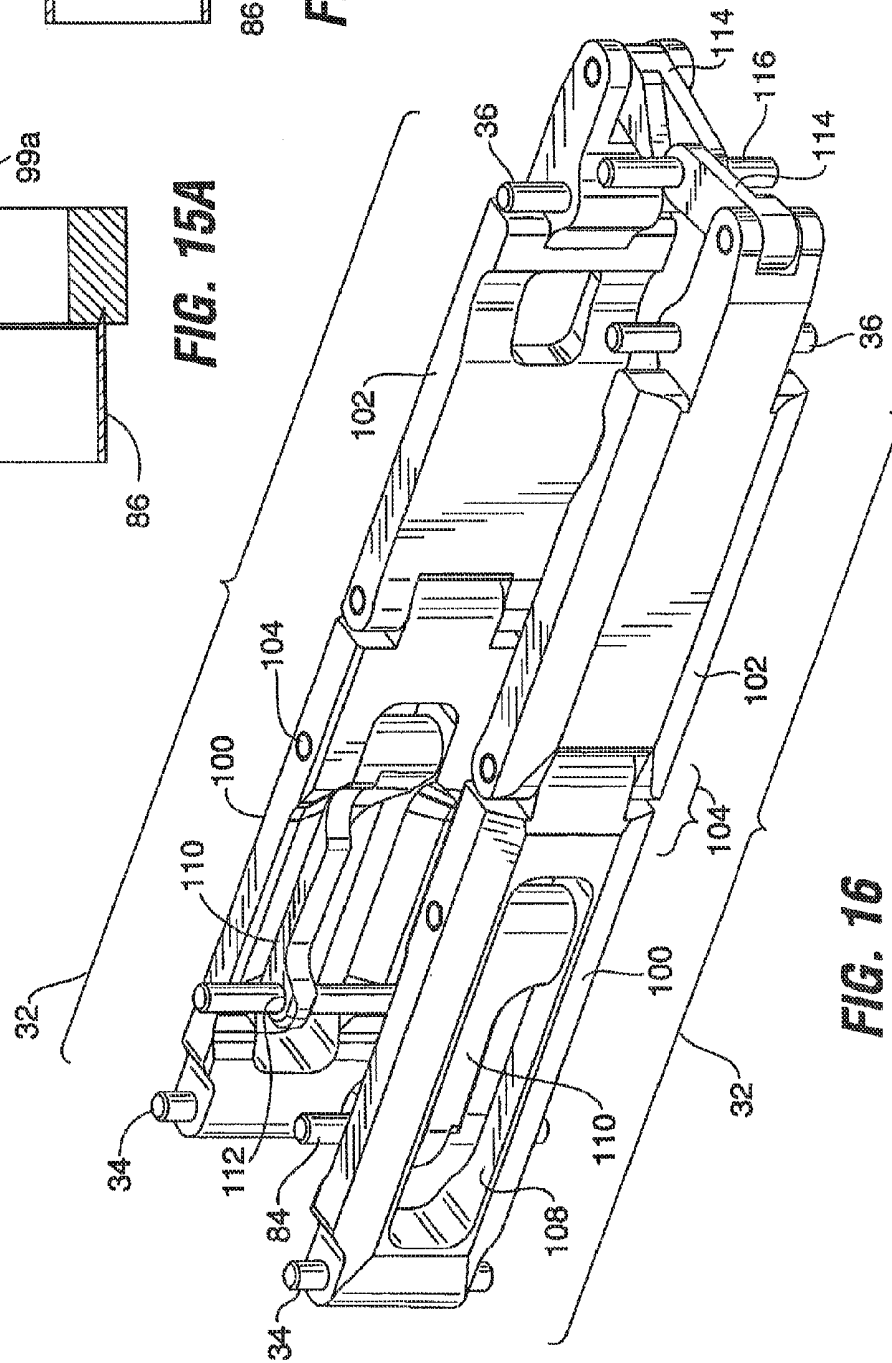


FIG. 16

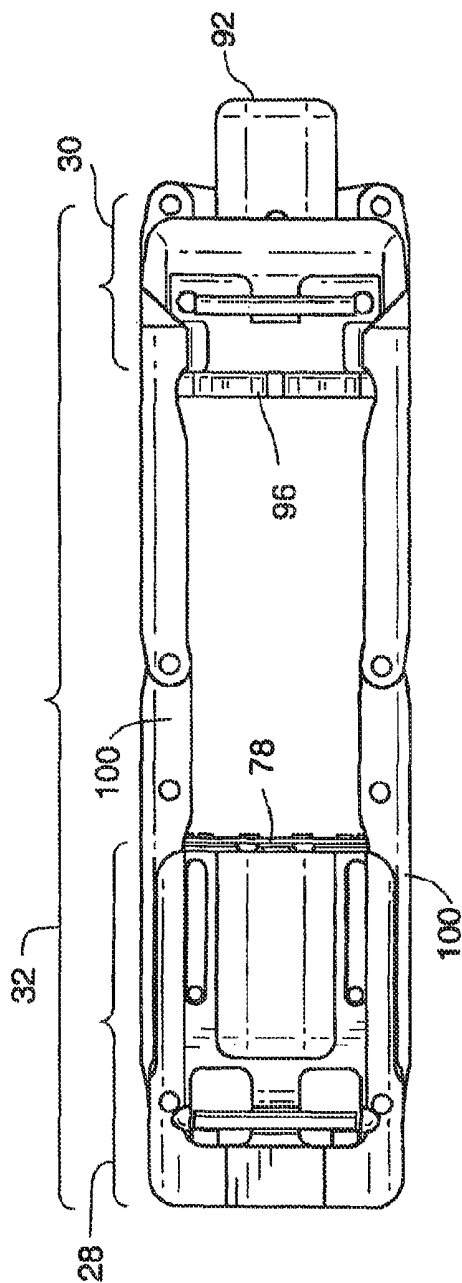


FIG. 17

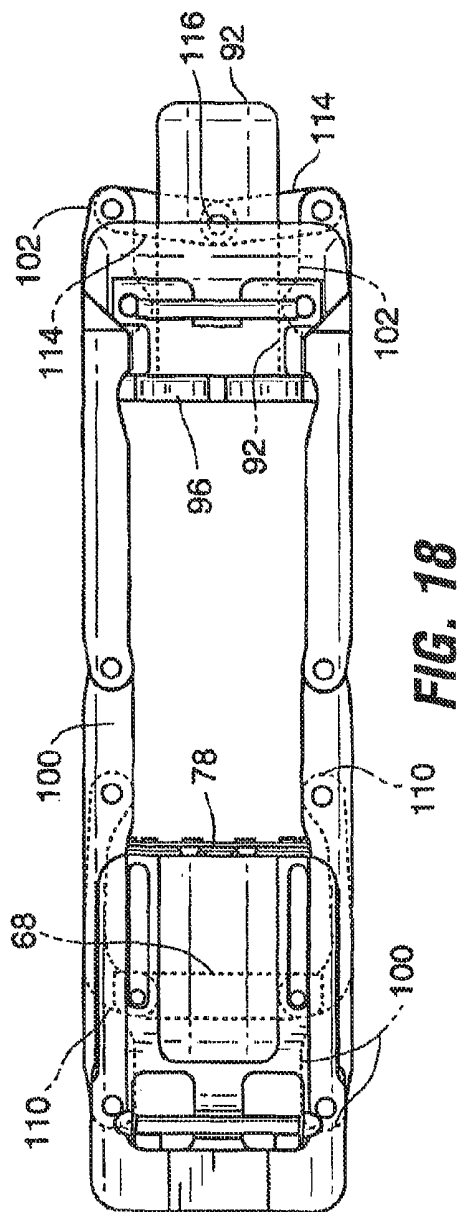


FIG. 18

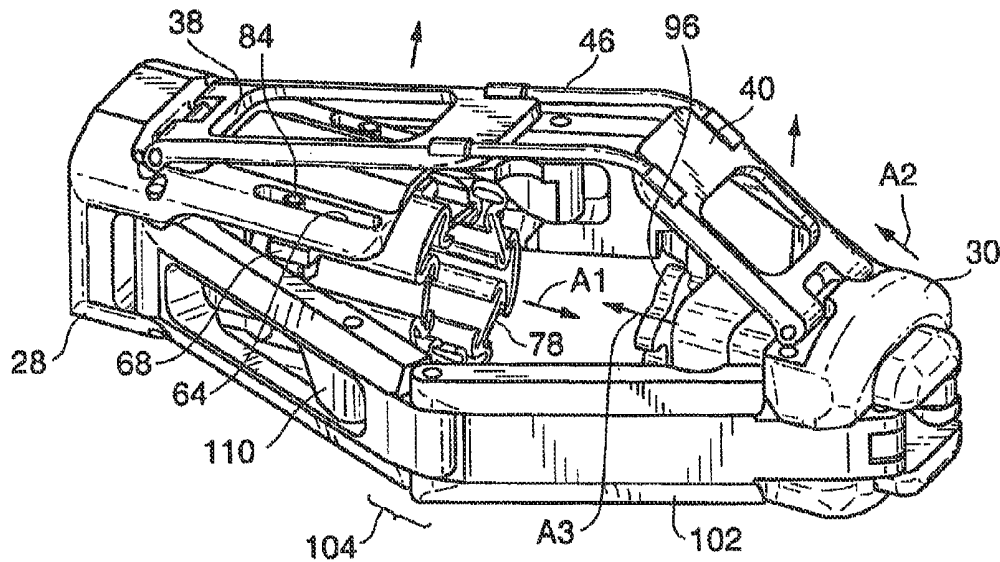


FIG. 19

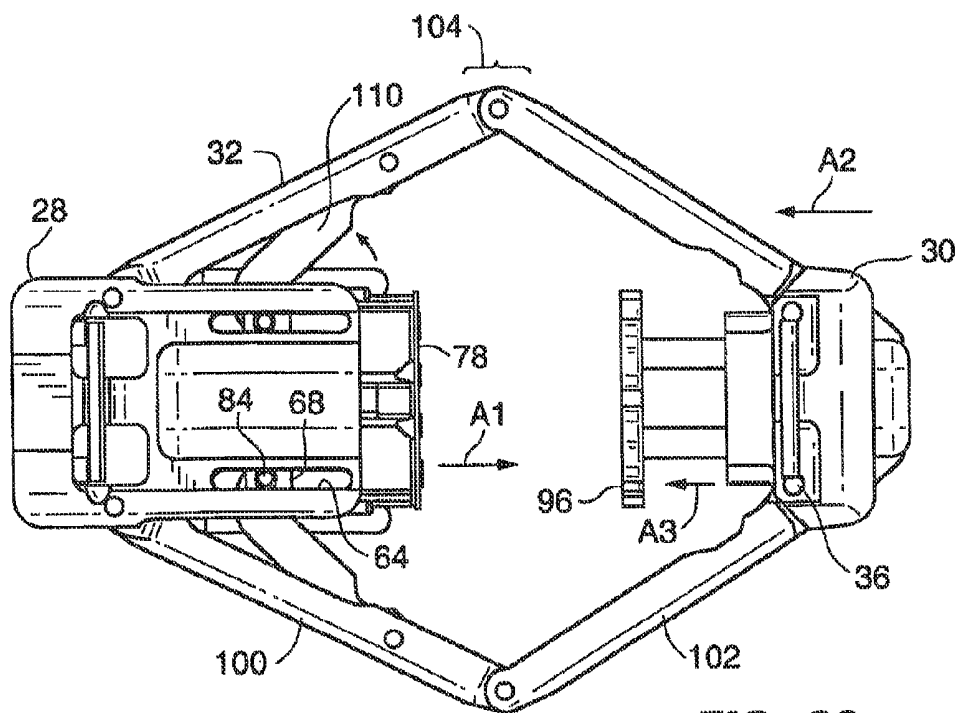


FIG. 20

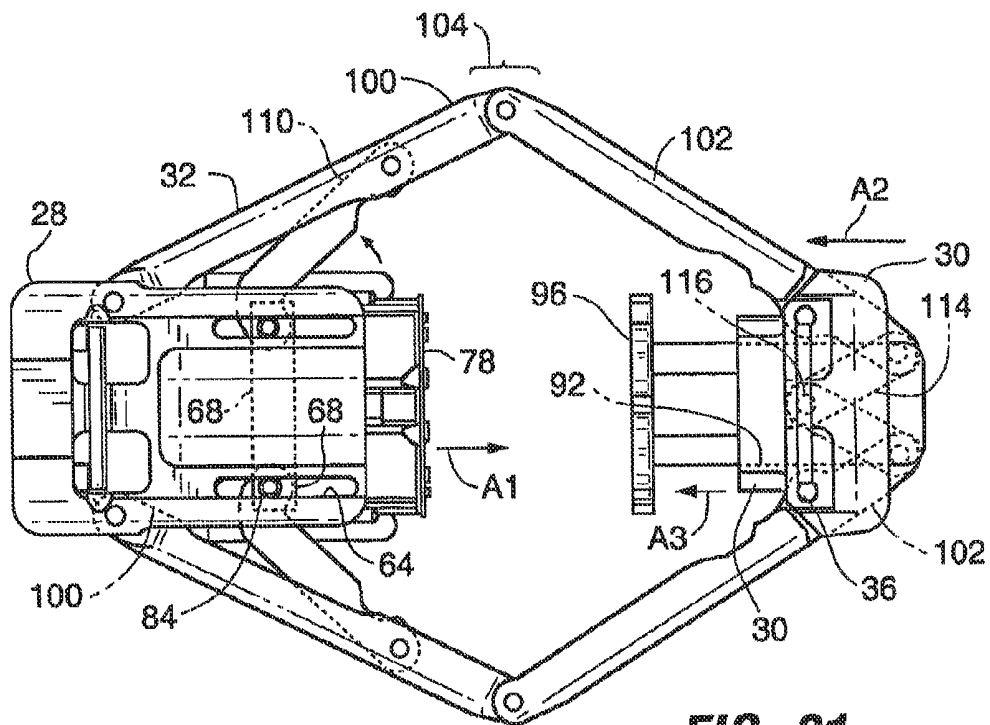


FIG. 21

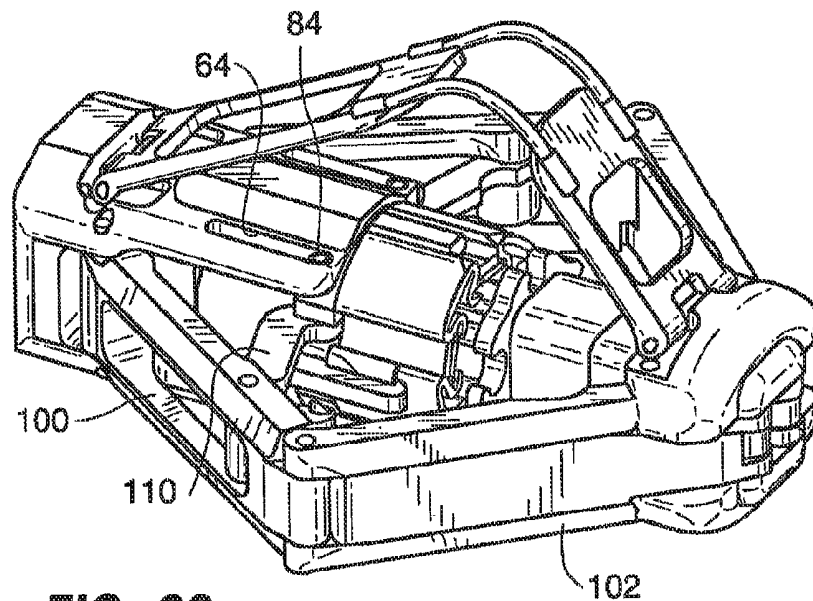


FIG. 22

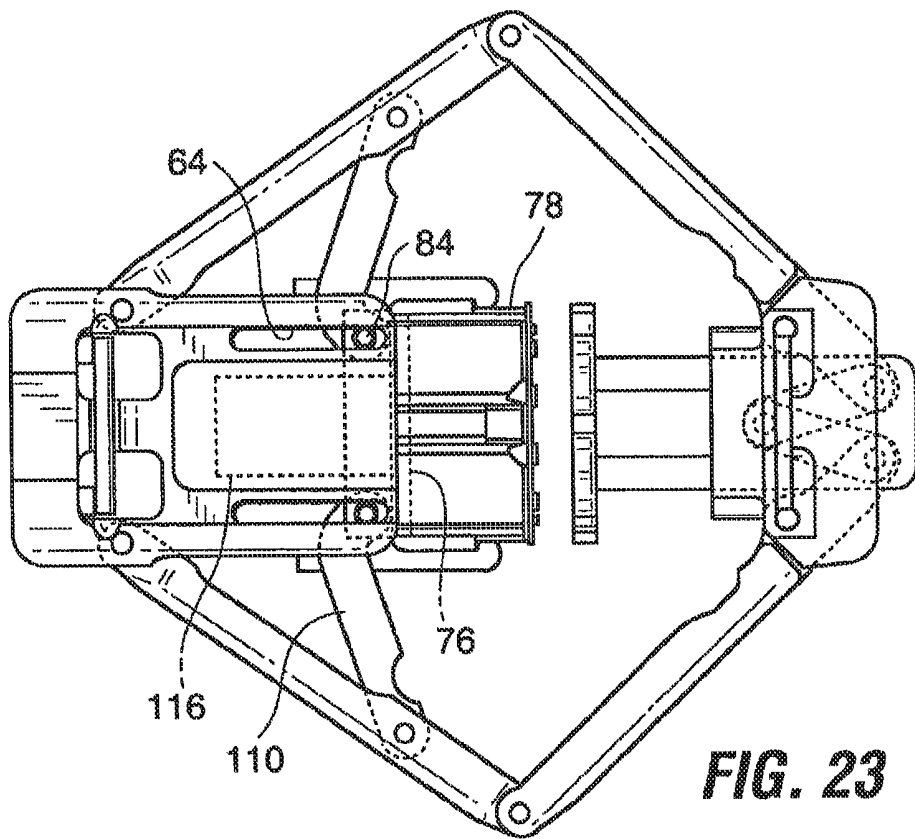


FIG. 23

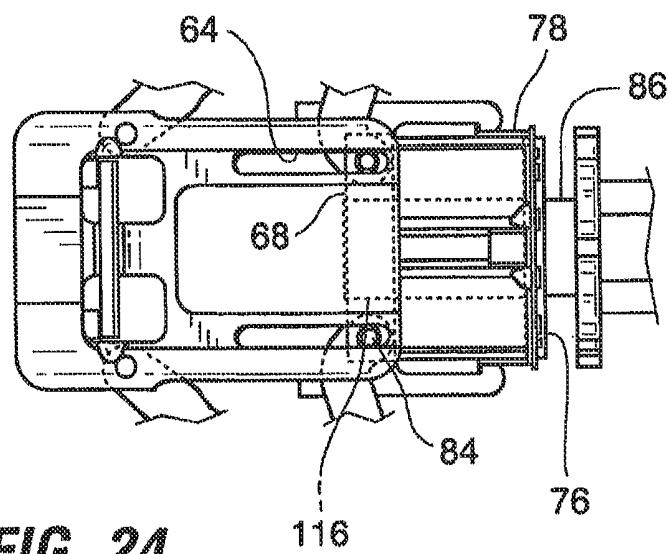
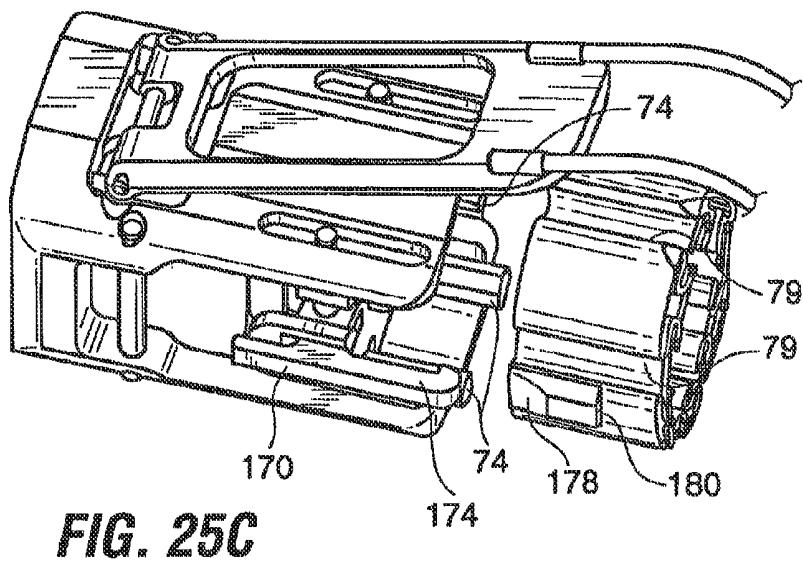
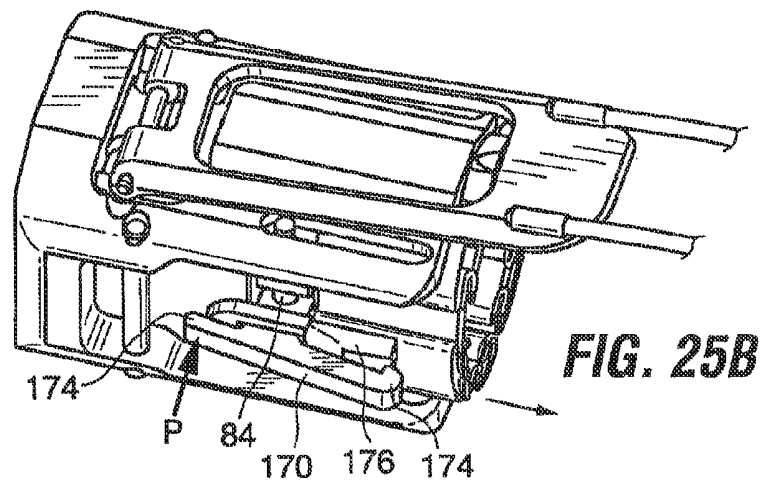
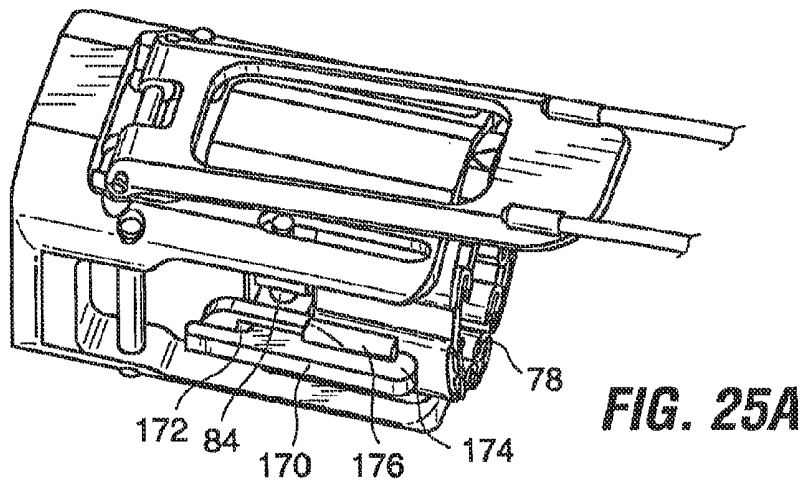
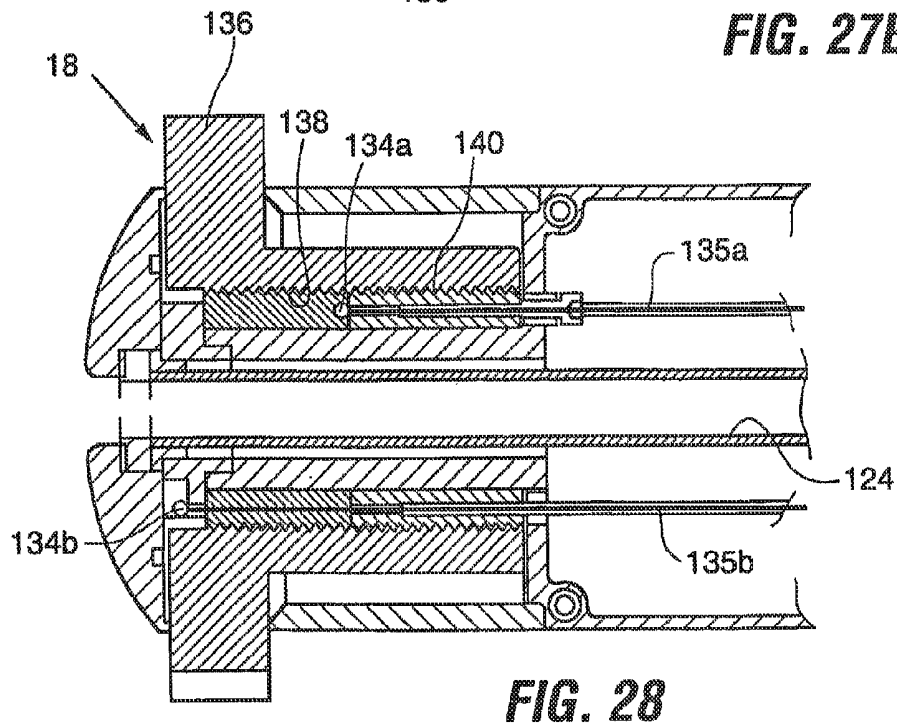
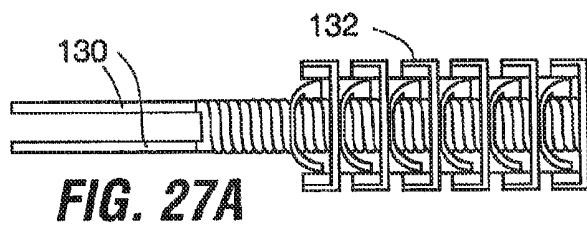
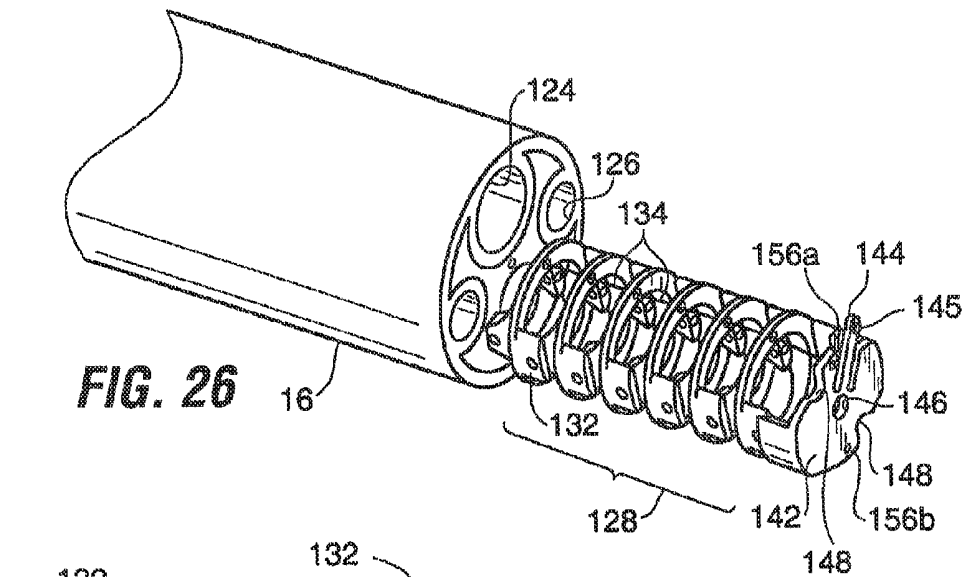


FIG. 24





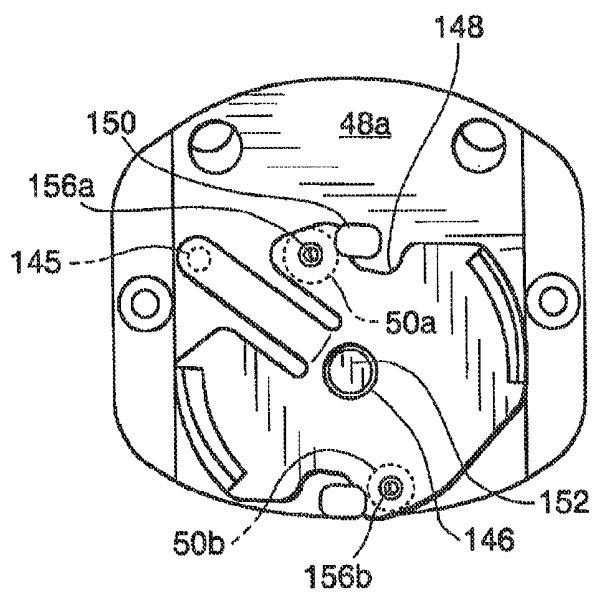
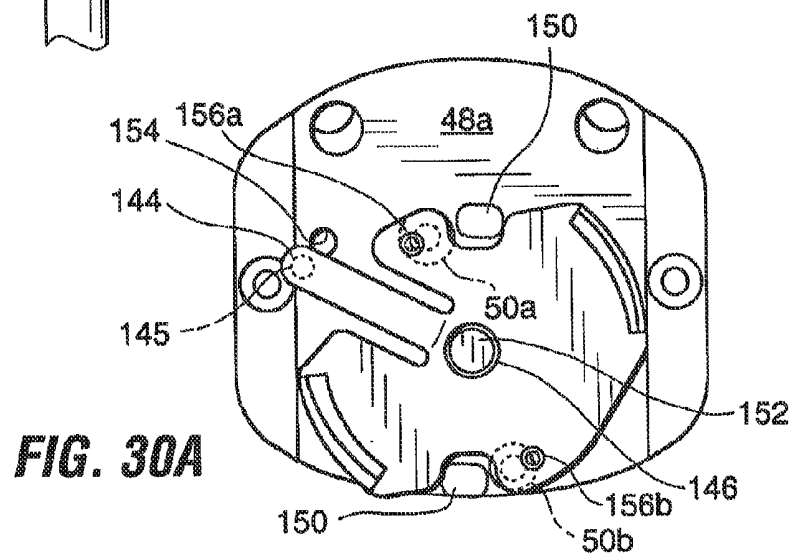
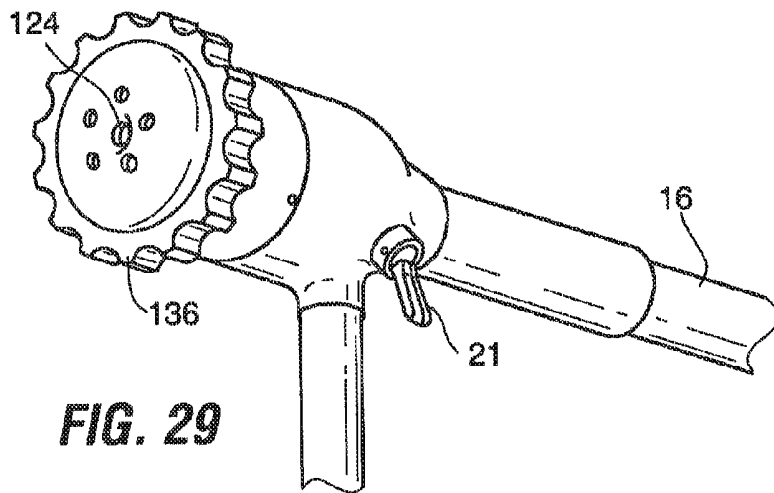


FIG. 30B

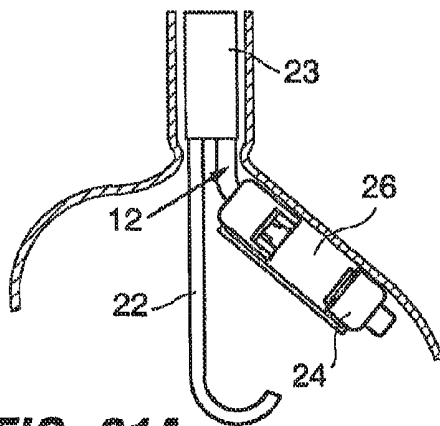


FIG. 31A

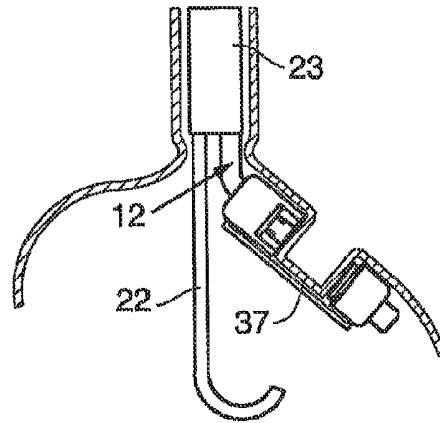


FIG. 31B

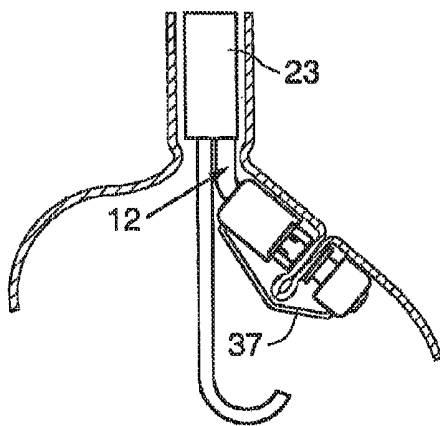


FIG. 31C

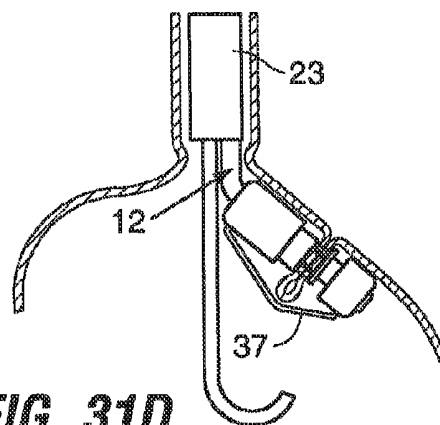


FIG. 31D

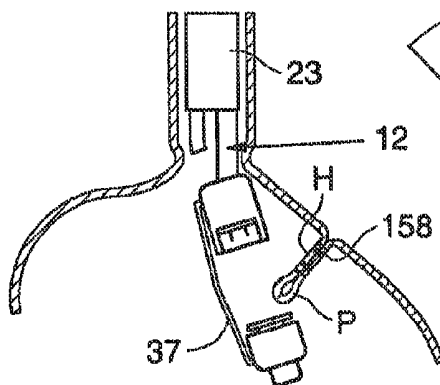


FIG. 31E

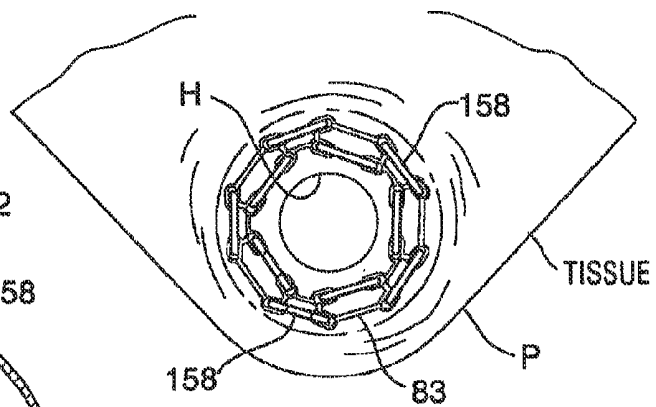
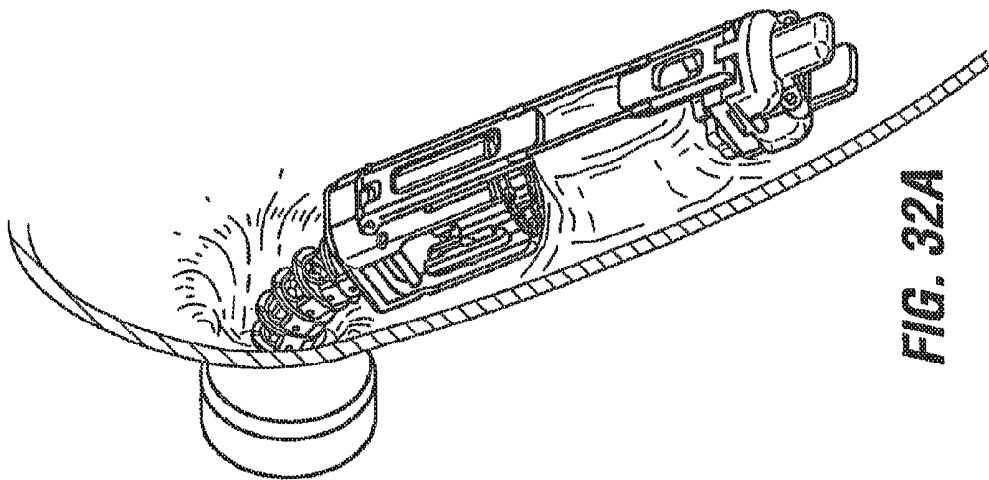
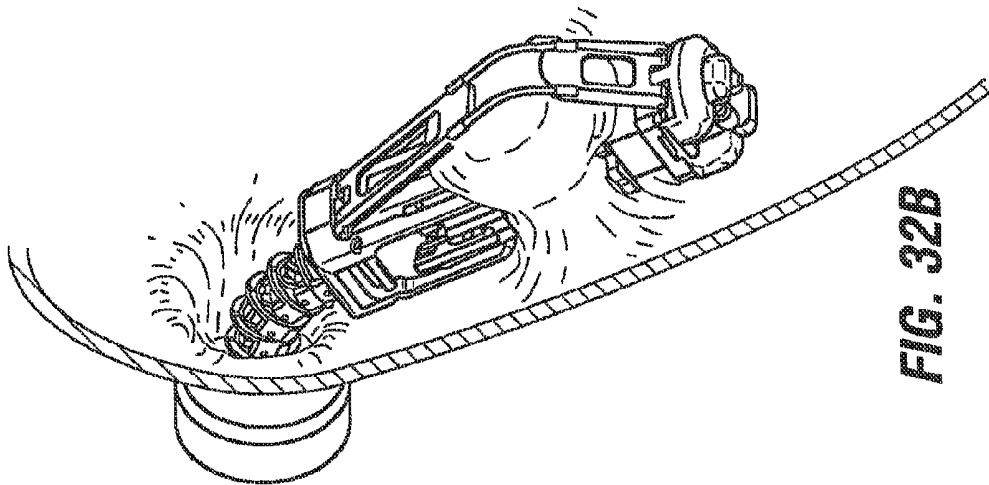
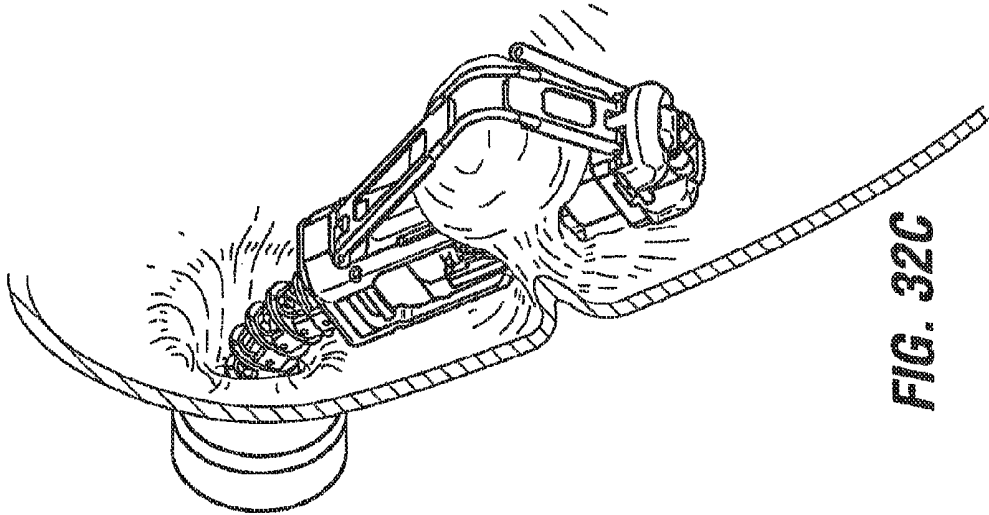
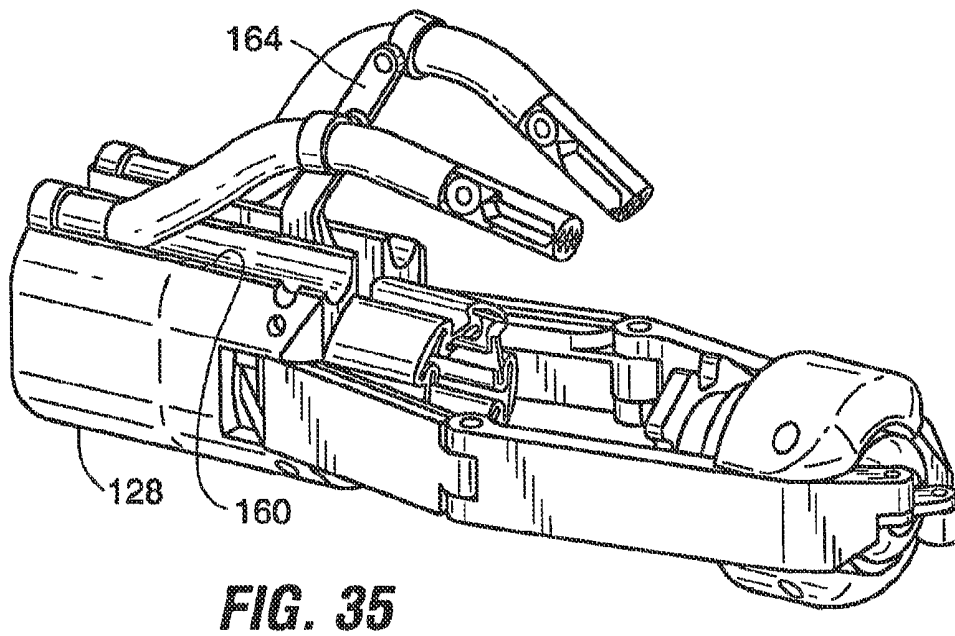
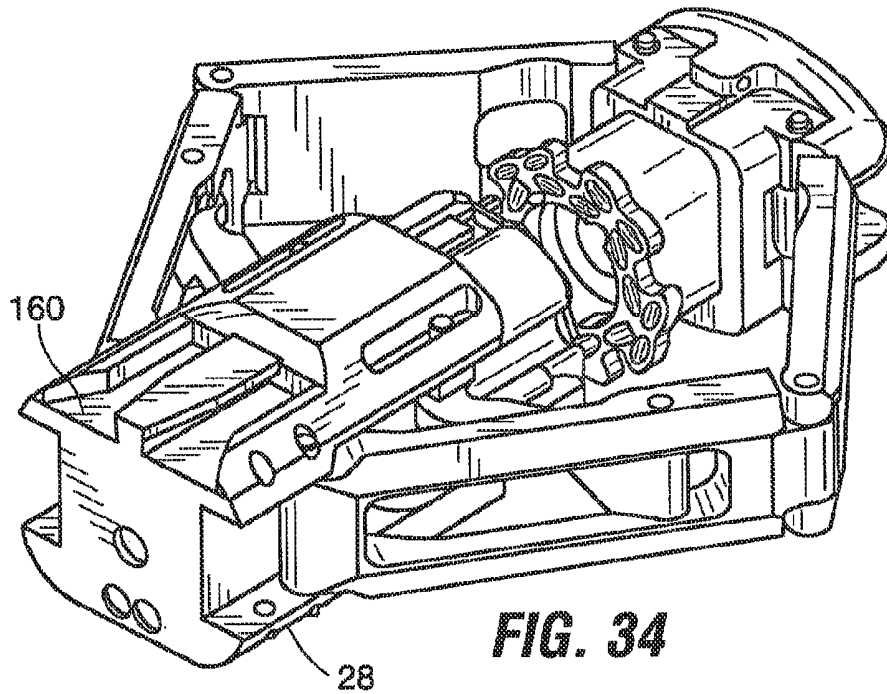


FIG. 33





ENDOSCOPIC STAPLING DEVICES

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation of U.S. patent application Ser. No. 13/053,133, filed Mar. 21, 2011, now which is a continuation of U.S. patent application Ser. No. 12/050,169, filed Mar. 18, 2008, entitled ENDOSCOPIC STAPLING DEVICES AND METHODS, now U.S. Pat. No. 8,020,741; U.S. patent application Ser. No. 13/053,133 is also a divisional of U.S. patent application Ser. No. 12/052,997, filed Mar. 21, 2008, now U.S. Pat. No. 7,909,222, which is a divisional of U.S. patent application Ser. No. 12/050,169, filed Mar. 18, 2008, now U.S. Pat. No. 8,020,741; U.S. patent application Ser. No. 13/053,133 is also a divisional of U.S. patent application Ser. No. 12/053,010, filed Mar. 21, 2008, now U.S. Pat. No. 7,922,062, which is a divisional of U.S. patent application Ser. No. 12/050,169, filed Mar. 18, 2008, now U.S. Pat. No. 8,020,741; U.S. patent application Ser. No. 13/053,133 is also a divisional of U.S. patent application Ser. No. 12/053,027, filed Mar. 21, 2008, now U.S. Pat. No. 7,909,219, which is a divisional of U.S. patent application Ser. No. 12/050,169, filed Mar. 18, 2008, now U.S. Pat. No. 8,020,741; U.S. patent application Ser. No. 13/053,133 is also a divisional of U.S. patent application Ser. No. 12/053,066, filed Mar. 21, 2008, now U.S. Pat. No. 7,909,223, which is a divisional of U.S. patent application Ser. No. 12/050,169, filed Mar. 18, 2008, now U.S. Pat. No. 8,020,741; U.S. patent application Ser. No. 13/053,133 is also a divisional of U.S. patent application Ser. No. 12/053,182, filed Mar. 21, 2008, now U.S. Pat. No. 7,913,892, which is a divisional of U.S. patent application Ser. No. 12/050,169, filed Mar. 18, 2008, now U.S. Pat. No. 8,020,741; all of which are incorporated herein by reference in their entirety.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

None

TECHNICAL FIELD

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

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.

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.

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 submucosa SM and the multi-layer muscularis M lie between the mucosa and the serosa.

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.

Several prior applications, including International Application No. WO 2005/037152 having an international filing date of Oct. 8, 2004 and U.S. application Ser. No. 11/439,461, filed May 23, 2006 (both incorporated herein by reference) 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.

For example, U.S. application Ser. No. 11/439,461 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.

In other instances, tissue plications may themselves 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 Applicants' co-pending application Ser. No. 11/542,457, filed Oct. 3, 2006, U.S. Publication No. 2007-0219571, which is incorporated herein by reference.

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.

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.

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.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1A is a schematic illustration of a human stomach and a portion of the small intestine.

FIG. 1B is a cross-sectional perspective view of a portion of a stomach wall, illustrating the layers of tissue forming the wall.

FIG. 2 illustrates an endoscopic stapling system.

FIGS. 3A-3C are perspective views showing the stapler head of the stapling system of FIG. 2 in three different positions.

FIG. 4 is a perspective view of the stapler head, with the membrane removed.

FIG. 5 is a perspective view of the proximal end of the staple housing of the stapler head of FIG. 4.

FIG. 6 is a perspective view of the distal end of the staple housing of the stapler head of FIG. 4.

FIG. 7 is an exploded perspective view showing elements advanceable within the staple housing during compression and stapling operations.

FIG. 8 is a plan view of a staple reinforcement device.

FIG. 9 is a side elevation view of a staple cartridge.

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.

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.

FIG. 11E is similar to FIG. 11D and shows an alternative piston configuration.

FIG. 12 is a perspective view of the anvil housing of the stapler head of FIG. 4.

FIG. 13 is a perspective view of the anvil support.

FIG. 14 is a plan view of the anvil.

FIG. 15A is a cross-sectional side view of the cutting device and a first embodiment of a cutting board.

FIG. 15B is a cross-sectional side view of the cutting device and a second embodiment of a cutting board.

FIG. 16 is a perspective view of the hinged arm assemblies of the stapler head of FIG. 4.

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.

FIG. 18 is similar to FIG. 17 and illustrates hidden features of FIG. 17.

FIG. 19 is a perspective view of the stapler head in an intermediate, partially expanded, position.

FIG. 20 is a plan view similar to FIG. 17 but showing the stapler head in the intermediate position.

FIG. 21 is similar to FIG. 20 and illustrates hidden features of FIG. 20.

FIG. 22 is a perspective view of the stapler head in a fully expanded, full compression position.

FIG. 23 is a plan view similar to FIG. 20 but showing the stapler head in the full compression position.

FIG. 24 is similar to FIG. 23 and illustrates hidden features of FIG. 24.

FIGS. 25A-25C 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.

FIG. 26 is a perspective view of the stapler of FIG. 2, with the staple head removed.

FIG. 27A is a plan view of the articulating section of the stapler of FIG. 2, showing the drive fluid lines.

FIG. 27B shows a drive fluid line having an alternate longitudinally expandable shape.

FIG. 28 is a cross-sectional side view of the handle of the stapler of FIG. 2.

FIG. 29 is a perspective view of the handle of the stapler of FIG. 2.

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.

FIGS. 31A-31E are a series of drawings schematically illustrating use of the system of FIG. 2 to form a plication in a stomach.

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.

FIG. 33 is a top plan view of a plication formed in body tissue.

FIGS. 34 and 35 are perspective views of an alternative stapler head equipped to carry additional tools.

DETAILED DESCRIPTION

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.

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 of 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.

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.

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. 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.

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

Generally speaking, system 10 includes a stapler 12 having a stapler head 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, 25 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.

The fluid source 25 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 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.

Referring to FIG. 3A, a covering or membrane 24 encloses the stapler head 14 to form a vacuum chamber 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 capable of forming a vacuum chamber that will expand in volume to accommodate tissue drawn into the chamber.

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 elements may be formed into or onto the membrane material, or embedded in the membrane material.

Stapler Head

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 a 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.

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 member comprising a proximal staple housing 28, a second member comprising a distal anvil housing 30, and at least one elongate member but preferably a pair of hinged arm assemblies 32.

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 and an anvil on the anvil housing.

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

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.

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.

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.

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.

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. 1A-11D.

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 (not shown). The drive member is coupled to the arm assemblies 32, anvil housing, and staple housing so that advancing the drive member distally effects tissue compression by bringing the contact surfaces of the staple housing and anvil housing relatively towards one another.

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 using pins 84 that extend through the housing's guide slots 64 and through mounting bores 70 in the disk 68.

A portion of the staple housing 28 contains staples to be fired into the tissue. The staples are contained within a staple holder 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.

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.

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 distally 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.

Again referring to FIG. 7, cartridge 78 includes a number of staple locations 80, each housing a staple. 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 Ser. No. 11/542,457, entitled ENDOSCOPIC PLICATION DEVICES AND METHODS, filed Oct. 3, 2006, and published Sep. 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.

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.

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 (not shown in FIG. 7) in the hydraulic chamber 66 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

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. 1A and 11B schematically show the fluid flow in the hydraulic chamber 66 of the staple 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 114 are formed through the side wall 110, between the o-ring seals 112.

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. 1A-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 114 in the compression piston.

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. 1A. Fluid pressure within the hydraulic chamber 66 advances the compression piston 106, with the stapler piston 116 within it, in a distal direction. FIG. 11B shows the compression piston 106 approaching the end of its travel. Once the compression piston reaches the end of its travel as shown in FIG. 11C, channel 114 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. 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. 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.

The anvil housing (identified by numeral 30 in FIG. 4) 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.

A central bore 90 extends longitudinally through the anvil housing 30. An anvil support 92 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.

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. A central opening 97 extends through the anvil 96 and is contiguous with a lumen in the anvil support 92.

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.

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.

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.

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.

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 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

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 pivotally coupled to the staple housing and a second arm section 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.

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 a spreader arm 110 pivotally mounted within the cutout 108. The distal end of each spreader arm 110 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 spreader arms 110 to pivot relative to the pins 84 and to thus drive the arm assemblies 32 out-

wardly. Additional specifics concerning movement of the arm assemblies 32 is set forth in the section entitled. Stapler Head. Operation.

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 spreader arms 110 drove 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.

Stapler Head Operation

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.

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.

Both the disk 68 and the arm spreaders 110 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 110 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.

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.

The relative motion of the two housings 28, 30 toward each other also drives upward links 38, 40 and their inter-connecting 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.

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 5.5 mm length) or 0.109 inches for 6.5 mm leg length staples. Application of additional pressure into the hydraulic circuit will not compress the tissue any further.

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.

At the fully compressed position, the arm spreaders 110 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.

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.

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 110 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.

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.

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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.

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 Shaft and Handle

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.

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.

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 articulating section 128.

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 1000 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).

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

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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.

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.

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 distalmost link 132 of the stapler housing 28.

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.

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.

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.

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.

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.

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.

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.

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 **148** along its edges. Hydraulic feed holes **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**.

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 **50a, 50b** are repositioned as shown. Additionally, the rear surface **48a** has been modified to include a pair of catches in the form of undercut bosses **150**, plus an aligning pin **152**, and a hole **154**.

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

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.

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 advance further into the stomach.

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. **3A**. The opening **26** in the membrane **24** is positioned against the target tissue. The endoscope **22** is placed in a retroflexed position as shown.

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 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.

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.

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. 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, allowing 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.

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.

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.

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

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.

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. Moreover, features of the disclosed embodiments may be combined with one another and with other features in varying ways to produce additional embodiments. Thus, the embodiments described herein should be treated as representative examples of systems useful for forming endoscopic tissue plications, and should not be used to limit the scope of the claimed invention.

Any and all patents, patent applications and printed publications referred to above, including those relied upon for purposes of priority, are incorporated herein by reference.

We claim:

1. A stapler, comprising:

a staple housing including a staple cartridge;

an anvil housing including an anvil; and

a member configured to (a) move the staple cartridge relative to the staple housing in the direction of the anvil, and (b) move the anvil in the direction of the staple cartridge.

2. The stapler of claim 1, wherein the staple cartridge includes a plurality of staples positioned therein.

3. The stapler of claim 1, wherein the member is configured to move the anvil relative to the anvil housing in the direction of the staple cartridge.

4. The stapler of claim 1, wherein the staple cartridge is spaced apart from the anvil along a longitudinal axis of the stapler, and wherein the stapler further includes one or more arm members coupling the staple housing and the anvil housing, and wherein each arm member of the one or more arm members includes a first arm coupled to the staple housing at a first pivot, and a second arm coupled to the anvil housing at a second pivot, and a hinge connecting the first arm and the second arm.

5. The stapler of claim 4, wherein the one or more arm members are coupled to the staple housing and the anvil housing such that (i) the first arm rotates about the first pivot away from the longitudinal axis, and (ii) the second arm

rotates about the second pivot away from the longitudinal axis, as the staple cartridge and the anvil move towards each other.

6. The stapler of claim 4, wherein the one or more arm members are coupled to the staple housing and the anvil housing such that the hinge moves away from the longitudinal axis as the staple cartridge and the anvil move towards each other.

7. The stapler of claim 4, further including a vacuum chamber coupled to and positioned between the staple cartridge and the anvil, wherein movement of the staple cartridge and the anvil towards each other is configured to expand the vacuum chamber in a direction transverse to the movement.

8. A stapler, comprising:

a staple cartridge;

an anvil spaced apart from the staple cartridge along a longitudinal axis, the anvil being coupled to an anvil housing;

an expandable vacuum chamber positioned between the staple cartridge and the anvil; and

a member configured to (a) move the staple cartridge towards the anvil, and (b) move the anvil relative to the anvil housing towards the staple cartridge, wherein movement of the staple cartridge and the anvil towards each other is configured to expand the vacuum chamber in a direction transverse to the longitudinal axis.

9. The stapler of claim 8, wherein the staple cartridge is coupled to a staple housing, and wherein the member is configured to move the staple cartridge relative to the staple housing towards the anvil.

10. The stapler of claim 8, wherein the staple cartridge includes a plurality of staples positioned therein.

11. The stapler of claim 8, wherein the stapler further includes one or more arm members coupling the staple cartridge and the anvil, and wherein each arm member of the one or more arm members includes a first arm coupled to the staple cartridge at a first pivot, and a second arm coupled to the anvil at a second pivot, and a hinge connecting the first arm and the second arm.

12. The stapler of claim 11, wherein the one or more arm members are coupled to the staple cartridge and the anvil such that (i) the first arm rotates about the first pivot away from the longitudinal axis, and (ii) the second arm rotates about the second pivot away from the longitudinal axis, as the staple cartridge and the anvil move towards each other.

13. The stapler of claim 11, wherein the one or more arm members are coupled to the staple cartridge and the anvil such that the hinge moves away from the longitudinal axis as the staple cartridge and the anvil move towards each other.

14. A stapler, comprising:

a staple cartridge coupled to a staple housing;

an anvil spaced apart from the staple cartridge along a longitudinal axis;

a member operatively connected to the staple cartridge and configured for movement from a retracted position to an extended position, wherein the member is configured to move the staple cartridge relative to the staple housing in the direction of the anvil; and

an arm assembly operatively coupled to the staple cartridge and the anvil, the arm assembly including a plurality of arm members coupled together by a hinge, such that movement of the member from the retracted position to the extended position pivots each arm member of the plurality of arm members outwardly to

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move the hinge away from the longitudinal axis and drive the staple cartridge and the anvil towards each other.

15. The stapler of claim 14, wherein the anvil is coupled to an anvil housing. 5

16. The stapler of claim 15, wherein the member is configured to move the anvil relative to the anvil housing towards the staple cartridge.

17. The stapler of claim 14, further including a vacuum chamber coupled to and positioned between the staple cartridge and the anvil, wherein movement of the staple cartridge and the anvil towards each other is configured to expand the vacuum chamber in a direction transverse to the longitudinal axis. 10

18. The stapler of claim 14, wherein the staple cartridge includes a plurality of staples positioned therein. 15

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专利名称(译)	内窥镜缝合装置		
公开(公告)号	US9636114	公开(公告)日	2017-05-02
申请号	US14/487936	申请日	2014-09-16
[标]申请(专利权)人(译)	波士顿科学西美德公司		
申请(专利权)人(译)	BOSTON SCIENTIFIC SCIMED , INC.		
当前申请(专利权)人(译)	BOSTON SCIENTIFIC SCIMED , INC.		
[标]发明人	COLE DAVID SMITH ANDREW		
发明人	COLE, DAVID SMITH, ANDREW		
IPC分类号	A61B17/072 A61B17/115 A61B17/30 A61B17/00 A61B17/3205 A61F5/00 A61B17/064		
CPC分类号	A61B17/072 A61B17/115 A61F5/0083 A61B17/064 A61B17/32053 A61B2017/00539 A61B2017/00818 A61B2017/07271 A61B2017/306 A61F5/0013		
其他公开文献	US20150001274A1		
外部链接	Espacenet USPTO		

摘要(译)

本文描述了内窥镜缝合器和用于将一个或多个紧固件施加到身体组织的方法。在一个实施例中，紧固件施加装置（优选地是订书机）经口部通过进入胃以通过从胃内部接合组织并将其向内拉入而使胃组织复杂化。在所公开的实施例中，组织被吸入组织室，使得浆膜组织的部分彼此面对地定位。所公开的订书机允许相对的组织部分移动到与另一个组织接触，并且优选地递送钉以维持组织切片之间的接触，至少直到形成浆膜结合。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。

