



(86) Date de dépôt PCT/PCT Filing Date: 2007/05/03

(87) Date publication PCT/PCT Publication Date: 2007/11/15

(45) Date de délivrance/Issue Date: 2015/11/17

(85) Entrée phase nationale/National Entry: 2008/10/23

(86) N° demande PCT/PCT Application No.: US 2007/068149

(87) N° publication PCT/PCT Publication No.: 2007/131112

(30) Priorité/Priority: 2006/05/03 (US60/797,346)

(51) Cl.Int./Int.Cl. *A61B 17/22* (2006.01),
A61B 1/00 (2006.01), *A61B 17/94* (2006.01),
A61B 17/128 (2006.01), *A61B 18/12* (2006.01)

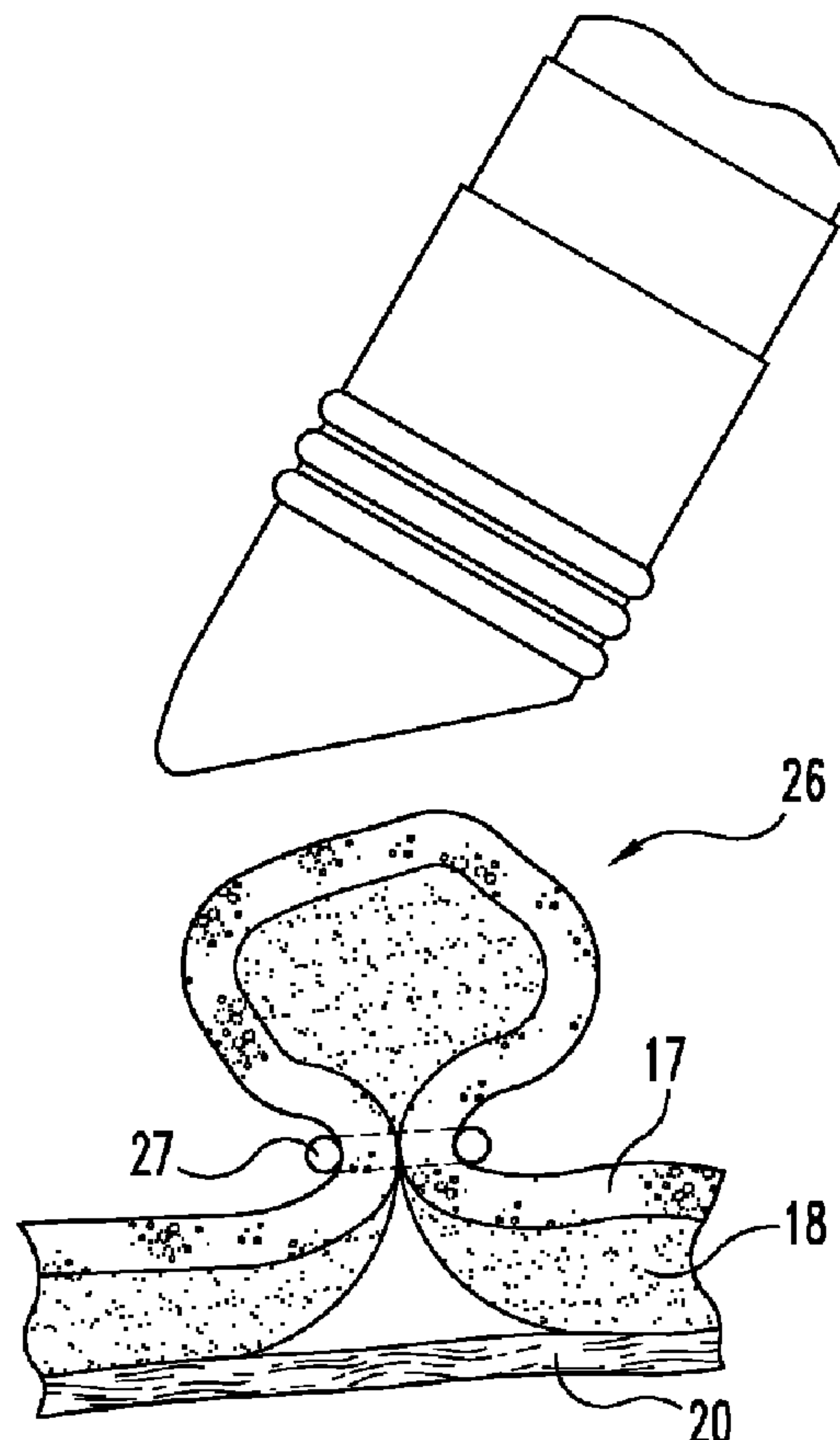
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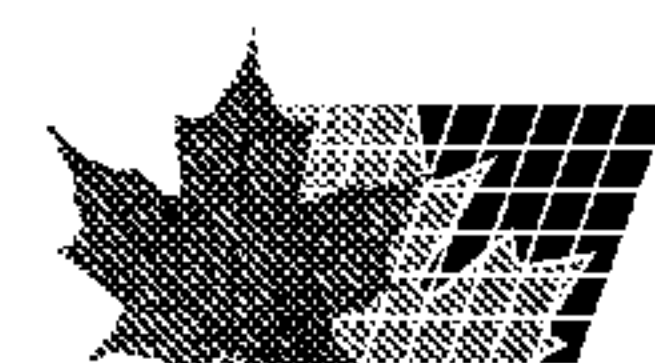
(54) Titre : PROCÉDES ET APPAREILS DESTINÉS A REFORMER L'ŒSOPHAGE ET D'AUTRES LUMIÈRES
CORPORELLES

(54) Title: METHODS AND APPARATUSES FOR RESHAPING THE ESOPHAGUS AND OTHER BODY LUMENS



(57) Abrégé/Abstract:

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(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
15 November 2007 (15.11.2007)

PCT

(10) International Publication Number
WO 2007/131112 A3(51) International Patent Classification:
A61B 17/10 (2006.01)(21) International Application Number:
PCT/US2007/068149

(22) International Filing Date: 3 May 2007 (03.05.2007)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/797,346 3 May 2006 (03.05.2006) US(71) Applicant (for all designated States except US): **INDIANA UNIVERSITY RESEARCH & TECHNOLOGY CORPORATION** [US/US]; 351 West 10th Street, Indianapolis, IN 46202 (US).

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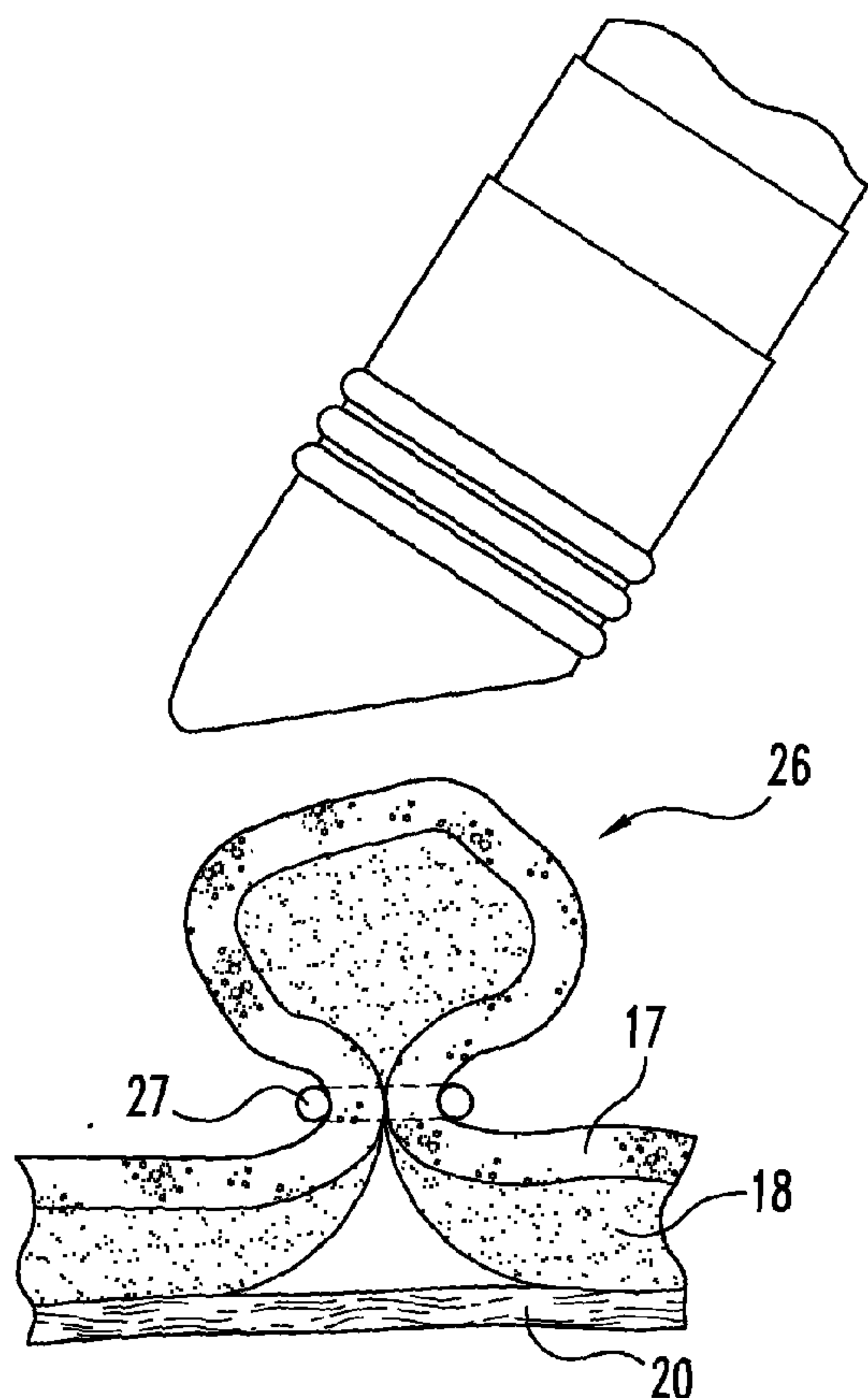
(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP,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

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

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**Fig. 3**

WO 2007/131112 A3

WO 2007/131112 A3



Published:

- *with international search report*
- *before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments*

(88) Date of publication of the international search report:
4 December 2008

53672-1

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**METHODS AND APPARATUSES FOR RESHAPING THE ESOPHAGUS AND OTHER
BODY LUMENS**

BACKGROUND

The present invention relates generally to the field
5 of medicine and in particular aspects to methods and
apparatuses for reshaping the esophagus or other body lumens.

Gastroesophageal reflux disease (GERD) has been
described as an improper functioning of the lower or inferior
esophageal sphincter (LES) in which stomach contents are able
10 to leak back, or reflux, abnormally into the esophagus. When
refluxed stomach acid contacts the lining of the esophagus, it
causes a burning sensation in the chest or throat called
heartburn. Reports show that as many as 10 percent of
Americans have daily episodes of heartburn and that up to 44
15 percent have symptoms at least once a month. While occasional
heartburn is common and does not necessarily mean that the
subject has GERD, chronic heartburn may signify GERD and can
lead to more serious health problems.

Esophagitis, a complication of GERD, can become a
20 relapsing chronic condition. Other complications of

GERD include strictures, ulcerations and Barrett's esophagus (progressive replacement of distal eroded squamous mucosa with metaplastic intestinal epithelium). Treatment of GERD may include medication
5 and/or surgery.

Morbid obesity (clinically severe obesity) has been described as a condition in which a person's weight creates major primary or threatened secondary
10 effects on the patient's health and well being. Unfortunately, morbid obesity often results in early death. A 12-year follow-up of 336,442 men and 419,060 women found that the mortality rate increased approximately two-fold for men having a 50% above-
15 average weight, while the mortality rate increased 5-fold for diabetics and 4-fold for those with digestive tract disease in this same group. In a similar group of women, the overall mortality rate also increased two-fold, while the mortality rate increased 8-fold for
20 diabetics and 3-fold for those with digestive tract disease.

Associations between the average weight of population groups and the prevalence of non-insulin-
25 dependent diabetes have also been repeatedly observed. Reports show that the risk of developing diabetes increases about two-fold in the mildly obese, five-fold in the moderately obese and ten-fold in severely obese persons. It is also known that the duration of obesity
30 is an important determinant of the risk for developing diabetes. Further, studies show increases in cancer mortality rates in severely obese females, e.g.,

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endometrium (5.4 times), gall bladder (3.6 times), uterine cervix (2.4 times), ovary (1.6 times) and breast (1.5 times). As well, studies show increases in cancer mortality rates in severely obese males, e.g., colorectal (1.7 times) and prostate
5 (1.3 times).

There remains a need for improved and/or alternative methods and apparatuses that are useful for treating GERD, obesity and other medical conditions. Embodiments of the present invention may be addressed to those needs.

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SUMMARY

There is provided a medical device for use to reshape a body passage of a patient, the medical device comprising: an access device for accessing a luminal surface of a body passage of a patient, the access device including a barrel having a proximal end and a beveled distal end, the beveled distal end including a first side that is generally opposite a second side, the first side extending distally of the second side in a generally longitudinal direction along the barrel; one or more bands received upon the barrel for banding tissue on the luminal surface; a single pull line operably associated with the one or more bands, the single pull line operable to deploy the one or more bands from the barrel, the single pull line extending through the barrel and exiting the barrel at or substantially at the first side of the barrel as the one or more bands are deployed; and a tissue resection device for resecting tissue from the luminal surface; wherein the medical device is for removing tissue from one or more areas of the luminal surface for the purpose of resulting in a healing response for reshaping the body passage.

There is also provided a medical device, comprising: an endoscopic ligator barrel having a proximal end and a beveled distal end, the beveled distal end including a first side that is generally opposite a second side, the first side extending distally of the second side in a generally longitudinal direction along the ligator barrel; one or more ligation bands received upon the ligation barrel; a single pull line operably associated with the one or more ligation bands, the single pull line operable to deploy the one or more

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ligation bands from the ligation barrel, the single pull line extending through the ligator barrel and exiting the barrel at or substantially at the first side of the barrel as the one or more ligation bands are deployed.

- 5 Another aspect provides a medical device, comprising:
an endoscopic ligator barrel having a proximal end and a
beveled distal end, the beveled distal end including a first
side that is generally opposite a second side, the first side
extending distally of the second side in a generally
10 longitudinal direction along the ligator barrel; one or more

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ligation bands received upon the ligation barrel, the one or more ligation bands being deployable from the beveled distal end; and a single pull line operably associated with the one or more ligation bands, the single pull line operable to deploy
5 the one or more ligation bands from the ligation barrel, the single pull line extending through a guide member coupled to the ligator barrel, the guide member configured to prevent circumferential migration of the single pull line around the barrel as the ligation bands are deployed.

10 In one aspect of the invention, methods for treating a patient are provided that include reshaping a body lumen or passageway of the patient by a procedure including selective tissue removal which, upon healing, results in the reshaping. In particular embodiments, the invention provides methods that
15 comprise an endoluminal mucosaplasty (ELM) procedure in which tissue from a wall defining the body passageway is removed to invoke a healing process that reshapes the passageway, and systems designed therefor. The resected tissue can be non-diseased tissue. In certain inventive procedures, these methods
20 may be used to modify an esophagus to a new condition that provides increased resistance to reflux of gastric contents, for instance in the treatment of a patient having gastrointestinal reflux disease (GERD). In other inventive procedures, these methods can be used to modify the stomach in
25 a bariatric surgery to contribute to a change in eating habits of the patient, e.g. in the treatment of obesity. In certain forms, the inventive methods lead to a narrowing of a body passageway. Such methods may include the steps of (i) identifying a body passageway within a patient presenting a

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medical condition in which a reduction in the luminal diameter of the body passageway will reduce flow therethrough in at least one direction such that the patient will experience at least a partial relief of the medical condition; (ii)

- 5 determining a degree of reduction in the diameter of the passageway expected to provide effective treatment for the medical condition in the patient; (iii) removing tissue from an inner portion of a wall defining the body passageway sufficient to elicit a healing response. The ensuing healing of the
- 10 tissue can cause contraction of tissues within the body passageway such that the luminal diameter thereof is reduced. In many typical treatment regimens, the patient will then be assessed post-operatively to determine if the body passageway reduction is effective for at least partial relief of the
- 15 medical condition.

In one embodiment, the present invention provides a method for treating a patient comprising resecting tissue from a luminal surface of a body passage of the patient for the purpose of inducing a healing response that will result in a

20 reshaping of the body passage.

In another embodiment, the invention provides a system for reshaping a body lumen. The system comprises an endoscope configured to generate an image of the body lumen and at least one reference of known dimension located in the body

25 lumen, and software for calculating a dimension of the body lumen based on a comparison of the reference and the body lumen in the image. The system further includes an endoluminal surgical device for removing tissue from the body lumen.

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In another embodiment, the invention provides a medical device comprising an endoscopic ligator barrel, one or more ligation bands received upon the ligation barrel, and a single pull line operably associated with

said one or more ligation bands, said single pull line operable to deploy said one or more ligation bands from said ligation barrel. The device can also include an endoscope upon which the barrel is received.

5

In another embodiment, the invention provides a medical device comprising an endoscopic ligator barrel having a proximal end and a beveled distal end, and one or more ligation bands received upon the ligation barrel, said one or more ligation bands being depoloyable from said beveled distal end. The device can further include an endoscope upon which the barrel is received.

15 In another embodiment, the present invention provides a method for narrowing a body passageway, comprising (a) identifying a body passageway within a patient presenting with a known medical condition in which the effective reduction in the nominal luminal diameter of said body passageway will lessen flow therethrough in at least one direction such that said patient will experience at least partial relief of said medical condition; (b) determining a degree of reduction in the diameter of said passageway expected to provide effective treatment for said medical condition in said patient; (c) using a medical device to cause injury to a portion of the luminal surface of said body passage sufficient to elicit a healing response that causes contraction of the tissues within said body passageway such that the nominal luminal diameter thereof is reduced without causing injury to the muscular layer of said body passageway; and (d)

assessing said patient post-operatively to determine if said body passageway reduction is effective for at least partial relief of said medical condition.

5 In another embodiment, the present invention provides a method for treating a patient that comprises reshaping the gastroesophageal junction of the patient with an endoscopic procedure that includes removing non-diseased tissue, wherein said reshaping is
10 effective to reduce gastroesophageal reflux in the patient.

 In another embodiment, the invention provides a method for treating a patient comprising reshaping the
15 stomach of the patient with an endoscopic procedure that includes removing non-diseased tissue, wherein said reshaping creates a narrowing in the stomach to modify satiety sensation of the patient.

20 In other embodiments, the invention provides method for treating a patient comprising operating through a channel of an endoscope to remove patient tissue (i) for the purpose of modifying the esophagus of the patient to reduce reflux of gastric contents
25 into the esophagus; or (ii) for the purpose of modifying the cardia of the patient to treat obesity.

 In another embodiment, the present invention provides a method for treating a patient comprising
30 responding to an identified need in the patient for treatment to reduce gastroesophageal reflux or obesity by conducting an endoscopic procedure that includes

removing tissue of the patient's esophagus and/or cardia.

In another embodiment, the invention provides a
5 method for treating a patient comprising (a) diagnosing
a patient as having GERD; and (b) based on said
diagnosing, conducting an endoscopic procedure that
includes removing patient tissue for the purpose of
modifying the shape of the gastroesophageal junction
10 of the patient.

In another embodiment, the present invention
provides a method for treating a patient comprising (a)
diagnosing a patient as needing treatment for obesity;
15 and (b) based on said diagnosing, conducting an
endoscopic procedure that includes removing patient
tissue for the purpose of modifying the shape of the
cardia of the patient.

20 In another embodiment, the invention provides a
method for narrowing a body passageway comprising the
steps of (a) endoscopically visualizing a portion of
said body passageway selected for modification to
provide a reduced luminal diameter thereof;
25 (b) determining the degree of reduction required to
provide the reduced luminal diameter; (c) identifying a
first treatment site located on a luminal surface of
the body passageway; (d) capturing a first portion of
tissue at the first treatment site; (e) placing a
30 tissue-securing element around the captured first
portion of tissue; (f) excising the first captured
portion of tissue; (g) identifying a second treatment

site located circumferentially adjacent to said first site; (h) capturing a second portion of tissue at the second treatment site; (i) placing a tissue-securing element around the captured second portion of tissue;
5 and (j) excising the second captured portion of tissue.

In another embodiment, the invention provides a method for narrowing a body passageway that is at least partially dysfunctional with respect to restriction of
10 materials flowing therethrough, comprising the steps of (a) selecting a first zone of treatment extending along a length of said body passageway, said first zone of treatment extending circumferentially around the inner surface of said body passageway; (b) establishing a
15 nominal diameter of the body passageway within the zone of treatment; (c) calculating an amount of resected tissue effective to reduce the body passageway to a desired second nominal diameter smaller than said first nominal diameter such that said dysfunctional condition
20 is at least partially corrected; (d) visualizing the zone of treatment using an imaging device; (e) isolating a first portion of the mucosa and submucosa from muscle layer within said first zone of treatment; (f) resecting the first portion of the mucosa and
25 submucosa above said muscle layer using a medical device; (g) moving circumferentially of said body passageway adjacent the first resected portion of tissue to a second portion of tissue spaced apart from said first portion; and (h) resecting said second
30 portion of tissue leaving an untreated area of tissue between said first portion and said second portion of resected tissue.

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In another embodiment, the invention provides a method for narrowing a body passageway comprising excising luminal tissue of the body passageway to create a resected area in a configuration that promotes healing of the luminal tissue
5 such that there is an overall contraction of the luminal tissue during the healing process whereby the nominal luminal diameter of the body passageway is reduced.

In another embodiment, the present invention provides a method for treating a patient to modify a body passage of the
10 patient, comprising removing tissue from a luminal surface of the body passage without substantial damage to underlying muscle tissue of the passage for the purpose of inducing a healing response that will result in a reshaping of the body passage.

15 In further embodiments, the present invention provides kits for reshaping a body passage such as an esophagous or stomach, as described herein, that include means or devices as described herein for resecting or otherwise removing tissue from the luman of the body passage, and written
20 materials regarding use of the means or devices to reshape a body passage, e.g. in the treatment of GERD or obesity as described herein.

Additional embodiments as well as features and advantages will be apparent from the further descriptions
25 herein.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1A is a schematic drawing of the stomach in cross-section.

5 FIG. 1B is a cross-sectional view of the wall of the stomach and esophagus.

FIG. 2 is a view showing a step in an inventive process.

10 FIGS. 3-5 are views showing further steps in an inventive process.

FIGS. 6-7 are sectional views of a body lumen wall showing healing response.

FIG. 8 is a sectional view of the junction of the stomach and the esophagus.

15 FIG. 9 is a cross-section of the gastroesophageal junction taken along the line 9-9 of FIG. 8 in the direction of the arrows showing a step in the process.

FIGS. 10-11 are cross-sectional views showing steps in an obesity treatment method.

20 FIG. 12 provides a partial cross-sectional view of an illustrative beveled ligating barrel disposed upon the end of an endoscope.

FIGS. 13-15 are graphs depicting results from an experimental study described in Example 1.

DETAILED DESCRIPTION

For the purposes of promoting an understanding of the principles of the disclosure, reference will now be made to the embodiment illustrated in the drawings and specific language will be used to describe the same. It will nevertheless be understood that no limitation of the scope of the claims is thereby intended, such alterations and further modifications in the illustrated device, and such further applications of the principles of the disclosure as illustrated therein, being contemplated as would normally occur to one skilled in the art to which the disclosure relates.

As disclosed above, aspects of the present invention relate to methods and systems for reshaping body lumens such as those of the esophagus or stomach. This reshaping may, for example, be effective to modify the flow of materials within the body lumen. Such reshaping can involve decreasing the resting diameter of the body lumen, creating folds or irregular surfaces within a body lumen wall, and/or altering the contour of the inner periphery of a body lumen wall. These and other modifications that can affect the flow of materials within the lumen are contemplated as being a part of the present invention.

As noted above, in particular inventive aspects, methods and systems are provided for reshaping a region of an esophagus of a patient, especially a region at or near the gastroesophageal junction. Such procedures can be undertaken, for example, to result in a

reduction of gastroesophageal reflux potentially in the treatment of a patient diagnosed with GERD. In addition or alternatively, procedures to reduce reflux can involve the treatment and reshaping of non-
5 esophageal tissues as well, for example cardia tissues occurring just below the termination of the esophagus.

GERD is typically diagnosed or at least suspected when a person complains of heartburn. Heartburn is
10 frequently described as a sub-sternal (under the middle of the chest) burning that occurs after meals and often worsens when lying down. When suspected, the treating physician will often conduct a therapeutic trial (at least initially) in which the patient is treated with
15 medications to suppress the production of acid by the stomach. The diagnosis is then confirmed on the basis of a response of the symptoms to the treatment. For example, if the heartburn is diminished to a large extent by the medication, the diagnosis of GERD is
20 generally considered confirmed. GERD can be diagnosed in a number of other suitable manners as well, for example, using diagnostic and other tests including but not limited to endoscopic procedures and esophageal acid testing. Oftentimes, the abnormal reflux
25 associated with GERD is due to the presence of a patient gastroesophageal junction that is distended or over-sized (e.g., has a diameter that is considered too large). In inventive variants, such an over-sized gastroesophageal junction is subjected to a reshaping
30 treatment in order to reduce its diameter and thereby reduce the amount of gastric reflux into the esophagus.

In additional embodiments of the invention, methods and systems are provided for modifying the lumen of the stomach of a patient, for example, in a bariatric surgery potentially in the treatment for obesity. Such procedures can, for example, be undertaken to treat patients that have been diagnosed as being morbidly obese. Typically, morbid obesity is diagnosed when a person's weight creates major primary or threatened secondary effects on the patient's health and well being. The usual diagnostic threshold for this condition is 100 pounds over what is considered ideal body weight, e.g., as determined by standard height versus weight charts. Examples of obesity-aggravated secondary health problems are arthritic symptoms on weight-bearing joints, adult-onset diabetes, sleep apnea/severe snoring, hygiene problems related to skin folds, depression and low self-esteem. In accordance with one aspect of the invention, a stricture or other narrowing or reshaping can be created in the stomach lumen of a patient who is morbidly obese or another patient for which modification of eating habits is considered to be beneficial. The stricture or other reshaping modifies the passage of food materials through the stomach lumen so as to facilitate a modification of eating habits in the patient. In addition, as all or a portion of the procedure, tissue from the fundus of the stomach can be removed, so as to reshape (e.g. reduce the size of) the stomach and potentially result in the removal of cells, such as ghrelin-secreting epithelial cells, that secrete hormonal or other signals related to hunger. Ghrelin is understood to be a hormonal signal for

hunger, and the reduction or removal of cells that secrete ghrelin or other hunger signals may be used in inventive embodiments disclosed herein, e.g. for the treatment of obesity.

5

Endoluminal mucosaplasty procedures can be used in the inventive reshaping methods disclosed herein. In this regard, "endoluminal mucosaplasty" as used herein refers to an endoluminally conducted procedure which results in a change in the mucosa tissue that lines the body lumen under treatment. This change may, for example, be a change in the diameter of the presented mucosal surface of the passageway, in the presence or absence of folds or ridges, and the like.

15

Some inventive reshaping methods described herein comprise resecting an amount of tissue from a body lumen wall and/or subjecting tissue at a treatment site to an injurious stimulus which results in cellular death within the tissue and an ensuing tissue healing response. In some forms of the invention, the tissue that is removed will include at least some non-diseased tissue for which there is no alternate indication (other than the desired reshaping) requiring its removal, and in certain instances all tissue that is removed will be normal non-diseased tissue for which there is no other indication for removal.

In advantageous inventive embodiments, the tissue that is removed will be essentially confined to tissue that occurs to the luminal side of a discernable muscular layer of the passageway wall. Lumen walls of

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body passageways such as the esophagus and stomach generally include a mucosal layer and an underlying submucosal connective tissue layer, commonly followed by a muscular layer. In aspects of the present
5 invention, amounts of mucosa and/or submucosa are removed at a location without causing any substantial damage to the underlying muscle layer. In this regard, it is known that the connective tissue of submucosal layers often interleaves with amounts of muscle tissue,
10 especially but not exclusively in areas transitioning from histologically-identifiable submucosa layers to underlying muscle layers. It will thus be understood that some muscular tissue can be present in the removed tissue without having caused any substantial damage to
15 the underlying muscle layer.

In some forms of the invention, the removal of both mucosal and submucosal tissues is sufficient to expose the underlying muscle layer. To protect against
20 significant damage to the muscle layer, amounts of mucosal and submucosal tissue can be drawn away from the underlying muscle layer, for example, using preferential suction into a chamber or using physical grasping devices such as forceps, after which the drawn
25 tissue is resected or otherwise removed. Illustratively, as discussed elsewhere herein, endoscopic equipment and techniques, such as but not limited to those that have traditionally been used in endoscopic mucosal resection (EMR) procedures which
30 remove unhealthy tissue, can be used to accomplish this type of tissue removal. Other endoscopic and non-endoscopic devices and techniques may be used as well.

In both endoscopic and non-endoscopic procedures, fluid (e.g., saline) injections may optionally be used to separate mucosal and submucosal tissue from an underlying muscle layer prior to tissue removal. These
5 and/or other devices, techniques or materials can be used in certain procedures of the invention to separate or isolate amounts of the tissue targeted for removal from tissue to remain, e.g. in order to protect or insulate the remaining tissue during removal of the
10 targeted tissue. Such selective tissue removal and protection can facilitate a more controlled pattern of injury to result in the desired shape modification while reducing the risk of undesired tissue damage and any potential complications associated therewith.

15

Medical procedures of the invention will involve the removal of amounts of tissue in at least one site of a body lumen wall and more typically at multiple sites contemplated as useful to achieve the desired
20 reshaping of the lumen. For instance, amounts of tissue can be removed from two to ten sites, and in certain treatments from two to eight sites. These sites may be discrete from one another, i.e., leaving amounts of undisturbed patient tissue in between the
25 sites, or they can overlap one another so as to create a larger contiguous region of removed tissue. Where multiple targeted removal sites are employed, at least some of them and often all of them will be positioned circumferentially relative to one another, although
30 sites that are longitudinally spaced from one another can also be used. The tissue removal sites will be selected so as to result in the desired reshaping of

the body lumen when a subsequent tissue healing process occurs. Such tissue healing will result in a contraction of the body lumen in the healed region, leading to the desired reshaping. In certain desired treatments, the tissue removal site or sites will result in a decrease in diameter of the lumen relative to the diameter prior to the treatment. In this regard, any appropriate decrease in diameter can be used in the treatment of the patient, including for example diameter decreases of at least about 10 percent, and oftentimes at least about 10 percent up to about 50 percent, or at least about 25 percent up to about 50 percent. In certain cases, it is expected that percent diameter decreases in the range of about 30 percent to about 35 percent will be achieved in the patient treatment.

Medical procedures of the invention are desirably performed in conjunction with a careful assessment of the characteristics of the body passageway to be treated, such as but not limited to its size, shape, lumen diameter, and/or tissue and cellular makeup, prior to removal of amounts of tissue. In this manner, the amount and pattern of tissue removal can be correlated to a desired and expected reduction in diameter and/or other reshaping characteristic. This assessment can be conducted in a variety of fashions, and may involve the use of any number of suitable devices and techniques that prove useful in providing a desired lumen reshaping. For example, some inventive apparatuses may include a device that can directly or indirectly measure one or more lumen wall dimensions.

In preferred aspects, systems especially adapted for performing inventive procedures as described herein include an overall apparatus having both an ability to measure the diameter of the body lumen and to undertake
5 tissue removal in the body lumen, so as to ultimately achieve the desired reduction in diameter and/or other reshaping. For these purposes, a video endoscope system can include an imaging means for creating a viewable image of the body lumen, at least one scaling
10 reference of known dimension, means for calculating the diameter of the body lumen based upon a reference to the scaling reference, and a device for removing amounts of tissue from a body lumen wall. In this manner, the body lumen can be viewed, a measurement of
15 the diameter of the lumen at a point of interest can be taken, and the apparatus can then be used to remove amounts of tissue in a desired pattern to achieve beneficial reshaping of the lumen. It should be noted that such an imaging means may be useful in other
20 regards, for example, in allowing the surgeon to view certain anatomical markers or other guidelines during an inventive procedure, which can help ensure that tissue removal occurs where desired.

25 A body lumen diameter can be calculated in a variety of fashions, and in some embodiments, is accomplished using software incorporated into the endoluminal operating system. Illustratively, inventive systems can incorporate software that is able
30 to indirectly measure the diameter of a lumen at a location along the length of a body passageway, for example, by comparing software-based measurements, such

as those based on vectors displayed on an imaging means, with a reference article of known diameter (e.g., an anatomical feature or mechanical article, such as a ligation barrel or another viewable portion of an endoscope). Then, this measurement can be used to determine an effective amount and pattern of tissue removal to achieve a desired result, such as but not limited to a desired reduction in the lumen diameter.

Reshaping procedures can include removal of tissue from a substantial percentage of the circumference of the inner surface of a body lumen wall at one or more desired locations along the length of the wall. For instance, removal of tissue from about 30% to 100% of such a circumference at a desired longitudinal location can be used, and in this regard, it will be understood that in inventive procedures involving tissue removal from the circumference of more than one longitudinal location along a body lumen wall, any suitable percentage of the body lumen wall circumference can be removed at each longitudinal location. For example, in an inventive procedure involving tissue removal from the circumference of a first region (e.g., above the lower esophageal sphincter) and a second region (e.g., below the lower esophageal sphincter), the percentage of tissue removed from the first region (in terms of the region's circumference) may be the same as or different from the percentage of tissue removed from the second region circumference. Illustratively, in some forms, the same general percentage of tissue is removed from the circumference of each of these regions, yet because the second region circumference is

typically greater than the first region circumference,
more actual tissue (in terms of circumference length)
is removed from the second region circumference. In
other forms, the percentage of tissue removed from the
5 second region circumference will be about 10% to about
200% greater than the amount of actual tissue remove
removed from the circumference of the first region,
although other percentage differences are contemplated
within the scope of the invention. It should be noted
10 that in measuring the percentage of tissue removed from
the circumference of the inner surface of a body lumen
wall, such a measurement may be practically based on an
average or nominal circumference length, for example,
to account for folds, ridges, bumps or other surface
15 irregularities in the body lumen wall inner surface.

For reshaping of the gastroesophageal junction to
reduce reflux of gastric contents into the esophagus,
certain inventive procedures will typically involve the
20 removal of tissue from about 30% to about 70% of the
circumference of the lumen wall at one or more
locations in and/or around the junction.
Illustratively, tissue can be removed directly at
and/or near the junction, slightly above the junction
25 and/or slightly below the junction, to achieve the
desired result, e.g. a reduction in the diameter of the
junction and/or a lengthening of the junction. For
creating a stricture in the stomach, e.g., to modify
the eating habits of the patient, about 60% to about
30 100% of the circumference of one or more regions of the
stomach, especially the gastric cardia, will typically
be subjected to tissue removal.

Tissue removal can also involve a significant longitudinal swath. For instance, longitudinal dimensions of the regions of tissue removal will typically be at least about 0.25 centimeters, and more typically in the range of about 0.5 centimeters to about 3 centimeters. It will be understood, however, that other longitudinal tissue removal lengths can be used within the broader aspects of the present invention.

Advantageously, the reshaping of a body lumen achieved in accordance with embodiments of the invention is largely or completely reversible, for instance, by dilating the treated region after it has partially or completely healed. Full or partial reversal may be desirable for a number of reasons, for example, where a patient has trouble swallowing or wants to resume pre-treatment eating habits. In some forms, the diameter of a gastroesophageal junction, which is determined to be too small after treatment and healing, can be dilated to a modified, enlarged diameter. Dilation of a body lumen segment can be accomplished in any suitable manner, e.g., using a balloon or other expandable device. Such an enlarged diameter may be the pre-treatment diameter, or a diameter between the pre-treatment diameter and the initial healed diameter. Similar principles apply to strictures created in the stomach.

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Moreover, should the diameter or other reshaping of the body lumen after an initial tissue removal and

healing phase be to an extent that is insufficient to result in the desired modified flow of material through the lumen, follow-up procedures similarly conducted to remove additional amounts of tissue can be undertaken
5 to result in further reshaping.

A variety of devices and techniques can be used to achieve tissue removal according to the invention. Suitable such devices can be effective for resecting or
10 otherwise removing tissues, especially essentially mucosal and submucosal tissues. In some embodiments, an inventive tissue removal apparatus is comprised of a tubular surgical access device equipped with (or otherwise usable in conjunction with) suitable direct
15 or indirect (e.g. imaging) visualization means. Although not necessary to broader aspects of the invention, endoscopic devices and techniques, including those that have traditionally been used in endoscopic mucosal resection techniques are useful in some
20 inventive embodiments. These include both non-ligated tissue capture/resection techniques, which may involve a preceding injection of saline or other fluid to lift the tissue to be removed from an underlying muscle layer, and ligated tissue capture/resection techniques
25 and associated instrumentation therefor. In ligating techniques, a ligating barrel can be affixed to the end of an endoscope. Suction can be used to draw mucosal and submucosal tissues into the ligating barrel, and a ligating band thereafter deployed to result in a
30 ligated tissue mass. An electrosurgical snare or other similar device can then be used to resect the ligated tissue. Desirably, the endoscopic apparatus will

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include an electrosurgical snare receivable through the operating channel of the endoscope and effective to resect the ligated tissue mass, although snares passed separately and/or outside of the endoscope may be employed as well. One suitable
5 multiple-band ligating apparatus that can be used in methods of the invention is the Duette® Multi-Band Mucosectomy device commercially available from Wilson-Cook Medical, Inc., dba Cook™ Endoscopy, Winston-Salem, North Carolina, USA.

The Duette® device is the subject of an
10 international patent application entitled "System and Method for Endoscopic Treatment of Tissue" (WO2005112797), published December 1, 2005, and a United States provisional patent application entitled "System and Method for Endoscopic Treatment of Tissue" filed March 31, 2006. The Duette® device
15 kit includes a 7FR Soft AcuSnare™ Mini Hexagonal Head disposable polypectomy snare (Wilson-Cook Medical, Inc.). This snare has a soft hexagonal snare loop (size 1.5 cm x 2.5 cm) made with braided stainless steel cable, and a catheter sheath size of 7.0 FR. The snare is connectible to an electrocautery
20 unit and a modified multi-band ligator. The braided stainless steel cable provides a hexagonal operating loop (or snare head) with a combination of flexibility, strength and resiliency that permits multiple resections of tissue without breaking or excessively deforming. The multi-band ligator which is
25 available with 4, 6, or 10 bands, has a relatively large operating channel bore so

that the snare can be advanced through the working lumen of the endoscopy device to slice away tissue that has been captured by a deployed ligation band. Multiple ligations and resections can be conveniently
5 accomplished with this device with no substantial deformation occurring to the snare.

Referring now to the drawings, illustrative, non-limiting systems and procedures of the invention will
10 be described. FIG. 1A illustrates the stomach and adjacent portions of the anatomy including the esophagus 11 and the duodenum 12. The lower esophageal sphincter is generally indicated at 15. At least one of the goals of some inventive procedures is to reshape
15 and narrow one or more portions of this body passageway such as, for example, between the esophagus 11 and the stomach 10. Illustratively, an inventive procedure may be used to treat GERD by narrowing the gastrointestinal junction to provide a more effective barrier against
20 the reflux of acid and food contents from the stomach back into the esophagus. Still another procedure involves treating the cardia 16 below the gastrointestinal junction to narrow the passageway into the stomach so as to reduce an obese patient's food
25 intake capacity for the purpose of contributing to weight loss. Although not necessary to broader aspects of the invention, in certain embodiments, such procedures include an endoluminal mucosaplasty procedure. Other suitable tissue removal and/or injury
30 techniques, whether performed endoscopically or not, may be used in accordance with the invention to reshape or otherwise treat these or other body lumens. Other

candidate body lumens include but not limited to the anal sphincter area which can be treated to remodel tissues of this area to improve compliance of the anal sphincter to treat fecal incontinence and/or other
5 medical problems.

Referring to FIG. 1B, there is illustrated a section of the wall between the esophagus and stomach and showing the mucosa layer 17, submucosa layer 18,
10 muscle layer 20 and adventitia or serosa layer 21. Referring to FIG. 2, the multi-band ligator 22 is shown with a beveled barrel 25. Beveled barrels such as that shown in FIG. 2, disposed upon an endoscope, can facilitate the capture of tissue at preferred angles,
15 although it will be understood that a variety of other beveled and non-beveled barrels may be utilized as well. Suction is provided in the beveled barrel 25 of the ligator, thereby causing tissue of the mucosa layer 17 and the submucosa layer 18 to pull away from muscle
20 20 and be drawn into the barrel 25. The multi-band ligator 22 is then operated to deploy an elastic band 27 from the barrel 25 onto the captured tissue 26 to produce the result shown in FIG. 3, wherein the elastic band 27 has the illustrated reduced size based upon its
25 elasticity, capturing a mass of tissue. Although FIG. 2 shows multi-band ligator 22 having six elastic bands, it will be understood that this and other similar multi-band ligators useful in the invention can include any suitable number of elastic bands, and in some
30 instances, will include one to ten or more elastic bands, more typically two to eight elastic bands. In the illustrated and preferred embodiment, the captured

tissue 26 includes essentially only mucosa and submucosa and does not include any significant amount of captured muscle layer 20. After the tissue is captured with the band, the suction can be released or
5 discontinued.

Referring to FIG. 4, the multi-band ligator 22 further includes a snare 30 which is connected to an electrocautery unit. The snare 30 is operated through
10 a catheter 31 to tighten the snare 30 about the base 32 of the captured tissue 26 as generally shown in FIG. 5. The tightened snare can be positioned above or below the band 27, and in some modes, will occur around the band 27 or traversing the band 27 (i.e., with portions
15 above and below the band 27). The snare 30 is then used to slice away and cauterize the tissue, causing the resulting resection edges 35 (FIG. 6) to extend down into the submucosa, exposing a portion of muscle layer 20 as shown. In doing so, an amount of muscular
20 tissue may also be removed and/or some non-removed muscular tissue may be damaged at the site.

In some forms, the deep resection of cut 35 through the mucosa and submucosa and down to the muscle
25 layer causes the healing of the cut area to occur primarily from the edges 36 of the wound as indicated by the arrows 37, more so than from the bottom of the wound as can occur, for example, in a typical ulceration of superficial cuts of the mucosa. Such
30 healing causes the sides or edges 36 of the wound to move toward one another as new tissue is formed in and around the wound, which in turn, can be effective to

pull tight tissues adjacent to sides 36, and in this regard, such generally lateral healing can be highly effective in reducing the passage diameter in regions in and/or around the inferior esophageal sphincter
5 (e.g., in lower regions of the esophagus and/or upper regions of the cardia). Additionally, such healing in certain forms can cause a beneficial thickening of underlying muscle tissue in such regions.

10 In general, the healing response initiated can depend on a number of factors including but not limited to the wounded tissue(s) involved, the devices and techniques used to remove and/or traumatize the tissue(s), and the size and shape of the wound(s)
15 created. In this regard, it will be understood that wounds resulting from procedures of the invention can exhibit any suitable size, shape and configuration to achieve the desired reshaping. Illustratively, FIG. 7 shows an alternative cut 35, which extends at least
20 partially down into the submucosa. The relatively deep extension of the cut 35 into the submucosa causes at least some of the healing of the cut area to generally occur from the sides 36 of the wound as indicated by the arrows 37.

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Inventive methods may be used in treating patients who have GERD. For example, some methods involve reshaping the alimentary canal about the gastroesophageal junction so as to reduce the size of
30 this junction such that the luminal opening has a final diameter (i.e., when healed) less than the original or pre-treatment diameter. In many adult humans, it will

be desirable to reduce the diameter of the gastroesophageal junction, but not below approximately 15mm to 18mm when in a relaxed condition, although it will be understood that the appropriate diameter will vary from patient to patient within, above, or below this range.

In certain individuals, dimensions below this level will increase the likelihood of dysphagia (difficulty in swallowing). It will be understood, however, that these and other similar considerations will be undertaken by the health care provider, and re-sizing and final sizes or shapes of the body lumen will depend upon a variety of factors specific to the patient.

FIG. 9 shows a cross-section of esophagus adjacent to the lower esophageal sphincter. In order to accomplish the reduction in size of the gastroesophageal junction, multiple cuts 38 are accomplished circumferentially around the esophageal wall above the gastroesophageal junction while leaving untreated tissue 39 between each of the cuts as shown in FIG. 9. Although FIG. 9 shows three cuts 38 occurring above the gastroesophageal junction, it will be understood that any suitable number of cuts may be made in this region for reshaping purposes including 1 to 10 or more cuts, although 3 or 4 cuts will be more typical. Because the area below the gastroesophageal junction typically widens out, in instances where tissue is removed from both above and below the junction, a greater number of cuts and/or larger sized

cuts will typically (but not necessarily) be made in this lower region compared to the relatively higher region. Nonetheless, regardless of whether tissue above the gastroesophageal junction is also being removed in a given procedure, any suitable number of cuts may be made in regions below the gastroesophageal junction for reshaping purposes including 1 to 10 or more cuts, although 3 or 4 cuts will be more typical. In one illustrative procedure as suggested by areas 40 and 41 in FIG. 8, a total of 7 cuts can be taken, with 3 above and 4 below the gastroesophageal junction. Although some treatment site locations have been described herein in relation to the gastroesophageal junction (GEJ, the most proximate part of gastric folds), it will be understood that treatment site locations useful in the present invention may also be described in relation to other anatomical markers or structures such as but not limited to the lower esophageal sphincter and the squamo-columnar junction (SCJ), also known as the Z-line. The SCJ (Z-line) is the endoscopically visible line formed by the juxtaposition of pale squamous epithelium and reddish columnar epithelium. In healthy individuals, this junction does not always precisely coincide with the GEJ, as it can also occur adjacent the GEJ, such as above the GEJ. The identification and use of these and other similar anatomical markers in the reshaping of body lumens will be well within the purview of skilled practitioners given the teachings herein.

In certain embodiments, the ligator barrel 25 is provided with calibrations inside of the barrel that

are visible to the physician endoscopist, so that the amount of captured tissue 26 can be determined. In this manner, the endoscopist can estimate the size of the wound that will be produced at each treated site
5 around the circumference of the lumen. In these and other embodiments, various forms of software may be utilized (whether incorporated directly into or otherwise used in conjunction with an inventive apparatus) to measure the diameter of the lumen and
10 potentially address the size of the cuts necessary to provide the desired amount of reduction of lumen size.

Illustratively, the software can operate based upon one or more captured images including both the
15 lumen region to be measured and at least one reference marker of known dimension positioned at the level of the lumen region to be measured. The image(s) can be imported into the software, and the user can generate an outline (e.g. circle) based on the reference marker,
20 and an outline (e.g. circle) of the periphery of the lumen region to be measured. Using the known dimension of the reference marker, the software calculates the circumference of the lumen region on the basis of the relative dimensions of the outlines (e.g. circles). In
25 some situations, the shape of the lumen will correspond precisely to the shape of the outline generated (e.g. circle). In these situations, multiple measurements can be taken, and a mean calculated based upon the multiple measurements, to serve as the circumference
30 value for the procedure. As one example, to measure a size representative of the circumference of a gastroesophageal junction, an endoscope can be passed

through the esophagus and into the stomach, and a retroflexed image can be captured of the endoscope shaft at the point where it enters the stomach (the "reference point"). This image can be imported into a software program as described above, and a first circle drawn to the outer dimension of the shaft at the reference point. A second circle can then be drawn to the inner periphery of the esophagus-stomach junction at the reference point. Using the known dimension of the first circle and a comparison to the second circle, the software calculates a value of the circumference of the second circle which serves as a representative value for the circumference of the junction. This process can be repeated multiple times if desired and a mean value calculated for the representative value. This representative value can then be used as a basis for deciding on a treatment regimen as described herein.

20 Since the healing process will take place after the cutting procedure so that the initial diameter of the treated portion of the esophagus will be somewhat larger than the expected final diameter, a software program can be beneficially used to guide the physician as to what percentage of the esophageal wall should be treated and/or what pattern of tissue removal utilized. As clinical experience is obtained, further correlations can be developed as to the amount and/or pattern of tissue removal required to produce a desired result. If the post-treatment gastroesophagel junction diameter reduction is not sufficient following healing, the endoscopist can perform one or more follow-up

surgeries to make further resections to achieve the target diameter. On the other hand, if the junction area is made too small such that dysphagia becomes a problem, the procedure can be partially or completely reversed using standard esophageal dilation techniques (e.g., balloon dilatation) inasmuch as the muscle layer(s) below remains essentially intact. With the improved acid reflux barrier created by the inventive treatment, measurable pH should be improved above the gastroesophageal junction (a measure of treatment effectiveness).

Methods of the invention may also be used in treating obesity. These methods can involve making a series of resections only below the gastroesophageal (GE) junction. As shown in FIG. 10, which is a cross-sectional view just below the GE junction(taken along line 10-10), the number of resections may be four in number, although any suitable number of cuts may be made including 1 to 10 or more cuts, with 3 to 6 cuts being more typical. FIG. 11 shows the cross-section of the esophagus 50 and stomach 51 in which there is a hiatal hernia 52. In such a situation, fewer cuts at location 55 below the GE junction may be necessary. By narrowing the cardia below the GE junction, a smaller passageway for receiving food intake is established and, as a result, the patient will feel the sensation of satiety earlier. This will cause or assist the patient to stop eating earlier than he or she would otherwise do so. Of course, the patient may defeat the beneficial results of the procedure by eating small, more frequent meals; however, patient screening and

counseling can be provided to patients undergoing bariatric treatment to increase the likelihood of a successful and long-term result.

5 Unlike the treatment for GERD, where in most cases no more than about 70-75% of the tissue defining the circumference of the inner surface of a body lumen wall is removed and/or traumatized, the obesity patient is expected to commonly benefit from removal of and/or
10 damage to about 75% to about 100% of the tissue defining this body lumen wall inner surface, with higher amounts such as 90% to 100% sometimes being necessary to achieve the desired narrowing or other reshaping. In some embodiments, tissue is removed from
15 100% of the circumference of a body lumen wall at one or more longitudinal locations (e.g., 2 or 3) along the wall so as to provide one or more continuous "bands" of removed tissue along the wall. In many adult bariatric surgeries, a target lumen diameter of about 10
20 millimeters may be established. An advantage of more desired inventive procedures as compared to other bariatric surgeries is that they can be accomplished so as to be partially or completely reversible or otherwise adjustable. Thus, should the patient decide
25 that he or she wants the procedure reversed or should reversal be necessitated for other reasons, standard dilatation techniques, e.g., using a dilatation balloon, can be undertaken.

30 In additional aspects of the invention, modified endoscopic ligator barrels and devices are provided. Referring to FIG. 12, such a device 60 may include

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multiple ligation bands 61 received upon a ligation barrel 62, potentially having a beveled barrel end 63, wherein the bands 61 are deployable by a single flexible pull line, such as string 64, extending through the barrel. In this regard, the
5 pull string 64 can include a plurality of engagement members 65 configured to reside against and facilitate movement of the ligations bands 61 as the string 64 is pulled. The engagement members 65 can, for instance, be knots or polymeric beads positioned upon the string. The string 61 can then be situated
10 in a looped pattern around the barrel 62 (see phantom lines illustrating loops) or linear or other pattern through the ligation bands in such a manner that pulling the string causes a controlled, sequential deployment of the bands. Illustrative pull string/ligation band looping arrangements that can be
15 adapted as single-string configurations for use in aspects of the present invention are disclosed, for example, in U.S. Patent Nos. 6,007,551, 6,730,101 and 6,974,466.

In certain embodiments in which the barrel end 63 defines a bevel, the string will be positioned to exit the
20 barrel 62 at or substantially at the long side 66 of the barrel so as to provide an enhanced deployment of the bands 61. As well, the ligation barrel 62 can include an internal or external adaptation to prevent migration of the string circumferentially around the barrel as it is pulled.
25 Illustratively, an internal hoop or ring 67 can be attached to the inner barrel wall 68 and the string 64 passed therethrough to

control the position of the string. Such single-string devices can be connected to the end of an endoscope and provide enhanced visibility through the barrel, while still allowing for effective band deployment. As well,
5 it has been found that ligation bands received against the outer surface of a ligation barrel having a beveled tip can be successfully and effectively deployed employing longitudinally-directed pull string(s), despite the presence of the bevel. Accordingly, in one
10 embodiment, a beveled-tip ligation barrel includes one or more tissue ligation bands positioned against its outer surface and one or more pull strings for deploying the band(s). Such barrels can be mounted to the end of an endoscope, and used for inventive
15 reshaping methods as described herein or for other purposes.

In additional embodiments, the present invention provides kits for reshaping a body passage such as an
20 esophagous or stomach, as described herein, that include means or devices as described herein for resecting or otherwise removing tissue from the luman of the body passage, and written materials including instructions for use of the means or devices to reshape
25 a body passage, e.g. in the treatment of GERD or obesity as described herein. The kits can include the means or devices packaged together with the instructions, e.g. in sterile medical packaging. Related embodiments of the invention include methods
30 for distributing such means or devices, or otherwise conducting business, which include distributing such means or devices for reshaping a body passage, and also

distributing information relating the use of such means or devices for reshaping a body passage. Such information can be distributed packaged with the means or device, or separately, e.g. including information or instructions available on a communication network, including a global computer communication network such as the internet.

For the purpose of promoting a further understanding of certain aspects of the present invention, the following Examples are provided. It will be understood that these Examples are illustrative, and not limiting, in nature.

EXAMPLE 1

In this Example, endoscopic endoluminal mucoplasty (ELM) was performed on domestic farm pigs in a fashion that reshaped the gastroesophageal junction (GEJ) of the pigs. A first pig (Pig #1) underwent ELM of the cardia to reduce the diameter of the lumen at the (GEJ) by about one-half. To accomplish this, an endoscopic system equipped with the Duette® Multi-Band Mucosectomy device (Wilson-Cook Medical, Inc., dba Cook™ Endoscopy, Winston-Salem, North Carolina, USA) was used to band and resect tissue at 3 sites taken circumferentially around the cardia, representing about 70% of the circumference of the cardia. A second pig (Pig #2) was similarly treated, except the procedure was reversed after 5 weeks using a dilation balloon. These experimental pigs, along with controls, had access to an unlimited amount of food and full access to water. The pigs were healthy and active throughout the study.

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The results of the study are presented graphically in Figures 13, 14 and 15. As can be seen, the treated pigs experienced significantly less weight gain (shown in pounds) than the controls. Additionally, Pig #2 experienced accelerated weight
5 gain following reversal of the procedure.

EXAMPLE 2

In this Example, endoscopic endoluminal mucoplasty (ELM) was used to perform a fundoplasty on a domestic farm pig. An endoscopic system was used, equipped with the Duette®
10 Multi-Band Mucosectomy device as in Example 1. Repeated banding and resection was used to remove 10 square centimeters of the mucosa and submucosa of the fundus of the stomach. The fundus, which is the most expansile part of the stomach, was shrunk and reshaped upon healing after the resection. This
15 pig along with a control had access to an unlimited amount of food and full access to water over five weeks. The pigs were healthy and active throughout the study, and the treated pig experienced significantly less weight gain than the control pig.

20 While the illustrated embodiments have been detailed in the drawings and foregoing description, the same is to be considered as illustrative and not restrictive in character, it being understood that only the preferred embodiment has been shown and described and that all changes and modifications that
25 come within the invention are desired to be protected. The articles "a", "an", "said" and "the" are not limited to a singular element, and include one or more such elements. The scope of the claims should not be limited by the examples

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herein, but should be given the broadest interpretation consistent with the description as a whole.

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CLAIMS:

1. A medical device for use to reshape a body passage of a patient, the medical device comprising:

an access device for accessing a luminal surface of a
5 body passage of a patient, the access device including a barrel having a proximal end and a beveled distal end, the beveled distal end including a first side that is generally opposite a second side, the first side extending distally of the second side in a generally longitudinal direction along the barrel;

10 one or more bands received upon the barrel for banding tissue on the luminal surface;

a single pull line operably associated with the one or more bands, the single pull line operable to deploy the one or more bands from the barrel, the single pull line extending
15 through the barrel and exiting the barrel at or substantially at the first side of the barrel as the one or more bands are deployed; and

a tissue resection device for resecting tissue from the luminal surface;

20 wherein the medical device is for removing tissue from one or more areas of the luminal surface for the purpose of resulting in a healing response for reshaping the body passage.

2. The medical device of claim 1, wherein the device is
25 for removing mucosa and submucosa tissue.

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3. The medical device of claim 1, wherein the device is for making a plurality of resections distributed circumferentially within a target area of said body passage.

4. The medical device of claim 1, wherein the device is
5 for use in an esophagus of the patient.

5. The medical device of claim 4, wherein the device is used to treat gastroesophageal reflux in the patient.

6. The medical device of claim 1, wherein the device is for use in a stomach of the patient.

10 7. The medical device of claim 6, wherein the reshaping of the body passage caused by the use of the device is for modifying a satiety sensation of the patient.

8. The medical device of claim 1, the medical device further comprising: an endoscope for visualizing the luminal
15 surface of the body passage during resection.

9. The medical device of claim 8, wherein said endoscope has an operating channel for conducting the resection through the channel.

10. The medical device of claim 9, the tissue resection
20 device further comprising: an electrocautery snare, a loop of the electrocautery snare being passable through the operating channel of the endoscope for capturing and cutting the tissue from the one or more areas of the luminal surface.

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11. The medical device of claim 10, wherein the band is for banding the tissue prior to capturing the tissue in the loop.

12. The medical device of claim 11, wherein the band is
5 for deploying around a pseudo-polyp of tissue created in a container with vacuum.

13. The medical device of claim 1, wherein the patient is diagnosed as having gastroesophageal reflux disorder; and the medical device is for removing the tissue based upon the
10 diagnosing.

14. The medical device of claim 1, wherein the patient is diagnosed as needing treatment for obesity; and the medical device is for removing the tissue based upon the diagnosing.

15. The medical device of claim 1, wherein the medical
15 device is for removing the tissue from the body passage without substantial damage to underlying muscle tissue of the luminal body passage.

16. The medical device of claim 14, wherein the device is for use in an esophagus.

20 17. The medical device of claim 14, wherein the device is for use in a stomach.

18. The medical device of claim 1, wherein the tissue is non-diseased tissue.

19. The medical device of claim 1, wherein the tissue
25 comprises mucosa and submucosa.

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20. The medical device of claim 1, wherein the device further comprises a measuring device for measuring a diameter of said body passage prior to using the tissue resection device.

5 21. A medical device, comprising:

an endoscopic ligator barrel having a proximal end and a beveled distal end, the beveled distal end including a first side that is generally opposite a second side, the first side extending distally of the second side in a generally
10 longitudinal direction along the ligator barrel;

one or more ligation bands received upon the ligation barrel;

a single pull line operably associated with the one or more ligation bands, the single pull line operable to deploy
15 the one or more ligation bands from the ligation barrel, the single pull line extending through the ligator barrel and exiting the barrel at or substantially at the first side of the barrel as the one or more ligation bands are deployed.

22. A medical device according to claim 21, also
20 comprising an endoscope, with the endoscopic ligator barrel received upon the endoscope.

23. A medical device according to any one of claims 21 and 22, wherein the ligator barrel is provided with calibrations for determining an amount of tissue received in
25 the ligator barrel.

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24. A medical device according to any one of claims 21 and 22, wherein the beveled distal end extends substantially across the width of the ligator barrel.

25. A medical device according to any one of claims 21
5 through 24, wherein said one or more ligation bands are positioned along a longitudinal segment of the ligator barrel, and wherein the circumference of the barrel is substantially constant along the length of said longitudinal segment.

26. A medical device according to any one of claims 21
10 through 24, further comprising a guide member coupled to the ligator barrel, wherein the pull line passes through the guide member.

27. A medical device according to claim 26, wherein the guide member is positioned on an inner barrel wall within the
15 ligator barrel.

28. A medical device according to any one of claims 26 and 27, wherein the guide member is positioned on the inner barrel wall substantially on the first side.

29. A medical device of any one of claims 21 through 28,
20 wherein the single pull line exits the barrel to engage the one or more ligation bands without reentering the barrel.

30. A medical device, comprising:

an endoscopic ligator barrel having a proximal end and a beveled distal end, the beveled distal end including a
25 first side that is generally opposite a second side, the first

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side extending distally of the second side in a generally longitudinal direction along the ligator barrel;

one or more ligation bands received upon the ligation barrel, the one or more ligation bands being deployable from
5 the beveled distal end; and

a single pull line operably associated with the one or more ligation bands, the single pull line operable to deploy the one or more ligation bands from the ligation barrel, the single pull line extending through a guide member coupled to
10 the ligator barrel, the guide member configured to prevent circumferential migration of the single pull line around the barrel as the ligation bands are deployed.

31. A medical device according to claim 30, wherein a portion of the pull line is situated outside the ligator
15 barrel.

32. A medical device according to claim 30 or 31, wherein the pull line extends through the ligator barrel and exits the ligator barrel at or substantially at the first side of the barrel.

20 33. A medical device according to any one of claims 30 through 32, wherein the ligator barrel includes calibrations for determining an amount of tissue received in the ligator barrel.

34. A medical device according to any one of 30
25 through 33, wherein said beveled distal end extends substantially across the width of the ligator barrel.

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35. A medical device according to any one of claims 30 through 34, wherein said one or more ligation bands are positioned along a longitudinal segment of the ligator barrel, and wherein the circumference of the barrel is substantially
5 constant along the length of said longitudinal segment.

36. A medical device according to any one of claims 30 through 35, wherein the guide member is positioned on an inner barrel wall within said ligator barrel.

37. A medical device of any one of claims 30 through 36,
10 wherein the single pull line exits the barrel to engage the one or more ligation bands without reentering the barrel.

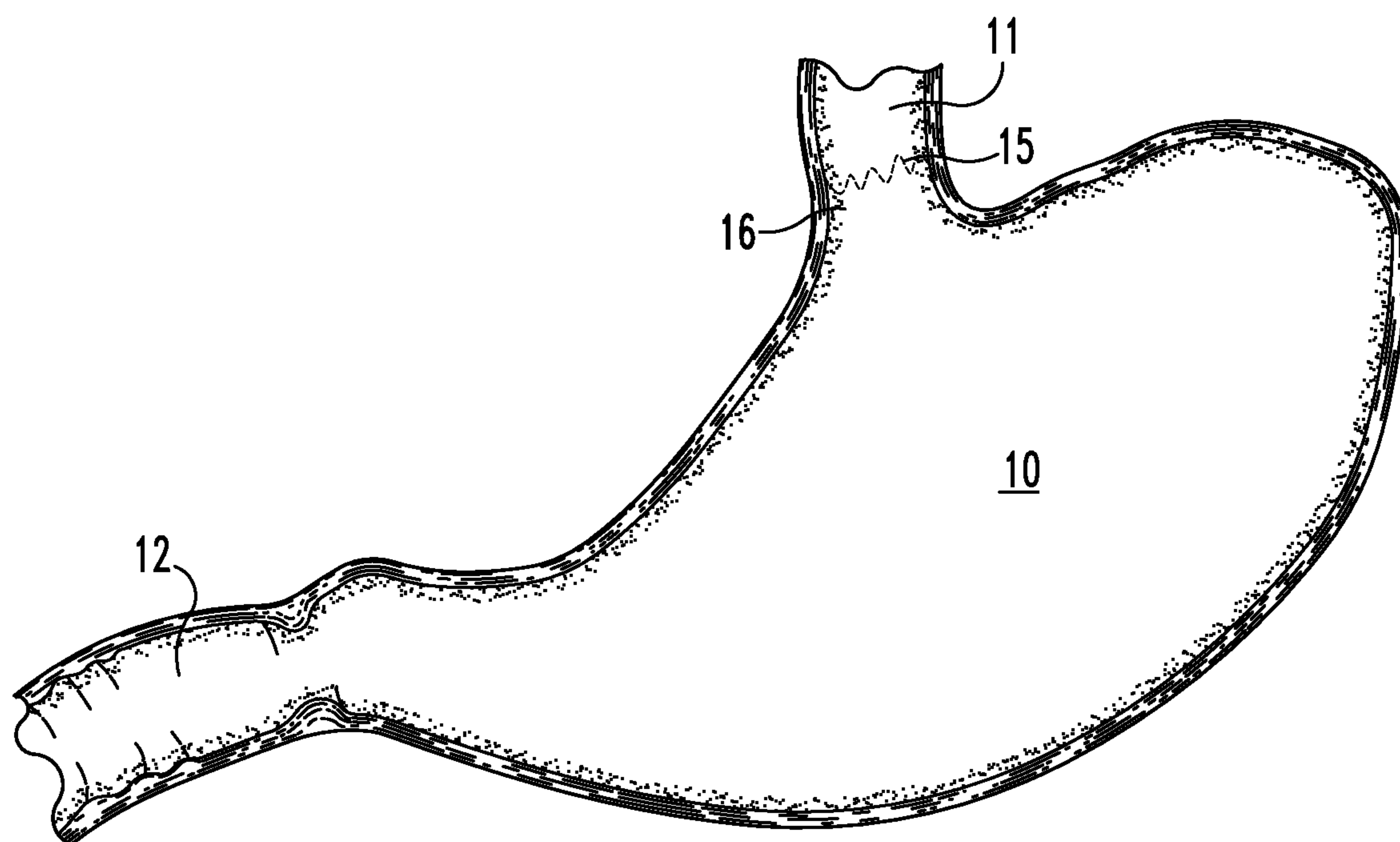


Fig. 1a

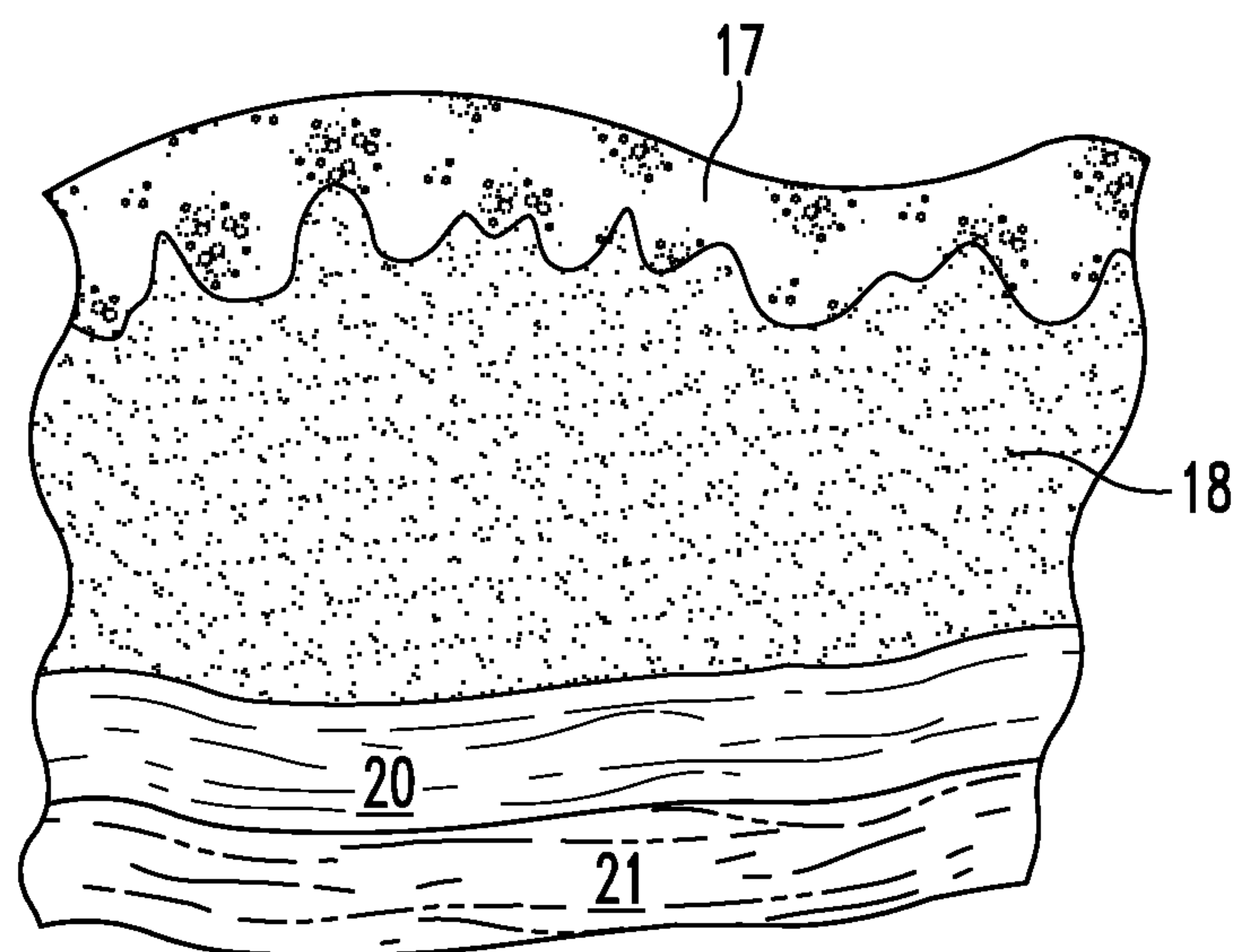
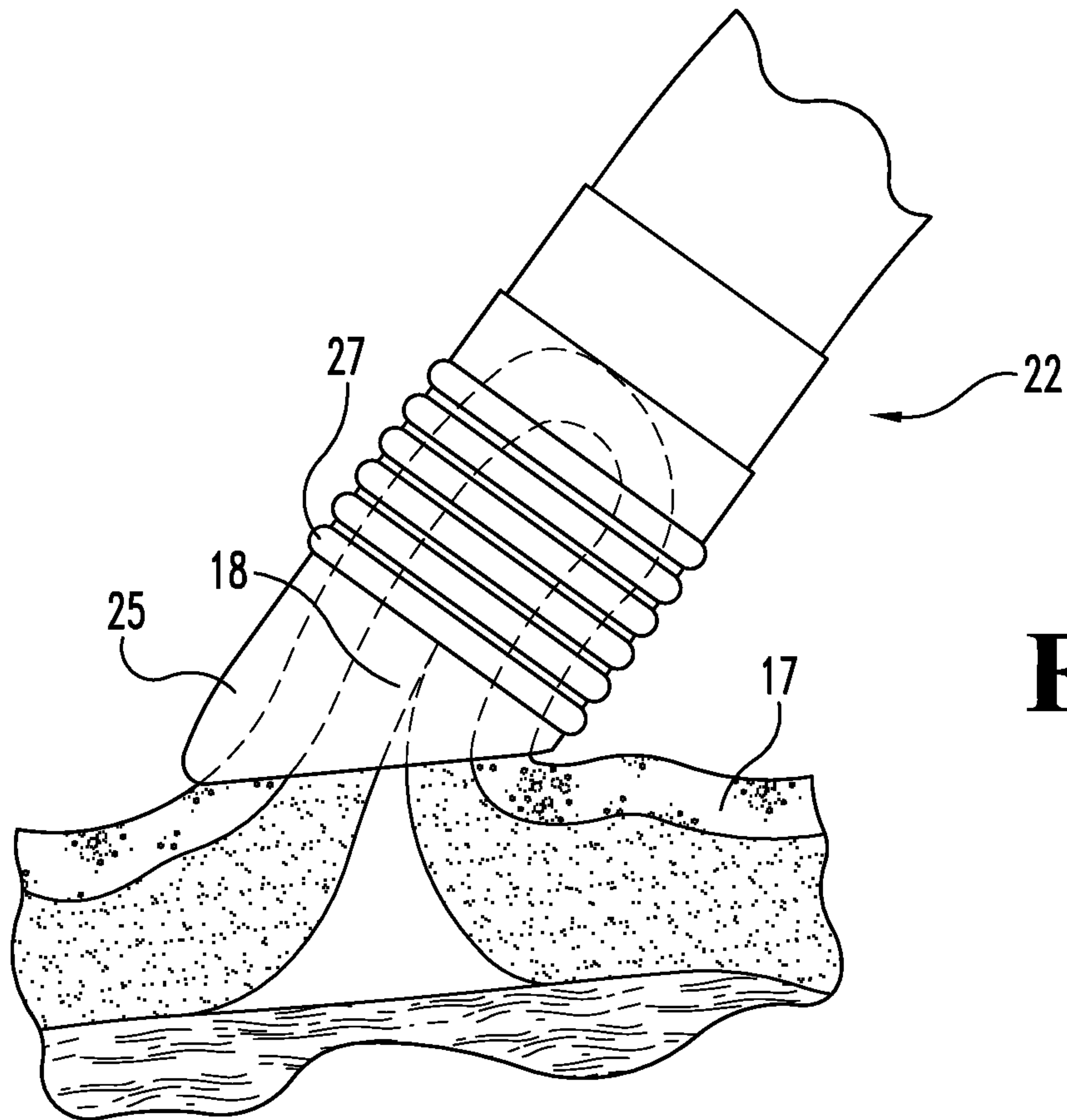
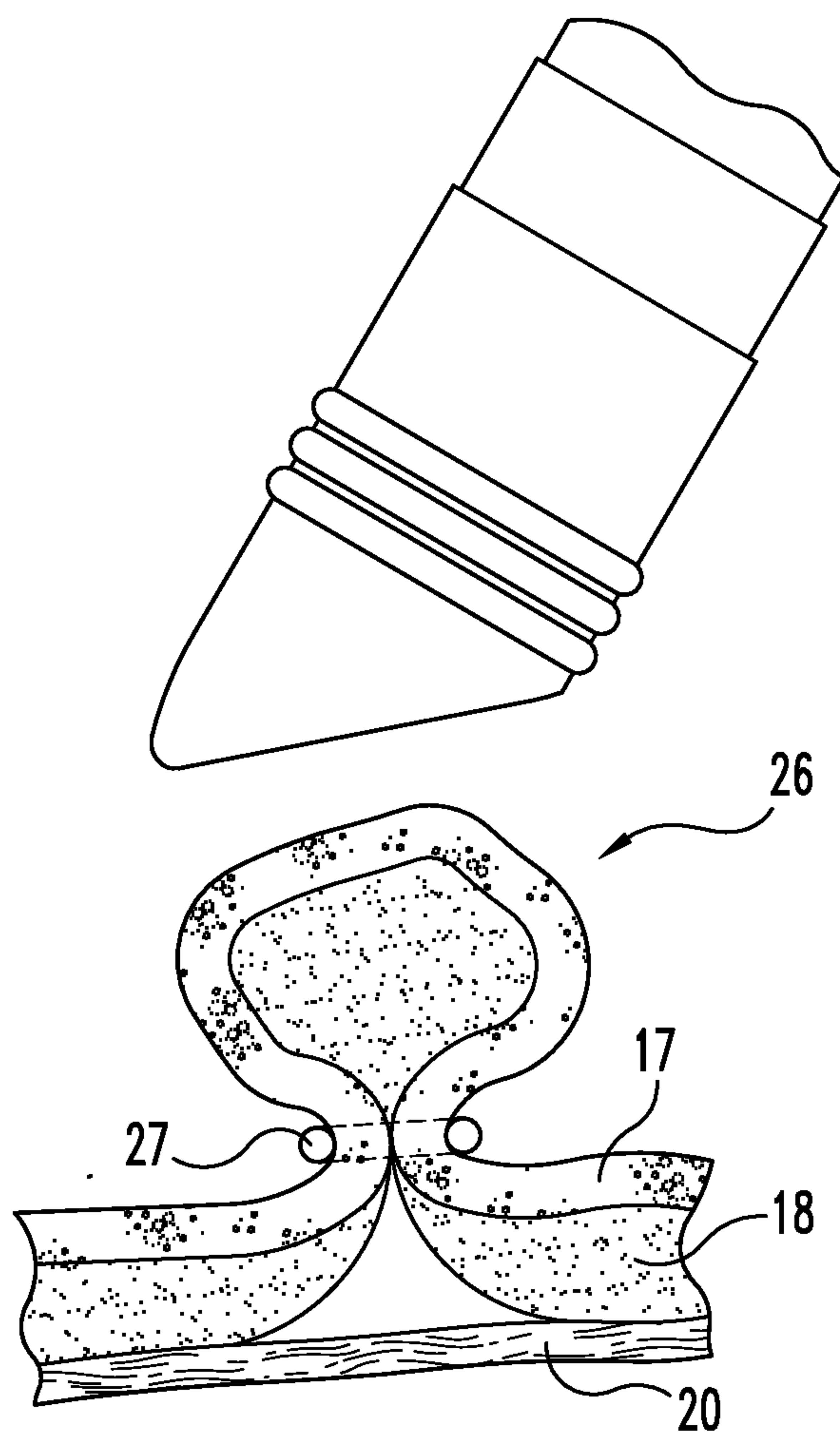
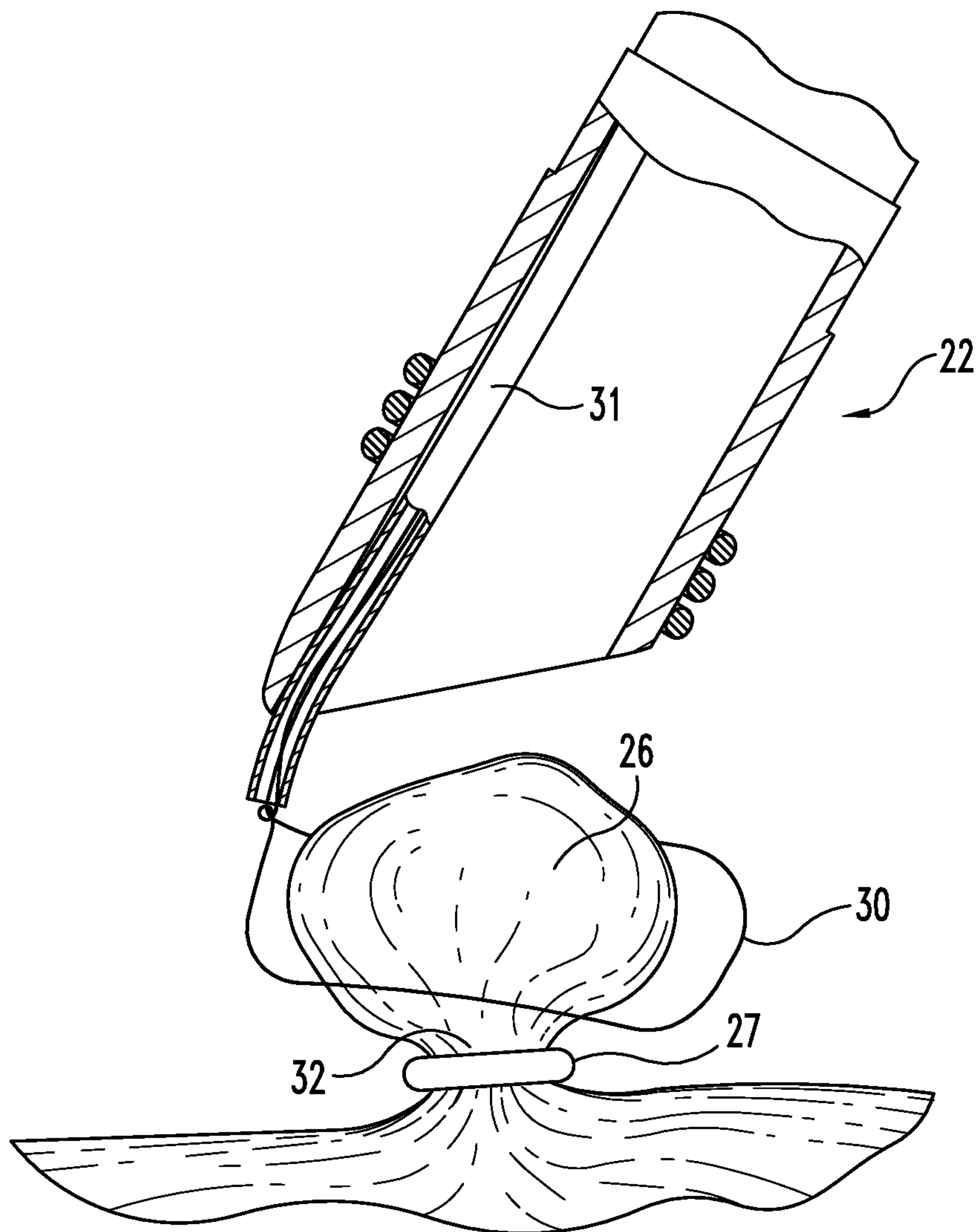


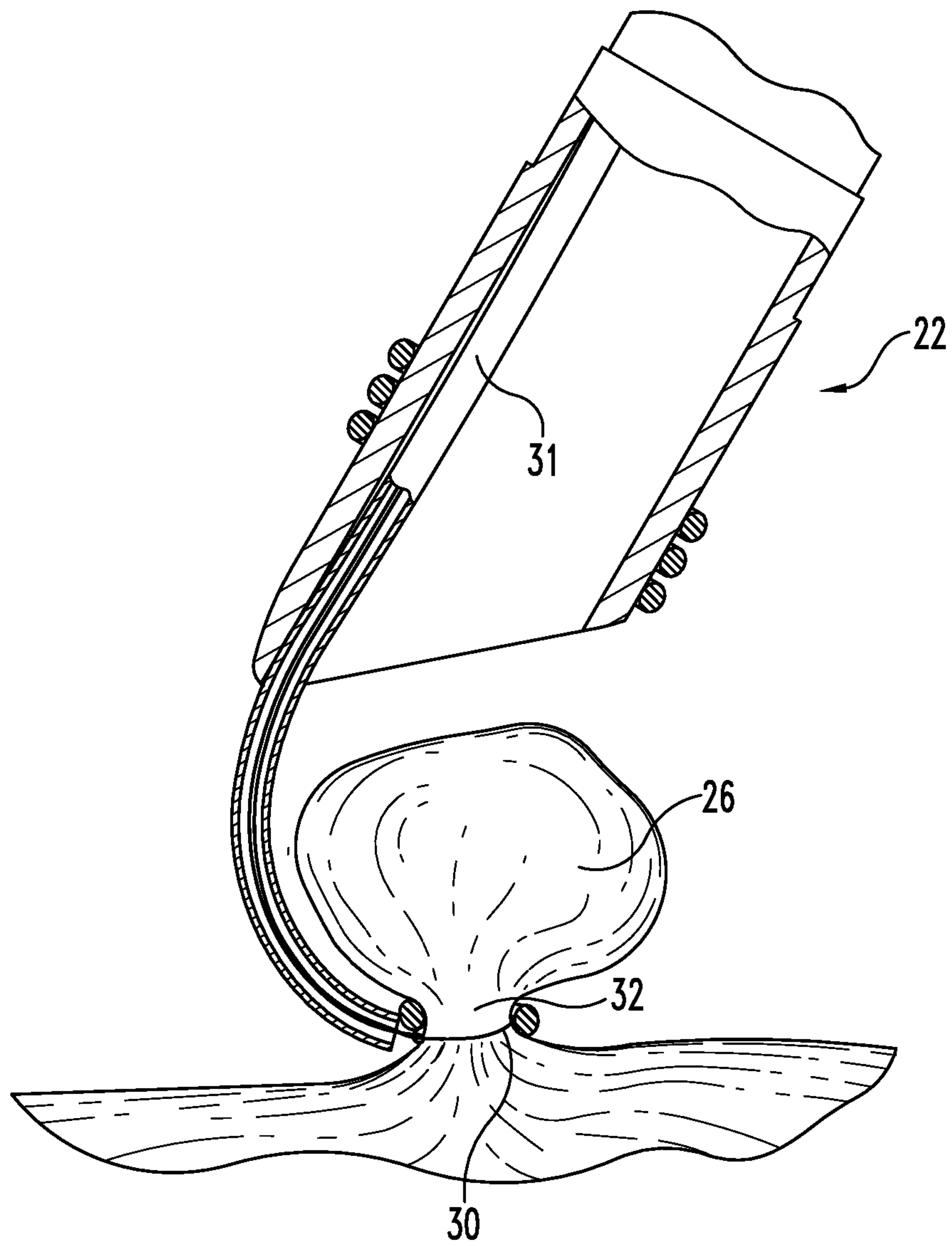
Fig. 1b

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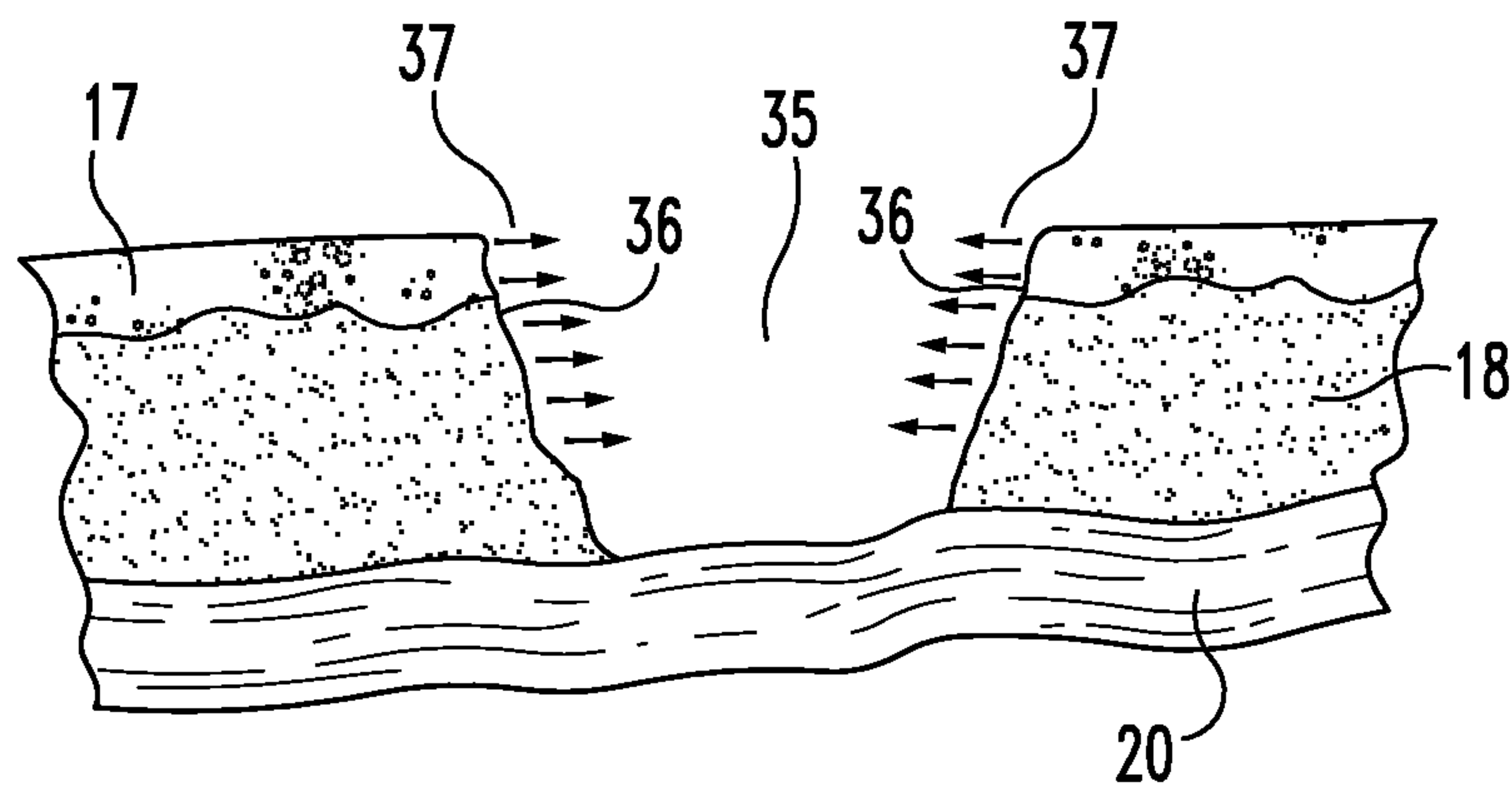
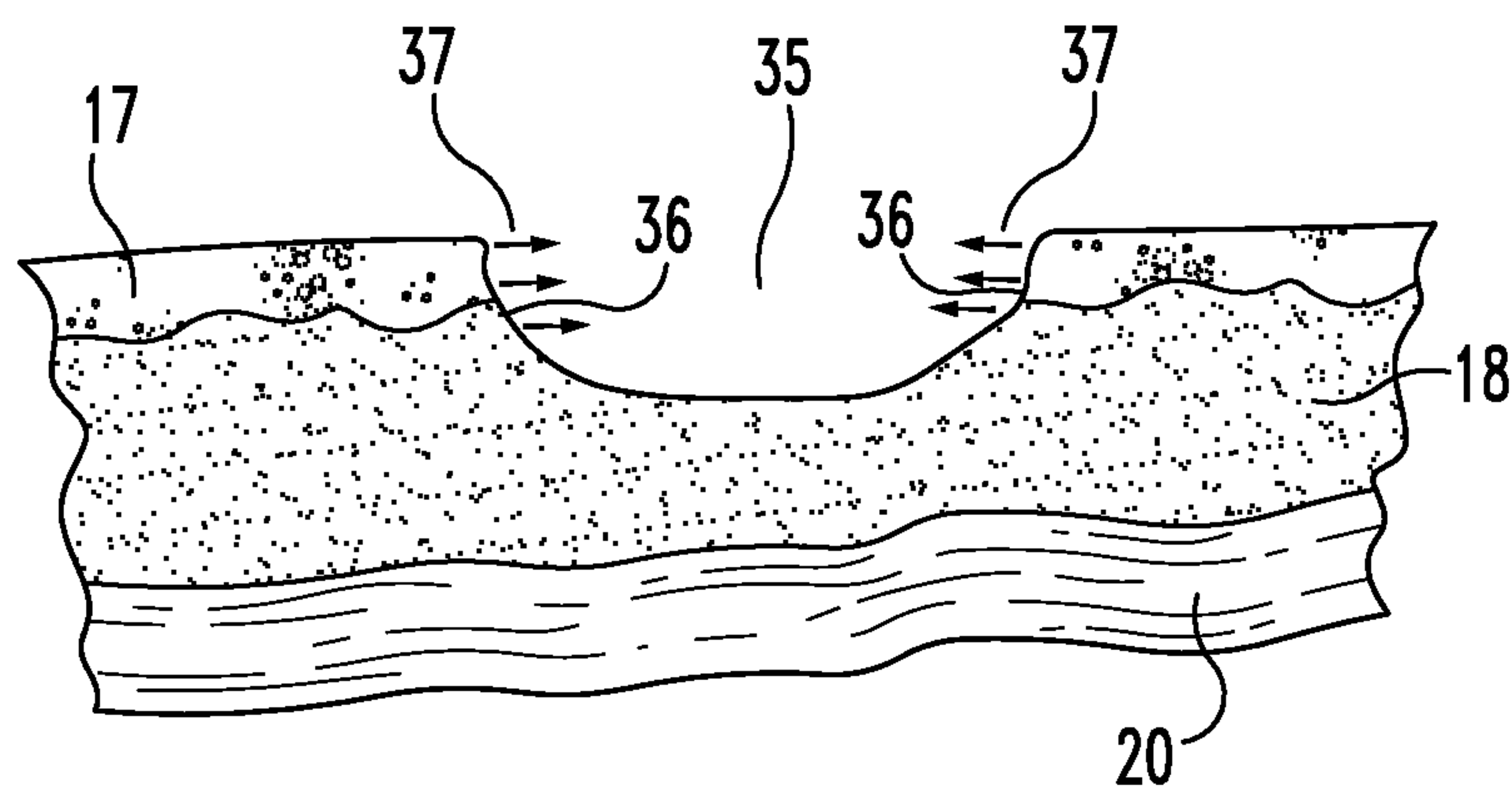
**Fig. 2****Fig. 3**

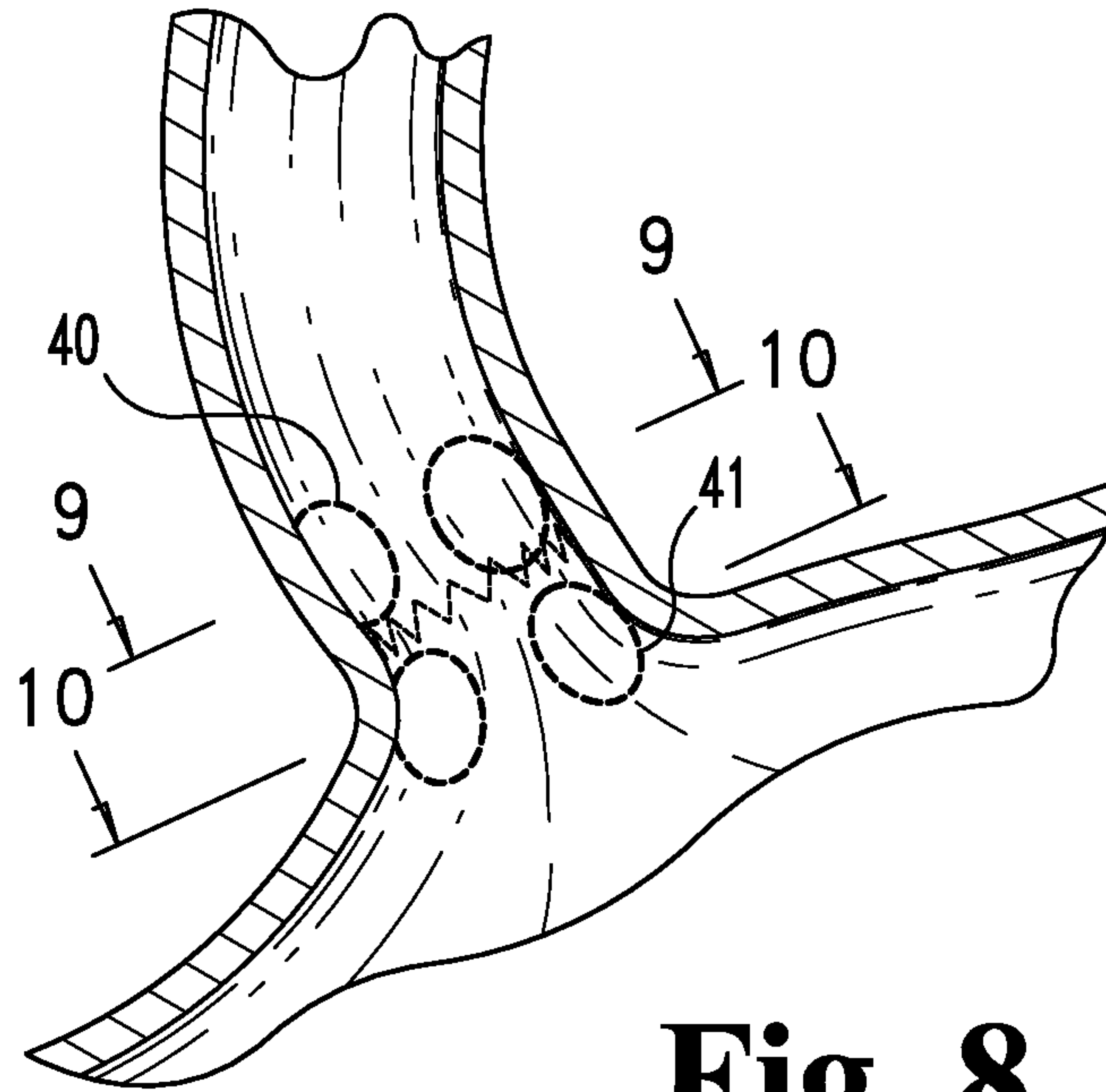
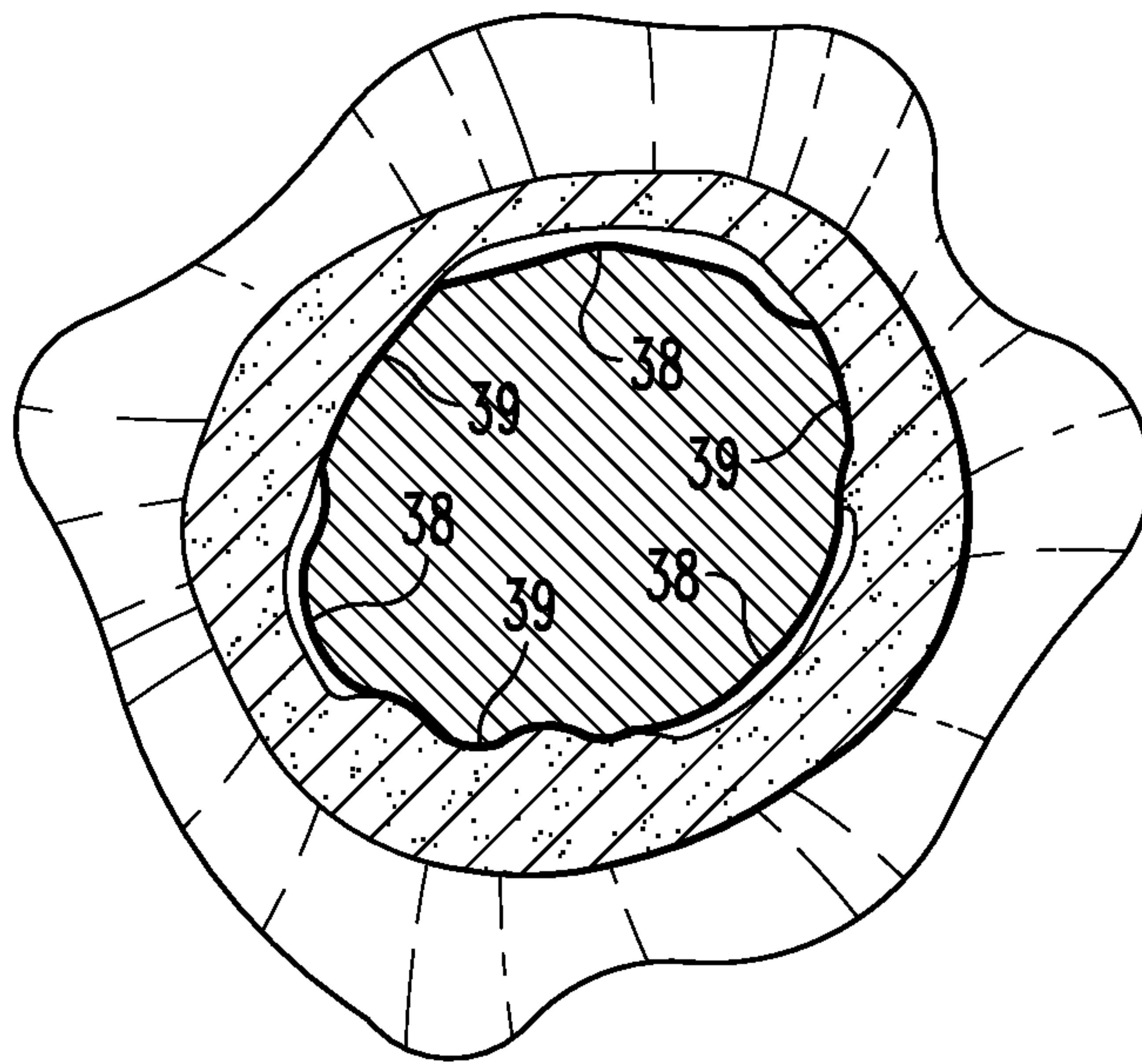
**Fig. 4**

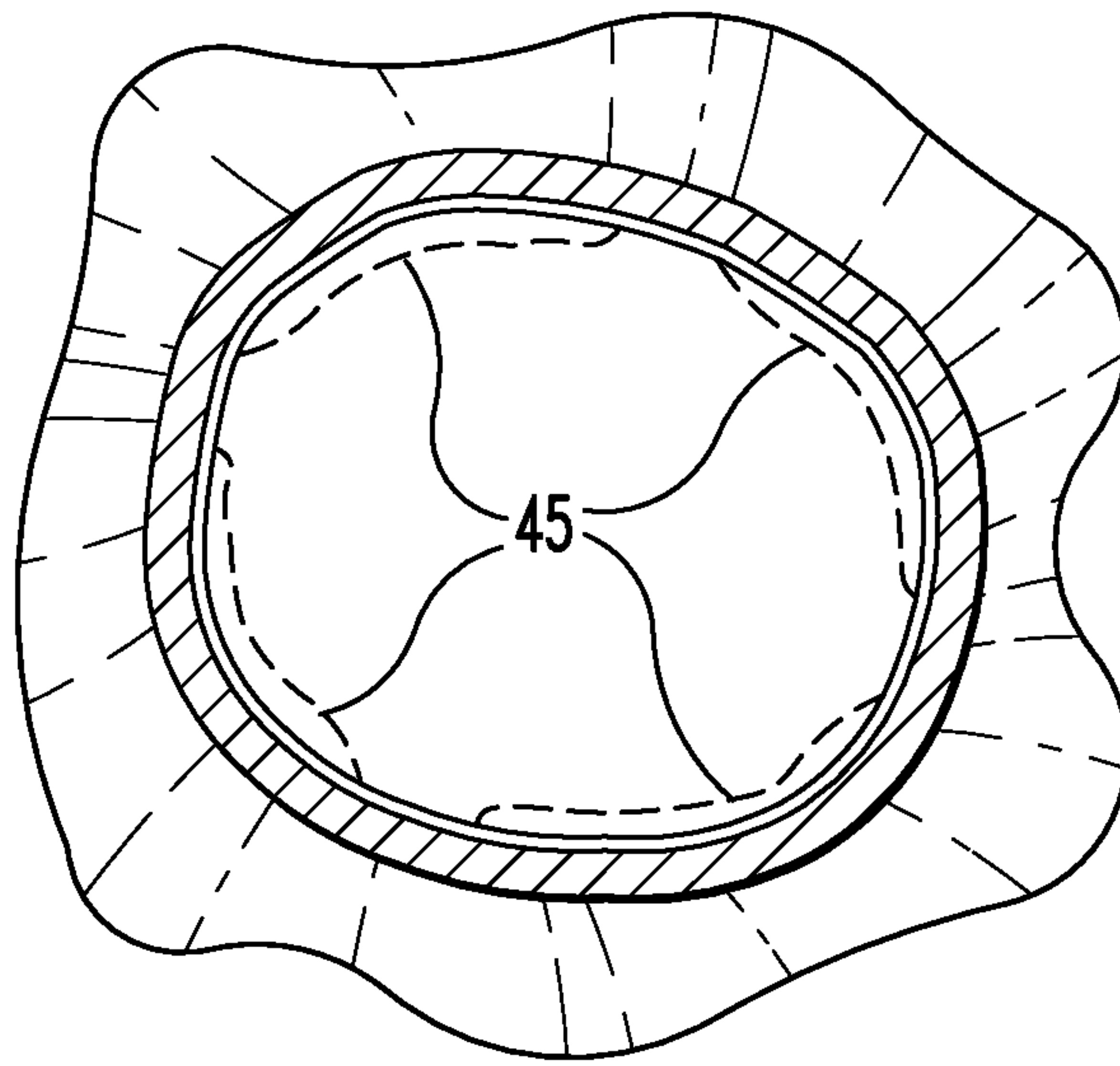
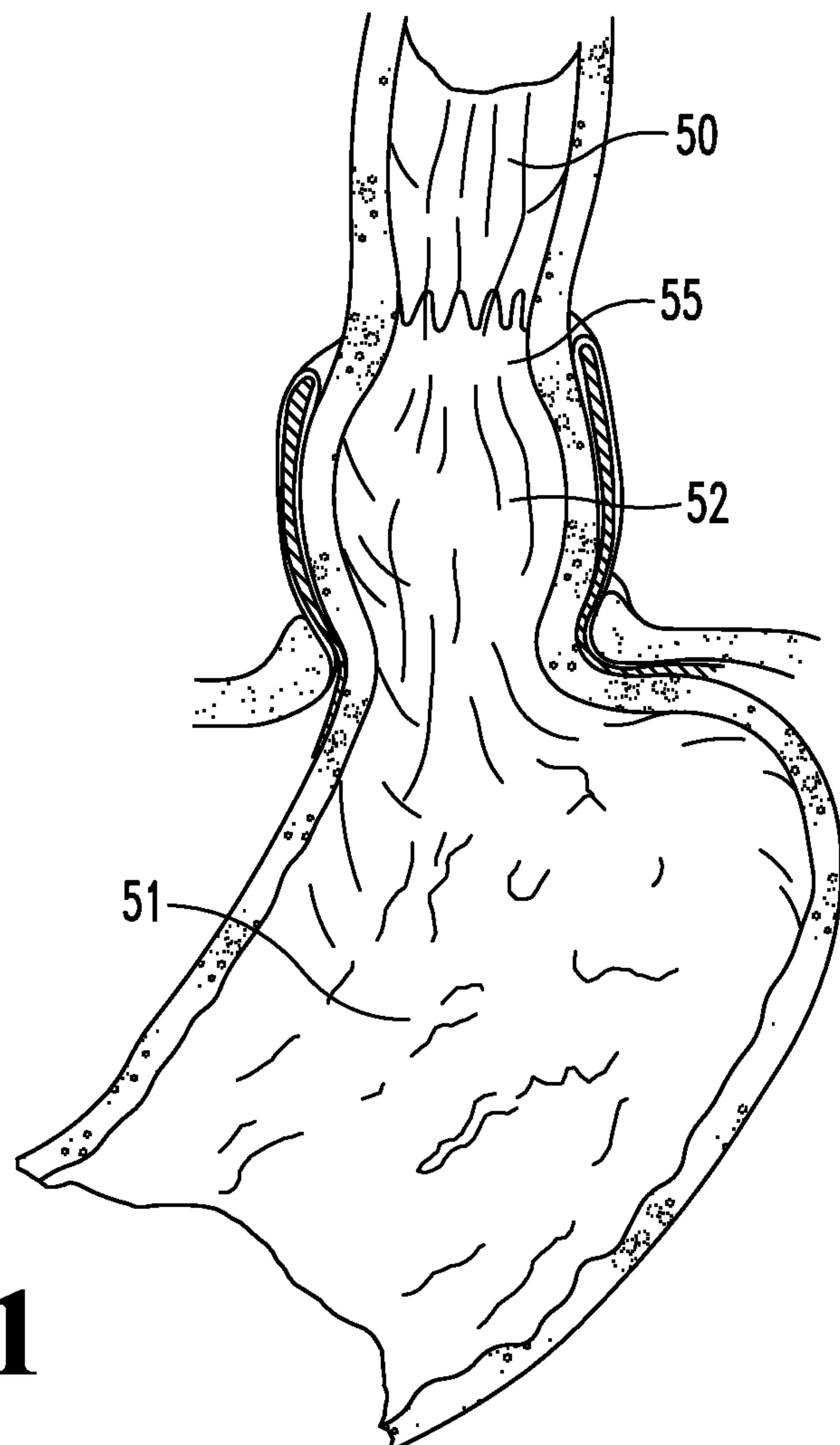
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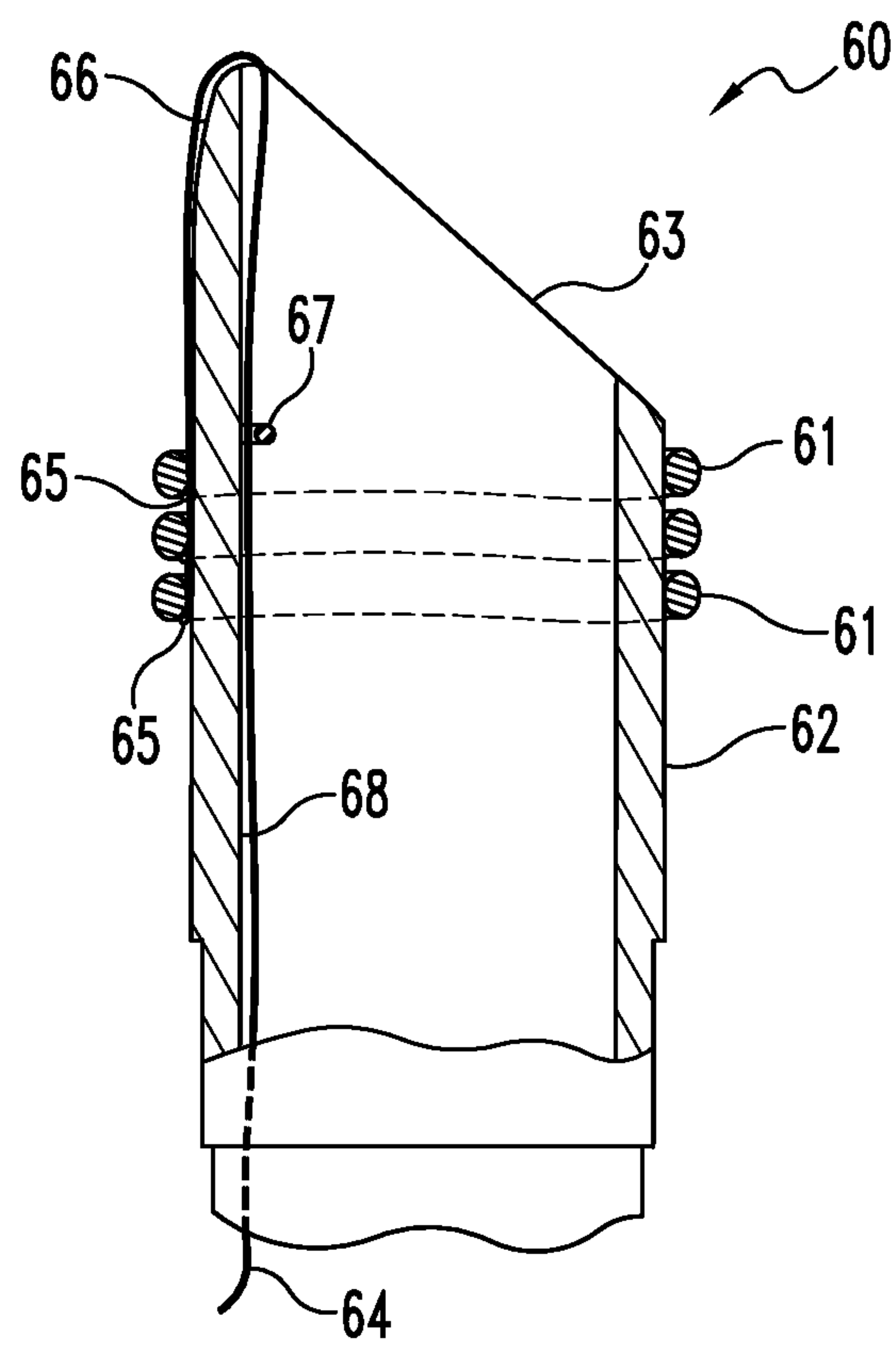
**Fig. 5**

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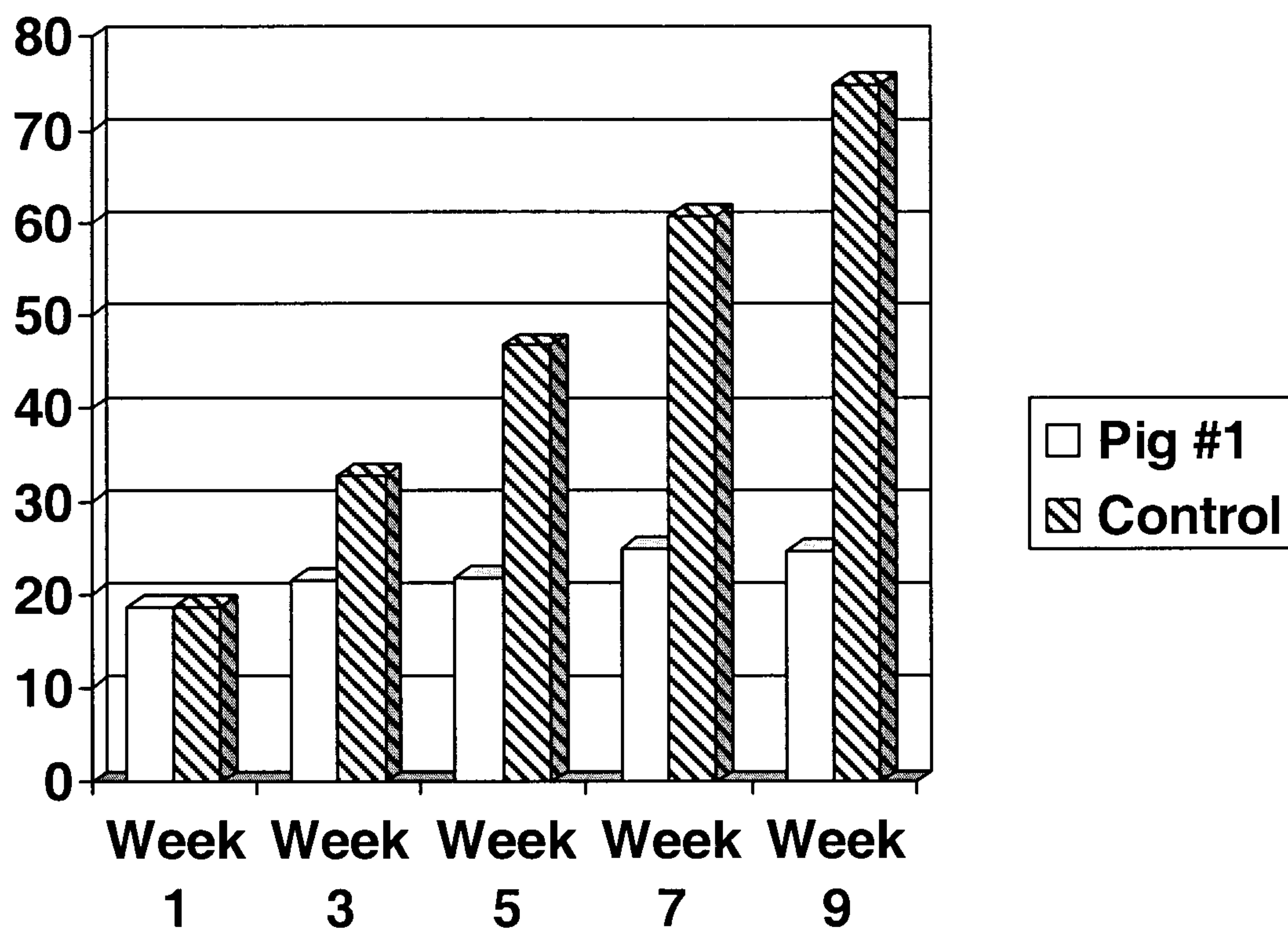
**Fig. 6****Fig. 7**

**Fig. 8****Fig. 9**

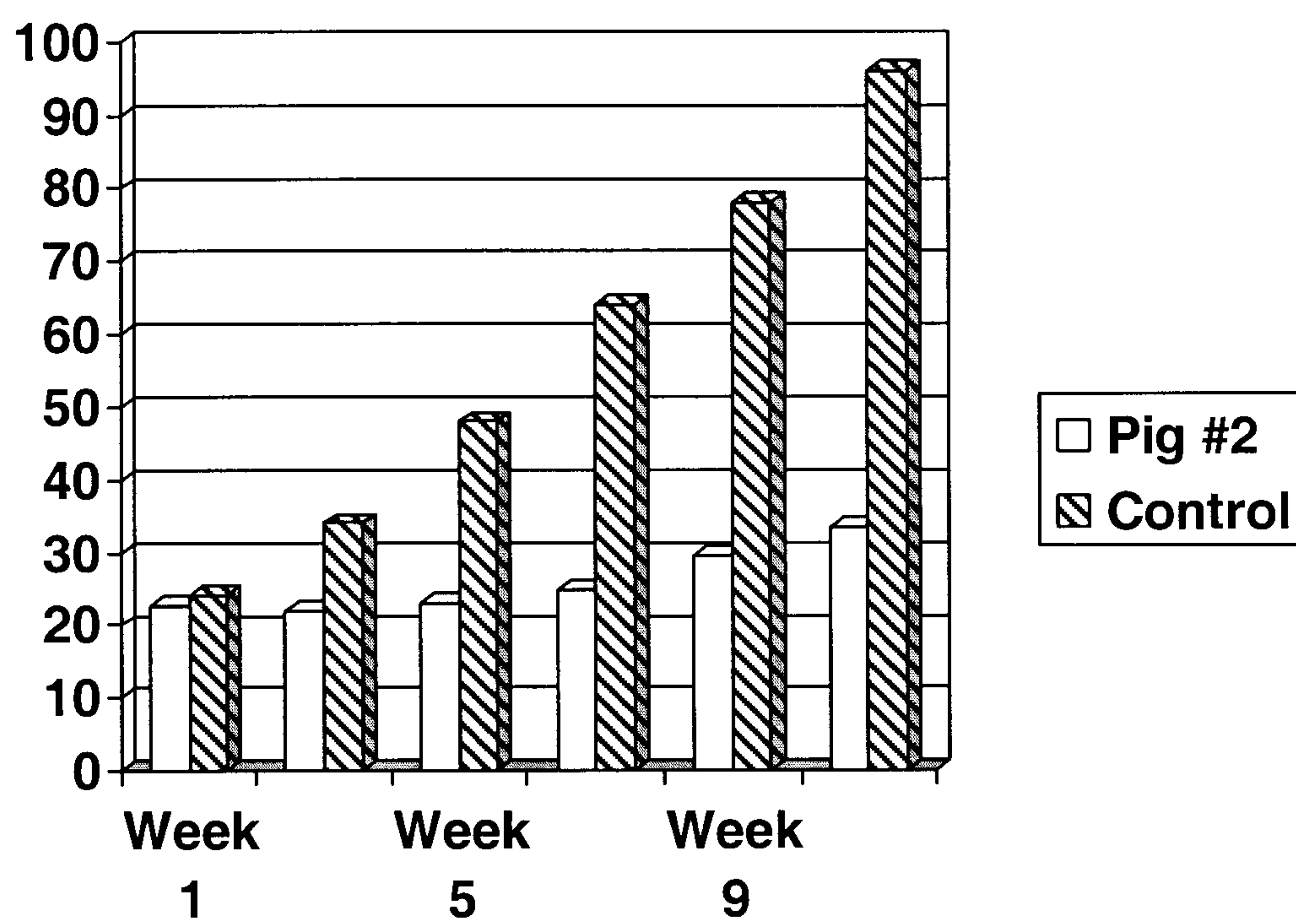
**Fig. 10****Fig. 11**

**Fig. 12**

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**FIGURE 13**

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**FIGURE 14**

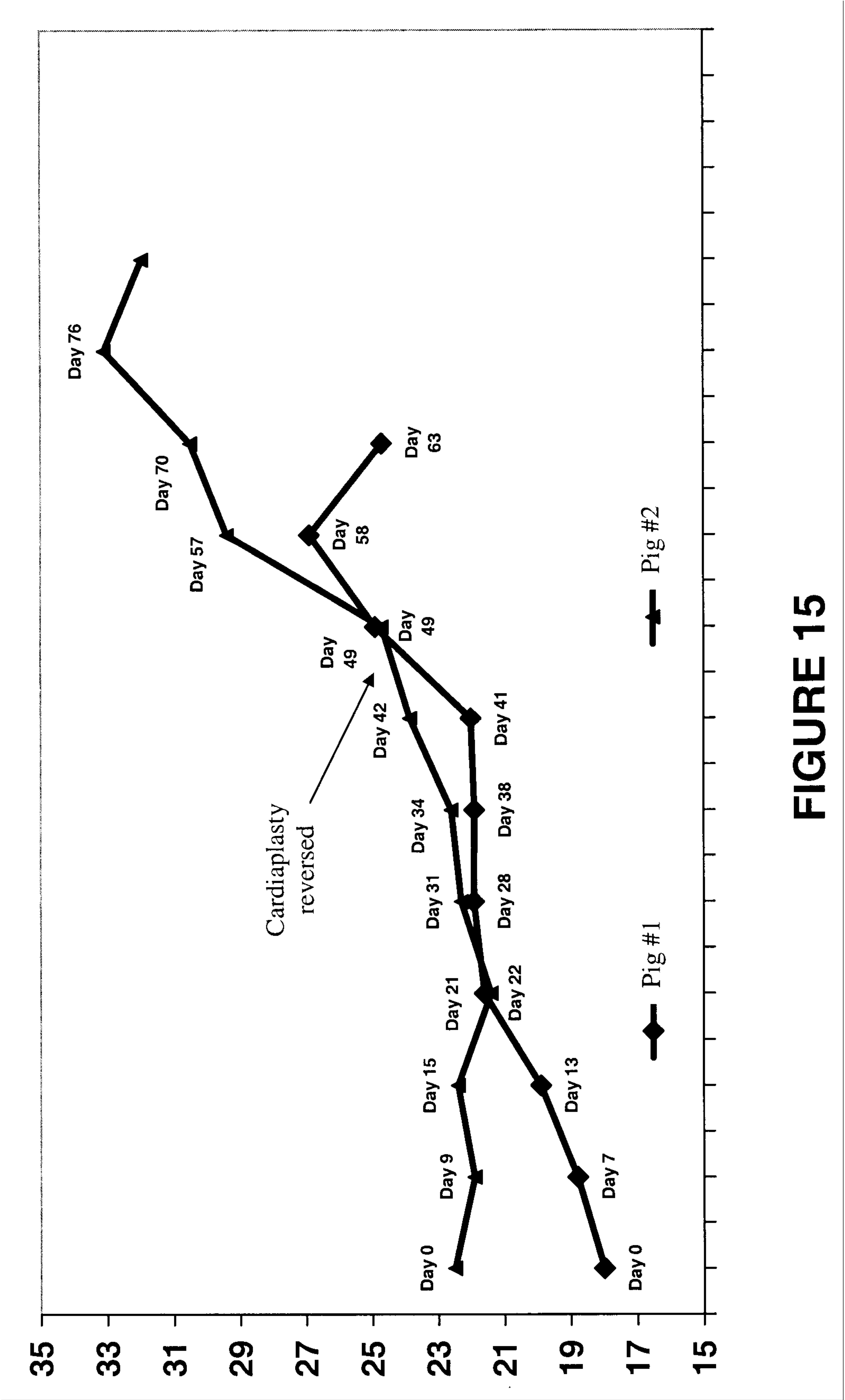


FIGURE 15

