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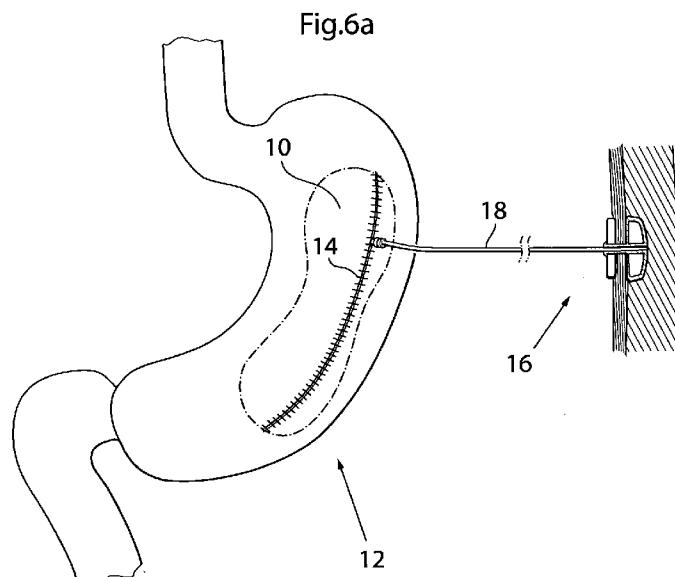
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(54) Title: APPARATUS FOR TREATING OBESITY



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(57) Abstract: An volume filling device for treatment of obesity is placed outside the stomach wall of the patient to reduce the inner volume of the stomach, thereby affecting the patient's appetite. By providing the volume filling device outside the stomach wall, contact with stomach acids is avoided, thereby increasing the life of the device.

APPARATUS FOR TREATING OBESITY

TECHNICAL FIELD

The present invention relates to an apparatus, a system, and a method for treating
5 obesity.

BACKGROUND

Obesity has been treated by gastric banding a band placed around the stomach to
create a stoma, a restricted opening, to restrict the flow of food down to below the
band. There has also been tried to use electrical stimulation of the stomach wall to
10 cause the patient to feel satiety.

When the stomach gets distended the patient gets a feeling that the stomach is full.

Another prior art way of treating obesity is to insert a balloon-like object into the
stomach of the patient. In this way, the patient is given the feeling of satiety much
more quickly when eating, preventing excessive intake of food. However, these prior
15 art balloon-like objects are subject to stomach acids, leading to their destruction
within a couple of months of use.

An example of a prior art inflatable gastric device for treating obesity is disclosed in
US patent No. 4,246,893 to Berson. In this document, it is disclosed an abdominal
method wherein an inflatable balloon is surgically implanted in the abdominal cavity
20 of the patient adjacent to the stomach. An adjusting port is provided subcutaneously
and the balloon is subsequently inflated by means of inserting a hypodermic needle
through the skin of the patient into the adjusting port and introducing a fluid under
pressure into the port for passage into the balloon to distend the upper abdomen,
compressing the stomach and thereby producing a sense of satiety.

25 SUMMARY

The object of the present invention to provide obesity treatment apparatus, system
and methods with improved long term properties.

This object and others are obtained by an apparatus described in the appended claims. Thus, by providing an apparatus comprising at least one volume filling device adapted to be at least substantially invaginated by a stomach wall portion of the patient and having an outer surface that includes a biocompatible material,

5 wherein the volume filling device is adapted to be placed outside of the stomach wall with the outer surface of the volume filling device resting against the outside of the stomach wall, such that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device, the volume filling device having a maximum circumference of at least 30 millimeters, an apparatus for

10 treating obesity is obtained. The present invention is based on the realization that by invaginating a volume filling device by the stomach wall on the outside thereof this device is protected from the stomach acids and will thus remain functioning for a very long time.

Preferably, the volume filling device is adapted to be completely invaginated by the

15 stomach wall of the patient and to be placed outside the stomach wall via a gastroscopic instrument. To this end the volume filling device may comprise an attachment device adapted to co-operate with a gripping instrument. Suitably, the volume filling device is adapted to be non-invasively adjustable postoperatively.

The apparatus may comprise a fixation device, suitably two or more fixation

20 devices, adapted to be involved in the fixation of the volume filling device to the stomach wall. The volume filling device may comprise a holding device adapted to be held by an instrument, suitably two or more holding devices, to simplify the implantation of the device.

At least a part of the volume filling device may be made of a material which is not

25 destructible by stomach acid. The volume filling device may be destructible by acids, for example hydrochloric acid.

In an embodiment, the volume filling device is inflatable to an expanded state and comprises an enclosure wall defining a chamber, wherein the volume filling device is inflated with a gel or fluid supplied into the chamber. At least one tube may be

30 connected to the volume filling device for supplying gel or fluid to the chamber. An

injection port connectible with the tube may be provided. Alternatively, the volume filling member may be provided with an inlet port for a fluid or a gel connectible to a gastroscopic instrument, wherein the inlet port comprises a fluid connection adapted to interconnect the inflatable device and the gastroscopic instrument.

- 5 The volume filling device may include a homogenous material, such as gel having a shore value of less than 15. The device may also be a solid body.

The volume filling device may comprise a rigid, elastic or flexible outer surface.

Where the outer surface is rigid, it is rigid enough to maintain non-deformed when subject to forces created by stomach movements. The volume filling device may

- 10 comprise a flexible non-elastic material.

In accordance with a first general design of the volume filling device, the device has a maximum circumference as seen in a plane perpendicular to an axis through the device. The circumferences of the device as seen in other planes perpendicular to said axis are equal to the maximum circumference or decrease as seen along said

- 15 axis in the direction from the maximum circumference. For example, the device may be substantially egg shaped, spherically shaped, or substantially shaped like an egg with an indented middle section or like a bent egg.

In accordance with a second general design of the device, the circumference of the device as seen in a plane perpendicular to an axis through the device increases and

- 20 decreases at least two times as the plane is displaced along said axis, or decreases and increases at least one time as the plane is displaced along said axis. For example, the device may be substantially shaped like a kidney.

The volume filling device have an elongated, rounded, bent and/or curved shape.

The volume filling device has a circumference of at least 120, 150, 180 or 220 mm.

- 25 The volume filling device has a volume in the range of 0.0001 to 0.001 m³ , or 0.00001 to 0.001 m³, or 0.00001 to 0.0002 m³. The volume of the volume filling device has a volume of less than 0.0002 m³.

The volume filling device may comprise at least two interconnectable portions adapted to be placed inside the stomach as separate portions.

The volume filling device may comprises an elastic material, a bio-compatible material and/or silicone.

- 5 Suitably, the volume filling device is provided with a coating. For example, a metal coating, a Parylene coating, a polytetrafluoroethylene coating or a polyurethane coating. The coating may be a multi-layer coating. Suitably, one of the layers may be made of made of metal, silicon or PTFE. The volume filling device may comprise an outer surface layer of polyurethane, Teflon®, or PTFE, or a combination thereof.
- 10 The volume filling device may comprise a fluid adapted to be transformed into solid state or fixed form. Such a fluid may be liquid polyurethane or iso-tonic. The fluid may comprises large molecules, such as iodine molecules, to prevent diffusion.

The volume filling device may have a maximum circumference of at least 50 millimeters, preferably at least 80 millimeters. Suitably, the volume filling device is deformable to a maximum diameter, so as to be insertable into a laparoscopic trocar.

15 Preferably, the volume filling device is adapted to be kept in place by stomach-to-stomach sutures or staples to invaginate the device in the stomach wall.

Advantageously, the volume filling device has varying circumference to better be kept in place invaginated in the stomach wall of the patient. The stomach-to-stomach

- 20 sutures or staples may be provided with fixation portions exhibiting a structure adapted to be in contact with the stomach wall to promote growth in of human tissue to secure long term placement of the volume filling device attached to the stomach wall. The structure may comprise a net like structure.

In embodiment of the invention, the apparatus comprises a stretching device placed

- 25 outside the stomach wall and adapted to stretch a portion of the stomach wall, thereby affecting the patient's appetite. Where the volume filling device is inflatable, the apparatus may comprise a fluid connection interconnecting the stretching device and the volume filling device.

In an embodiment, the apparatus comprises a stretching device comprising at least one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall and an operation device for operating the stretching device when implanted to stretch the stomach wall portion such that
5 satiety is created.

In an embodiment, the apparatus comprises at least one operable stretching device implantable in the patient and adapted to stretch a portion of the patient's stomach wall, and an implantable control unit for automatically controlling the operable stretching device, when the control unit and stretching device are implanted, to
10 stretch the stomach wall portion in connection with the patient eating such that satiety is created.

In an embodiment, the apparatus comprises a stretching device comprising at least one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall, wherein said stretching device comprising an
15 expandable stretching reservoir and an operation device for operating the stretching device when implanted to stretch the stomach wall portion, wherein the volume filling device is inflatable and in fluid connection with said stretching reservoir, wherein said operation device comprises a pump for pumping fluid between said main reservoir and said stretching reservoir to stretch said stomach wall portion such
20 that satiety is created. A control device may be provided for controlling said stretching device including said pump. The control device may comprise a wireless remote control adapted to control the stretching device from the outside of the patient's body, or an implantable control unit for controlling said stretching device. Alternatively, the control device may comprise a subcutaneously placed switch or
25 reservoir adapted to control the stretching device from the outside of the patient's body. A sensor or sensing device to be implanted in the patient body may be provided, wherein the implantable control unit is adapted to control the stretching device from the inside of the patient's body using information from said a sensor or sensing device, adapted to sense, direct or indirect, the food intake of the patient.

30 In an embodiment, the volume filling device comprises a main volume filling reservoir, a stretching device comprising at least one operable stretching device

implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall, wherein said stretching device comprising an expandable reservoir, adapted to be invaginated in the stomach wall at the upper part of the stomach, higher up than the inflatable main volume filling device when the patient is standing,

5 wherein the volume filling device is inflatable and in fluid connection with said stretching reservoir, wherein normal contractions of the stomach wall, related to food intake, cause fluid to flow from said invaginated main volume filling reservoir lower placed onto the stomach wall adapted to cause said stretching reservoir to stretch said stomach wall portion such that satiety is created. The fluid connection between the

10 main volume filling device reservoir and the stretching reservoir comprises a non-return valve. The fluid connection between the main volume filling device reservoir and the stretching reservoir comprises a release function adapted to release the volume in the stretching reservoir back to the main volume filling device reservoir. Said release function may comprise a fluid return connection of a substantially

15 smaller area than said fluid connection, to slowly release back fluid to said main volume filling device reservoir from the stretching reservoir to release said stretching of the stomach wall portion. A further manual control device comprising a subcutaneously placed reservoir adapted to control the stretching device from the outside of the patient's body may be provided to further affect the stretching device

20 to stretch the stomach wall portion.

In an embodiment, the a main volume filling device reservoir adapted to be inflatable may be provided, wherein the volume filling device further comprises an expandable structure, adapted to expand, when the device is invaginated in the stomach wall, wherein said structure comprising a bellow adapted to take into account the fibrosis surrounding the device when implanted, such that the movement of the bellow is substantially un-affected of said fibrosis.

In an embodiment, the apparatus comprises a stretching device comprising at least one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall and wherein the stretching device comprising a expandable structure, adapted to expand and stretch the stomach wall portion, when the device is invaginated in the stomach wall, wherein said structure comprising a

special bellow adapted to take into account the fibrosis surrounding the device when implanted, such that the movement of the bellow is substantially un-affected of said fibrosis. An operation device for operating the stretching device may be provided to stretch the stomach wall portion such that satiety is created. The apparatus may

5 comprise an implantable control unit for automatically controlling the operable stretching device, when the control unit and stretching device are implanted, to stretch the stomach wall portion in connection with the patient eating such that satiety is created.

In an embodiment, the apparatus comprises a stretching device comprising at least

10 one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall such that satiety is created. The control device may comprise a wireless remote control adapted to control the stretching device from the outside of the patient's body or an implantable control unit for controlling said stretching device. Alternatively, said control device may comprise a subcutaneously

15 placed switch or reservoir adapted to control the stretching device from the outside of the patient's body. A sensor or sensing device adapted to be implanted in the patient body may be provided, wherein the implantable control unit is adapted to control the stretching device from the inside of the patient's body using information from said sensor or sensing device, adapted to sense, direct or indirect, the food intake of the

20 patient.

In an embodiment, the apparatus is further adapted to treat reflux disease. To this end, it further comprises an implantable movement restriction device adapted to be at least partly invaginated by the patient's stomach fundus wall and having an outer surface that includes a biocompatible material, wherein a substantial part of the outer

25 surface of the movement restriction device is adapted to rest against the stomach wall without injuring the latter in a position between the patient's diaphragm and at least a portion of the lower part of the invaginated stomach fundus wall, such that movement of the cardiac notch of the patient's stomach towards the patient's diaphragm is restricted, when the movement restriction device is invaginated, to

30 thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's

cardia sphincter muscle exerted from the patient's abdomen, the movement restriction device having a size of at least 125 mm³ and a circumference of at least 15 mm.

In another embodiment, the apparatus is further adapted to treat reflux disease. To 5 this end, it further comprises an implantable movement restriction device having an outer surface including a biocompatible material, wherein the movement restriction device is adapted to rest with at least a part of its outer surface against the patient's stomach fundus wall, in a position between the patient's diaphragm and the fundus wall, such that movement of the cardiac notch of the patient's stomach towards the 10 patient's diaphragm is restricted, when the movement restriction device is implanted in the patient, to thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's cardia sphincter muscle exerted from the patient's abdomen, wherein the movement restriction device having a size of at least 125 mm³ 15 and a circumference of at least 15 mm, and an a fixation device adapted to secure the movement restriction device in said position, when the movement restriction device is implanted.

In another embodiment, the apparatus is further adapted to treat reflux disease. To this end, it further comprises an implantable movement restriction device adapted to 20 be at least partly invaginated by the patient's stomach fundus wall and having an outer surface that includes a biocompatible material, wherein a substantial part of the outer surface of the movement restriction device is adapted to rest against the stomach wall without injuring the latter in a position between the patient's diaphragm and at least a portion of the lower part of the invaginated stomach fundus 25 wall, such that movement of the cardiac notch of the patient's stomach towards the patient's diaphragm is restricted, when the movement restriction device is invaginated, to thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's cardia sphincter muscle exerted from the patient's 30 abdomen, the movement restriction device having a size of at least 125 mm³ and a circumference of at least 15 mm, further comprising a stretching device comprising

at least one operable stretching device implantable in the obese patient and adapted to stretch a portion of the patient's stomach wall such that satiety is created.

In another embodiment, the apparatus is further adapted to treat reflux disease. To this end, it further comprises an implantable movement restriction device having an 5 outer surface including a biocompatible material, wherein the movement restriction device is adapted to rest with at least a part of its outer surface against the patient's stomach fundus wall, in a position between the patient's diaphragm and the fundus wall, such that movement of the cardiac notch of the patient's stomach towards the patient's diaphragm is restricted, when the movement restriction device is implanted 10 in the patient, to thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's cardia sphincter muscle exerted from the patient's abdomen, wherein the movement restriction device having a size of at least 125 mm³ and a circumference of at least 15 mm, and a fixation device adapted to secure the 15 movement restriction device in said position, when the movement restriction device is implanted, further comprising a stretching device comprising at least one operable stretching device implantable in the obese patient and adapted to stretch a portion of the patient's stomach wall such that satiety is created.

In an embodiment, the apparatus further comprises a stretching device comprising 20 three or more mechanical parts engaged with different parts of the stomach wall, one part each, wherein said engagement includes suturing or stapling to the stomach wall or invaginating the mechanical parts in the stomach wall part with stomach to stomach sutures, wherein the three or more mechanical parts are adapted to move in relation to each other adapted to stretch three different wall portions, the stretching 25 device further adapted to having said wall portions stretched independently from each other both regarding force used for stretching the stomach wall portion as well as, time periods the stretching is applied, and when the stretching is applied.

In an embodiment, the apparatus further comprises a stretching device comprising 30 two or more hydraulic parts engaged with different parts of the stomach wall, one part each, wherein said engagement includes suturing or stapling to hydraulic part to the stomach wall or invaginating the hydraulic parts in the stomach wall part, with

stomach to stomach sutures, wherein the two or more hydraulic parts are adapted to move in relation to each other adapted to stretch three different wall portions, the stretching device further adapted to having said wall portions stretched independently from each other both regarding force used for stretching the stomach 5 wall portion as well as, time periods the stretching is applied, and when the stretching is applied.

In an embodiment, the apparatus further comprises a stretching device is engaged with a part of the stomach wall, including suturing or stapling the stretching device to the stomach wall or invaginating the stretching device in the stomach wall part, with 10 stomach to stomach sutures, wherein the stretching device is further adapted to stretch a stomach wall portion controlling force used for stretching the stomach wall portion as well as, time periods the stretching is applied, and when the stretching is applied.

In an embodiment, the apparatus further comprises a stretching device comprising 15 two parts engaged with different parts of the stomach wall, one part each, wherein said engagement includes suturing or stapling the parts to the stomach wall or invaginating the parts in the stomach wall part, with stomach to stomach sutures, wherein the stretching device further adapted to have different wall portions stretched independently from each other controlling force used for stretching the 20 stomach wall portion as well as, time periods the stretching is applied, and when the stretching is applied.

In an embodiment, the apparatus further comprises an external control unit for controlling the volume filling device from the outside of the patient's body. The external control unit may comprise a wireless remote control adapted to control the 25 device from the outside of the patient's body. Alternatively, the external control unit may comprise a subcutaneously placed switch or reservoir adapted to control the device from the outside of the patient's body.

In an embodiment, the apparatus further comprises a sensor or sensing device adapted to be implanted in the patient body, wherein the implantable control unit is 30 adapted to control the device from the inside of the patient's body using information

from said a sensor or sensing device, adapted to sense, direct or indirect, the food intake of the patient.

In accordance with another aspect of the present invention, there is provided an apparatus for treating obesity of an obese patient having a stomach with a food cavity, the apparatus comprising at least one volume filling device adapted to be at least substantially invaginated by a stomach wall portion of the patient and having an outer surface that includes a biocompatible material, wherein the volume filling device is adapted to be placed inside the stomach with the outer surface of the volume filling device resting against the inside of the stomach wall, such that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device, the volume filling device having a maximum circumference of at least 30 millimeters.

In a preferred embodiment, the system comprises at least one switch implantable in the patient for manually and non-invasively controlling the apparatus.

15 In another preferred embodiment, the system comprises a wireless remote control for non-invasively controlling the apparatus.

In a preferred embodiment, the system comprises a hydraulic operation device for operating the apparatus.

20 In one embodiment, the system comprises comprising a motor or a pump for operating the apparatus.

BRIEF DESCRIPTION OF THE DRAWINGS

The present invention will now be described in more detail by way of non-limiting examples and with reference to the accompanying drawings, in which:

Fig. 1 is an overall view of a patient showing the outlines of the stomach,

25 Fig. 2a is a view of a first embodiment of an apparatus for treating obesity implanted in a human patient,

Fig. 2b is a sectional view taken along line IIb – IIb of Fig. 2a,

Figs. 3a-k, 3m, 3n, 3p show different shapes and features of a volume filling device comprised in an apparatus according to the invention,

Figs. 4a-d show a deflated inflatable volume filling device comprised in an apparatus according to the invention and an instrument for placing the volume filling device,

5 Figs. 5a-i illustrate different steps of invaginating the inflatable device of Fig. 4a on the outside of a stomach wall of a patient,

Figs. 6-8 show alternative embodiments wherein the volume filling device is adapted to be non-invasively adjustable postoperatively,

10 Figs. 9 and 10 show embodiments wherein the volume filling device is adapted to be invaginated in the fundus region of the patient's stomach,

Fig. 11 shows an embodiment wherein the volume filling device is also adapted to treat reflux,

15 Fig. 12 shows an embodiment wherein the volume filling device adapted also for treating reflux is combined with stretching devices for stretching part of the stomach fundus wall,

Figs. 13-16 show alternative embodiments wherein a combination of a volume filling device and a stretching device is used,

Figs. 17a and 17b show an embodiment wherein the volume filling device is provided on the inside of the stomach wall,

20 Figs. 18a-h illustrate different steps of invaginating the inflatable device of Fig. 4a on the inside of a stomach wall of a patient,

Figs. 19a-j illustrate different steps of invaginating the inflatable device of Fig. 4a on the inside of a stomach wall of a patient,

25 Figs. 20a-f illustrate different steps of invaginating the inflatable device of Fig. 4a on the inside of a stomach wall of a patient,

Figs. 21a and 21b

Fig. 22 is an overall view of a patient with an implanted apparatus for treating obesity,

Figs. 23-41 show various ways of powering an apparatus for treating obesity

5 implanted in a human patient,

Figs. 42-47 show various ways of arranging hydraulic or pneumatic powering of an apparatus for treating obesity implanted in a human patient,

Fig. 48 illustrate the invagination of a plurality volume filling devices, and

Figs. 49a and 49b illustrate an abdominal method.

10 DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

Preferred embodiments of the invention will now be described in detail with reference to the drawing figures.

Fig. 1 shows a human patient 1, who is being treated for obesity. A volume filling device 10 is provided so that it reduces the inner volume of the stomach 12 – the food cavity of the stomach, thereby affecting the patient's appetite. The function and the operation of this volume filling device will be described and explained in detail in the following description.

Figs. 2a and 2b show in detail a first embodiment of an apparatus to treat obesity according to the invention, wherein Fig. 2a show a side view of the stomach while

20 Fig. 2b is a sectional view taken along line IIb – IIb of Fig 2a. The apparatus comprises a volume filling device 10 implanted in a human patient. More specifically, in the embodiment of Fig. 2a the volume filling device 10 is invaginated in the wall 12a of the patient's stomach 12 on the outside of the stomach wall. The body of the volume filling device 10 is elongated and shaped to rest against the wall 12a of the stomach 12 and further has an outer surface suitable to rest against this wall.

By invaginating the volume filling device 10 in part of the stomach wall, the size of the food cavity, generally designated 12b in Fig. 2b, will be reduced, resulting in a more rapid feeling of satiety after food intake.

5 The volume filling device 10 preferably comprises an elastic material, such as silicone. In this way, the volume filling device can adapt to the movements of the stomach, the degree of food intake etc.

By providing the volume filling device from a bio-compatible material, the risk of the patient's body rejecting the implant is to a very large extent reduced.

10 The volume filling device 10 can be fixed to the wall 12a of the stomach 12 in a number of different ways. In the embodiment shown in Fig. 2b, the volume filling device 10 is invaginated in the stomach wall 12a. After invagination, a number of stomach-to-stomach sutures or staples 14 are applied to keep the invagination in the short term. This allows growth of human tissue, keeping the invagination in the long term.

15 The volume filling device 10 preferably has an essentially round shape to not damage the stomach wall. An example thereof is shown in Fig. 3a, wherein the volume filling device is essentially egg-shaped. In another preferred embodiment, the volume filling device is slightly bent, such as the embodiment shown in Fig. 3b. However, since the stomach wall is strong many different shapes, forms, and dimensions may be used. In 20 one embodiment, the volume filling device has a diameter of about 40 millimeters and a length of about 120 millimeters, resulting in a volume that is about half the volume of the patient's stomach. However, it is preferred that the maximum circumference of the volume filling device is at least 30 millimeters, more preferably at least 50 millimeters, and even more preferably at least 80 millimeters.

25 It is not necessary that the volume filling device is elongated. In the embodiment shown in Fig. 3c, the volume filling device 10 is essentially spherical or ball-shaped. In order to fill out the stomach, two or more such volume filling devices may be combined to achieve the desired decrease of the food cavity of the patient's stomach.

It has been mentioned that the volume filling device is secured by the stomach-to-stomach sutures or staples. In order to further improve the fixation, the volume filling device may be provided with a waist portion having smaller diameter than the maximum diameter of the volume filling device. Such volume filling device having a 5 waist portion 10a is shown in Fig. 3d.

The volume filling device 10 may consist of at least two interconnectable portions so that each portion is easier to insert into the stomach and further through a hole in the stomach wall. Thus, Fig. 3e shows a volume filling device comprising two more or less spherical sub-parts 10b, 10c interconnected by a portion with which preferably 10 has smaller diameter. The portion with smaller diameter may comprise an interconnection means with a reversible function allowing subsequent disconnection of the two interconnected sub-parts 10b, 10c. Such means may comprise a bayonet socket, a screw connection or the like, designated 10d in the figure. Alternatively, the portion with smaller diameter may comprise a fixed interconnection, such as resilient 15 locking hooks provided on one of the sub-parts 10b, 10c and engaging the rim of a hole provided in the other one of the sub-parts 10b, 10c.

The configuration of the volume filling device 10 is not limited to one waist portion 10a. Thus, in Fig. 3f a volume filling device with two waist portions is shown.

In order to facilitate positioning of the volume filling device, an attachment means in 20 the form of a handle or the like may be provided on the outer surface of the volume filling device. One example thereof is shown in Fig. 3g, wherein also a detail view of a handle 10e is shown. In a preferred embodiment, the attachment means is provided at an end portion of the volume filling device 10. In order to avoid protruding portion 25 on the surface of the volume filling device 10, the handle 10e is provided flush with the outer surface of the volume filling device 10 and a recess 10f is arranged to allow a gripping tool or instrument (not shown in Fig. 3g) to achieve firm gripping around the handle 10e.

The volume filling device may comprise a tube for filling or emptying the volume filling device of a fluid or gel. By injecting fluid or gel into the volume filling device 30 10, the volume filling device is inflated to an inflated state, as will be described

below. The size of the volume filling device can also be adjusted by moving fluid or gel therefrom to a different reservoir.

A volume filling device 10 adapted for this is shown in Fig. 3h. A tube 10g is fixedly attached to the volume filling device. This tube can be attached to a suitable

5 instrument (not shown) or an injection port, which will be explained in detail below.

Instead of having a fixedly attached tube, the volume filling device 10 may comprise an inlet port 10h adapted for connection of a separate tube (not shown in this figure).

It is important that the implanted volume filling device is firmly kept in place in the stomach wall in which it is invaginated. To this end, the volume filling device can be

10 provided with one or more through holes adapted for receiving sutures or staples used for fixation of the invagination. Such an embodiment is shown in Fig. 3j, where the volume filling device 10 is provided with a row of holes 10i provided on a protruding flange-like protrusion on the volume filling device. In this embodiment, the row of holes extend along the longitudinal axis of the volume filling device.

15 Fig. 3k illustrates how sutures 14 are provided so that they run through the stomach wall 12a and through the holes 10i. In this way, the volume filling device is fixed in place in the pouch created from the stomach wall and will thus be prevented from sliding.

20 Although a plurality of holes is illustrated in the Fig. 3j, it will be appreciated that one single hole is sufficient to obtain improved fixation of the volume filling device 10.

Fig. 3m illustrates a volume filling device provided with an inlet port 10h. The volume filling device is invaginated in the stomach wall and the inlet port 10h is available for connection to a tube or the like from the abdominal area of the patient.

25 Fig. 3n illustrates an invaginated volume filling device wherein, instead of an inlet port, a fixed tube 10g extends into the abdominal area of the patient.

Fig. 3p is a figure similar to Fig. 3m but also illustrating tunneling of a connection tube 10g in the stomach wall between the inlet port 10h and the volume filling device 10.

It has been shown that the shape of the volume filling device can take many different

5 forms. It will be appreciated that also the material of the volume filling device can vary. It is preferred that the volume filling device is provided with a coating, such as a Parylene, polytetrafluoroethylene (PTFE), or polyurethane coating, or a combination of such coatings, i.e., a multi-layer coating. This coating or multi-layer coating improves the properties of the volume filling device, such as its resistance to
10 wear.

In one embodiment, the volume filling device comprises an inflatable device expandable to an expanded state. In this case, the inflatable device is provided with an inlet port for a fluid and is adapted to be connected to a gastroscopic instrument. This embodiment will now be described in detail with reference to Figs. 4a-4d.

15 An inflatable volume filling device in its non-expanded state is shown in Fig. 4a. It is essentially a balloon-like, deflated device 10 having an inlet port 10h. In this state, the inflatable device has a diameter of a few millimeters at the most, allowing it to be inserted into the stomach through the esophagus of the patient by means of a gastroscopic, tube-like instrument 600, depicted in figure 4b. The instrument
20 comprises an outer sleeve 600a and an inner sleeve 600b which can be displaced longitudinally relatively to the outer sleeve. The inner sleeve is provided with a cutter in the form of a cutting edge 615 at the distal end thereof. This cutting edge can be used for cutting a hole in the stomach wall, as will be explained in detail in the following.

25 When the instrument reaches a stomach wall, see Fig. 4c, the inner sleeve is brought forward from its position in the outer sleeve and into contact with the stomach wall 12a. The cutting edge 615 of the inner sleeve then cuts a hole in the stomach wall so as to allow subsequent insertion of the volume filling device 10 into and through this hole, see Fig. 4d. In order to push the volume filling device through the hole, a piston
30 602 may be provided in the instrument. Thus, the instrument further comprises a

piston 602 adapted for pushing a deflated volume filling device 10 out from a position in the inner sleeve, this position being shown in Fig. 4b, to a position outside of the inner sleeve, this being shown in Fig. 4d.

5 In order to protect the deflated volume filling device 10 from the cutting edge 615 of the inner sleeve, a further protective sleeve (not shown) can be provided around the volume filling device.

An intraluminar method of invaginating a volume filling device 10 on the outside of the stomach wall 12a will now be described with reference to Figs. 5a-i. Initially, an 10 instrument 600, preferably a gastroscopic instrument, is inserted into the mouth of the patient, see Fig. 5a. The instrument comprises an injection device 601, 602 for injecting either fluid or a device into the stomach of the patient. The instrument 600 further comprises a control unit 606 adapted for controlling the operation of the instrument. To this end, the control unit 606 comprises one or more steering devices, in the embodiment shown in the figure in the form of two joysticks 603 and two 15 control buttons 604. A display 605 is provided for displaying the image provided by an optical device for viewing inside the stomach, such as a camera (not shown) arranged at the outer end of the elongated member 607, see Figs. 5e-i. The camera, which may comprise connecting electrical wires extending along the elongated member, may be assisted by a light source (not shown) placed distally on the 20 elongated member for illuminating the inside of the stomach. The optical device may also comprise optical fibers placed along the elongated member and leading out from the patient's body for external viewing of the inside of the stomach.

The instrument is further inserted into the esophagus and into the stomach of the patient, see Fig. 5b. By means of the instrument 600, a hole 12b is created in the wall 25 of the stomach 12, . To this end, the instrument is provided with one or more cutters 615 at the distal end thereof, for example in the way described above with reference to Figs. 4a-d. These cutters can of course be designed in different ways, such as a toothed drum cutter rotating about the center axis of the tube-like instrument.

After cutting a hole in the stomach wall, the distal end of the instrument 600 is 30 inserted into and through the hole 12b so that it ends up outside the stomach wall

12a. This is shown in Fig. 5c, showing a side view of the stomach 12, and Fig. 5d, which is a sectional view through the stomach of Fig. 5c taken along the lines Vd – Vd.

5 The instrument 600 is adapted to create a “pocket” or “pouch” on the outside of the stomach 12 around the hole 12b in the stomach wall. Such an instrument and the method of providing the pouch will now be described.

10 Figs. 5e-i show a gastroscopic or laparoscopic instrument for invaginating a volume filling device 10 in the stomach wall 12a of the patient by creating a pouch of stomach wall 12a material in which the volume filling device is placed. The instrument, generally designated 600, and which may comprise the features described above with reference to Figs. 4a-d, comprises an elongated member 607 having a proximal end and a distal end, the elongated member 607 having a diameter less than that of the patient’s esophagus and being flexible such as to allow introduction of the flexible elongated member 607 with its distal end first through the 15 patient’s throat, esophagus and into the stomach 12 to the stomach wall 12a.

20 The stomach penetration device or cutter 615 is provided on the elongated member 607 at the distal end thereof for penetrating the stomach wall 12a so as to create a hole in the stomach wall 12a, to allow introduction of the elongated member 607 through the hole. The stomach penetration device 615 could be adapted to be operable for retracting said stomach penetration device 615 after the stomach fundus wall 12a has been penetrated, for not further damaging tissue within the body. The instrument further comprises a special holding device 609 provided on the elongated member 607 on the proximal side to the penetration device 615.

25 The elongated member further comprises an expandable member 611 which is adapted to be expanded after the elongated member has penetrated the stomach wall 12a and thereby assist in the creation of a cavity or pouch adapted to hold the volume filling device 610. The expandable member 611 may comprise an inflatable circular balloon provided circumferentially around the distal end portion of the flexible elongated member 607.

The method steps when invaginating the volume filling device will now be described in detail. After the instrument 600 has been inserted into the stomach 12, the stomach penetration device 615 is placed into contact with the stomach wall 12a, see Fig. 5e. The stomach penetration device or cutter 615 is then brought to create the hole 12b in the stomach wall, whereafter at least the expandable member 611 is brought through the hole 12b in the stomach wall. The special holding device 609 is in this step brought to a holding state wherein it expands radially so as to form an essentially circular abutment surface to the stomach wall 12a, see Fig. 5f. In this way, the insertion of the stomach penetration device 615 and the expandable member 611 through the hole 12a in the stomach wall is limited to the position shown in Fig. 5f.

The expandable member 611 is then expanded. In the case the expandable member comprises a balloon or the like, air or other fluid is injected into it.

The part of the elongated member 607 comprising the expandable member 611 is then retracted in the proximal direction, as indicated by the arrow in Fig. 5g, thereby pulling the stomach wall 612 into a basket or cup like structure created by the special holding device 609.

A suturing or stapling device 608 is further provided, either as a device connected to the elongated member 607 or as a separate instrument. The suturing or stapling member comprises a suturing or stapling end 613 which is adapted to close the cavity or pouch by means of stomach to stomach sutures or staples 14.

In a further step, illustrated in Fig. 5h, an inflatable volume filling device 10 is placed in its deflated state in the cup like structure. The volume filling device 10 is then inflated to its inflated or expanded state, see Fig. 5i. This inflation of the volume filling device 10 can be accomplished by injecting a fluid or a gel into the deflated volume filling device. It can also be accomplished by injecting a material which is allowed to cure, thereby forming a solid device 10. Thus, the volume filling device 10 shown in Figs. 5h and 5i can illustrate either a balloon-like device which is subsequently filled with fluid or gel or alternatively a material which is simply injected into the cup like structure formed by the stomach wall 12a.

The fluid which is used to fill the volume filling device 10 could be any suitable fluid suitable to fill the inflatable device 10, such as a salt solution. In another embodiment, when this fluid is a fluid which is adapted to be transformed into solid state, the fluid could be liquid polyurethane.

- 5 In order to minimize or entirely eliminate leakage, the fluid is iso-tonic, i.e., it has the same osmolarity as human body fluids. Another way of preventing diffusion is to provide a fluid which comprises large molecules, such as iodine molecules.

The stomach-to-stomach sutures or staples are preferably provided with fixation portions exhibiting a structure, such as a net like structure, adapted to be in contact

- 10 with the stomach wall to promote growth in of human tissue to secure the long term placement of the volume filling device attached to the stomach wall.

After the inflatable device 10 has been inflated, partly or fully, the inlet port 10b (not shown in Figs. 5h and 5i) of the volume filling device 10, is sealed and the instrument 600 is retracted from the hole 12b, which is subsequently closed in some 15 suitable way, such as by means of the instrument 600. The instrument is then removed from the stomach 600 and the inflatable device 10 in its inflated or expanded state is invaginated by a stomach wall portion of the patient on the outside of the stomach wall. This reduces the inner volume of the stomach, thereby affecting the patent's appetite.

- 20 During one or more of the above described steps, the stomach may be inflated with gas, preferably by means of the gastroscopic instrument.

The volume filling device 10 described above with reference to Figs. 5a-i has been described as an inflatable volume filling device. It will be appreciated that is also can be an elastic volume filling device with an elasticity allowing compression so as to 25 be inserted into a gastroscopic instrument and which expands to an expanded state after leaving the instrument.

An alternative embodiment of an apparatus for treating obesity will now be described with reference to Figs. 6a and 6b, showing a sectional view of a stomach in which a volume filling device is invaginated in the stomach wall on the outside thereof

together with a system for regulating the size of the volume filling device. The volume filling device is an inflatable device as described above with reference to Figs. 5a-h and thus comprises a fluid. The inflatable device 10 thus forms a fluid chamber, in which fluid is allowed to flow. The inflatable device thereby forms an 5 expandable chamber that can change the volume it occupies in the stomach wall, thereby forming a hydraulically or pneumatically regulated inflatable device.

In Fig. 6a, an injection port 16 for fluids is connected to the inflatable volume filling device 10 by means of a conduit 18 in the form of a tube. The inflatable device 10 is thereby adapted to be regulated, preferably non-invasively, by moving liquid or air 10 from the injection port 16 to the chamber formed by the inflatable device. By using a hypodermic needle or the like, the amount of fluid in the inflatable device 10 can thus be adjusted, thereby adjusting the size of the adjustable device. The injection port 16 can also be used simply for refilling the volume filling device 10.

The regulation reservoir 17 can be regulated in several ways. In an alternative 15 embodiment, the regulation reservoir 17 is regulated by manually pressing a regulation reservoir. In other words, the regulation reservoir is regulated by moving a wall of the reservoir. It is then preferred that the regulation reservoir is placed subcutaneously and non-invasive regulation is thereby achieved.

A similar embodiment is shown in Fig. 6b. However, in this embodiment the 20 injection port 16 has been replaced by an adjustable regulation reservoir 17 in fluid connecting with the volume filling device 10 via a tube 18. When the regulation reservoir 17 is pressed, the volume thereof decreases and hydraulic fluid is moved from the reservoir to the chamber formed by the inflatable device 10 via the conduit or tube 18, enlarging or expanding the inflatable device 10. In this way, the volume 25 filling device is non-invasively adjustable postoperatively.

It will be appreciated that instead of hydraulic operation, pneumatic operation can be used, wherein air instead of hydraulic fluid is moved between the regulation reservoir and the chamber formed by the inflatable device 10. Preferable the regulation reservoir has a locking position to keep it in the desired position. If the patient

compresses the reservoir it preferably stays compressed and releases after pressing again.

Any kind of hydraulic solution may be used for the inflatable device. The hydraulic solution may be driven by both mechanically and be powered with any motor or 5 pump as well as manually.

In another embodiment, shown in Fig. 7, a motor 40 is adapted to move a wall of the regulation reservoir 17. The powered regulation reservoir 17 is then preferably placed in the abdomen of the patient. In this embodiment, a wireless external remote control unit forming part of the external energy transmission device 34 can be

10 provided to perform non-invasive regulation of the motor via an energy transforming device 30, which is adapted to supply an energy consuming operation device, in the present example the motor 40, with energy via a power supply line 32.

The remote control may comprise a wireless energy transmitter, whereby the non-invasive regulation is performed by the energy transmitter. When the regulation is 15 performed by means of a remote control an internal power source for powering the regulating device is provided. The internal energy source can for example be a chargeable implanted battery or a capacitor or a device for receiving wireless energy transmitted from outside the body of the patient. Different ways of regulating the inflatable device 10 will be described below with reference to Figs. 22-41.

20 In yet an alternative embodiment, shown in Fig. 8, the apparatus for treating obesity comprises a pump 44, wherein the reservoir is regulated by the pump 44 pumping fluid or air from the reservoir to the chamber formed by the inflatable device.

Different configurations of this pump will be described below with reference to Figs. 22-41.

25 Yet an alternative embodiment of an apparatus for treating obesity will now be described with reference to Fig. 9, which shows a stomach 12 of a patient who is treated for obesity. This embodiment is similar to the one described above with reference to Fig. 7 and the apparatus comprises a volume filling device in the form of an inflatable device 10 which is invaginated in the wall 12a of the patient's stomach

12. However, in this case the invagination has been performed in the fundus, i.e., the upper portion of the stomach, where the number of receptors in the stomach wall is large, and the inflatable device functions as a stretching device for part of the stomach fundus wall.

5 A regulation reservoir 17 for fluids is connected to the inflatable device by means of a conduit 18 in the form of a tube. The inflatable device 10 is thereby adapted to be regulated, preferably non-invasively, by moving liquid or air from the regulation reservoir 17 to the chamber formed by the inflatable device 10. The regulation of the inflatable device 10 preferably comprises a reversed servo, i.e., a small volume is actuated for example by the patient's finger and this small volume is in connection with a larger volume, i.e., the regulation reservoir 17.

10 Thus, the inflatable device 10 is placed outside the stomach wall and is adapted to stretch a part of the stomach fundus wall, thereby affecting the patient's appetite. By enlarging the size of the stretching device, the stomach fundus wall 12a surrounding the inflatable stretching device 10 is stretched since the circumference of the inflatable stretching device 10 is increased. By this stretching, the receptors in the stomach wall indicate that the stomach is full, thereby creating a feeling of satiety to the patient. Correspondingly, when the stretching device 10 is contracted, the receptors indicate that the stomach is not full, thereby returning the feeling of hunger.

15 20 It will be appreciated that this embodiment combines the effects of both reducing the volume of the stomach food cavity and stretching part of the stomach wall, thereby increasing the treatment effect.

25 The expansion and contraction of the stretching device 10 can be performed under direct control of the patient. Alternatively, the expansion and contraction can be performed according to a pre-programmed schedule.

30 In a preferred embodiment, shown in Fig. 10, a sensor 19 is provided at a suitable position, such as at the esophagus. The volume filling device 10 in the form of the inflatable stretching device is similar to the one shown in Fig. 9. By providing one or more sensors, the apparatus for treating obesity can be automated in that the size of the volume filling device 10 in the form of the inflatable stretching device is adjusted

depending on the amount of food entering the food cavity of the stomach. The fluid is thereby moved between the inflatable volume filling device 10 and a fluid reservoir 15.

The apparatus for treating obesity can have the additional functionality of treating

5 reflux. An embodiment having this function is shown in Fig. 11, wherein the volume filling device 10 is invaginated in the stomach wall close to and at least partially above the patient's cardia 26 when the patient is in a standing position and is fixed to a position above the cardia area 26 by a fixation, such as sutures or staples 14a. For example a direct or indirect fixation to the diaphragm muscle or associated muscles 10 may be provided. As an alternative a direct or indirect fixation to the esophagus above and close to the angle of His can be provided. In this alternative embodiment, the volume filling device 10 rests in a position against stomach wall of the fundus when implanted and which also fills a volume above the cardia area 26 between the cardia and the diaphragm muscle so that the cardia is prevented from slipping up into 15 the thorax cavity, whereby reflux disease is prevented.

Such a volume filling device 10 may be used for keeping electronics and/or an energy source and/or hydraulic fluid. Hydraulic fluid from that device may be distributed to several smaller inflatable device areas to vary the stretching area from time to time avoiding any possible more permanent stretching effect of the stomach 20 wall. Even mechanically several stretching areas may be used.

In an alternative embodiment, which is shown in Fig. 12, the volume of an inflatable volume filling device 10 may be in fluid connection with one or more preferably smaller inflatable devices or chambers 50. These chambers are adapted to communicate with fluid or air being moved between the chambers.

25 Thus, the large chamber 10 is adapted to, with its main volume to be a volume filling device for reducing the size of the food cavity and for treating reflux disease and the one or several small chambers are adapted to function as the inflatable devices to treat obesity, wherein the main chamber is adapted to communicate with fluid or air to the small chambers causing a stretching effect in the stomach wall thereby further 30 treating obesity.

In Figs. 13-16, different embodiments embodying a combination of a volume filling device invaginated in the central or lower portion of the stomach and a stretching device invaginated in the upper portion or fundus of the patient's stomach. Thus, in Fig. 13 there is shown an adjustable volume filling device 10, which is invaginated in the stomach wall of a patient's stomach 12. Additionally, an adjustable stretching device 50 with the previously described function is invaginated in the stomach fundus wall of the patient. It is preferred that the volume filling device 10 is substantially larger than the stretching device 50.

The volume filling device 10 and the stretching device 50 are in fluid communication with each other via a fluid communication device comprising a first fluid tube 52, in which a pump 54 is provided. The pump 54 is under the control from an energy transforming device 30, which is adapted to supply the pump 50 with energy via a power supply line 56. The energy transforming device 30 is also connected to a sensor 19 provided in the esophagus of the patient so that food intake can be detected.

The volume filling device 10 and the stretching device 50 are also in fluid communication with each other via a second fluid tube 58, which preferably has a smaller cross-sectional area than the first fluid tube 52.

The operation of this arrangement is as follows. The volume filling device 10 functions as in the above described embodiments, i.e., it reduces the size of the food cavity of the patient's stomach 12. Additionally, when the stretching device 50 is enlarged by pumping fluid from the volume filling device 10 and to the stretching device 50 by means of the pump 54, the stomach fundus wall is stretched, creating a feeling of satiety for the patient. Thus, for example when food intake is detected by means of the sensor 19, fluid is automatically pumped into the stretching device 50 to increase the feeling of satiety and thereby limit the food intake.

When fluid has been injected into the stretching device 50, the internal pressure therein is higher than the internal pressure in the volume filling device 10. This difference in pressure will create a flow of fluid in the second, preferably narrower tube 58 from the stretching device 50 to the volume filling device 10. The flow rate

will be determined by among other things the difference in pressure and the cross-sectional area of the second tube 58. It is preferred that the second tube is so dimensioned, that the pressures in the volume filling device 10 and the stretching device 50 will return to equilibrium after 3 hours after fluid has been injected into the 5 stretching device 50 to create the feeling of satiety.

In this embodiment, the function of the second tube 58 is to allow fluid to return from the stretching device 50 to the volume filling device 10. It will be appreciated that this function also can be performed by the pump 54 in the first tube 52 and that the second tube 58 then can be omitted.

10 Fig. 14 illustrates an embodiment similar to the one illustrated in Fig. 13. Thus, there is provided an adjustable volume filling device 10, which is invaginated in the stomach wall of a patient's stomach 12. Additionally, an adjustable stretching device 50 with the previously described function is invaginated in the stomach fundus wall of the patient. It is preferred that the volume filling device 10 is substantially larger 15 than the stretching device 50.

The volume filling device 10 and the stretching device 50 are in fluid communication with each other via a first fluid tube 52, and a second fluid tube, which preferably has a smaller cross-sectional area than the first tube. However, instead of a pump, there is provided a non-return valve 60 in the first fluid tube 52 instead of an energized 20 pump. This non-return valve 60 allows fluid to flow in the direction from the volume filling device 10 and to the stretching device 10 but not vice versa. This means that this embodiment may be entirely non-energized. Instead, it operates according to the following principles.

When the food cavity of the stomach 12 is essentially empty, there is a state of 25 equilibrium between the internal pressure of the volume filling device 10 and the stretching device 50. In this state, the stretching device is in a non-stretch state, i.e., it does not stretch a part of the stomach fundus wall and thus does not create a feeling of satiety.

When the patient starts to eat, food will enter the food cavity of the stomach 12. This will create increased pressure on the stomach wall in which the volume filling device 10 is invaginated and the internal pressure therein will increase. Also, the stomach wall muscles will begin to process the food in the food cavity by contraction, which 5 also contributes to an increased internal pressure in the volume filling device 10.

Since the internal pressure in the stretching device 50 will remain essentially unchanged, because it is located in the upper part of the stomach 12 where no food is exerting a pressure on the stomach wall, a fluid flow will be created through the first and second fluid tubes 52, 58 in the direction from the volume filling device 10 and 10 to the stretching device 50. This in turn will increase the volume of the stretching device 50, which, by stretching the stomach fundus wall, will provide a feeling of satiety to the patient.

A fluid flow from the stretching device 50 to the volume filling device 10 through the second tube 58 will return the pressure of these devices to equilibrium as 15 described above with reference to Fig. 13.

Fig. 15 illustrates an embodiment, which is similar to the one shown in Fig. 14 but with the addition of an injection port 16, which is used for refilling the fluid system comprising the volume filling device 10 and the stretching device 50 or alternatively for actively adjusting the size thereof.

20 Similarly, Fig. 16 illustrates an embodiment wherein the stretching device 50 can be actively regulated by manually pressing an adjustment reservoir which is provided subcutaneously below the patient's skin, similar to the embodiment shown in Fig. 9. Thus, a regulation reservoir 17 for fluids is connected to the inflatable device by means of a conduit 18 in the form of a tube. The stretching device 50 is thereby 25 adapted to be regulated, non-invasively, by moving liquid or air from the regulation reservoir 17 to the chamber formed by the inflatable device. The regulation of the stretching device 50 preferably comprises a reversed servo, i.e., a small volume is actuated for example by the patient's finger and this small volume is in connection with a larger volume.

An alternative placement of the volume filling device 10 is shown in Figs. 17a and 17b, wherein Fig. 17b shows a sectional view through the stomach shown in Fig. 17a along the line XVIIb - XVIIb. There, the volume filling device 10 is adapted to be placed inside the wall of the stomach 12, such as via a gastroscope or similar 5 intraluminar instrument, and resting against the inside of the stomach wall 12a. The inflatable device can be kept invaginated by means of sutures or staples 14, like in the embodiment of Figs. 2a and 2b. In this embodiment, no hole is required in the stomach wall. Instead, a method of providing the volume filling device 10 can comprise the following steps, which will be explained with reference to Figs. 18a-i 10 showing an invagination instrument

The invagination instrument, generally designated 630, comprises an elongated tube member 632 similar to the elongated member 607 described above with reference to Figs. 5a-i. Thus, it can be connected to a control unit 606, see Fig. 5a. The invagination instrument 630 further comprises a perforated suction portion 634, 15 which preferably is elongated. The suction portion 634 exhibits a plurality of small holes 636, into which air will be sucked by providing suction in the tube member 632. This suction effect will be used to create a “pocket” or “pouch” in a part of a stomach wall, generally designated 12a.

In other words, when the tip of the suction portion 634 is pressed against the stomach 20 wall 12a, see Fig. 18a, a small recess will be formed therein. When the suction portion 634 is further pressed against the stomach wall 12a, see Fig. 18b, a larger recess will be formed. The part of the stomach wall 12a that forms the recess will, due to the suction effect, adhere to the suction portion 634 of the invagination instrument 630. As the suction portion 634 is further pressed into the stomach wall 25 12a, see Fig. 18c, a deeper recess will be formed until the entire suction portion 634 is embedded in the recess, see Fig. 18d.

The rim of the recess will at this stage be fixated by means of fixation elements 638 and the suction portion be removed from the instrument, see Fig. 18e. A compressed elastic volume filling device 10 will subsequently be inserted into the recess, see Fig. 30 18f, for example in the way described above with reference to Fig. 4d. This compressed volume filling device is then expanded to its final shape, see Fig. 18g,

where after the pouch is sealed by suturing or stapling by means of the fixations elements, see Fig. 18h.

All the alternatives described above with reference to Figs. 2-16 are also applicable to the embodiment described with reference to Figs. 17 and 18, i.e., to the 5 embodiment where the volume filling device is invaginated on the inside of the stomach wall.

Figs. 19a-j show an instrument for use in a method of engaging a volume filling device 10 to the stomach wall 12 of a patient. The instrument is adapted to be inserted through a narrow tube shaped object such as a gastroscope, used in an 10 intraluminar procedure, or a laparoscopic trocar used in a laparoscopic procedure. The instrument comprises an elongated member 650 which is adapted to be flexible by means of a construction comprising multiple ring shaped members, however it is equally conceivable that said elongated member 650 is adapted to be flexible by means of said elongated member 650 being made of a flexible or adjustable material. 15 The elongated member 650 is inserted into the body and placed in proximity to the stomach wall 12 of the patient, from the outside or inside thereof. The elongated member 650 has a special holding device 651 adapted to hold the stomach by means of mechanical grabbing members or vacuum. The special holding device 651 comprises a first joint 652 and a second joint 653, which enable the special holding 20 device 651 be operable in relation to the elongated member 650 and thereby place the part of the holding device 651 comprising the mechanical grabbing members or vacuum elements into contact with the stomach wall 12 of the patient. Fig. 19b shows the special holding device 651 when placed in contact with the stomach wall 12 of the human patient, after which the special holding member 651 connects to the 25 stomach wall 12, for holding the stomach wall 12. Fig. 19c shows the instrument when the step of advancing a pushing rod 654 from the elongated member 650 is performed. The pushing rod 654 pushes the stomach wall 12 to create a cavity or pouch thereof. Fig. 19d shows the instrument turned 90° in relation to figs. 19a-c. This view shows the special holding members 651a,b operably attached to two sides 30 of the elongated member 650 and being in contact with the stomach wall 12, holding the stomach wall 12 as the pushing rod 654 pushes to create a cavity or pouch. When

the pushing rod 654 has pushed the stomach wall 12 to a desired position the special holding devices 651a,b moves towards the pushing rod 654 and thereby closes the cavity or pouch.

After the cavity or pouch has been created it needs to be sealed. Fig. 19f shows the

5 advancement of a suturing or stapling device 655 from the elongated member 650. The suturing or stapling device 655 is positioned in connection with the stomach wall after which the suturing or stapling device commences with the suturing or stapling of the stomach wall 12, creating a seal of stomach to stomach sutures or staples 14. The instrument is moved along the stomach wall 12 of the patient and thereby a
10 cavity or pouch is created and sealed using the instrument, as shown in fig. 19g and 19h. When a cavity or pouch or desired size has been created and sealed an inserting member 656 is advanced from the elongated member 650. The inserting member 656 is adapted to insert a volume filling device 10 being inflatable, as described earlier in this application. After the inserting member 656 has been positioned in the cavity or
15 pouch the volume filling device 10 is inserted through the inserting member 656 and into the cavity or pouch by means of a pressurized fluid or gas, or a mechanical advancement member pushing said inflatable volume filling device 10 into the cavity or pouch. The insertion member then inflates the inflatable volume filling device with a fluid or gas and seals of the final section of the pouch using stomach to
20 stomach sutures or staples 14. The embodiment described explains the process of inserting an inflatable volume filling device, however it is equally conceivable that the volume filling device 10 is expandable by means of the volume filling device 10 being made of an elastic material.

Figs. 20a-f show an instrument for use in a method of engaging a volume filling

25 device 10 to the stomach wall 12 of a patient. The instrument is adapted to be inserted through a narrow tube shaped object such as a gastroscope, used in an intraluminar procedure, or a laparoscopic trocar used in a laparoscopic procedure. The instrument comprises an elongated member 660 which is adapted to be flexible by means of a construction comprising multiple ring shaped members, however it is
30 equally conceivable that said elongated member 660 is adapted to be flexible by means of said elongated member 660 being made of a flexible or adjustable material.

The elongated member 660 is inserted into the body and placed in proximity to the stomach wall 12 of the patient, from the outside or inside thereof. The elongated member 660 has multiple special holding devices 661 adapted to hold the stomach by means of mechanical grabbing members or vacuum. The special holding devices 661 are locked in a position alongside the elongated member 660 by means of a locking ring 662. The special holding devices are made of a flexible material and pre-bent to expand into a funnel-shaped device when said locking ring 662 is removed. The special holding device in its funnel shaped expandable state is shown in fig. 20b. Fig. 20b further shows the special holding device 661 when placed in contact with the 5 stomach wall 12 of the human patient, after which the special holding member 661 connects to the stomach wall 12, for holding the stomach wall 12. Fig. 20c shows the instrument when the step of advancing a pushing rod 664 from the elongated member 660 is performed. The pushing rod 664 pushes the stomach wall 12 to create a cavity or pouch thereof. When the pushing rod 664 has pushed the stomach wall 12 to a 10 desired position the special holding devices 661 moves towards the pushing rod 664 and thereby closes the cavity or pouch.

After the cavity or pouch has been created it needs to be sealed. Fig. 20d shows the advancement of a suturing or stapling device 665 from the elongated member 660.

The suturing or stapling device 665 is positioned in connection with the stomach wall

12 after which the suturing or stapling device 665 commences with the suturing or 20 stapling of the stomach wall 12, creating a seal of stomach to stomach sutures or staples 14. Thereafter an inserting member 666 is advanced from the elongated member 660 and the special holding devices 661 are retracted. The inserting member 666 is adapted to insert a volume filling device 10 being inflatable, as described

25 earlier in this application. After the inserting member 666 has been positioned in the cavity or pouch the volume filling device 10 is inserted through the inserting member 666 and into the cavity or pouch by means of a pressurized fluid or gas, or a

mechanical advancement member pushing said inflatable volume filling device 10 into the cavity or pouch. The insertion member 656 then inflates the inflatable 30 volume filling device with a fluid or gas and seals of the final section of the pouch using stomach to stomach sutures or staples 14. The embodiment described explains the process of inserting an inflatable volume filling device 10, however it is equally

conceivable that the volume filling device 10 is expandable by means of the volume filling device 10 being made of an elastic material. Fig.20 f shows the volume filling device 10 as the volume filling device 10 is invaginated in the stomach wall 12, in a cavity or pouch sealed with stomach sutures or staples 14.

5 Fig. 21a shows an instrument used in a method of engaging the volume filling device according to any of the embodiments of the application to the stomach wall 12. The instrument comprises an elongated member 670 which is adapted to be flexible by means of a construction comprising multiple ring shaped members, however it is equally conceivable that said elongated member 670 is adapted to be flexible by 10 means of said elongated member 670 being made of a flexible or adjustable material. The elongated member 670 is inserted into the body and placed in proximity to the stomach wall 12 of the patient, from the inside thereof. A stomach penetrating member 672 is placed in the distal end of the elongated member 670, retractably fixated to a protective sleeve 673 adapted to protect the tissue of the body from the 15 sharp penetrating member 672 or cutter 672 after the cutting operation has been performed.

Fig. 21b shows the instrument comprising the elongated member 670 after the cutting operation has been performed and the stomach penetrating member or cutter 672 has been retracted into the protective sleeve 673. A guiding wire 671 is pushed 20 through the elongated member 670, through the hole made in the stomach wall 12 and out through the abdomen and placed on the inside of the patients skin, which is penetrated from the outside to enable the guiding wire 671 to exit the abdomen. The guiding wire 671 can then be used to guide a conduit 18 or a lead attached to the volume filling device 10 being placed in the stomach from the inside thereof. The 25 volume filling device 10 with the conduit 18 or electrical lead being a volume filling device 10 according to any of the embodiments of this application. The guiding of the conduit 18 or electrical lead enables the attachment of the conduit 18 or electrical lead to a control unit 42 placed subcutaneously in the patient from the outside of the abdomen.

30 Figs. 20a-f show an instrument for use in a method of engaging a volume filling device 10 to the stomach wall 12 of a patient. The instrument is adapted to be

inserted through a narrow tube shaped object such as a gastroscope, used in an intraluminar procedure, or a laparoscopic trocar used in a laparoscopic procedure. The instrument comprises an elongated member 660 which is adapted to be flexible by means of a construction comprising multiple ring shaped members, however it is equally conceivable that said elongated member 660 is adapted to be flexible by means of said elongated member 660 being made of a flexible or adjustable material. The elongated member 660 is inserted into the body and placed in proximity to the stomach wall 12 of the patient, from the outside or inside thereof. The elongated member 660 has multiple special holding devices 661 adapted to hold the stomach by means of mechanical grabbing members or vacuum. The special holding devices 661 are locked in a position alongside the elongated member 660 by means of a locking ring 662. The special holding devices are made of a flexible material and pre-bent to expand into a funnel-shaped device when said locking ring 662 is removed. The special holding device in its funnel shaped expandable state is shown in fig. 20b. Fig. 20b further shows the special holding device 661 when placed in contact with the stomach wall 12 of the human patient, after which the special holding member 661 connects to the stomach wall 12, for holding the stomach wall 12. Fig. 20c shows the instrument when the step of advancing a pushing rod 664 from the elongated member 660 is performed. The pushing rod 664 pushes the stomach wall 12 to create a cavity or pouch thereof. When the pushing rod 664 has pushed the stomach wall 12 to a desired position the special holding devices 661 moves towards the pushing rod 664 and thereby closes the cavity or pouch.

After the cavity or pouch has been created it needs to be sealed. Fig. 20d shows the advancement of a suturing or stapling device 665 from the elongated member 660. The suturing or stapling device 665 is positioned in connection with the stomach wall 12 after which the suturing or stapling device 665 commences with the suturing or stapling of the stomach wall 12, creating a seal of stomach to stomach sutures or staples 14. Thereafter an inserting member 666 is advanced from the elongated member 660 and the special holding devices 661 are retracted. The inserting member 666 is adapted to insert a volume filling device 10 being inflatable, as described earlier in this application. After the inserting member 666 has been positioned in the cavity or pouch the volume filling device 10 is inserted through the inserting member

666 and into the cavity or pouch by means of a pressurized fluid or gas, or a mechanical advancement member pushing said inflatable volume filling device 10 into the cavity or pouch. The insertion member 656 then inflates the inflatable volume filling device with a fluid or gas and seals off the final section of the pouch

5 using stomach to stomach sutures or staples 14. The embodiment described explains the process of inserting an inflatable volume filling device 10, however it is equally conceivable that the volume filling device 10 is expandable by means of the volume filling device 10 being made of an elastic material. Fig.20 f shows the volume filling device 10 as the volume filling device 10 is invaginated in the stomach wall 12, in a

10 cavity or pouch sealed with stomach to stomach sutures or staples 14.

Fig. 21a shows an instrument used in a method of engaging the volume filling device according to any of the embodiments of the application to the stomach wall 12. The instrument comprises an elongated member 670 which is adapted to be flexible by means of a construction comprising multiple ring shaped members, however it is equally conceivable that said elongated member 670 is adapted to be flexible by means of said elongated member 670 being made of a flexible or adjustable material. The elongated member 670 is inserted into the body and placed in proximity to the stomach wall 12 of the patient, from the inside thereof. A stomach penetrating member 672 is placed in the distal end of the elongated member 670, retractably fixated to a protective sleeve 673 adapted to protect the tissue of the body from the sharp penetrating member 672 or cutter 672 after the cutting operation has been performed.

Fig. 21b shows the instrument comprising the elongated member 670 after the cutting operation has been performed and the stomach penetrating member or cutter 672 has been retracted into the protective sleeve 673. A guiding wire 671 is pushed through the elongated member 670, through the hole made in the stomach wall 12 and out through the abdomen and placed on the inside of the patient's skin, which is penetrated from the outside to enable the guiding wire 671 to exit the abdomen. The guiding wire 671 can then be used to guide a conduit 18 or a lead attached to the volume filling device 10 being placed in the stomach from the inside thereof. The volume filling device 10 with the conduit 18 or electrical lead being a volume filling

device 10 according to any of the embodiments of this application. The guiding of the conduit 18 or electrical lead enables the attachment of the conduit 18 or electrical lead to a control unit 42 placed subcutaneously in the patient from the outside of the abdomen.

5 Fig. 22 illustrates a system for treating a disease comprising an apparatus 10 comprising a volume filling device of the present invention placed in the abdomen of a patient. An implanted energy-transforming device 1002 is adapted to supply energy consuming components of the apparatus with energy via a power supply line 1003. An external energy-transmission device 1004 for non-invasively energizing the 10 apparatus 10 transmits energy by at least one wireless energy signal. The implanted energy-transforming device 1002 transforms energy from the wireless energy signal into electric energy which is supplied via the power supply line 1003.

The implanted energy-transforming device 1002 may also comprise other components, such as: a coil for reception and/or transmission of signals and energy, 15 an antenna for reception and/or transmission of signals, a microcontroller, a charge control unit, optionally comprising an energy storage, such as a capacitor, one or more sensors, such as temperature sensor, pressure sensor, position sensor, motion sensor etc., a transceiver, a motor, optionally including a motor controller, a pump, and other parts for controlling the operation of a medical implant.

20 The wireless energy signal may include a wave signal selected from the following: a sound wave signal, an ultrasound wave signal, an electromagnetic wave signal, an infrared light signal, a visible light signal, an ultra violet light signal, a laser light signal, a micro wave signal, a radio wave signal, an x-ray radiation signal and a gamma radiation signal. Alternatively, the wireless energy signal may include an 25 electric or magnetic field, or a combined electric and magnetic field.

The wireless energy-transmission device 1004 may transmit a carrier signal for carrying the wireless energy signal. Such a carrier signal may include digital, analogue or a combination of digital and analogue signals. In this case, the wireless energy signal includes an analogue or a digital signal, or a combination of an 30 analogue and digital signal.

Generally speaking, the energy-transforming device 1002 is provided for transforming wireless energy of a first form transmitted by the energy-transmission device 1004 into energy of a second form, which typically is different from the energy of the first form. The implanted apparatus 10 is operable in response to the 5 energy of the second form. The energy-transforming device 1002 may directly power the apparatus with the second form energy, as the energy-transforming device 1002 transforms the first form energy transmitted by the energy-transmission device 1004 into the second form energy. The system may further include an implantable accumulator, wherein the second form energy is used at least partly to charge the 10 accumulator.

Alternatively, the wireless energy transmitted by the energy-transmission device 1004 may be used to directly power the apparatus, as the wireless energy is being transmitted by the energy-transmission device 1004. Where the system comprises an operation device for operating the apparatus, as will be described below, the wireless 15 energy transmitted by the energy-transmission device 1004 may be used to directly power the operation device to create kinetic energy for the operation of the apparatus.

The wireless energy of the first form may comprise sound waves and the energy-transforming device 1002 may include a piezo-electric element for transforming the 20 sound waves into electric energy. The energy of the second form may comprise electric energy in the form of a direct current or pulsating direct current, or a combination of a direct current and pulsating direct current, or an alternating current or a combination of a direct and alternating current. Normally, the apparatus comprises electric components that are energized with electrical energy. Other 25 implantable electric components of the system may be at least one voltage level guard or at least one constant current guard connected with the electric components of the apparatus.

Optionally, one of the energy of the first form and the energy of the second form may comprise magnetic energy, kinetic energy, sound energy, chemical energy, radiant 30 energy, electromagnetic energy, photo energy, nuclear energy or thermal energy.

Preferably, one of the energy of the first form and the energy of the second form is non-magnetic, non-kinetic, non-chemical, non-sonic, non-nuclear or non-thermal.

The energy-transmission device may be controlled from outside the patient's body to release electromagnetic wireless energy, and the released electromagnetic wireless

5 energy is used for operating the apparatus. Alternatively, the energy-transmission device is controlled from outside the patient's body to release non-magnetic wireless energy, and the released non-magnetic wireless energy is used for operating the apparatus.

The external energy-transmission device 1004 also includes a wireless remote

10 control having an external signal transmitter for transmitting a wireless control signal for non-invasively controlling the apparatus. The control signal is received by an implanted signal receiver which may be incorporated in the implanted energy-transforming device 1002 or be separate there from.

The wireless control signal may include a frequency, amplitude, or phase modulated signal or a combination thereof. Alternatively, the wireless control signal includes an analogue or a digital signal, or a combination of an analogue and digital signal.

15 Alternatively, the wireless control signal comprises an electric or magnetic field, or a combined electric and magnetic field.

The wireless remote control may transmit a carrier signal for carrying the wireless

20 control signal. Such a carrier signal may include digital, analogue or a combination of digital and analogue signals. Where the control signal includes an analogue or a digital signal, or a combination of an analogue and digital signal, the wireless remote control preferably transmits an electromagnetic carrier wave signal for carrying the digital or analogue control signals.

25 Fig. 23 illustrates the system of Fig. 22 in the form of a more generalized block

diagram showing the apparatus 10, the energy-transforming device 1002 powering the apparatus 10 via power supply line 1003, and the external energy-transmission device 1004, The patient's skin 1005, generally shown by a vertical line, separates

the interior of the patient to the right of the line from the exterior to the left of the line.

Fig. 24 shows an embodiment of the invention identical to that of Fig. 23, except that a reversing device in the form of an electric switch 1006 operable for example by 5 polarized energy also is implanted in the patient for reversing the apparatus 10.

When the switch is operated by polarized energy the wireless remote control of the external energy-transmission device 1004 transmits a wireless signal that carries polarized energy and the implanted energy-transforming device 1002 transforms the wireless polarized energy into a polarized current for operating the electric switch 10 1006. When the polarity of the current is shifted by the implanted energy-transforming device 1002 the electric switch 1006 reverses the function performed by the apparatus 10.

Fig. 25 shows an embodiment of the invention identical to that of Fig. 23, except that an operation device 1007 implanted in the patient for operating the apparatus 10 is 15 provided between the implanted energy-transforming device 1002 and the apparatus 10. This operation device can be in the form of a motor 1007, such as an electric servomotor. The motor 1007 is powered with energy from the implanted energy-transforming device 1002, as the remote control of the external energy-transmission device 1004 transmits a wireless signal to the receiver of the implanted energy-transforming device 1002.

Fig. 26 shows an embodiment of the invention identical to that of Fig. 23, except that it also comprises an operation device in the form of an assembly 1008 including a motor/pump unit 1009 and a fluid reservoir 1010 is implanted in the patient. In this case the apparatus 10 is hydraulically operated, i.e. hydraulic fluid is pumped by the 25 motor/pump unit 1009 from the fluid reservoir 1010 through a conduit 1011 to the apparatus 10 to operate the apparatus, and hydraulic fluid is pumped by the motor/pump unit 1009 back from the apparatus 10 to the fluid reservoir 1010 to return the apparatus to a starting position. The implanted energy-transforming device 1002 transforms wireless energy into a current, for example a polarized current, for 30 powering the motor/pump unit 1009 via an electric power supply line 1012.

Instead of a hydraulically operated apparatus 10, it is also envisaged that the operation device comprises a pneumatic operation device. In this case, the hydraulic fluid can be pressurized air to be used for regulation and the fluid reservoir is replaced by an air chamber.

- 5 In all of these embodiments the energy-transforming device 1002 may include a rechargeable accumulator like a battery or a capacitor to be charged by the wireless energy and supplies energy for any energy consuming part of the system.

As an alternative, the wireless remote control described above may be replaced by manual control of any implanted part to make contact with by the patient's hand

- 10 most likely indirect, for example a press button placed under the skin.

Fig. 27 shows an embodiment of the invention comprising the external energy-transmission device 1004 with its wireless remote control, the apparatus 10, in this case hydraulically operated, and the implanted energy-transforming device 1002, and further comprising a hydraulic fluid reservoir 1013, a motor/pump unit 1009 and an 15 reversing device in the form of a hydraulic valve shifting device 1014, all implanted in the patient. Of course the hydraulic operation could easily be performed by just changing the pumping direction and the hydraulic valve may therefore be omitted.

The remote control may be a device separated from the external energy-transmission device or included in the same. The motor of the motor/pump unit 1009 is an electric

- 20 motor. In response to a control signal from the wireless remote control of the external energy-transmission device 1004, the implanted energy-transforming device 1002 powers the motor/pump unit 1009 with energy from the energy carried by the control signal, whereby the motor/pump unit 1009 distributes hydraulic fluid between the hydraulic fluid reservoir 1013 and the apparatus 10. The remote control of the

- 25 external energy-transmission device 1004 controls the hydraulic valve shifting device 1014 to shift the hydraulic fluid flow direction between one direction in which the fluid is pumped by the motor/pump unit 1009 from the hydraulic fluid reservoir 1013 to the apparatus 10 to operate the apparatus, and another opposite direction in which the fluid is pumped by the motor/pump unit 1009 back from the apparatus 10 to the 30 hydraulic fluid reservoir 1013 to return the apparatus to a starting position.

Fig. 28 shows an embodiment of the invention comprising the external energy-transmission device 1004 with its wireless remote control, the apparatus 10, the implanted energy-transforming device 1002, an implanted internal control unit 1015 controlled by the wireless remote control of the external energy-transmission device

5 1004, an implanted accumulator 1016 and an implanted capacitor 1017. The internal control unit 1015 arranges storage of electric energy received from the implanted energy-transforming device 1002 in the accumulator 1016, which supplies energy to the apparatus 10. In response to a control signal from the wireless remote control of the external energy-transmission device 1004, the internal control unit 1015 either
10 releases electric energy from the accumulator 1016 and transfers the released energy via power lines 1018 and 1019, or directly transfers electric energy from the implanted energy-transforming device 1002 via a power line 1020, the capacitor 1017, which stabilizes the electric current, a power line 1021 and the power line 1019, for the operation of the apparatus 10.

15 The internal control unit is preferably programmable from outside the patient's body. In a preferred embodiment, the internal control unit is programmed to regulate the apparatus 10 according to a pre-programmed time-schedule or to input from any sensor sensing any possible physical parameter of the patient or any functional parameter of the system.

20 In accordance with an alternative, the capacitor 1017 in the embodiment of Fig. 28
10 may be omitted. In accordance with another alternative, the accumulator 1016 in this embodiment may be omitted.

25 Fig. 29 shows an embodiment of the invention identical to that of Fig. 23, except that a battery 1022 for supplying energy for the operation of the apparatus 10 and an electric switch 1023 for switching the operation of the apparatus 10 also are implanted in the patient. The electric switch 1023 may be controlled by the remote control and may also be operated by the energy supplied by the implanted energy-transforming device 1002 to switch from an off mode, in which the battery 1022 is not in use, to an on mode, in which the battery 1022 supplies energy for the operation
30 of the apparatus 10.

Fig. 30 shows an embodiment of the invention identical to that of Fig. 29, except that an internal control unit 1015 controllable by the wireless remote control of the external energy-transmission device 1004 also is implanted in the patient. In this case, the electric switch 1023 is operated by the energy supplied by the implanted energy-transforming device 1002 to switch from an off mode, in which the wireless remote control is prevented from controlling the internal control unit 1015 and the battery is not in use, to a standby mode, in which the remote control is permitted to control the internal control unit 1015 to release electric energy from the battery 1022 for the operation of the apparatus 10.

Fig. 31 shows an embodiment of the invention identical to that of Fig. 30, except that an accumulator 1016 is substituted for the battery 1022 and the implanted components are interconnected differently. In this case, the accumulator 1016 stores energy from the implanted energy-transforming device 1002. In response to a control signal from the wireless remote control of the external energy-transmission device 1004, the internal control unit 1015 controls the electric switch 1023 to switch from an off mode, in which the accumulator 1016 is not in use, to an on mode, in which the accumulator 1016 supplies energy for the operation of the apparatus 10. The accumulator may be combined with or replaced by a capacitor.

Fig. 32 shows an embodiment of the invention identical to that of Fig. 31, except that a battery 1022 also is implanted in the patient and the implanted components are interconnected differently. In response to a control signal from the wireless remote control of the external energy-transmission device 1004, the internal control unit 1015 controls the accumulator 1016 to deliver energy for operating the electric switch 1023 to switch from an off mode, in which the battery 1022 is not in use, to an on mode, in which the battery 1022 supplies electric energy for the operation of the apparatus 10.

Alternatively, the electric switch 1023 may be operated by energy supplied by the accumulator 1016 to switch from an off mode, in which the wireless remote control is prevented from controlling the battery 1022 to supply electric energy and is not in use, to a standby mode, in which the wireless remote control is permitted to control the battery 1022 to supply electric energy for the operation of the apparatus 10.

It should be understood that the switch 1023 and all other switches in this application should be interpreted in its broadest embodiment. This means a transistor, MCU, MCPU, ASIC, FPGA or a DA converter or any other electronic component or circuit that may switch the power on and off. Preferably the switch is controlled from 5 outside the body, or alternatively by an implanted internal control unit.

Fig. 33 shows an embodiment of the invention identical to that of Fig. 29, except that a motor 1007, a mechanical reversing device in the form of a gear box 1024, and an internal control unit 1015 for controlling the gear box 1024 also are implanted in the patient. The internal control unit 1015 controls the gear box 1024 to reverse the 10 function performed by the apparatus 10 (mechanically operated). Even simpler is to switch the direction of the motor electronically. The gear box interpreted in its broadest embodiment may stand for a servo arrangement saving force for the operation device in favour of longer stroke to act.

Fig. 34 shows an embodiment of the invention identical to that of Fig. 40 except that 15 the implanted components are interconnected differently. Thus, in this case the internal control unit 1015 is powered by the battery 1022 when the accumulator 1016, suitably a capacitor, activates the electric switch 1023 to switch to an on mode. When the electric switch 1023 is in its on mode the internal control unit 1015 is permitted to control the battery 1022 to supply, or not supply, energy for the 20 operation of the apparatus 10.

Fig. 35 schematically shows conceivable combinations of implanted components of the apparatus for achieving various communication options. Basically, there are the apparatus 10, the internal control unit 1015, motor or pump unit 1009, and the external energy-transmission device 1004 including the external wireless remote 25 control. As already described above the wireless remote control transmits a control signal which is received by the internal control unit 1015, which in turn controls the various implanted components of the apparatus.

A feedback device, preferably comprising a sensor or measuring device 1025, may be implanted in the patient for sensing a physical parameter of the patient. The 30 physical parameter may be at least one selected from the group consisting of

pressure, volume, diameter, stretching, elongation, extension, movement, bending, elasticity, muscle contraction, nerve impulse, body temperature, blood pressure, blood flow, heartbeats and breathing. The sensor may sense any of the above physical parameters. For example, the sensor may be a pressure or motility sensor.

5 Alternatively, the sensor 1025 may be arranged to sense a functional parameter. The functional parameter may be correlated to the transfer of energy for charging an implanted energy source and may further include at least one selected from the group of parameters consisting of; electricity, any electrical parameter, pressure, volume, diameter, stretch, elongation, extension, movement, bending, elasticity, temperature

10 and flow.

The feedback may be sent to the internal control unit or out to an external control unit preferably via the internal control unit. Feedback may be sent out from the body via the energy transfer system or a separate communication system with receiver and transmitters.

15 The internal control unit 1015, or alternatively the external wireless remote control of the external energy-transmission device 1004, may control the apparatus 10 in response to signals from the sensor 1025. A transceiver may be combined with the sensor 1025 for sending information on the sensed physical parameter to the external wireless remote control. The wireless remote control may comprise a signal transmitter or transceiver and the internal control unit 1015 may comprise a signal receiver or transceiver. Alternatively, the wireless remote control may comprise a signal receiver or transceiver and the internal control unit 1015 may comprise a signal transmitter or transceiver. The above transceivers, transmitters and receivers may be used for sending information or data related to the apparatus 10 from inside 20 the patient's body to the outside thereof.

25

30 Where the motor/pump unit 1009 and battery 1022 for powering the motor/pump unit 1009 are implanted, information related to the charging of the battery 1022 may be fed back. To be more precise, when charging a battery or accumulator with energy feed back information related to said charging process is sent and the energy supply is changed accordingly.

Fig. 36 shows an alternative embodiment wherein the apparatus 10 is regulated from outside the patient's body. The system 1000 comprises a battery 1022 connected to the apparatus 10 via a subcutaneous electric switch 1026. Thus, the regulation of the apparatus 10 is performed non-invasively by manually pressing the subcutaneous

5 switch, whereby the operation of the apparatus 10 is switched on and off. It will be appreciated that the shown embodiment is a simplification and that additional components, such as an internal control unit or any other part disclosed in the present application can be added to the system. Two subcutaneous switches may also be used. In the preferred embodiment one implanted switch sends information to the 10 internal control unit to perform a certain predetermined performance and when the patient press the switch again the performance is reversed.

Fig. 37 shows an alternative embodiment, wherein the system 1000 comprises a hydraulic fluid reservoir 1013 hydraulically connected to the apparatus. Non-invasive regulation is performed by manually pressing the hydraulic reservoir 15 connected to the apparatus. Alternatively, the hydraulic fluid reservoir 1013 is adapted to work with an injection port for the injection of hydraulic fluid, preferably for calibration of hydraulic fluid.

The system may include an external data communicator and an implantable internal data communicator communicating with the external data communicator. The 20 internal communicator feeds data related to the apparatus or the patient to the external data communicator and/or the external data communicator feeds data to the internal data communicator.

Fig. 38 schematically illustrates an arrangement of the system that is capable of sending information from inside the patient's body to the outside thereof to give 25 feedback information related to at least one functional parameter of the apparatus or system, or related to a physical parameter of the patient, in order to supply an accurate amount of energy to an implanted internal energy receiver 1002 connected to implanted energy consuming components of the apparatus 10. Such an energy receiver 1002 may include an energy source and/or an energy-transforming device. 30 Briefly described, wireless energy is transmitted from an external energy source 1004a located outside the patient and is received by the internal energy receiver 1002

located inside the patient. The internal energy receiver is adapted to directly or indirectly supply received energy to the energy consuming components of the apparatus 10 via a switch 1026. An energy balance is determined between the energy received by the internal energy receiver 1002 and the energy used for the apparatus 5 10, and the transmission of wireless energy is then controlled based on the determined energy balance. The energy balance thus provides an accurate indication of the correct amount of energy needed, which is sufficient to operate the apparatus 10 properly, but without causing undue temperature rise.

In Fig. 38 the patient's skin is indicated by a vertical line 1005. Here, the energy 10 receiver comprises an energy-transforming device 1002 located inside the patient, preferably just beneath the patient's skin 1005. Generally speaking, the implanted energy-transforming device 1002 may be placed in the abdomen, thorax, muscle fascia (e.g. in the abdominal wall), subcutaneously, or at any other suitable location. The implanted energy-transforming device 1002 is adapted to receive wireless 15 energy E transmitted from the external energy-source 1004a provided in an external energy-transmission device 1004 located outside the patient's skin 1005 in the vicinity of the implanted energy-transforming device 1002.

As is well known in the art, the wireless energy E may generally be transferred by 20 means of any suitable Transcutaneous Energy Transfer (TET) device, such as a device including a primary coil arranged in the external energy source 1004a and an adjacent secondary coil arranged in the implanted energy-transforming device 1002. When an electric current is fed through the primary coil, energy in the form of a 25 voltage is induced in the secondary coil which can be used to power the implanted energy consuming components of the apparatus, e.g. after storing the incoming energy in an implanted energy source, such as a rechargeable battery or a capacitor. However, the present invention is generally not limited to any particular energy transfer technique, TET devices or energy sources, and any kind of wireless energy may be used.

The amount of energy received by the implanted energy receiver may be compared 30 with the energy used by the implanted components of the apparatus. The term "energy used" is then understood to include also energy stored by implanted

components of the apparatus. A control device includes an external control unit 1004b that controls the external energy source 1004a based on the determined energy balance to regulate the amount of transferred energy. In order to transfer the correct amount of energy, the energy balance and the required amount of energy is

5 determined by means of a determination device including an implanted internal control unit 1015 connected between the switch 1026 and the apparatus 10. The internal control unit 1015 may thus be arranged to receive various measurements obtained by suitable sensors or the like, not shown, measuring certain characteristics of the apparatus 10, somehow reflecting the required amount of energy needed for
10 proper operation of the apparatus 10. Moreover, the current condition of the patient may also be detected by means of suitable measuring devices or sensors, in order to provide parameters reflecting the patient's condition. Hence, such characteristics and/or parameters may be related to the current state of the apparatus 10, such as power consumption, operational mode and temperature, as well as the patient's
15 condition reflected by parameters such as; body temperature, blood pressure, heartbeats and breathing. Other kinds of physical parameters of the patient and functional parameters of the device are described elsewhere.

Furthermore, an energy source in the form of an accumulator 1016 may optionally be connected to the implanted energy-transforming device 1002 via the control unit

20 1015 for accumulating received energy for later use by the apparatus 10. Alternatively or additionally, characteristics of such an accumulator, also reflecting the required amount of energy, may be measured as well. The accumulator may be replaced by a rechargeable battery, and the measured characteristics may be related to the current state of the battery, any electrical parameter such as energy
25 consumption voltage, temperature, etc. In order to provide sufficient voltage and current to the apparatus 10, and also to avoid excessive heating, it is clearly understood that the battery should be charged optimally by receiving a correct amount of energy from the implanted energy-transforming device 1002, i.e. not too little or too much. The accumulator may also be a capacitor with corresponding
30 characteristics.

For example, battery characteristics may be measured on a regular basis to determine the current state of the battery, which then may be stored as state information in a suitable storage means in the internal control unit 1015. Thus, whenever new measurements are made, the stored battery state information can be updated 5 accordingly. In this way, the state of the battery can be “calibrated” by transferring a correct amount of energy, so as to maintain the battery in an optimal condition.

Thus, the internal control unit 1015 of the determination device is adapted to determine the energy balance and/or the currently required amount of energy, (either energy per time unit or accumulated energy) based on measurements made by the 10 above-mentioned sensors or measuring devices of the apparatus 10, or the patient, or an implanted energy source if used, or any combination thereof. The internal control unit 1015 is further connected to an internal signal transmitter 1027, arranged to transmit a control signal reflecting the determined required amount of energy, to an external signal receiver 1004c connected to the external control unit 1004b. The 15 amount of energy transmitted from the external energy source 1004a may then be regulated in response to the received control signal.

Alternatively, the determination device may include the external control unit 1004b. In this alternative, sensor measurements can be transmitted directly to the external control unit 1004b wherein the energy balance and/or the currently required amount 20 of energy can be determined by the external control unit 1004b, thus integrating the above-described function of the internal control unit 1015 in the external control unit 1004b. In that case, the internal control unit 1015 can be omitted and the sensor measurements are supplied directly to the internal signal transmitter 1027 which sends the measurements over to the external signal receiver 1004c and the external 25 control unit 1004b. The energy balance and the currently required amount of energy can then be determined by the external control unit 1004b based on those sensor measurements.

Hence, the present solution according to the arrangement of Fig. 38 employs the feed back of information indicating the required energy, which is more efficient than 30 previous solutions because it is based on the actual use of energy that is compared to the received energy, e.g. with respect to the amount of energy, the energy difference,

or the energy receiving rate as compared to the energy rate used by implanted energy consuming components of the apparatus. The apparatus may use the received energy either for consuming or for storing the energy in an implanted energy source or the like. The different parameters discussed above would thus be used if relevant and 5 needed and then as a tool for determining the actual energy balance. However, such parameters may also be needed per se for any actions taken internally to specifically operate the apparatus.

The internal signal transmitter 1027 and the external signal receiver 1004c may be implemented as separate units using suitable signal transfer means, such as radio, IR 10 (Infrared) or ultrasonic signals. Alternatively, the internal signal transmitter 1027 and the external signal receiver 1004c may be integrated in the implanted energy-transforming device 1002 and the external energy source 1004a, respectively, so as to convey control signals in a reverse direction relative to the energy transfer, basically using the same transmission technique. The control signals may be 15 modulated with respect to frequency, phase or amplitude.

Thus, the feedback information may be transferred either by a separate communication system including receivers and transmitters or may be integrated in the energy system. In accordance with the present invention, such an integrated information feedback and energy system comprises an implantable internal energy receiver for receiving wireless energy, the energy receiver having an internal first coil and a first electronic circuit connected to the first coil, and an external energy transmitter for transmitting wireless energy, the energy transmitter having an external second coil and a second electronic circuit connected to the second coil. The external second coil of the energy transmitter transmits wireless energy which is received by 20 the first coil of the energy receiver. This system further comprises a power switch for switching the connection of the internal first coil to the first electronic circuit on and off, such that feedback information related to the charging of the first coil is received by the external energy transmitter in the form of an impedance variation in the load of the external second coil, when the power switch switches the connection of the 25 internal first coil to the first electronic circuit on and off. In implementing this system in the arrangement of Fig. 38, the switch 1026 is either separate and controlled by the 30 internal first coil to the first electronic circuit on and off. In implementing this system in the arrangement of Fig. 38, the switch 1026 is either separate and controlled by the

internal control unit 1015, or integrated in the internal control unit 1015. It should be understood that the switch 1026 should be interpreted in its broadest embodiment. This means a transistor, MCU, MCPU, ASIC FPGA or a DA converter or any other electronic component or circuit that may switch the power on and off.

5

To conclude, the energy supply arrangement illustrated in Fig. 38 may operate basically in the following manner. The energy balance is first determined by the internal control unit 1015 of the determination device. A control signal reflecting the required amount of energy is also created by the internal control unit 1015, and the

10 control signal is transmitted from the internal signal transmitter 1027 to the external signal receiver 1004c. Alternatively, the energy balance can be determined by the external control unit 1004b instead depending on the implementation, as mentioned above. In that case, the control signal may carry measurement results from various sensors. The amount of energy emitted from the external energy source 1004a can 15 then be regulated by the external control unit 1004b, based on the determined energy balance, e.g. in response to the received control signal. This process may be repeated intermittently at certain intervals during ongoing energy transfer, or may be executed on a more or less continuous basis during the energy transfer.

The amount of transferred energy can generally be regulated by adjusting various

20 transmission parameters in the external energy source 1004a, such as voltage, current, amplitude, wave frequency and pulse characteristics.

This system may also be used to obtain information about the coupling factors between the coils in a TET system even to calibrate the system both to find an optimal place for the external coil in relation to the internal coil and to optimize 25 energy transfer. Simply comparing in this case the amount of energy transferred with the amount of energy received. For example if the external coil is moved the coupling factor may vary and correctly displayed movements could cause the external coil to find the optimal place for energy transfer. Preferably, the external coil is adapted to calibrate the amount of transferred energy to achieve the feedback 30 information in the determination device, before the coupling factor is maximized.

This coupling factor information may also be used as a feedback during energy transfer. In such a case, the energy system of the present invention comprises an implantable internal energy receiver for receiving wireless energy, the energy receiver having an internal first coil and a first electronic circuit connected to the first coil, and an external energy transmitter for transmitting wireless energy, the energy transmitter having an external second coil and a second electronic circuit connected to the second coil. The external second coil of the energy transmitter transmits wireless energy which is received by the first coil of the energy receiver. This system further comprises a feedback device for communicating out the amount of energy received in the first coil as a feedback information, and wherein the second electronic circuit includes a determination device for receiving the feedback information and for comparing the amount of transferred energy by the second coil with the feedback information related to the amount of energy received in the first coil to obtain the coupling factor between the first and second coils. The energy transmitter may regulate the transmitted energy in response to the obtained coupling factor.

With reference to Fig. 39, although wireless transfer of energy for operating the apparatus has been described above to enable non-invasive operation, it will be appreciated that the apparatus can be operated with wire bound energy as well. Such an example is shown in Fig. 39, wherein an external switch 1026 is interconnected between the external energy source 1004a and an operation device, such as an electric motor 1007 operating the apparatus 10. An external control unit 1004b controls the operation of the external switch 1026 to effect proper operation of the apparatus 10.

Fig. 40 illustrates different embodiments for how received energy can be supplied to and used by the apparatus 10. Similar to the example of Fig. 38, an internal energy receiver 1002 receives wireless energy E from an external energy source 1004a which is controlled by a transmission control unit 1004b. The internal energy receiver 1002 may comprise a constant voltage circuit, indicated as a dashed box "constant V" in the figure, for supplying energy at constant voltage to the apparatus 10. The internal energy receiver 1002 may further comprise a constant current

circuit, indicated as a dashed box "constant C" in the figure, for supplying energy at constant current to the apparatus 10.

The apparatus 10 comprises an energy consuming part 10a, which may be a motor, pump, restriction device, or any other medical appliance that requires energy for its 5 electrical operation. The apparatus 10 may further comprise an energy storage device 10b for storing energy supplied from the internal energy receiver 1002. Thus, the supplied energy may be directly consumed by the energy consuming part 10a, or stored by the energy storage device 10b, or the supplied energy may be partly consumed and partly stored. The apparatus 10 may further comprise an energy 10 stabilizing unit 10c for stabilizing the energy supplied from the internal energy receiver 1002. Thus, the energy may be supplied in a fluctuating manner such that it may be necessary to stabilize the energy before consumed or stored.

The energy supplied from the internal energy receiver 1002 may further be accumulated and/or stabilized by a separate energy stabilizing unit 1028 located 15 outside the apparatus 10, before being consumed and/or stored by the apparatus 10. Alternatively, the energy stabilizing unit 1028 may be integrated in the internal energy receiver 1002. In either case, the energy stabilizing unit 1028 may comprise a constant voltage circuit and/or a constant current circuit.

It should be noted that Fig. 38 and Fig. 40 illustrate some possible but non-limiting 20 implementation options regarding how the various shown functional components and elements can be arranged and connected to each other. However, the skilled person will readily appreciate that many variations and modifications can be made within the scope of the present invention.

Fig. 41 schematically shows an energy balance measuring circuit of one of the 25 proposed designs of the system for controlling transmission of wireless energy, or energy balance control system. The circuit has an output signal centered on 2.5V and proportionally related to the energy imbalance. The derivative of this signal shows if the value goes up and down and how fast such a change takes place. If the amount of received energy is lower than the energy used by implanted components of the 30 apparatus, more energy is transferred and thus charged into the energy source. The

output signal from the circuit is typically feed to an A/D converter and converted into a digital format. The digital information can then be sent to the external energy-transmission device allowing it to adjust the level of the transmitted energy. Another possibility is to have a completely analog system that uses comparators comparing 5 the energy balance level with certain maximum and minimum thresholds sending information to external energy-transmission device if the balance drifts out of the max/min window.

The schematic Fig. 41 shows a circuit implementation for a system that transfers 10 energy to the implanted energy components of the apparatus of the present invention from outside of the patient's body using inductive energy transfer. An inductive energy transfer system typically uses an external transmitting coil and an internal receiving coil. The receiving coil, L1, is included in the schematic Fig. 24; the transmitting parts of the system are excluded.

The implementation of the general concept of energy balance and the way the 15 information is transmitted to the external energy transmitter can of course be implemented in numerous different ways. The schematic Fig. 41 and the above described method of evaluating and transmitting the information should only be regarded as examples of how to implement the control system.

CIRCUIT DETAILS

20 In Fig. 41 the symbols Y1, Y2, Y3 and so on symbolize test points within the circuit. The components in the diagram and their respective values are values that work in this particular implementation which of course is only one of an infinite number of possible design solutions.

25 Energy to power the circuit is received by the energy receiving coil L1. Energy to implanted components is transmitted in this particular case at a frequency of 25 kHz. The energy balance output signal is present at test point Y1.

Those skilled in the art will realize that the above various embodiments of the system could be combined in many different ways. For example, the electric switch 1006 of Fig. 24 could be incorporated in any of the embodiments of Figs. 27-33, the

hydraulic valve shifting device 1014 of Fig. 27 could be incorporated in the embodiment of Fig. 26, and the gear box 1024 could be incorporated in the embodiment of Fig. 25. Please observe that the switch simply could mean any electronic circuit or component.

- 5 The embodiments described in connection with Figs. 38, 40 and 41 identify a method and a system for controlling transmission of wireless energy to implanted energy consuming components of an electrically operable apparatus. Such a method and system will be defined in general terms in the following.

A method is thus provided for controlling transmission of wireless energy supplied to

- 10 implanted energy consuming components of an apparatus as described above. The wireless energy E is transmitted from an external energy source located outside the patient and is received by an internal energy receiver located inside the patient, the internal energy receiver being connected to the implanted energy consuming components of the apparatus for directly or indirectly supplying received energy thereto. An energy balance is determined between the energy received by the internal energy receiver and the energy used for the apparatus. The transmission of wireless energy E from the external energy source is then controlled based on the determined energy balance.

The wireless energy may be transmitted inductively from a primary coil in the

- 20 external energy source to a secondary coil in the internal energy receiver. A change in the energy balance may be detected to control the transmission of wireless energy based on the detected energy balance change. A difference may also be detected between energy received by the internal energy receiver and energy used for the medical device, to control the transmission of wireless energy based on the detected energy difference.

When controlling the energy transmission, the amount of transmitted wireless energy may be decreased if the detected energy balance change implies that the energy balance is increasing, or vice versa. The decrease/increase of energy transmission may further correspond to a detected change rate.

The amount of transmitted wireless energy may further be decreased if the detected energy difference implies that the received energy is greater than the used energy, or vice versa. The decrease/increase of energy transmission may then correspond to the magnitude of the detected energy difference.

- 5 As mentioned above, the energy used for the medical device may be consumed to operate the medical device, and/or stored in at least one energy storage device of the medical device.

When electrical and/or physical parameters of the medical device and/or physical parameters of the patient are determined, the energy may be transmitted for

- 10 consumption and storage according to a transmission rate per time unit which is determined based on said parameters. The total amount of transmitted energy may also be determined based on said parameters.

When a difference is detected between the total amount of energy received by the internal energy receiver and the total amount of consumed and/or stored energy, and

- 15 the detected difference is related to the integral over time of at least one measured electrical parameter related to said energy balance, the integral may be determined for a monitored voltage and/or current related to the energy balance.

When the derivative is determined over time of a measured electrical parameter related to the amount of consumed and/or stored energy, the derivative may be

- 20 determined for a monitored voltage and/or current related to the energy balance.

The transmission of wireless energy from the external energy source may be controlled by applying to the external energy source electrical pulses from a first electric circuit to transmit the wireless energy, the electrical pulses having leading and trailing edges, varying the lengths of first time intervals between successive

- 25 leading and trailing edges of the electrical pulses and/or the lengths of second time intervals between successive trailing and leading edges of the electrical pulses, and transmitting wireless energy, the transmitted energy generated from the electrical pulses having a varied power, the varying of the power depending on the lengths of the first and/or second time intervals.

In that case, the frequency of the electrical pulses may be substantially constant when varying the first and/or second time intervals. When applying electrical pulses, the electrical pulses may remain unchanged, except for varying the first and/or second time intervals. The amplitude of the electrical pulses may be substantially constant 5 when varying the first and/or second time intervals. Further, the electrical pulses may be varied by only varying the lengths of first time intervals between successive leading and trailing edges of the electrical pulses.

A train of two or more electrical pulses may be supplied in a row, wherein when applying the train of pulses, the train having a first electrical pulse at the start of the 10 pulse train and having a second electrical pulse at the end of the pulse train, two or more pulse trains may be supplied in a row, wherein the lengths of the second time intervals between successive trailing edge of the second electrical pulse in a first pulse train and leading edge of the first electrical pulse of a second pulse train are varied.

15 When applying the electrical pulses, the electrical pulses may have a substantially constant current and a substantially constant voltage. The electrical pulses may also have a substantially constant current and a substantially constant voltage. Further, the electrical pulses may also have a substantially constant frequency. The electrical pulses within a pulse train may likewise have a substantially constant frequency.

20 The circuit formed by the first electric circuit and the external energy source may have a first characteristic time period or first time constant, and when effectively varying the transmitted energy, such frequency time period may be in the range of the first characteristic time period or time constant or shorter.

A system comprising an apparatus as described above is thus also provided for 25 controlling transmission of wireless energy supplied to implanted energy consuming components of the apparatus. In its broadest sense, the system comprises a control device for controlling the transmission of wireless energy from an energy-transmission device, and an implantable internal energy receiver for receiving the transmitted wireless energy, the internal energy receiver being connected to 30 implantable energy consuming components of the apparatus for directly or indirectly

supplying received energy thereto. The system further comprises a determination device adapted to determine an energy balance between the energy received by the internal energy receiver and the energy used for the implantable energy consuming components of the apparatus, wherein the control device controls the transmission of wireless energy from the external energy-transmission device, based on the energy balance determined by the determination device.

In one embodiment at least one battery may be a part of or replace the energy-transforming device 1002 to supply energy to the apparatus 10 over a power supply line. In one embodiment the battery is not rechargeable. In an alternative 10 embodiment the battery is rechargeable. The battery supply may of course be placed both remote to and incorporated in the device.

Further, the system may comprise any of the following:

- A primary coil in the external energy source adapted to transmit the wireless energy inductively to a secondary coil in the internal energy receiver.
- 15 - The determination device is adapted to detect a change in the energy balance, and the control device controls the transmission of wireless energy based on the detected energy balance change
- The determination device is adapted to detect a difference between energy received by the internal energy receiver and energy used for the implantable energy 20 consuming components of the apparatus, and the control device controls the transmission of wireless energy based on the detected energy difference.
- The control device controls the external energy-transmission device to decrease the amount of transmitted wireless energy if the detected energy balance change implies that the energy balance is increasing, or vice versa, wherein the decrease/increase of 25 energy transmission corresponds to a detected change rate.
- The control device controls the external energy-transmission device to decrease the amount of transmitted wireless energy if the detected energy difference implies that the received energy is greater than the used energy, or vice versa, wherein the

decrease/increase of energy transmission corresponds to the magnitude of said detected energy difference.

- The energy used for the apparatus is consumed to operate the apparatus, and/or stored in at least one energy storage device of the apparatus.

5 - Where electrical and/or physical parameters of the apparatus and/or physical parameters of the patient are determined, the energy-transmission device transmits the energy for consumption and storage according to a transmission rate per time unit which is determined by the determination device based on said parameters. The determination device also determines the total amount of transmitted energy based on 10 said parameters.

- When a difference is detected between the total amount of energy received by the internal energy receiver and the total amount of consumed and/or stored energy, and the detected difference is related to the integral over time of at least one measured electrical parameter related to the energy balance, the determination device

15 determines the integral for a monitored voltage and/or current related to the energy balance.

- When the derivative is determined over time of a measured electrical parameter related to the amount of consumed and/or stored energy, the determination device determines the derivative for a monitored voltage and/or current related to the energy 20 balance.

- The energy-transmission device comprises a coil placed externally to the human body, and an electric circuit is provided to power the external coil with electrical pulses to transmit the wireless energy. The electrical pulses have leading and trailing edges, and the electric circuit is adapted to vary first time intervals between

25 successive leading and trailing edges and/or second time intervals between successive trailing and leading edges of the electrical pulses to vary the power of the transmitted wireless energy. As a result, the energy receiver receiving the transmitted wireless energy has a varied power.

- The electric circuit is adapted to deliver the electrical pulses to remain unchanged except varying the first and/or second time intervals.
 - The electric circuit has a time constant and is adapted to vary the first and second time intervals only in the range of the first time constant, so that when the lengths of the first and/or second time intervals are varied, the transmitted power over the coil is varied.
 - The electric circuit is adapted to deliver the electrical pulses to be varied by only varying the lengths of first time intervals between successive leading and trailing edges of the electrical pulses.
- 10 - The electric circuit is adapted to supplying a train of two or more electrical pulses in a row, said train having a first electrical pulse at the start of the pulse train and having a second electrical pulse at the end of the pulse train, and
- the lengths of the second time intervals between successive trailing edge of the second electrical pulse in a first pulse train and leading edge of the first electrical pulse of a second pulse train are varied by the first electronic circuit.
 - The electric circuit is adapted to provide the electrical pulses as pulses having a substantially constant height and/or amplitude and/or intensity and/or voltage and/or current and/or frequency.
 - The electric circuit has a time constant, and is adapted to vary the first and second time intervals only in the range of the first time constant, so that when the lengths of the first and/or second time intervals are varied, the transmitted power over the first coil are varied.
 - The electric circuit is adapted to provide the electrical pulses varying the lengths of the first and/or the second time intervals only within a range that includes the first time constant or that is located relatively close to the first time constant, compared to the magnitude of the first time constant.

Figs. 42-45 show in more detail block diagrams of four different ways of hydraulically or pneumatically powering an implanted apparatus according to the invention.

Fig. 42 shows a system as described above with. The system comprises an implanted 5 apparatus 10 and further a separate regulation reservoir 1013, a one way pump 1009 and an alternate valve 1014.

Fig. 43 shows the apparatus 10 and a fluid reservoir 1013. By moving the wall of the regulation reservoir or changing the size of the same in any other different way, the adjustment of the apparatus may be performed without any valve, just free passage of 10 fluid any time by moving the reservoir wall.

Fig. 44 shows the apparatus 10, a two way pump 1009 and the regulation reservoir 1013.

Fig. 45 shows a block diagram of a reversed servo system with a first closed system controlling a second closed system. The servo system comprises a regulation 15 reservoir 1013 and a servo reservoir 1050. The servo reservoir 1050 mechanically controls an implanted apparatus 10 via a mechanical interconnection 1054. The apparatus has an expandable/contactable cavity. This cavity is preferably expanded or contracted by supplying hydraulic fluid from the larger adjustable reservoir 1052 in fluid connection with the apparatus 10. Alternatively, the cavity contains 20 compressible gas, which can be compressed and expanded under the control of the servo reservoir 1050.

The servo reservoir 1050 can also be part of the apparatus itself.

In one embodiment, the regulation reservoir is placed subcutaneous under the patient's skin and is operated by pushing the outer surface thereof by means of a 25 finger. This system is illustrated in Figs 46a-c. In Fig. 46a, a flexible subcutaneous regulation reservoir 1013 is shown connected to a bulge shaped servo reservoir 1050 by means of a conduit 1011. This bellow shaped servo reservoir 1050 is comprised in a a flexible apparatus 10. In the state shown in Fig. 46a, the servo reservoir 1050 contains a minimum of fluid and most fluid is found in the regulation reservoir 1013.

Due to the mechanical interconnection between the servo reservoir 1050 and the apparatus 10, the outer shape of the apparatus 10 is contracted, i.e., it occupies less than its maximum volume. This maximum volume is shown with dashed lines in the figure.

- 5 Fig. 46b shows a state wherein a user, such as the patient in with the apparatus is implanted, presses the regulation reservoir 1013 so that fluid contained therein is brought to flow through the conduit 1011 and into the servo reservoir 1050, which, thanks to its bellow shape, expands longitudinally. This expansion in turn expands the apparatus 10 so that it occupies its maximum volume, thereby stretching the
10 stomach wall (not shown), which it contacts.

The regulation reservoir 1013 is preferably provided with means 1013a for keeping its shape after compression. This means, which is schematically shown in the figure, will thus keep the apparatus 10 in a stretched position also when the user releases the regulation reservoir. In this way, the regulation reservoir essentially operates as an
15 on/off switch for the system.

An alternative embodiment of hydraulic or pneumatic operation will now be described with reference to Figs. 47 and 48a-c. The block diagram shown in Fig. 47 comprises with a first closed system controlling a second closed system. The first system comprises a regulation reservoir 1013 and a servo reservoir 1050. The servo
20 reservoir 1050 mechanically controls a larger adjustable reservoir 1052 via a mechanical interconnection 1054. An implanted apparatus 10 having an expandable/contactable cavity is in turn controlled by the larger adjustable reservoir 1052 by supply of hydraulic fluid from the larger adjustable reservoir 1052 in fluid connection with the apparatus 10.

25 An example of this embodiment will now be described with reference to Fig. 48a-c. Like in the previous embodiment, the regulation reservoir is placed subcutaneous under the patient's skin and is operated by pushing the outer surface thereof by means of a finger. The regulation reservoir 1013 is in fluid connection with a bellow shaped servo reservoir 1050 by means of a conduit 1011. In the first closed system

1013, 1011, 1050 shown in Fig. 48a, the servo reservoir 1050 contains a minimum of fluid and most fluid is found in the regulation reservoir 1013.

The servo reservoir 1050 is mechanically connected to a larger adjustable reservoir 1052, in this example also having a bellow shape but with a larger diameter than the

5 servo reservoir 1050. The larger adjustable reservoir 1052 is in fluid connection with the apparatus 10. This means that when a user pushes the regulation reservoir 1013, thereby displacing fluid from the regulation reservoir 1013 to the servo reservoir 1050, the expansion of the servo reservoir 1050 will displace a larger volume of fluid from the larger adjustable reservoir 1052 to the apparatus 10. In other words, in this 10 reversed servo, a small volume in the regulation reservoir is compressed with a higher force and this creates a movement of a larger total area with less force per area unit.

Like in the previous embodiment described above with reference to Figs. 46a-c, the regulation reservoir 1013 is preferably provided with means 1013a for keeping its

15 shape after compression. This means, which is schematically shown in the figure, will thus keep the apparatus 10 in a stretched position also when the user releases the regulation reservoir. In this way, the regulation reservoir essentially operates as an on/off switch for the system.

One single volume filling device has been described as invaginated in the stomach

20 wall. Alternatively, two or more volume filling devices 10 may be invaginated to obtain the desired reduction of the food cavity. One such example is illustrated in Fig. 48, wherein three ball-shaped volume filling devices 10 are invaginated in the wall of the patient's stomach 12.

It has been described how the volume filling device 10 is invaginated in the stomach

25 wall by means of a gastroscopic instrument. It will be appreciated that abdominal operation methods can be used as well. Such methods will now be described in detail with reference to Figs. 49a and 49b.

In a first alternative embodiment, the volume filling device is implanted using a laparoscopic method instead of the intraluminal method described above. According

to this embodiment, a needle or a tube-like instrument is inserted into the abdomen of the patient's body, and said needle or tube-like instrument is then used to fill the patient's abdomen with gas. Subsequently, at least two laparoscopic trocars are inserted into the patient's body; and a camera is inserted through one of said at least 5 two laparoscopic trocars. Then, at least one dissecting tool through one of said at least two laparoscopic trocars, and an area of the stomach is dissected. The volume filling device is then introduced into the abdominal cavity, and placed on the outside of the stomach wall. A pouch in the stomach wall for the device is created, and the device invaginated in said pouch by providing sutures or staples to the stomach wall, 10 thereby positioning the volume filling device so that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device.

The above first alternative preferably further comprises affixing the device to the stomach wall by providing sutures or staples.

15 The above embodiment preferably further comprises providing an apparatus for regulating the obesity treatment device from the outside of the patient's body; and operating said apparatus to regulate the obesity treatment device. Further, regulation of the obesity treatment device includes changing the volume of a filling body of the volume filling device when implanted.

20 The above embodiment preferably further comprises providing an injection type syringe comprising a fluid for injection into an implanted filling body; and injecting volume of fluid into said filling body.

According to an embodiment, the device is enclosed in the pouch or partially enclosed in that the pouch is left at least partly open. Further, the pouch can be 25 designed to exhibit only one opening. Alternatively the pouch is designed to exhibit two openings and to extend non-circumferentially around the stomach.

Preferably the pouch has a volume of more than 15 milliliters.

In a second alternative, also using a laparoscopic method instead of the intraluminal method, the initial steps are the same as described in the first alternative, but

following dissection of the stomach, a hole is created in the stomach wall and a volume filling device introduced into the abdominal cavity and through said hole into the stomach. The device is placed on the inside of the stomach wall, and a pouch is created on the outside of the stomach cavity for the device placed on the inside of the 5 stomach wall, and the device is invaginated in the pouch by providing sutures or staples to the stomach wall, thereby positioning the volume filling device so that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device.

10 The above embodiment preferably further comprises affixing the device to the stomach wall by providing sutures or staples. According to one embodiment, the stomach wall is affixed to the lower part of the patient's esophagus by providing sutures or staples.

15 The above second alternative preferably further comprises providing an apparatus for regulating the obesity treatment device from the outside of the patient's body; and operating said apparatus to regulate the obesity treatment device. Further, regulation of the obesity treatment device includes changing the volume of a filling body of the volume filling device when implanted.

20 The above embodiment preferably further comprises providing an injection type syringe comprising a fluid for injection into an implanted filling body; and injecting volume of fluid into said filling body.

According to an embodiment, the device is enclosed in the pouch or partially enclosed in that the pouch is left at least partly open. Further, the pouch can be designed to exhibit only one opening. Alternatively the pouch is designed to exhibit two openings and to extend non-circumferentially around the stomach.

25 Preferably the pouch has a volume of more than 15 milliliters.

A third alternative involves a surgical incision instead of the either the intraluminal or the laparoscopic method. Here, an opening in the patient's abdominal wall is made by surgical incision, and an area of the patient's stomach is dissected. The volume filling device is introduced through said abdominal incision, and attached to the

stomach wall, thereby positioning the volume filling device so that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device.

In an alternative embodiment of the above, third alternative, the initial steps are the

- 5 same including the dissection of an area of the stomach. Following this, a pouch in the stomach wall is created for the device, and the device invaginated in the pouch by providing sutures or staples to the stomach wall, thereby positioning the volume filling device so that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device.

10 In yet another alternative embodiment of the above, third alternative, the initial steps

are the same including the dissection of an area of the stomach. Following this, a hole in the stomach wall is created and the volume filling device introduced through the hole and into the stomach. The device is then placed on the inside of the stomach wall, and a pouch on the stomach wall created for the device. The device is then

15 invaginated in the pouch by providing sutures or staples to the stomach wall, thereby positioning the volume filling device so that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device.

The above embodiments of the third alternative further comprise affixing the device to the stomach wall by providing sutures or staples.

20 The above embodiment preferably further comprises providing an apparatus for regulating the obesity treatment device from the outside of the patient's body; and operating said apparatus to regulate the obesity treatment device. Further, regulation of the obesity treatment device includes changing the volume of a filling body of the volume filling device when implanted.

25 The above embodiment preferably further comprises providing an injection type syringe comprising a fluid for injection into an implanted filling body; and injecting volume of fluid into said filling body.

According to an embodiment, the device is enclosed in the pouch or partially enclosed in that the pouch is left at least partly open. Further, the pouch can be

designed to exhibit only one opening. Alternatively the pouch is designed to exhibit two openings and to extend non-circumferentially around the stomach.

Preferably the pouch has a volume of more than 15 milliliters.

A fourth alternative embodiment is a method comprising the steps of inserting a

5 needle or a tube-like instrument into the abdomen of the patient's body; using said
needle or tube-like instrument to fill the patient's abdomen with gas; placing at least
two laparoscopic trocars in the patient's body; inserting a camera through one of said
at least two laparoscopic trocars into the patient's abdomen; inserting at least one
dissecting tool through one of said at least two laparoscopic trocars; dissecting an
10 area of the stomach; creating a pouch from the stomach wall for the device; closing
the pouch by providing sutures and staples; introducing a injecting member
comprising an injectable filling material; and injecting the filling material into the
pouch, thereby creating a filling body that fills a volume in the patient's stomach,
reducing the food cavity in size by a volume substantially exceeding the volume of
15 the volume filling device.

Instead of the above disclosed laparoscopic method, a surgical incision or opening is
cut in the skin to enter the patients abdomen; an area of the stomach dissected; a
pouch created from the stomach wall for the device; and said pouch closed by
providing sutures and staples. An injecting member comprising an injectable filling
20 material is then introduced; and the filling material injected into the pouch, thereby
creating a filling body that reduces the food cavity in size by a volume substantially
exceeding the volume of the volume filling device.

According to an alternative embodiment of the above, the pouch is created on the
outside of the stomach wall, with the filling body placed against the inside of the
25 stomach wall.

The method according to either of the two previous embodiments comprises creating
a hole in the stomach wall wherein the pouch is created on the inside of the stomach
wall, with the filling body placed against the outside of the stomach wall.

The method according to either of the two previous embodiments may further comprise affixing the stomach wall to the lower part of the patient's esophagus by providing sutures or staples or affixing the stomach wall to the patient's diaphragm muscle or associated muscles.

- 5 Preferably the pouch has a volume of more than 15 milliliters.

In a method according to either of the two previous embodiments the filling material is preferably capable of undergoing a curing process from a fluid state to a semi-solid or solid state. Preferably said curing process is triggered by an increase in temperature from ambient temperature to body temperature.

- 10 The invention also makes available a method of treating obesity in a patient by implanting a volume filling device that, when implanted in a patient, reduces the food cavity in size by a volume substantially exceeding the volume of the volume filling device, the method comprising the steps of:

inserting a needle or a tube-like instrument into the abdomen of the patient's body;

- 15 using said needle or tube-like instrument to fill the patient's abdomen with gas;

placing at least two laparoscopic trocars in the patient's body;

inserting a camera through one of said at least two laparoscopic trocars into the patient's abdomen;

- 15 inserting at least one dissecting tool through one of said at least two laparoscopic trocars;

dissecting an area of the stomach;

creating a hole in the stomach wall;

introducing a device into the abdominal cavity;

introducing the device through the hole and into the stomach;

- 25 placing the device on the outside of the stomach wall;

fixating the device placed on the outside of the stomach wall, and thereby creating a filling body that reduces the food cavity in size by a volume substantially exceeding the volume of the volume filling device.

In the above method, the device is preferably affixed to the stomach wall by 5 providing sutures or staples.

The invention also comprises a laparoscopic instrument for providing a volume filling device to be invaginated in the stomach wall of a human patient to treat obesity, suitable for use with any of the laparoscopic methods described above, the instrument comprising:

- 10 an elongated member having a proximal end and a distal end, the elongated member having a diameter less than that of a laparoscopic trocar to be introduced into the patients abdomen during a laparoscopic operation;
- 15 a stomach pushing device for pushing the stomach wall to create a tube-like shaped portion of the stomach wall protruding into the normal stomach cavity, said pushing device comprising the volume filling device to be invaginated by the stomach wall in the tube-like shaped portion thereof;
- 20 wherein the pushing device comprises a vacuum device to suck the stomach fundus to assist the instrument in forming the tube-like shaped portion of the stomach wall together with the pushing device, and wherein the vacuum device comprises a vacuum passageway leading from the proximal to the distal end of the instrument and at the end portion of the instrument, which includes the pushing device, said vacuum passageway is divided up in multiple small openings adapted to suck the stomach wall portion to become adherent to the pushing device to further form the tube-like stomach wall portion; and
- 25 wherein the instrument comprises an insertion device adapted to introduce the volume filling device into the tube-like shaped stomach portion.

This instrument preferably comprises at least one clamping device for holding the opening of the tube-like portion substantially closed by clamping together stomach to

stomach in said opening, wherein the instrument is adapted to place the at least one clamping device at the opening in such a way that it allows later suturing of the opening.

Further, the instrument preferably comprises an inflation device for inflating the volume filling device before or after the suturing. Further still, the instrument preferably comprises a suturing device adapted to suture the opening of the tube-like portion with stomach to stomach sutures for creating at least partly a closed space enclosing the volume filling device, wherein the instrument is adapted to be withdrawn leaving the volume filling device at least partly invaginated in the stomach wall.

Said suturing device preferably comprises a first and second suture positioning member provided on the elongated member situated in the stomach at the distal end thereof, and wherein the instrument further comprises an operation device adapted to adjust the first and second suturing member in a position in which the first and second suture positioning members are in front of each other with the stomach wall on both sides of the open end of the cup like portion, and adapted to suture the open end of the cup like portion of the wall with a row of stomach to stomach sutures.

Preferably said suturing device comprises an operable re-loadable multi-suturing device, which is reloadable with sutures from outside of the patient's body and which is adapted to suture the open end of the cup like portion of the wall with said row of stomach to stomach sutures, wherein the row of sutures comprises two or more sutures or staples to be sutured simultaneously.

More preferably, said suturing device comprises multiple sutures for suturing two or more sutures simultaneously.

It is understood that a skilled person is in the position of combining steps, changing the order of steps, and combining elements of the different embodiments of the invention without inventive effort, and without departing from the scope of the invention as defined in the description and claims.

CLAIMS

1. An apparatus for treating obesity of an obese patient having a stomach with a food cavity, the apparatus comprising:
 - 5 at least one volume filling device adapted to be at least substantially invaginated by a stomach wall portion of the patient, and
 - 10 wherein the volume filling device is adapted to be placed on the outside of the stomach wall with the outer surface of the volume filling device resting against the outside of the stomach wall, such that the volume of the food cavity is reduced in size by a volume substantially exceeding the volume of the volume filling device, the volume filling device having a maximum circumference of at least 30 millimeters.
2. The apparatus according to claim 1, wherein the volume filling device comprises an inflatable device expandable to an expanded state.
- 15 3. The apparatus according to claim 2, wherein the inflatable device has an inlet port for a fluid or a gel and is adapted to be connected to a gastroscopic instrument.
4. The apparatus according to claim 3, wherein the inlet port comprises a fluid connection adapted to interconnect the inflatable device and the gastroscopic 20 instrument.
5. The apparatus according to claim 1, wherein the volume filling device has an elongated shape.
6. The apparatus according to claim 5, wherein the volume filling device has a rounded shape.
- 25 7. The apparatus according to claim 5, wherein the volume filling device has a bent or curved shape.

8. The apparatus according to claim 1, wherein the volume filling device comprises an elastic material.

9. The apparatus according to claim 1, wherein the volume filling device comprises a bio-compatible material.

5 10. The apparatus according to claim 1, wherein the volume filling device comprises silicone.

11. The apparatus according to claim 1, wherein the volume filling device is provided with a coating or a multilayer coating.

10 12. The apparatus according to claim 11, wherein the coating is a Parylene coating.

13. The apparatus according to claim 11, wherein the coating is a polytetra-fluoroethylene coating.

14. The apparatus according to claim 11, wherein the coating is a polyurethane coating.

15 15. The apparatus according to claim 11, wherein the coating is a multi-layer coating.

16. The apparatus according to claim 11, wherein the coating is a polyurethane coating.

17. The apparatus according to claim 1, wherein the volume filling device 20 comprises a fluid.

18. The apparatus according to claim 17, wherein the fluid is adapted to be transformed into solid state or fixed form.

19. The apparatus according to claim 17, wherein the fluid is liquid polyurethane.

25 20. The apparatus according to claim 17, wherein the fluid is iso-tonic.

21. The apparatus according to claim 17, wherein the fluid comprises large molecules to prevent diffusion.
22. 21 The apparatus according to claim 17, wherein the fluid comprises iodine molecules.
- 5 23. The apparatus according to claim 1, wherein the volume filling device comprises a homogenous material.
24. The apparatus according to claim 1, wherein the volume filling device is a solid body.
- 10 25. The apparatus according to claim 1, wherein the volume filling device comprises an enclosure wall defining a chamber.
26. The apparatus according to claim 1, wherein the volume filling device comprises a rigid outer surface.
27. The apparatus according to claim 1, wherein the volume filling device comprises an elastic outer surface.
- 15 28. The apparatus according to claim 1, wherein the volume filling device comprises a flexible outer surface.
29. The apparatus according to claim 1, wherein the volume filling device as a maximum circumference of at least 50 millimeters.
- 20 30. The apparatus according to claim 29, wherein the volume filling device has a circumference of at least 80 millimeters.
31. The apparatus according to claim 1, wherein the volume filling device has a volume of between 0.00001 and 0.001 m³.
32. The apparatus according to claim 31, wherein the volume filling device has a volume of between 0.00001 and 0.0002 m³.

33. The apparatus according to claim 1, wherein the volume filling device is deformable to a maximum diameter, so as to be inserted into a laparoscopic trocar.

34. The apparatus according to claim 1, wherein the volume filling device is adapted to be kept in place by stomach-to-stomach sutures or staples to invaginate 5 the device in the stomach wall.

35. The apparatus according to claim 34, wherein the stomach-to-stomach sutures or staples are provided with fixation portions exhibiting a structure adapted to be in contact with the stomach wall to promote growth in of human tissue to secure long term placement of the volume filling device attached to the stomach wall.

10 36. The apparatus according to claim 35, wherein the structure comprises a net like structure.

37. The apparatus according to claim 1, wherein the volume filling device is adapted to be non-invasively adjustable postoperatively.

15 38. The apparatus according to claim 1, wherein the volume filling device has varying circumference to be better adapted to be kept in place invaginated in the stomach wall of the patient.

20 39. The apparatus according to claim 1, comprising a stretching device placed outside the stomach wall and adapted to stretch a part of the stomach wall, thereby affecting the patient's appetite, the apparatus further comprising a fluid connection interconnecting the stretching device and the volume filling device.

40. The apparatus according to claim 38, wherein the volume filling device is adapted to be placed outside the stomach wall via a gastroscopic instrument.

25 41. The apparatus according to claim 1, wherein the volume filling device comprises at least two interconnectable portions adapted to be placed outside the stomach wall as separate portions.

42. The apparatus according to claim 1, wherein the volume filling device comprises an outer surface layer of polyurethane, Teflon®, or PTFE, or a combination thereof.

43. The apparatus according to claim 1, wherein the volume filling device 5 is destructible by acid, preferably hydrochloric acid.

44. The apparatus according to claim 1, wherein the volume filling device comprises gel.

45. The apparatus according to claim 43, wherein the gel has a shore value of less than 15.

10 46. The apparatus according to claim 1, wherein the volume filling device comprises an attachment device adapted to co-operate with a gripping instrument.

47. The apparatus according to claim 1, wherein the volume filling device is adapted to be completely invaginated by the stomach wall of the patient.

15 48. The apparatus according to claim 1, comprising a fixating device adapted to fixate the volume filling device to the stomach wall to keep the volume filling device in place, when the volume filling device is implanted.

49. The apparatus according to claim 31, wherein the volume filling device has a volume of less than 0.0002 m³.

50. The apparatus according to claim 31, wherein the volume filling device 20 has a volume of between 0.0001 and 0.001 m³.

51. The apparatus according to claim 1, wherein at least a part of the volume filling device is made of a material which is not destructible by acid.

52. The apparatus according to claim 29, wherein the volume filling device has a circumference of at least 120 mm.

25 53. The apparatus according to claim 29, wherein the volume filling device has a circumference of at least 150 mm.

54. The apparatus according to claim 29, wherein the volume filling device has a circumference of at least 180 mm.

55. The apparatus according to claim 29, wherein the volume filling device has a circumference of at least 220 mm.

5 56. The apparatus according to claim 1, wherein the volume filling device comprises a flexible non-elastic material.

57. The apparatus according to claim 1, further comprising a fixation device adapted to be involved in the fixation of the volume filling device to the stomach wall.

10 58. The apparatus according to claim 1, further comprising two or more fixation devices adapted to be involved in the fixation of the volume filling device to the stomach wall.

59. The apparatus according to claim 1, wherein the volume filling device comprises a holding device adapted to be held by an instrument and simplify the 15 implantation of the device.

60. The apparatus according to claim 1, wherein the volume filling device comprises two or more holding devices adapted to be held by an instrument and simplify the implantation of the device.

61. The apparatus according to claim 1, wherein the volume filling device 20 comprises an inflatable chamber, further comprising at least one tube connected to the device for supplying fluid to the chamber.

62. The apparatus according to claim 61, further comprising an injection port connectible with the tube.

63. The apparatus according to claim 1, wherein the volume filling device 25 has a maximum circumference as seen in a plane perpendicular to an axis through the device, and wherein the circumferences of the device as seen in other planes

perpendicular to said axis are equal to the maximum circumference or decrease as seen along said axis in the direction from the maximum circumference.

64. The apparatus according to claim 1, wherein the circumference of the volume filling device as seen in a plane perpendicular to an axis through the device increases and decreases at least two times as the plane is displaced along said axis, or decreases and increases at least one time as the plane is displaced along said axis.

65. The apparatus according to claim 11, wherein one layer is made of silicon.

66. The apparatus according to claim 11, wherein one layer is made of 10 PTFE.

67. The apparatus according to claim 11, wherein the coating is a metal coating.

68. The apparatus according to claim 11, wherein one layer is made of metal.

15 69. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 36, and 58, wherein the volume filling device comprising; - a stretching device comprising at least one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall and - an operation device for operating the stretching device when implanted to stretch 20 the stomach wall portion such that satiety is created.

70. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37, and 57, wherein the volume filling device comprising; at least one operable stretching device implantable in the patient and adapted to stretch a portion of the patient's stomach wall, and an implantable control unit for automatically controlling 25 the operable stretching device, when the control unit and stretching device are implanted, to stretch the stomach wall portion in connection with the patient eating such that satiety is created.

71. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37, and 57, wherein the volume filling device comprising;- a stretching device comprising at least one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall, wherein said stretching device comprising an expandable stretching reservoir and
- 5 - an operation device for operating the stretching device when implanted to stretch the stomach wall portion, wherein the volume filling device is inflatable and in fluid connection with said stretching reservoir, wherein said operation device comprises a pump for pumping fluid between said main reservoir and said stretching reservoir to stretch said stomach wall portion such that satiety is created.
- 10
72. An apparatus for treating obesity according to claim 71, comprising a control device for controlling said stretching device including said pump.
73. An apparatus for treating obesity according to claim 72, wherein said control device comprising a wireless remote control adapted to control the stretching device from the outside of the patient's body.
- 15
74. An apparatus for treating obesity according to claim 72, wherein said control device comprising a subcutaneously placed switch or reservoir adapted to control the stretching device from the outside of the patient's body.
75. An apparatus for treating obesity according to claim 72, wherein said 20 control device comprising an implantable control unit for controlling said stretching device.
76. The device according to claim 75, comprising a sensor or sensing device adapted to be implanted in the patient body, wherein the implantable control unit is adapted to control the stretching device from the inside of the patient's body 25 using information from said a sensor or sensing device, adapted to sense, direct or indirect, the food intake of the patient.
77. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37, and 57, wherein the volume filling device comprising;- a main volume filling reservoir, and- a stretching device comprising at least one operable stretching device

implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall, wherein said stretching device comprising an expandable reservoir, adapted to be invaginated in the stomach wall at the upper part of the stomach, higher up than the inflatable main volume filling device when the patient is standing,
5 wherein the volume filling device is inflatable and in fluid connection with said stretching reservoir, wherein normal contractions of the stomach wall, related to food intake, cause fluid to flow from said invaginated main volume filling reservoir lower placed onto the stomach wall adapted to cause said stretching reservoir to stretch said stomach wall portion such that satiety is created.

10 78. An apparatus for treating obesity according to claim 77, wherein the fluid connection between the main volume filling device reservoir and the stretching reservoir comprises a non-return valve.

15 79. An apparatus for treating obesity according to claim 77, wherein the fluid connection between the main volume filling device reservoir and the stretching reservoir comprises a release function adapted to release the volume in the stretching reservoir back to the main volume filling device reservoir.

20 80. apparatus for treating obesity according to claim 77, wherein said release function comprises a fluid return connection of a substantially smaller area than said fluid connection, to slowly release back fluid to said main volume filling device reservoir from the stretching reservoir to release said stretching of the stomach wall portion.

25 81. An apparatus for treating obesity according to anyone of claim 77, comprising a further manual control device comprising a subcutaneously placed reservoir adapted to control the stretching device from the outside of the patient's body to further affect the stretching device to stretch the stomach wall portion.

82. The device according to anyone of claim 1, 2, 11, 37, and 57, comprising a main volume filling device reservoir adapted to be inflatable, the device further comprising an expandable structure, adapted to expand, when the device is invaginated in the stomach wall, wherein said structure comprising a bellow adapted

to take into account the fibrosis surrounding the device when implanted, such that the movement of the bellow is substantially un-affected of said fibrosis.

83. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37, and 57, wherein the volume filling device comprising;

5 a stretching device comprising at least one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall and wherein the stretching device comprising a expandable structure, adapted to expand and stretch the stomach wall portion, when the device being invaginated in the stomach wall, wherein said structure comprising a special bellow adapted to take into
10 account the fibrosis surrounding the device when implanted, such that the movement of the bellow is substantially un-affected of said fibrosis.

84. An apparatus for treating obesity according to claim 83, comprising;

- an operation device for operating the stretching device when implanted to stretch the stomach wall portion such that satiety is created.

15 85. An apparatus for treating obesity according to claim 83, wherein the volume filling device comprising; an implantable control unit for automatically controlling the operable stretching device, when the control unit and stretching device are implanted, to stretch the stomach wall portion in connection with the patient eating such that satiety is created.

20 86. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37, and 57, wherein the volume filling device comprising: a stretching device comprising at least one operable stretching device implantable in an obese patient and adapted to stretch a portion of the patient's stomach wall such that satiety is created.

25 87. An apparatus for treating obesity according to claim 86, wherein said control device comprising a wireless remote control adapted to control the stretching device from the outside of the patient's body.

88. An apparatus for treating obesity according to claim 86, wherein said control device comprising a subcutaneously placed switch or reservoir adapted to control the stretching device from the outside of the patient's body.

89. An apparatus for treating obesity according to claim 86, wherein said 5 control device comprising an implantable control unit for controlling said stretching device.

90. The device according to anyone of claim 89, comprising a sensor or sensing device adapted to be implanted in the patient body, wherein the implantable control unit is adapted to control the stretching device from the inside of the patient's 10 body using information from said a sensor or sensing device, adapted to sense, direct or indirect, the food intake of the patient.

91. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further is adapted to treat reflux disease and further comprising an implantable movement restriction device adapted 15 to be at least partly invaginated by the patient's stomach fundus wall and having an outer surface that includes a biocompatible material, wherein a substantial part of the outer surface of the movement restriction device is adapted to rest against the stomach wall without injuring the latter in a position between the patient's diaphragm and at least a portion of the lower part of the invaginated stomach fundus 20 wall, such that movement of the cardiac notch of the patient's stomach towards the patient's diaphragm is restricted, when the movement restriction device is invaginated, to thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's cardia sphincter muscle exerted from the patient's 25 abdomen, the movement restriction device having a size of at least 125 mm³ and a circumference of at least 15 mm.

92. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further is adapted to treat reflux disease and further comprising: - an implantable movement restriction device having an outer 30 surface including a biocompatible material, wherein the movement restriction device

- is adapted to rest with at least a part of its outer surface against the patient's stomach fundus wall, in a position between the patient's diaphragm and the fundus wall, such that movement of the cardiac notch of the patient's stomach towards the patient's diaphragm is restricted, when the movement restriction device is implanted in the
- 5 patient, to thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's cardia sphincter muscle exerted from the patient's abdomen, wherein the movement restriction device having a size of at least 125 mm³ and a circumference of at least 15 mm, and
- 10 a fixation device adapted to secure the movement restriction device in said position, when the movement restriction device is implanted.
93. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further is adapted to treat reflux disease and further comprising;
- 15 an implantable movement restriction device adapted to be at least partly invaginated by the patient's stomach fundus wall and having an outer surface that includes a biocompatible material, wherein a substantial part of the outer surface of the movement restriction device is adapted to rest against the stomach wall without injuring the latter in a position between the patient's diaphragm and at least a portion
- 20 of the lower part of the invaginated stomach fundus wall, such that movement of the cardiac notch of the patient's stomach towards the patient's diaphragm is restricted, when the movement restriction device is invaginated, to thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's cardia sphincter muscle
- 25 exerted from the patient's abdomen, the movement restriction device having a size of at least 125 mm³ and a circumference of at least 15 mm, further comprising a stretching device comprising at least one operable stretching device implantable in the obese patient and adapted to stretch a portion of the patient's stomach wall such that satiety is created.

94. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further is adapted to treat reflux disease and further comprising;

an implantable movement restriction device having an outer surface including a

5 biocompatible material, wherein the movement restriction device is adapted to rest with at least a part of its outer surface against the patient's stomach fundus wall, in a position between the patient's diaphragm and the fundus wall, such that movement of the cardiac notch of the patient's stomach towards the patient's diaphragm is restricted, when the movement restriction device is implanted in the patient, to

10 thereby prevent the cardia from sliding through the patient's diaphragm opening into the patient's thorax, so as to maintain the supporting pressure against the patient's cardia sphincter muscle exerted from the patient's abdomen, wherein the movement restriction device having a size of at least 125 mm³ and a circumference of at least 15 mm, and

15 a fixation device adapted to secure the movement restriction device in said position, when the movement restriction device is implanted, further comprising a stretching device comprising at least one operable stretching device implantable in the obese patient and adapted to stretch a portion of the patient's stomach wall such that satiety is created.

20 95. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further comprising; a stretching device comprising three or more mechanical parts engaged with different parts of the stomach wall, one part each, wherein said engagement includes suturing or stapling to the stomach wall or invaginating the mechanical parts in the stomach wall part

25 with stomach to stomach sutures, wherein the three or more mechanical parts are adapted to move in relation to each other adapted to stretch three different wall portions, the stretching device further adapted to having said wall portions stretched independently from each other both regarding;

force used for stretching the stomach wall portion as well as,

30 time periods the stretching is applied, and

when the stretching is applied.

96. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further comprising; a stretching device comprising two or more hydraulic parts engaged with different parts of the stomach wall, one part each, wherein said engagement includes suturing or stapling to hydraulic part to the stomach wall or invaginating the hydraulic parts in the stomach wall part, with stomach to stomach sutures, wherein the two or more hydraulic parts are adapted to move in relation to each other adapted to stretch three different wall portions, the stretching device further adapted to having said wall portions stretched independently from each other both regarding;

force used for stretching the stomach wall portion as well as,

time periods the stretching is applied, and

when the stretching is applied.

97. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further comprising; a stretching device is engaged with a part of the stomach wall, including suturing or stapling the stretching device to the stomach wall or invaginating the stretching device in the stomach wall part, with stomach to stomach sutures, wherein the stretching device is further adapted to stretch a stomach wall portion controlling;

force used for stretching the stomach wall portion as well as,

time periods the stretching is applied, and

when the stretching is applied.

98. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further comprising; a stretching device comprising two parts engaged with different parts of the stomach wall, one part each, wherein said engagement includes suturing or stapling the parts to the stomach wall or invaginating the parts in the stomach wall part, with stomach to stomach sutures,

wherein the stretching device further adapted to have different wall portions stretched independently from each other controlling;

force used for stretching the stomach wall portion as well as,

time periods the stretching is applied, and

5 when the stretching is applied.

99. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device further comprising; an external control unit for controlling the volume filling device from the outside of the patient's body.

100. The device according to claim 99, wherein the external control unit 10 comprising a wireless remote control adapted to control the device from the outside of the patient's body.

101. The device according to claim 99, wherein the external control unit comprising a subcutaneously placed switch or reservoir adapted to control the device from the outside of the patient's body.

102. An apparatus for treating obesity according to anyone of claim 1, 2, 11, 37 and 57, wherein the volume filling device, further comprising; comprising a sensor or sensing device adapted to be implanted in the patient body, wherein the implantable control unit is adapted to control the device from the inside of the patient's body using information from said a sensor or sensing device, adapted to sense, direct or indirect, the food intake of 20 the patient.

103. An apparatus for treating obesity of an obese patient having a stomach with a food cavity, the apparatus comprising:

at least one volume filling device adapted to be at least substantially invaginated by a stomach wall portion of the patient and having an outer surface that includes a

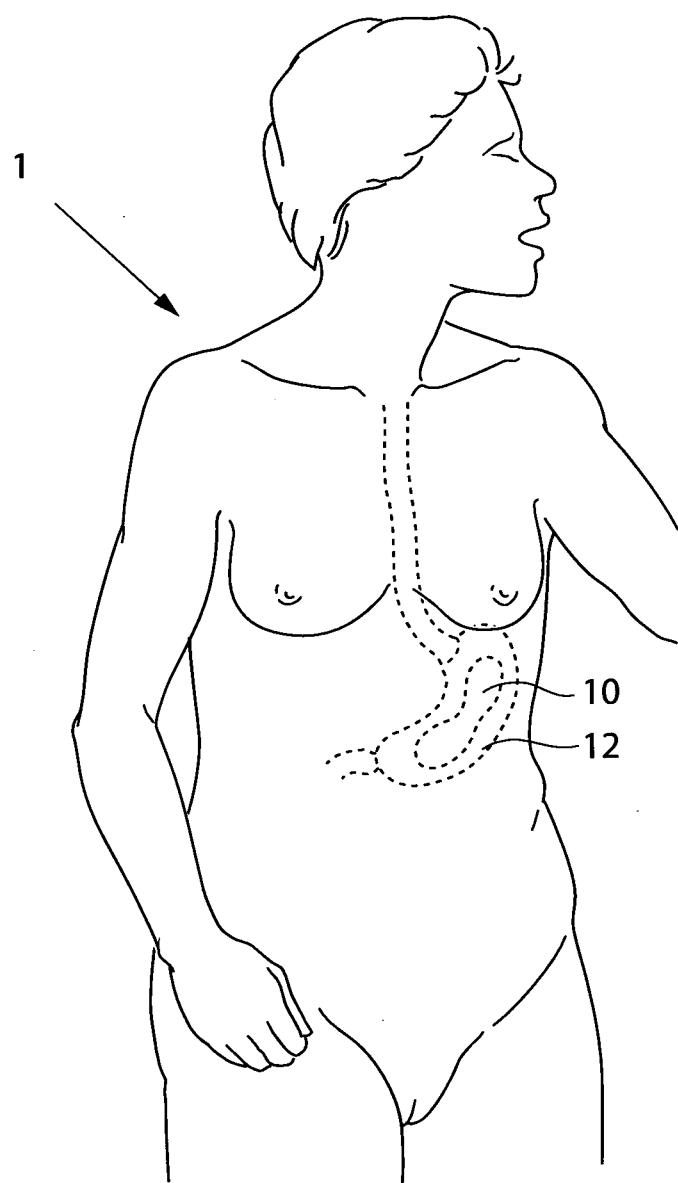
25 biocompatible material, and

wherein the volume filling device is adapted to be placed inside the stomach with the outer surface of the volume filling device resting against the inside of the stomach wall, such that the volume of the food cavity is reduced in size by a volume

substantially exceeding the volume of the volume filling device, the volume filling device having a maximum circumference of at least 30 millimeters.

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Fig.1



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Fig.2a

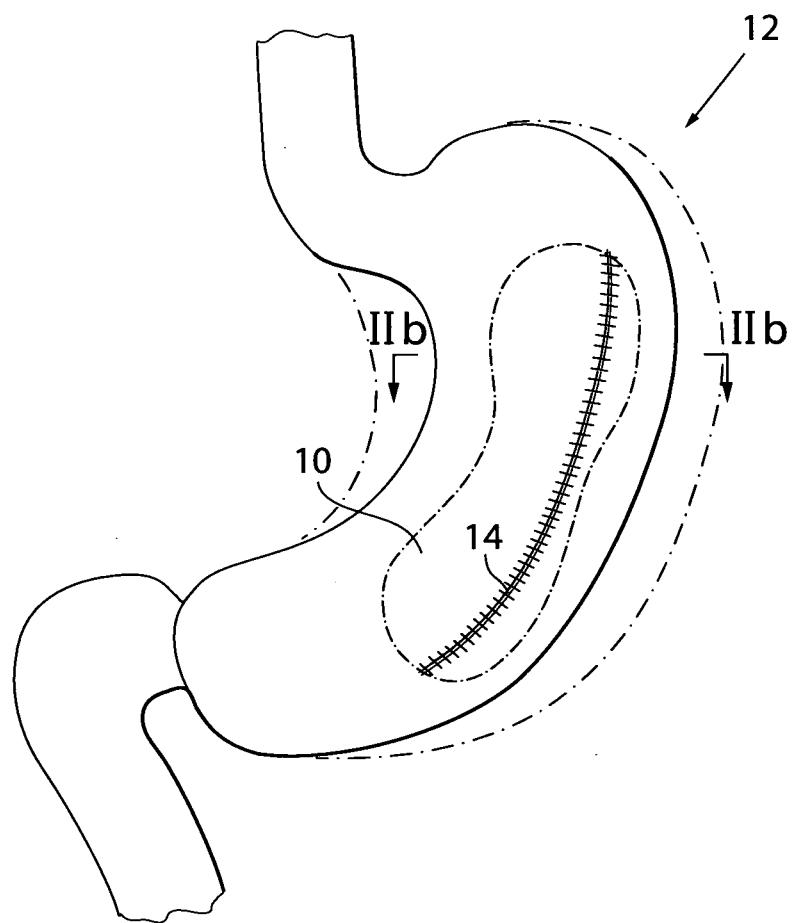
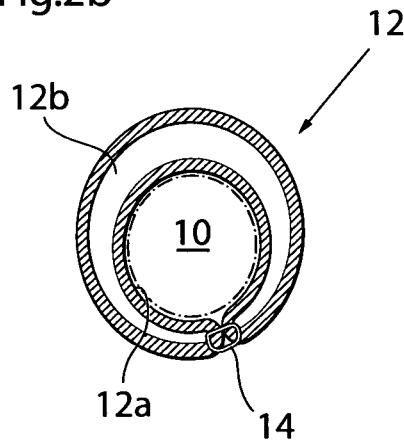


Fig.2b



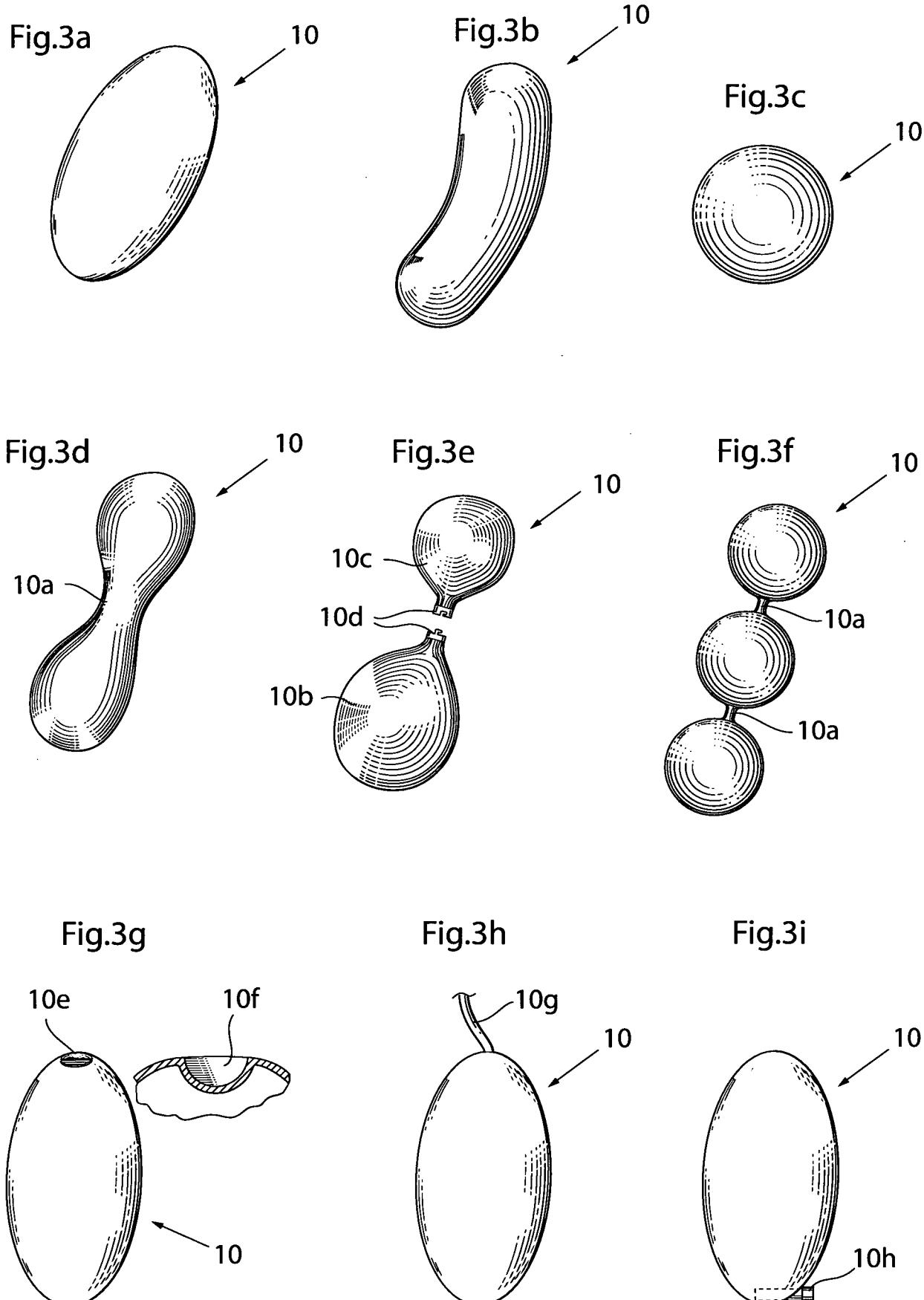


Fig.3j

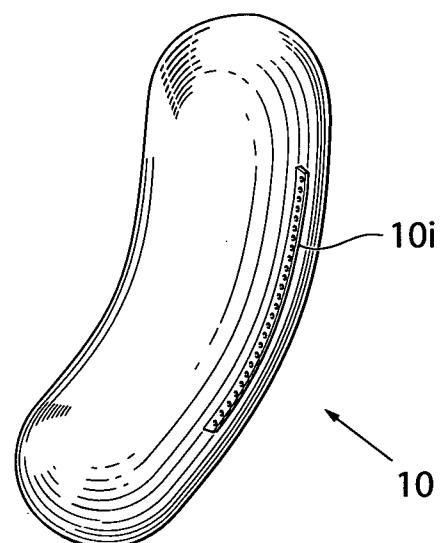


Fig.3k

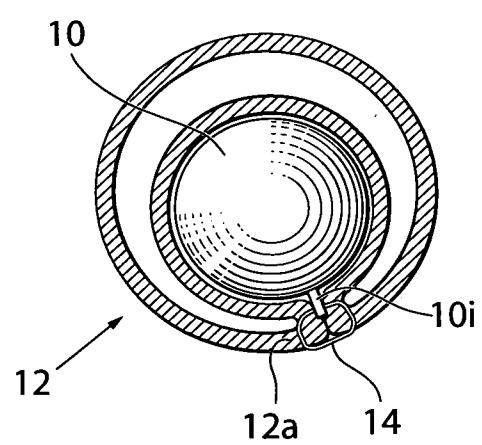


Fig.3m

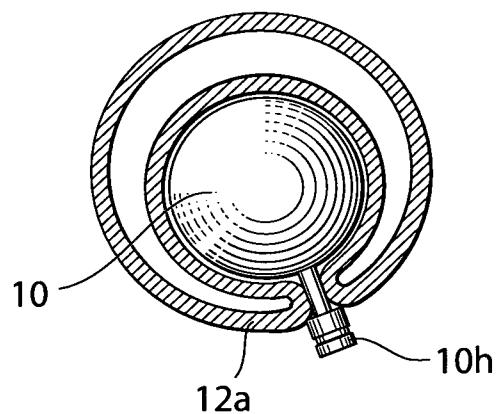


Fig.3n

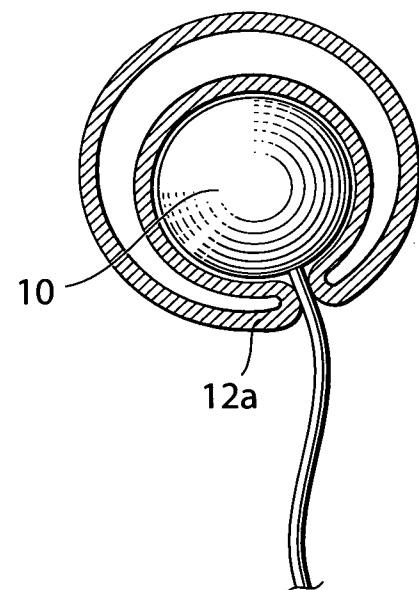


Fig.3p

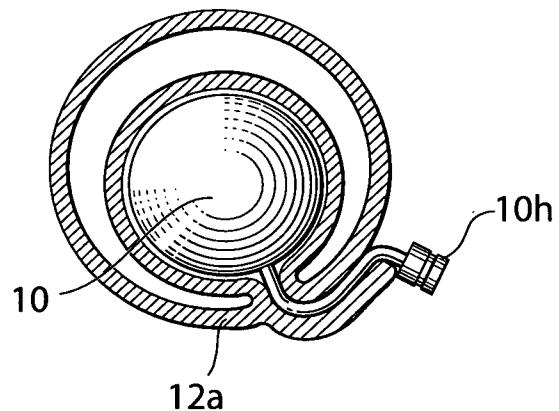


Fig.4a

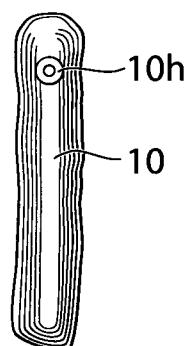


Fig.4b

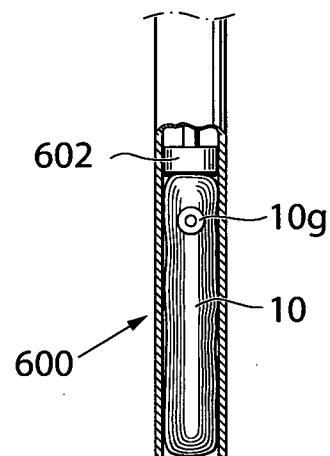
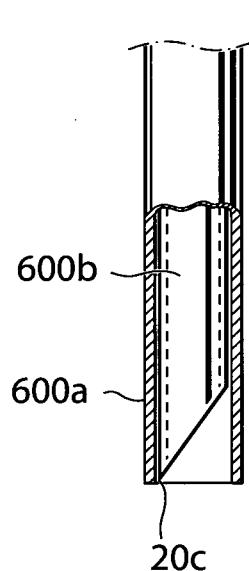


Fig.4c

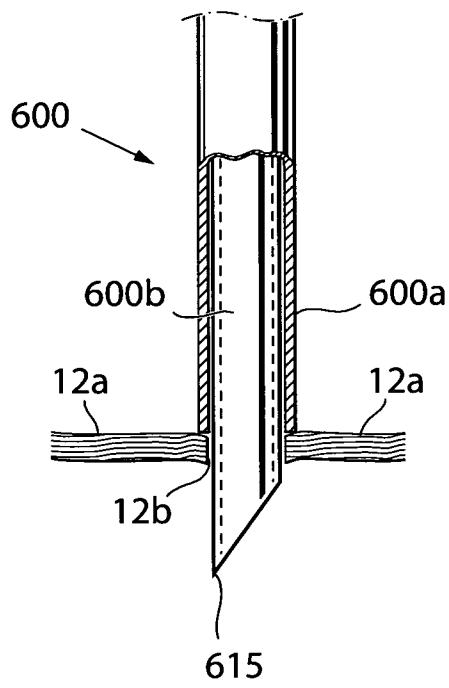


Fig.4d

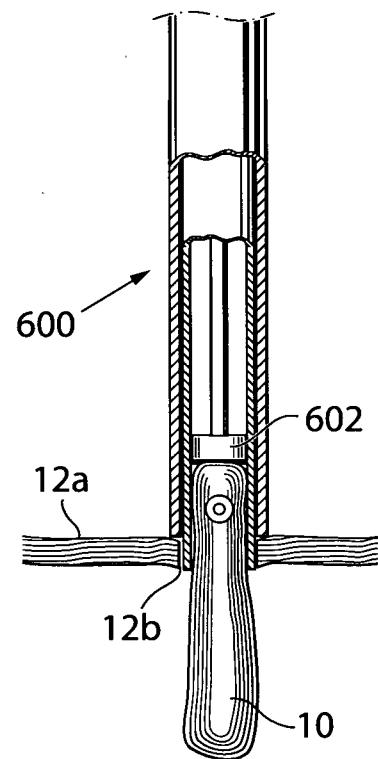


Fig.5a

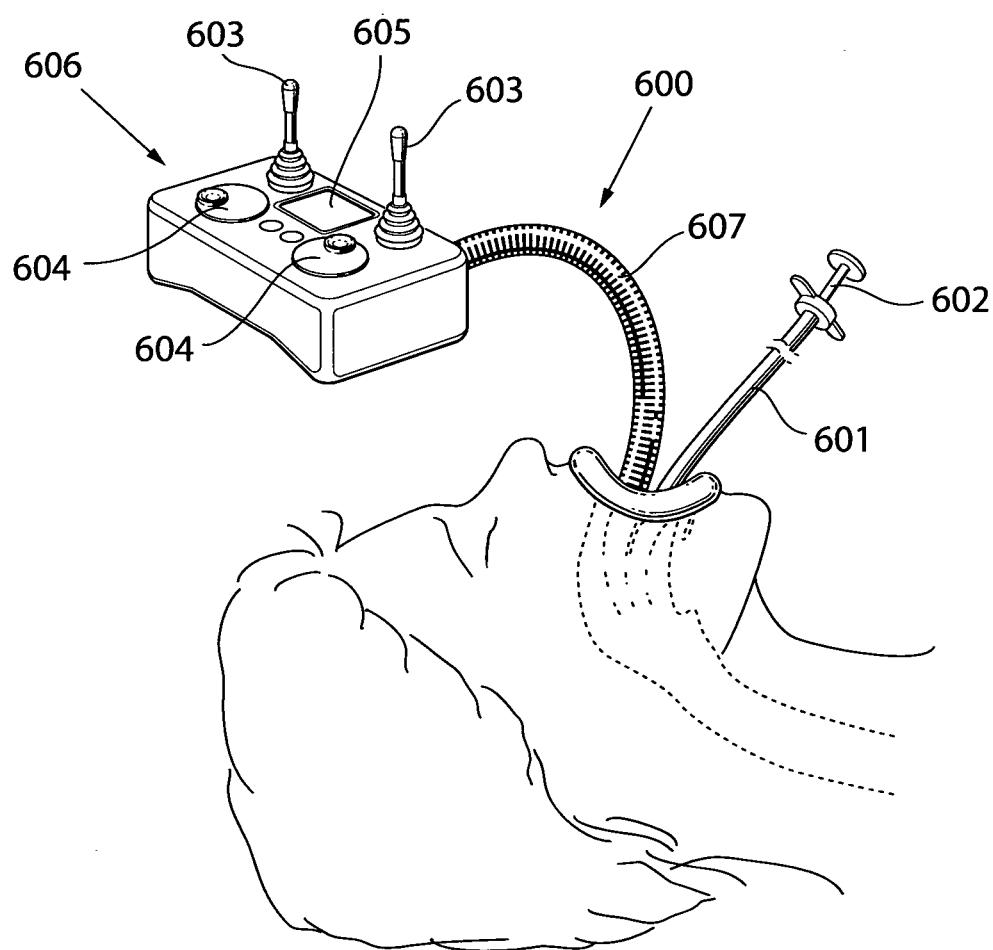


Fig.5b

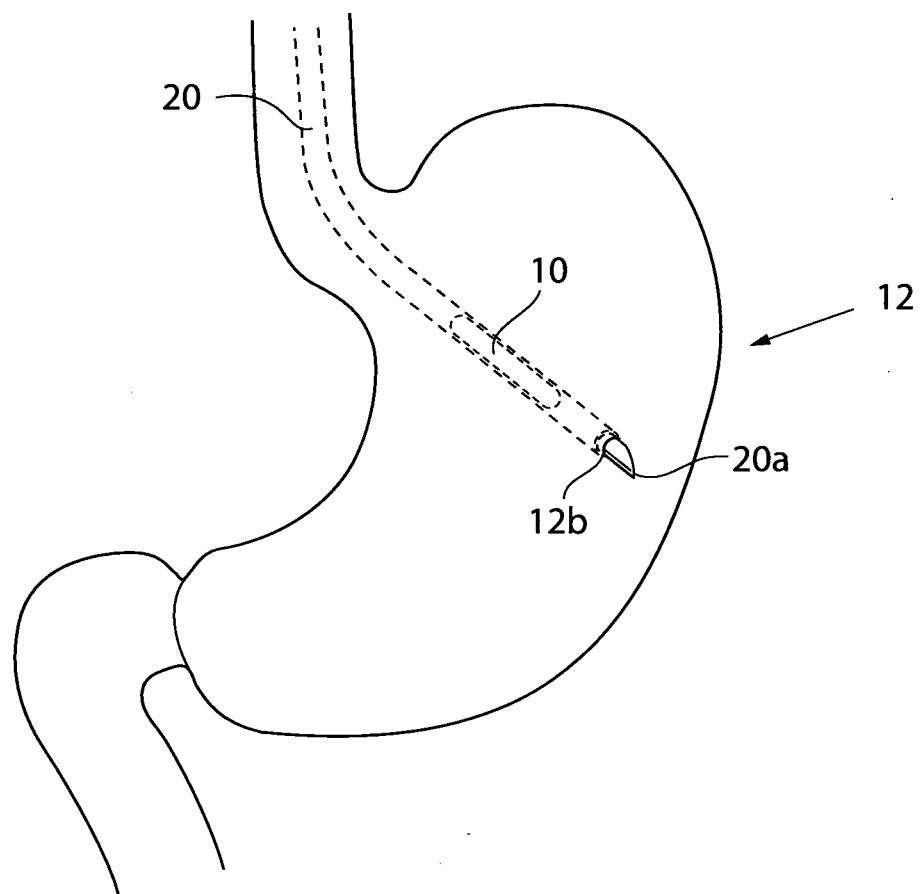


Fig.5c

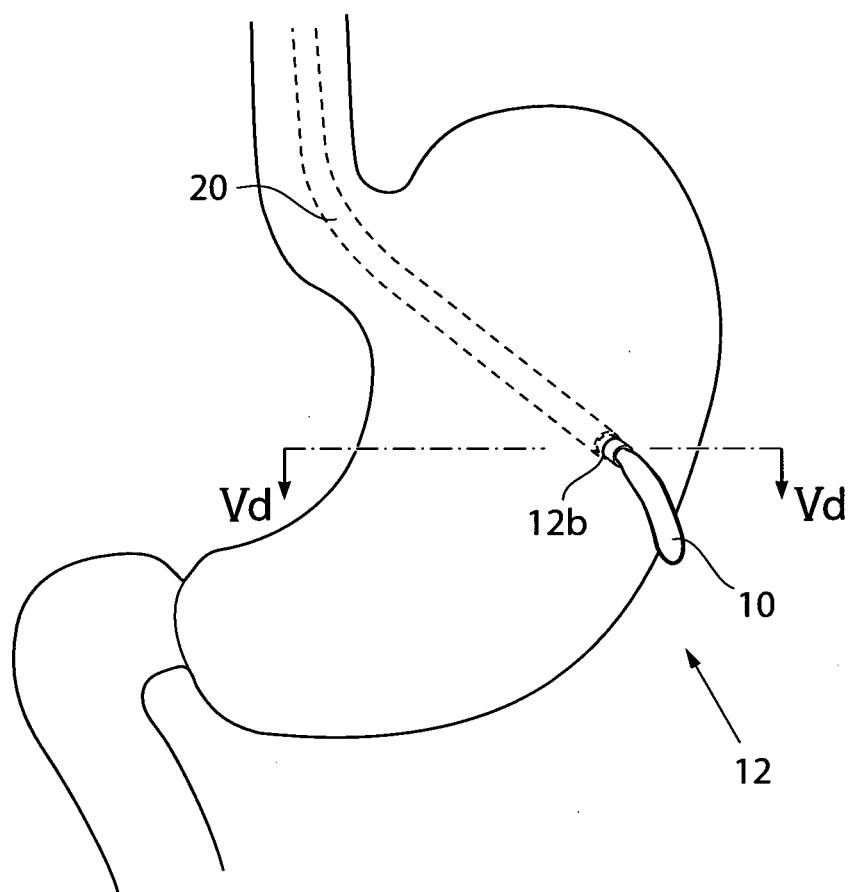
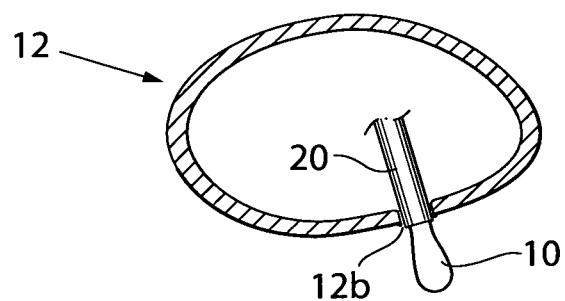


Fig.5d



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Fig.5e

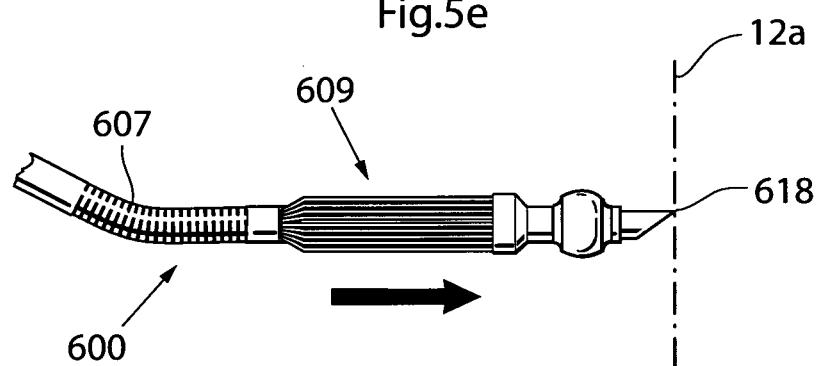


Fig.5f

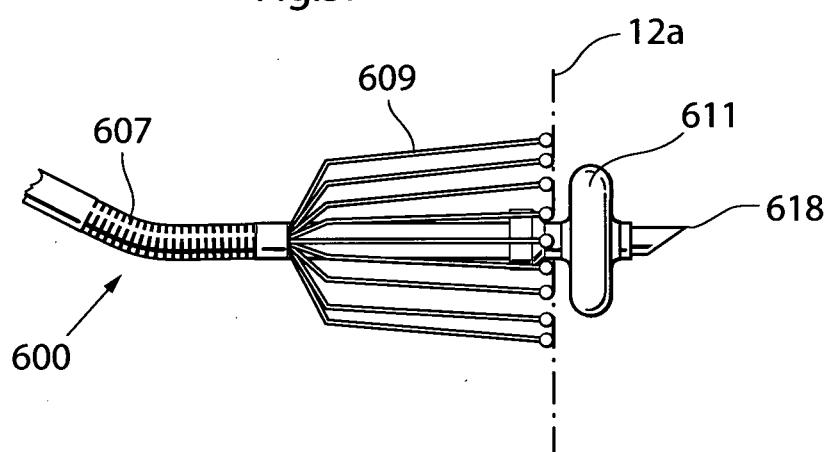


Fig.5g

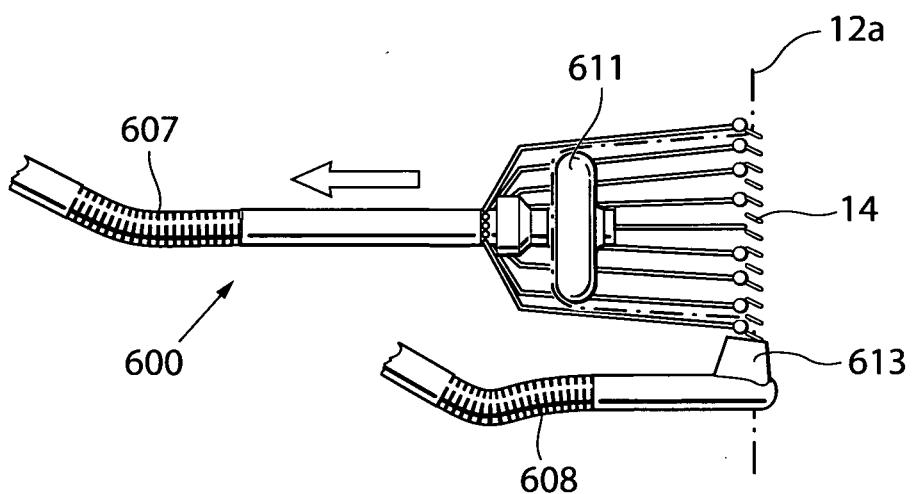


Fig.5h

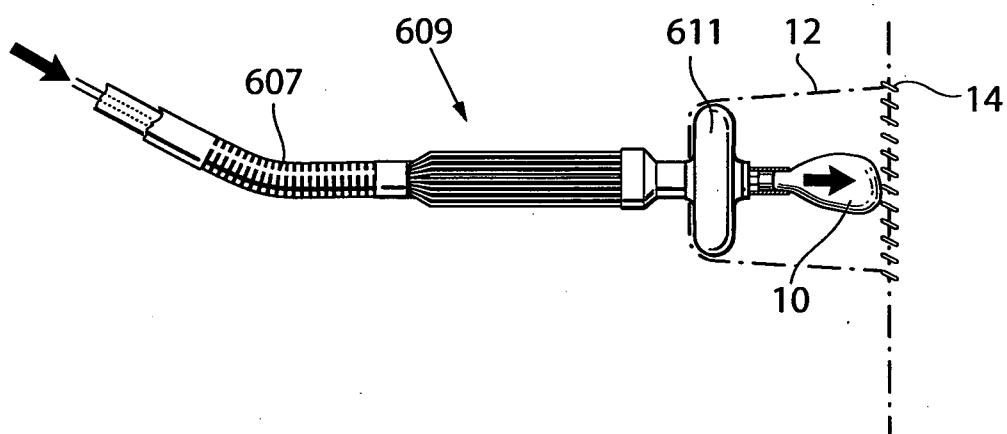
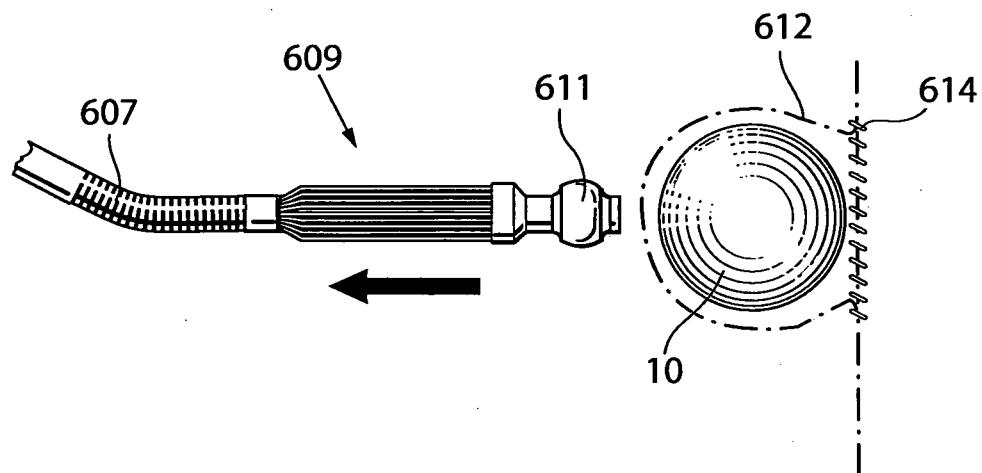


Fig.5i



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Fig.6a

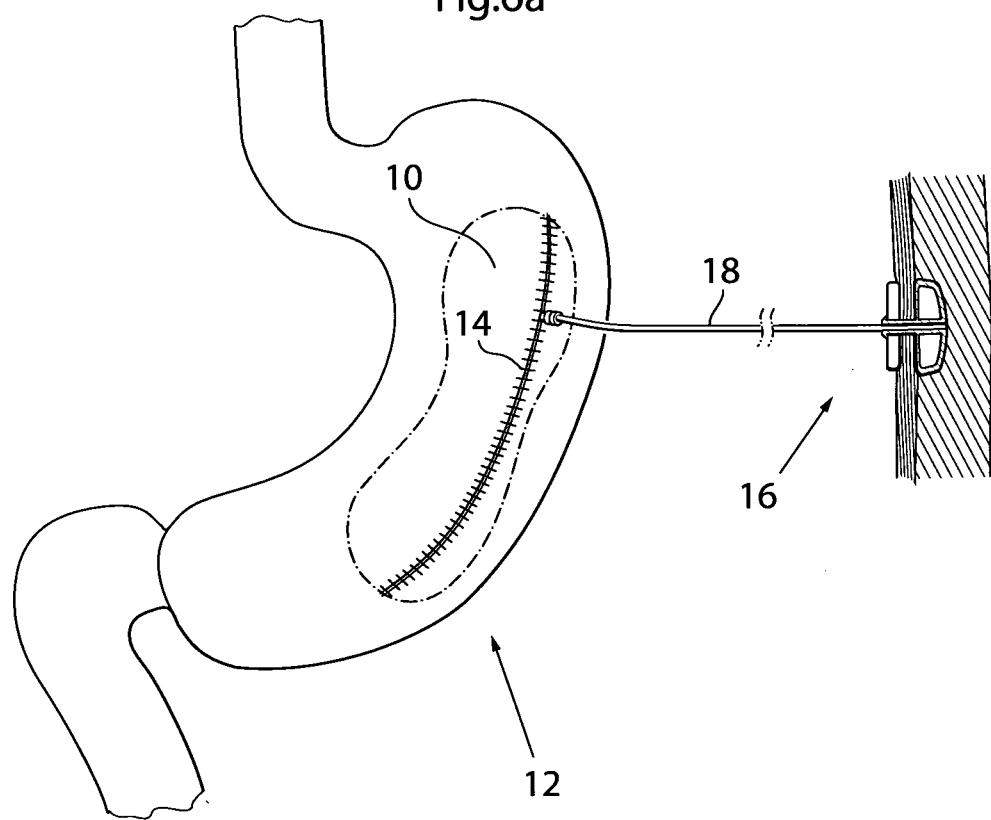


Fig.6b

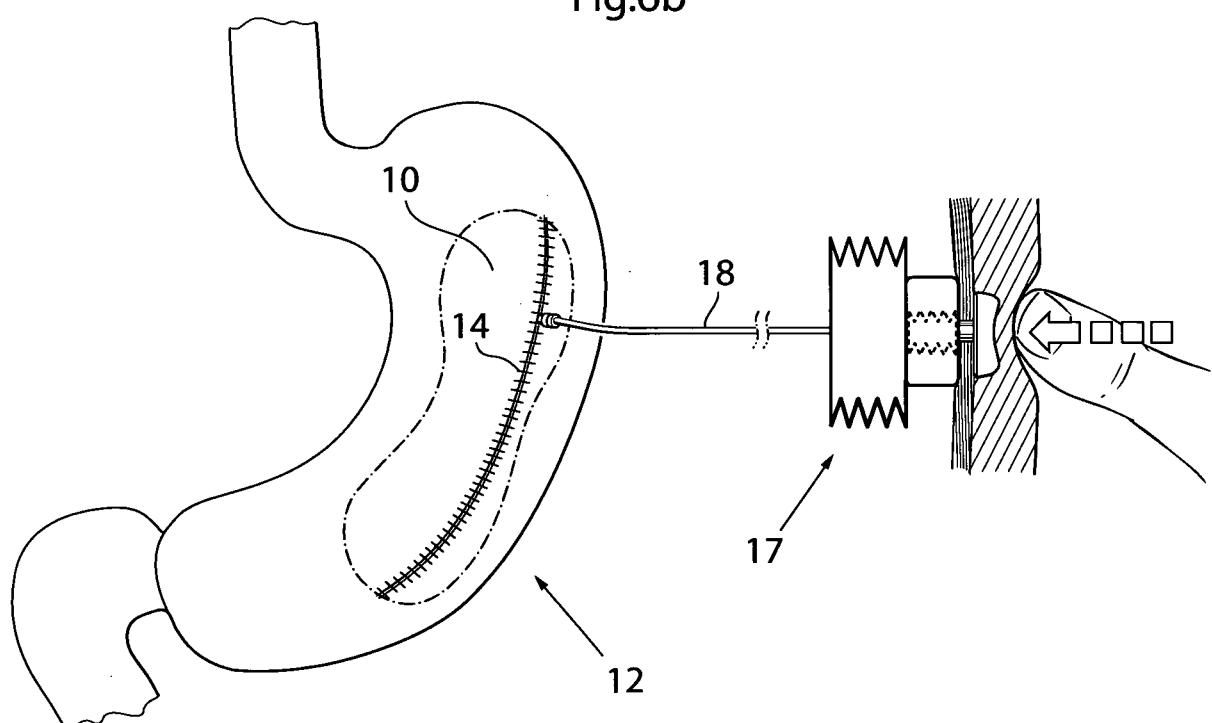


Fig.7

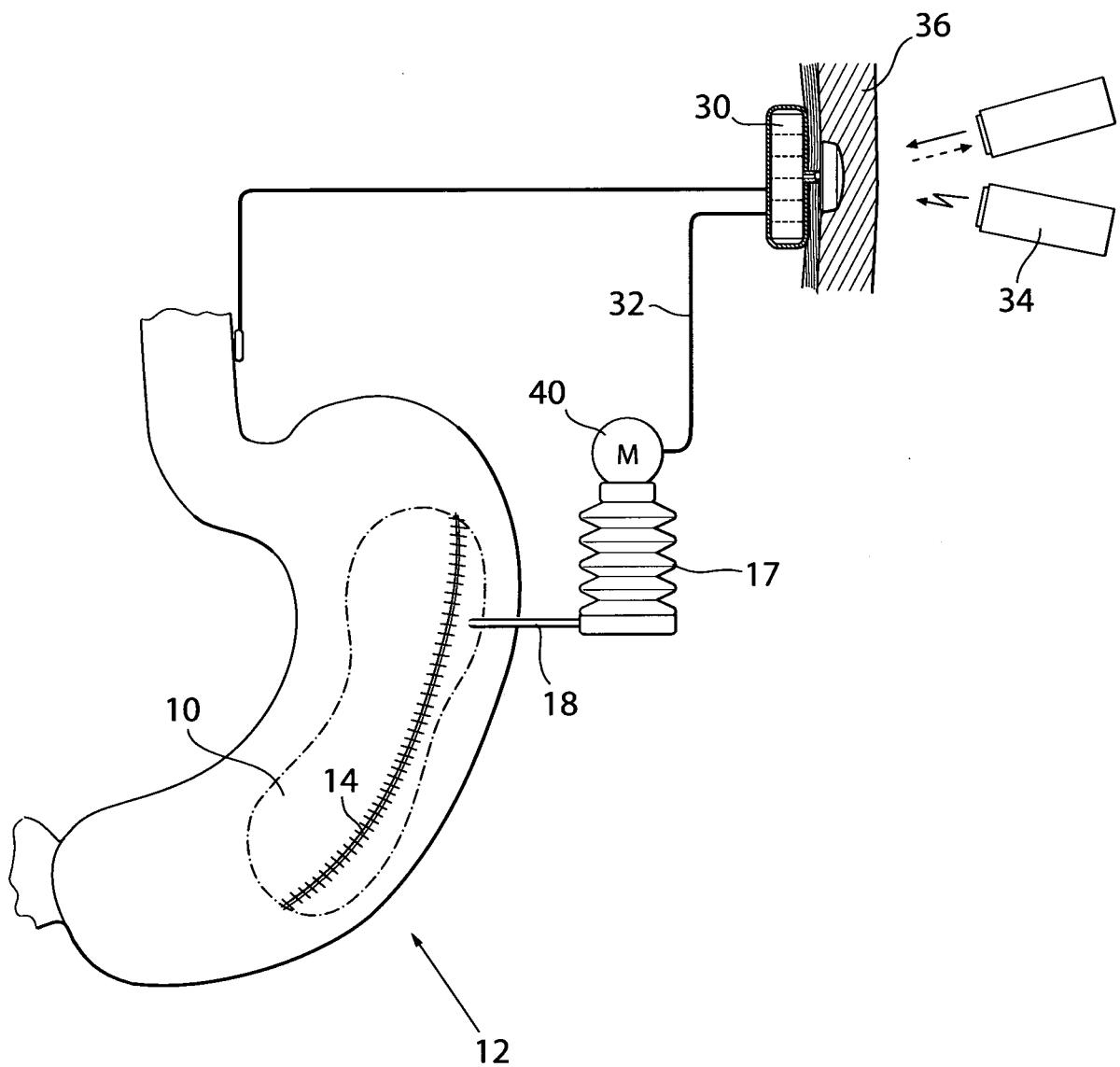


Fig.8

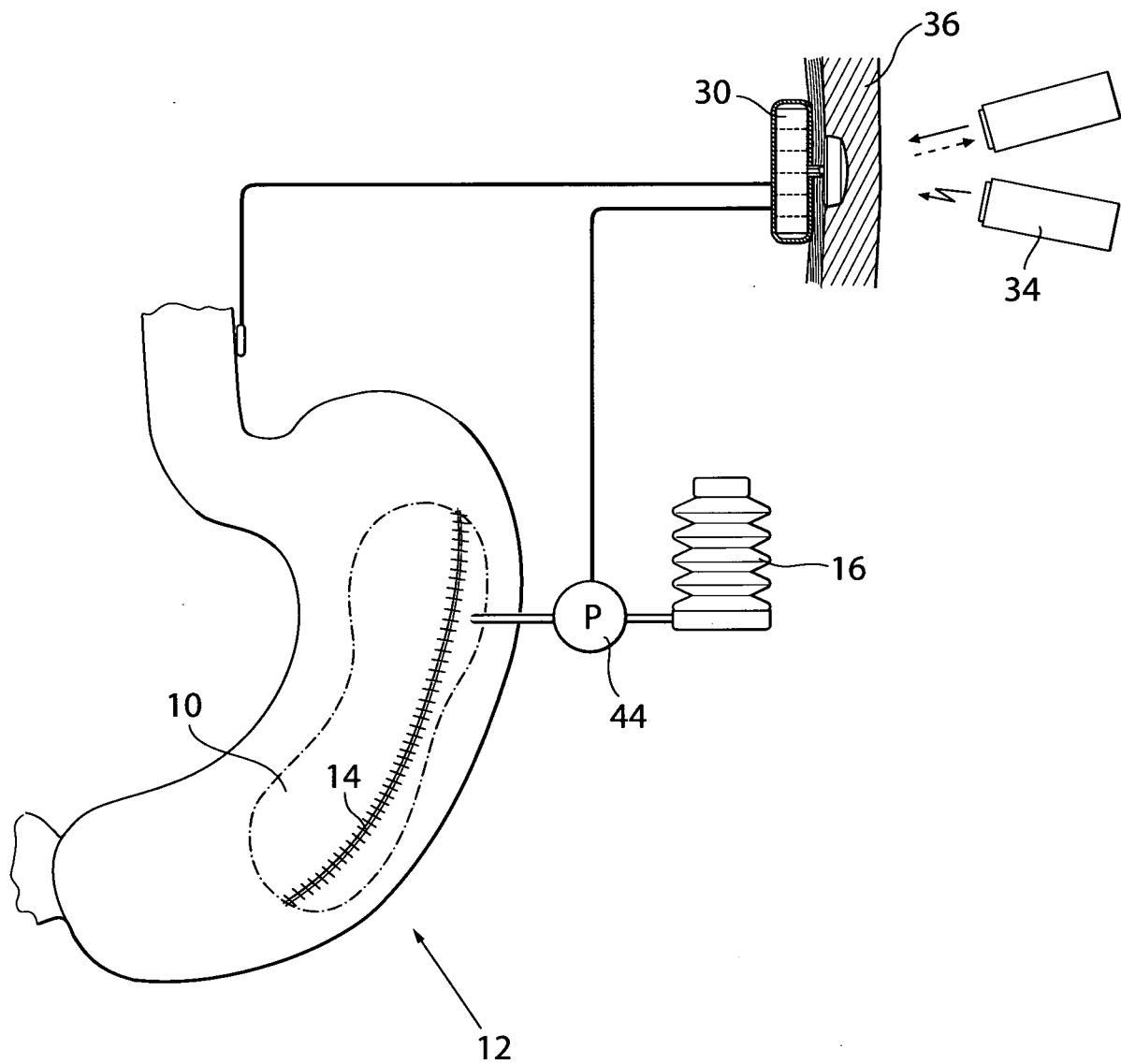


Fig.9

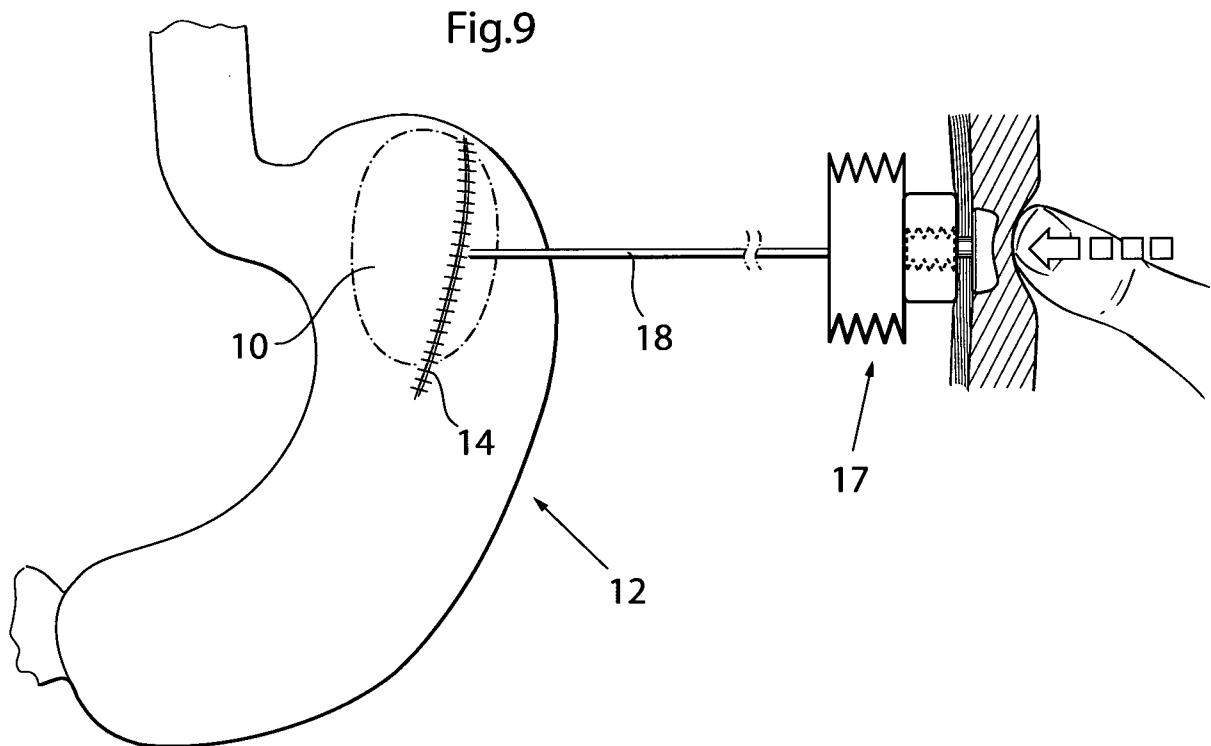


Fig.10

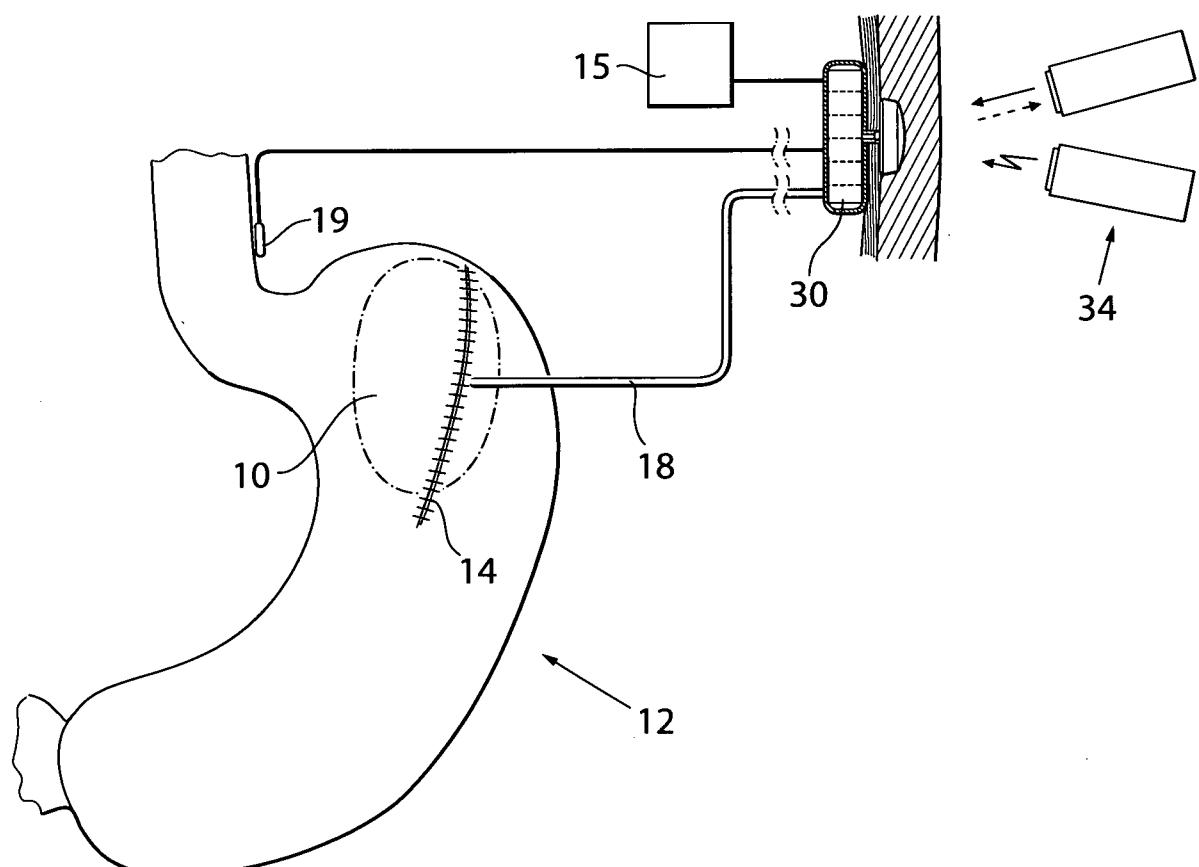


Fig.11

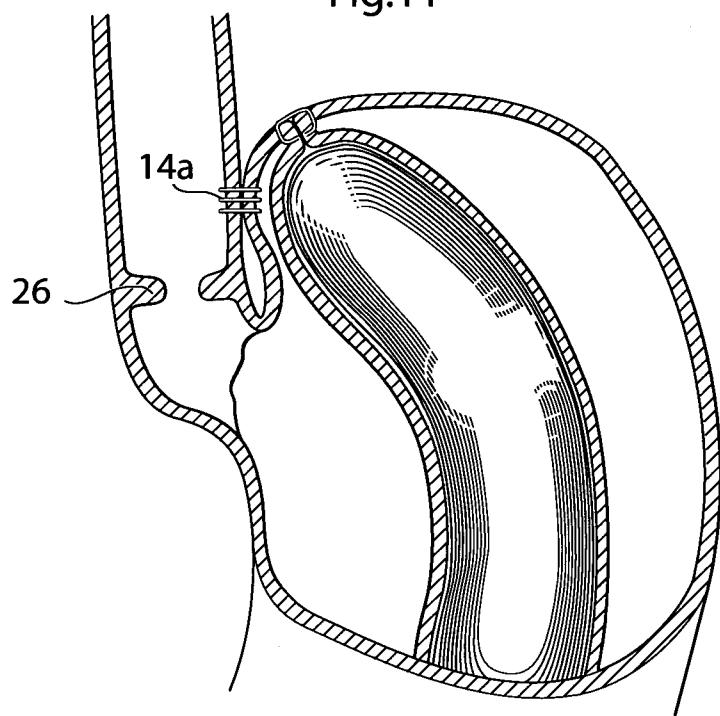
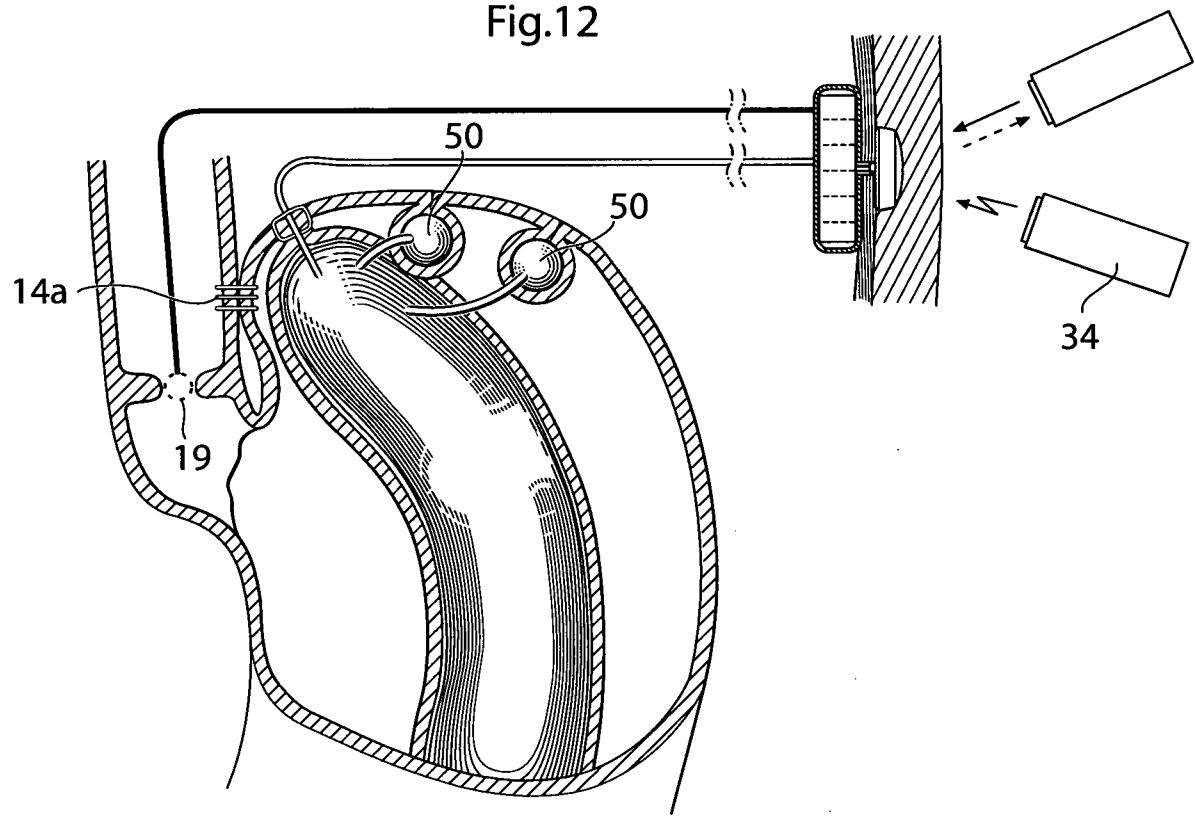


Fig.12



SUBSTITUTE SHEET (RULE 26)

Fig.13

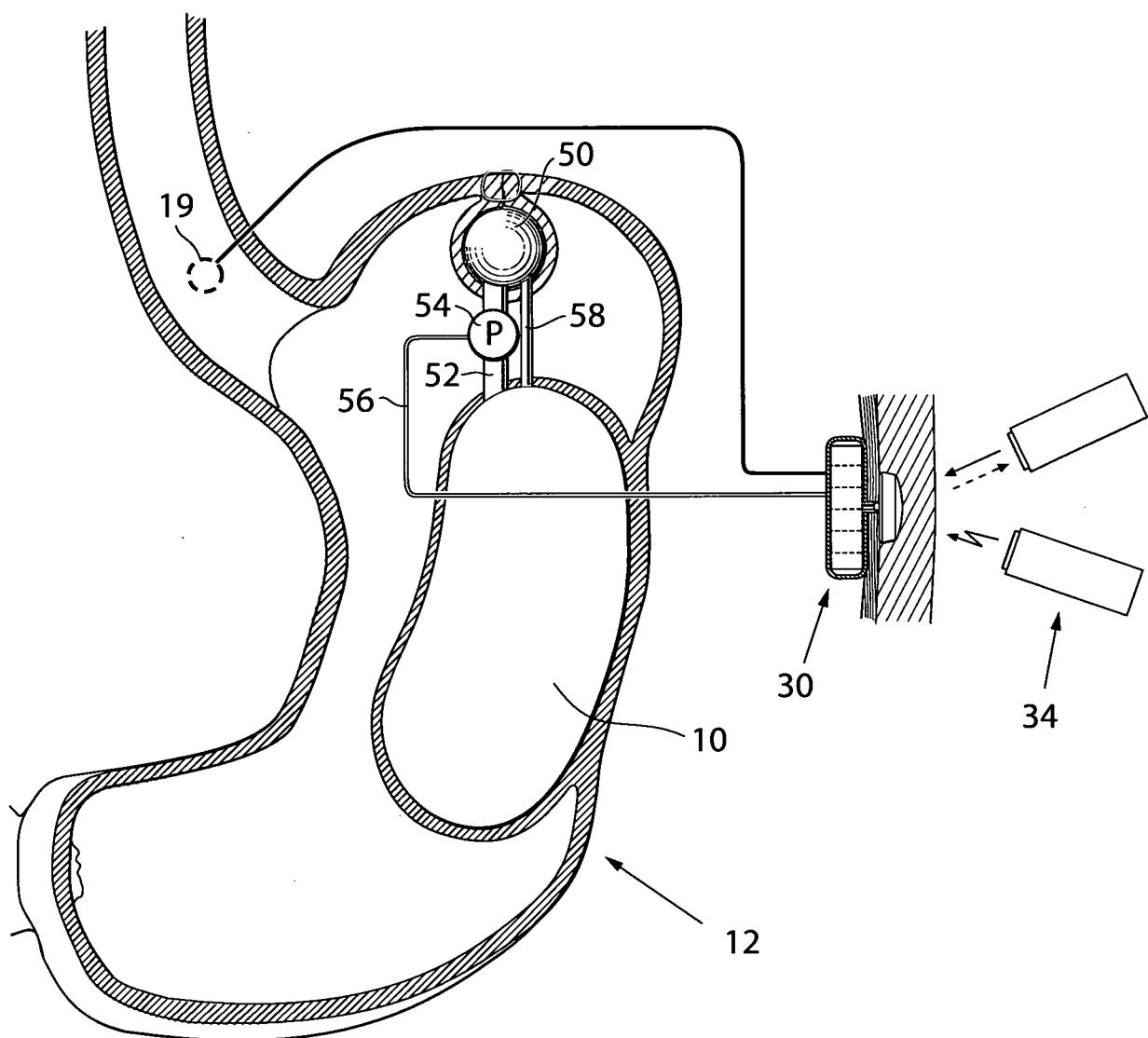


Fig.14

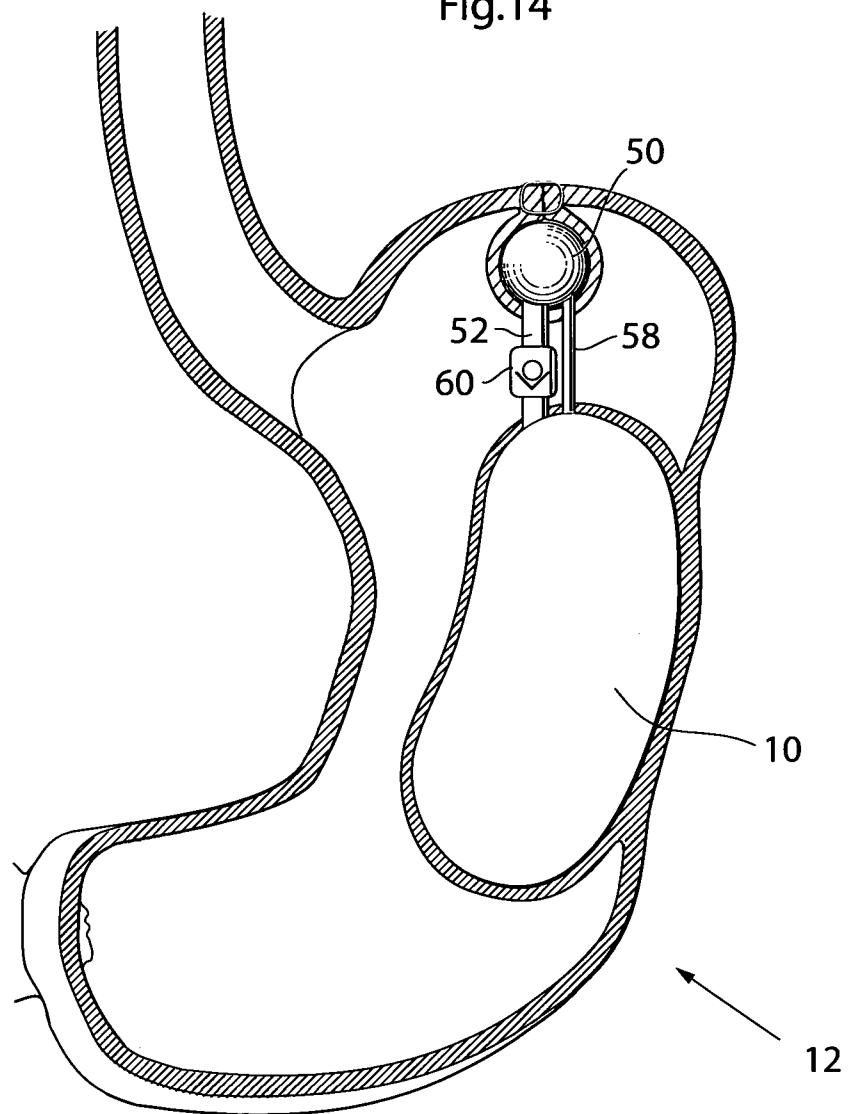


Fig.15

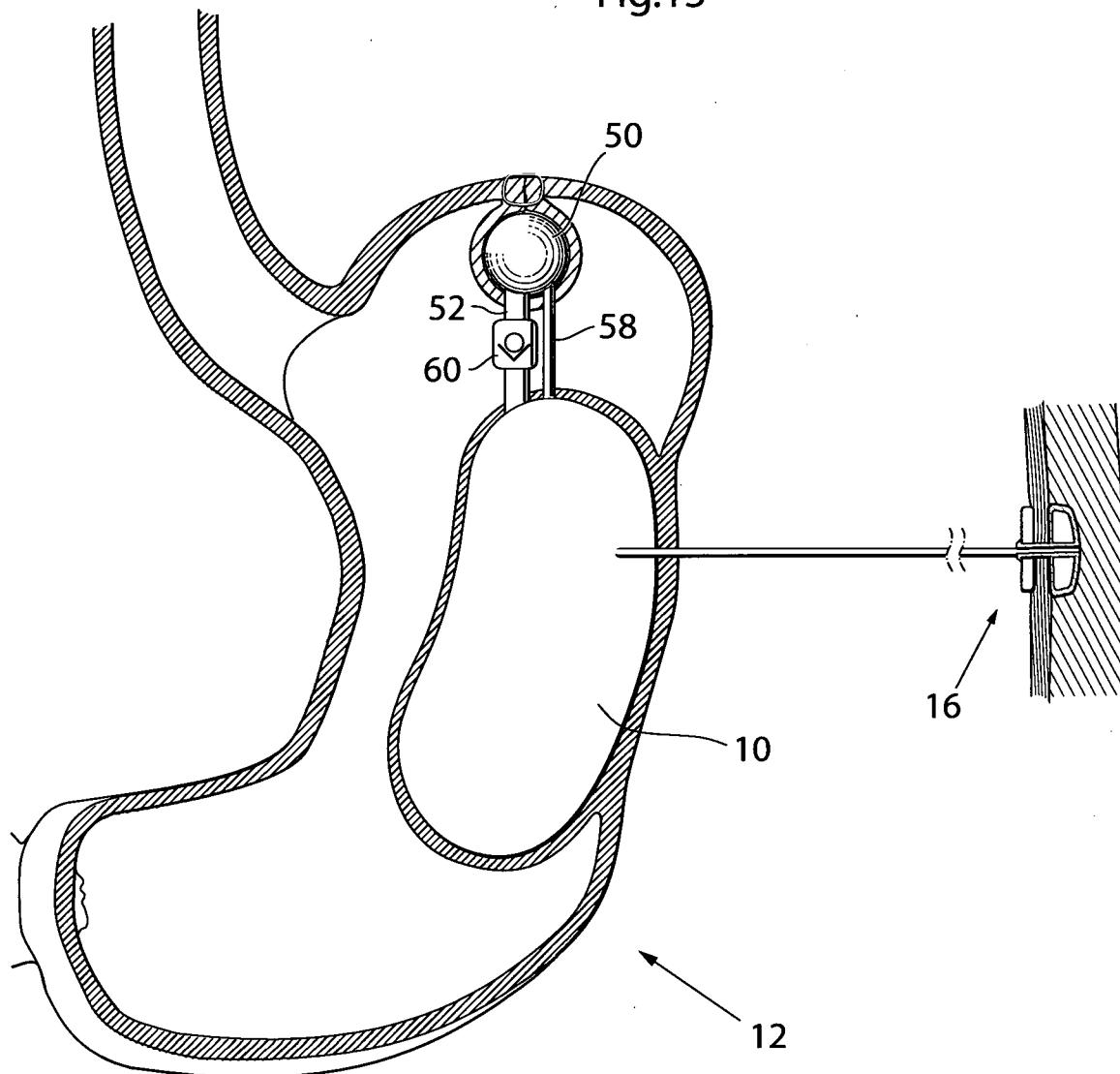


Fig.16

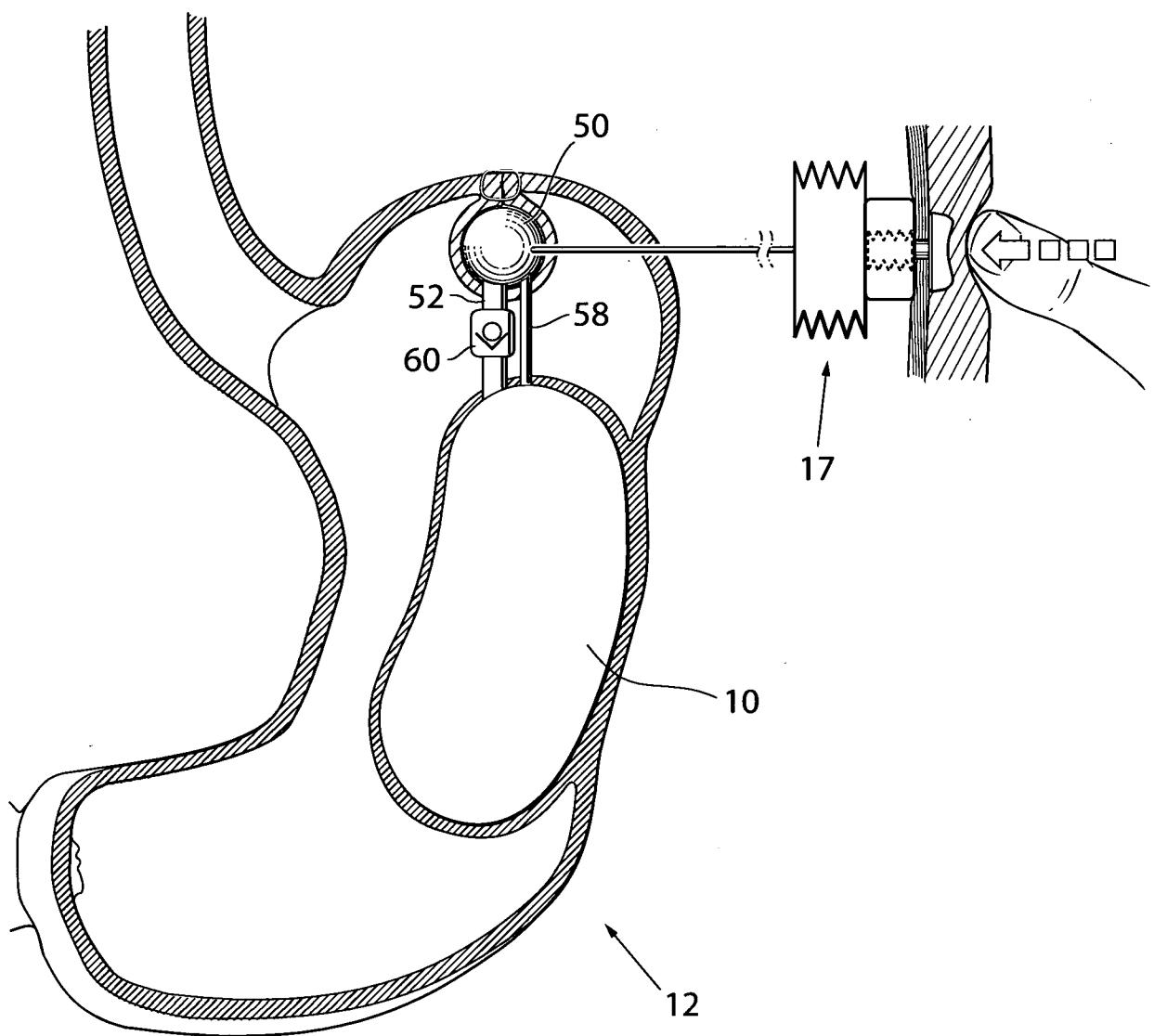


Fig.17a

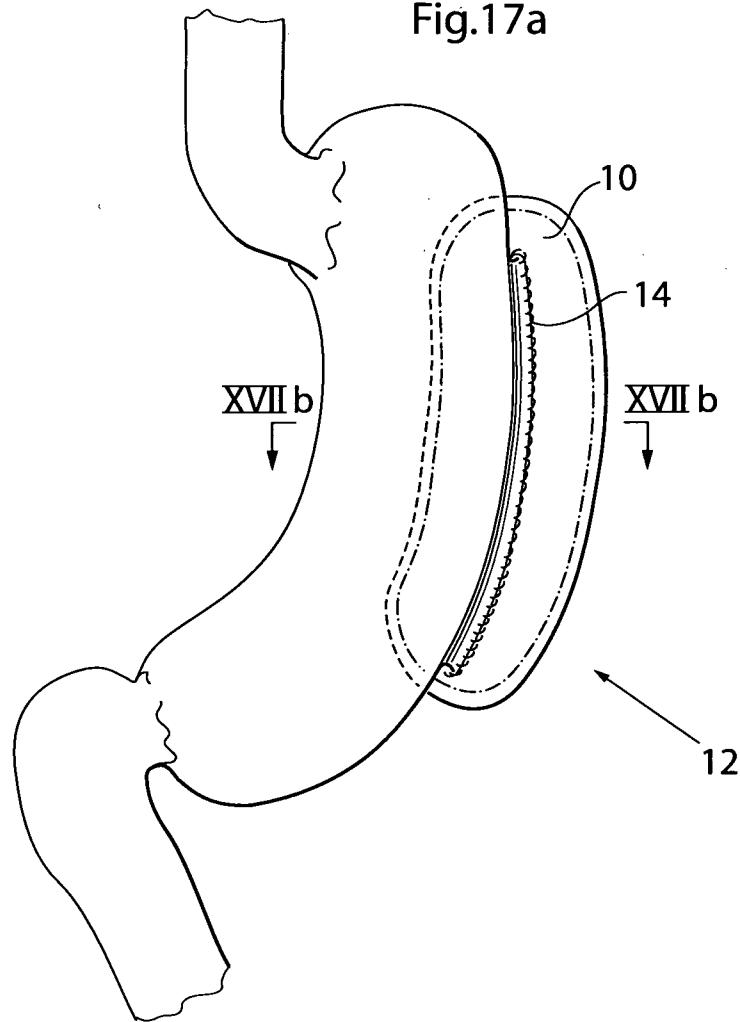
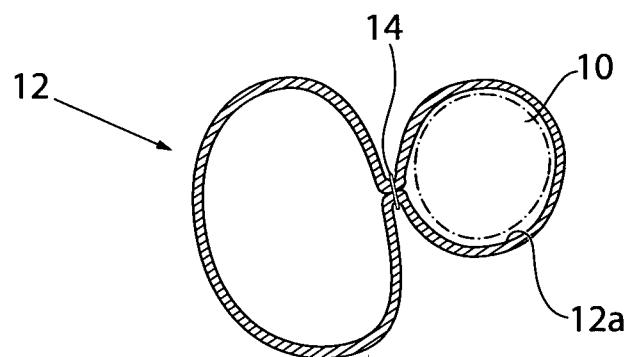


Fig.17b



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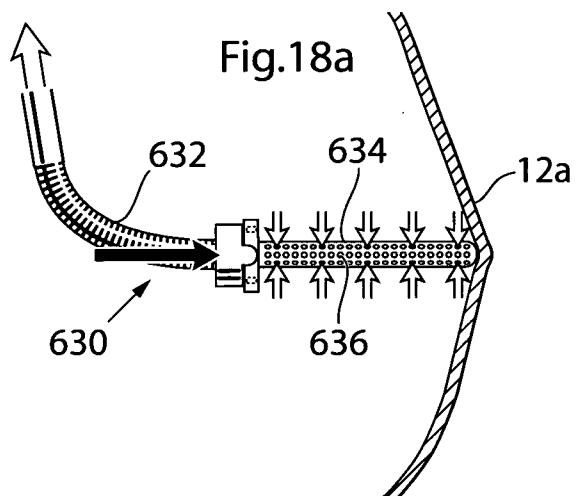


Fig.18a

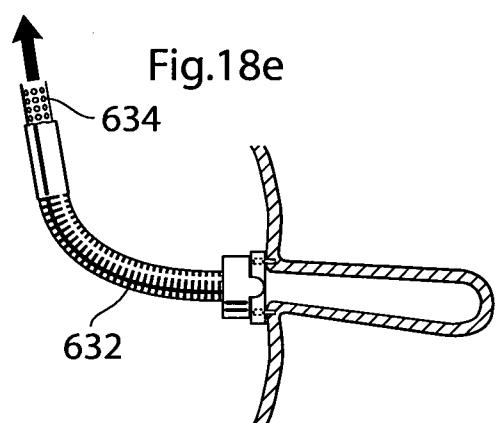


Fig.18e

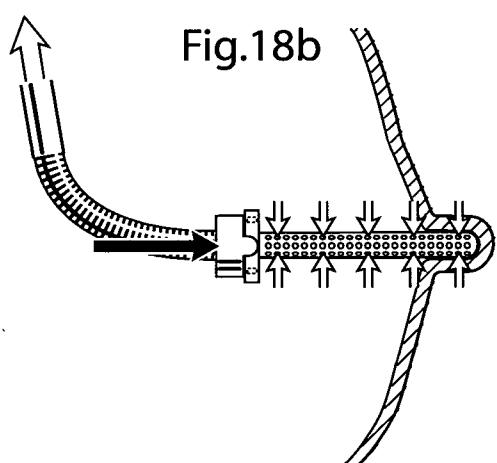


Fig.18b

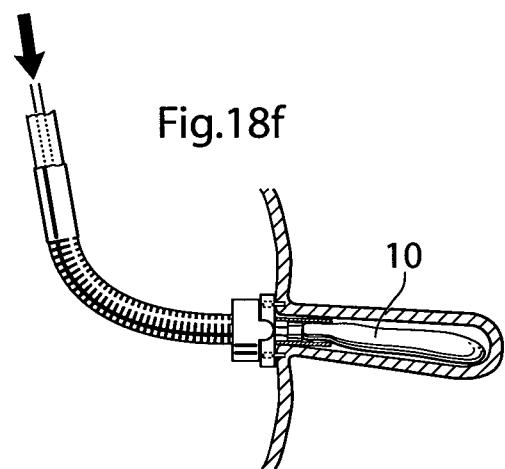


Fig.18f

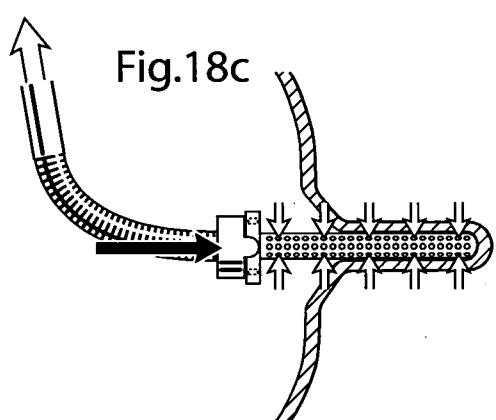


Fig.18c

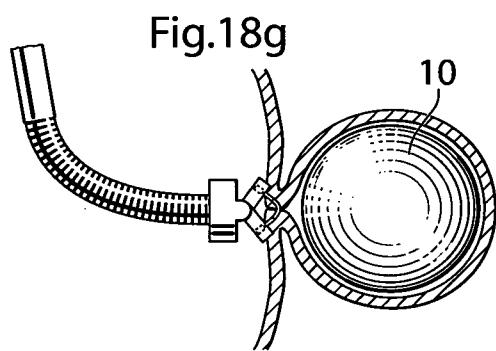


Fig.18g

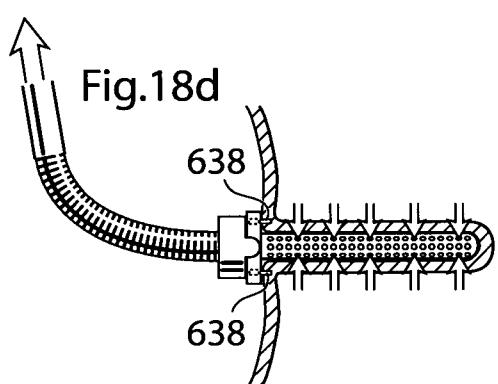


Fig.18d

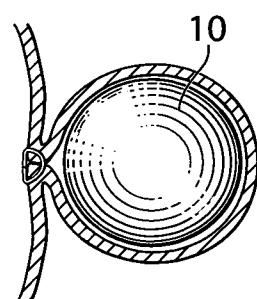


Fig.18h

Fig. 19a

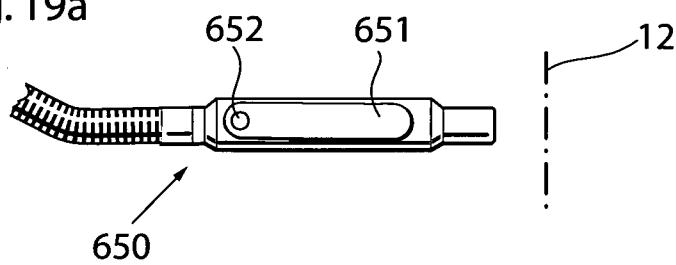


Fig. 19b

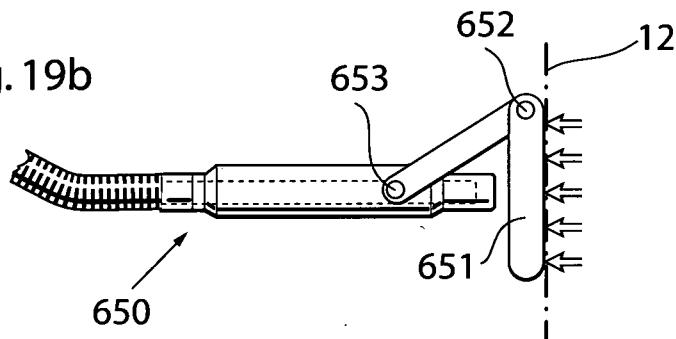


Fig. 19c

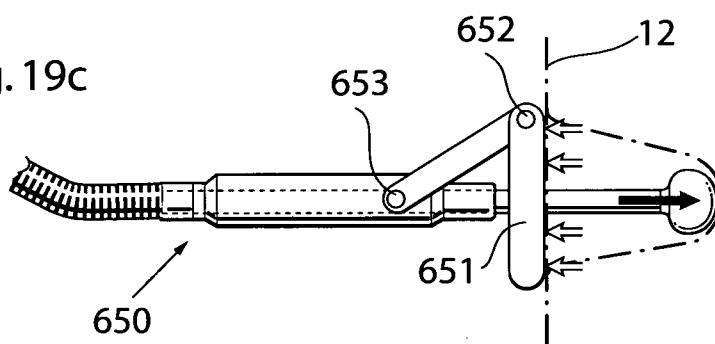


Fig. 19d

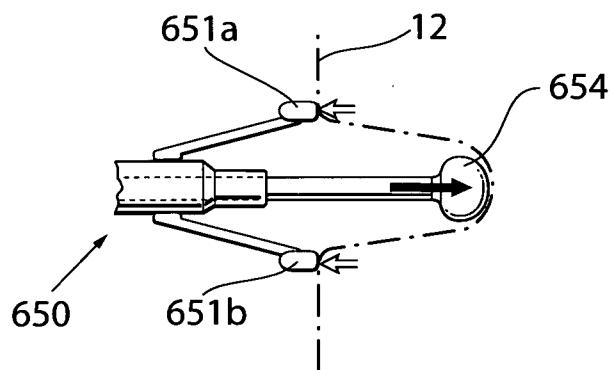
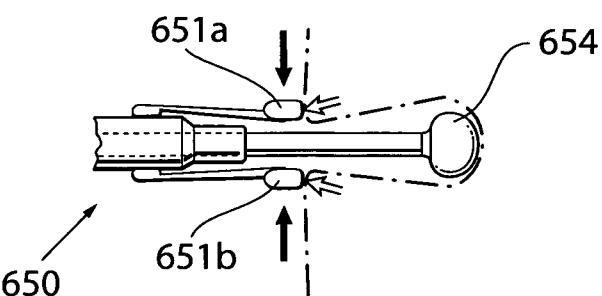


Fig. 19e



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Fig. 19f

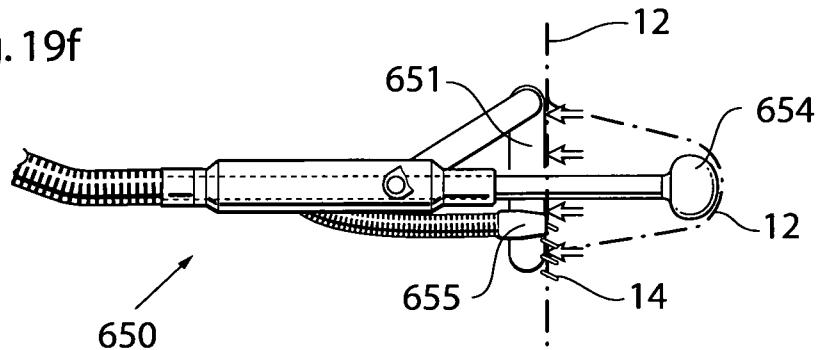


Fig. 19g

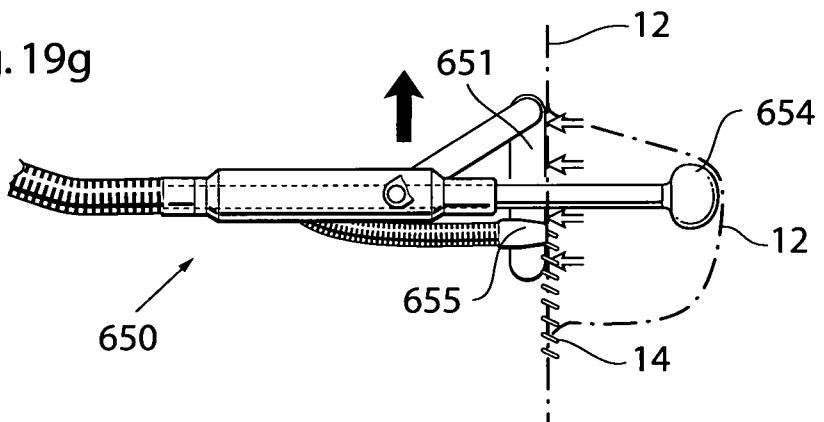


Fig. 19h

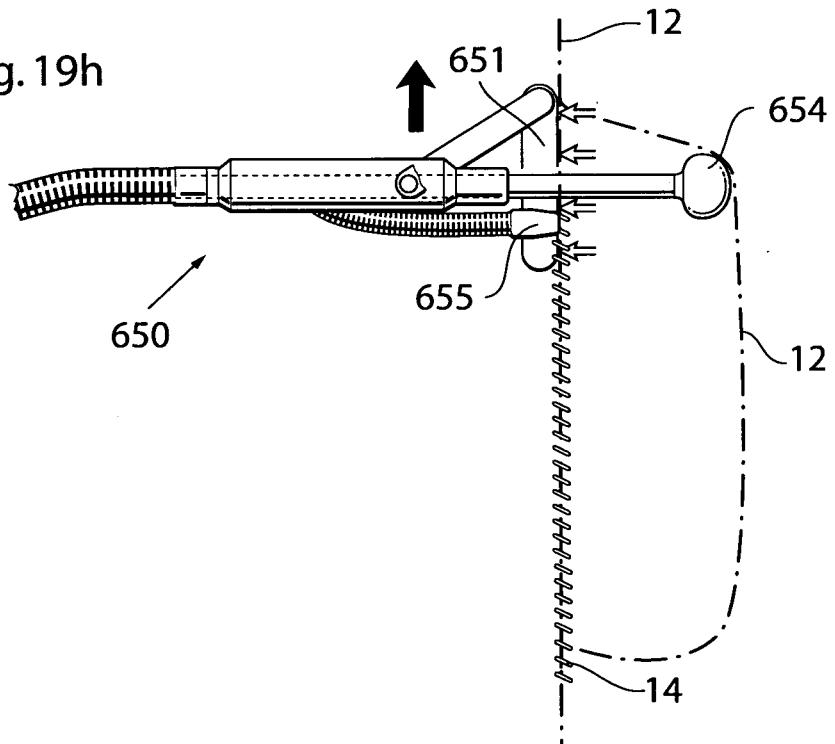


Fig. 19i

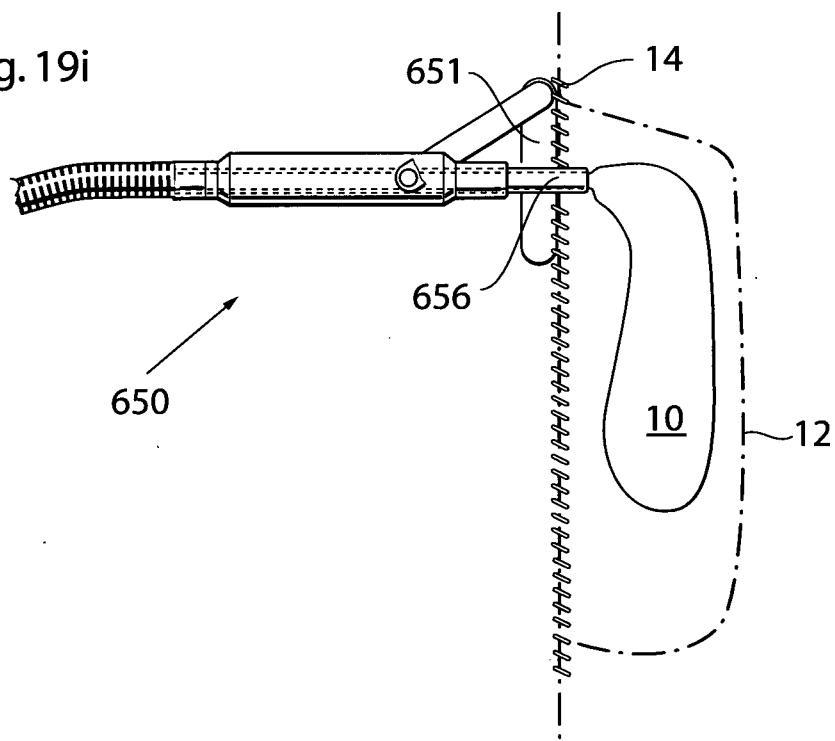


Fig. 19j

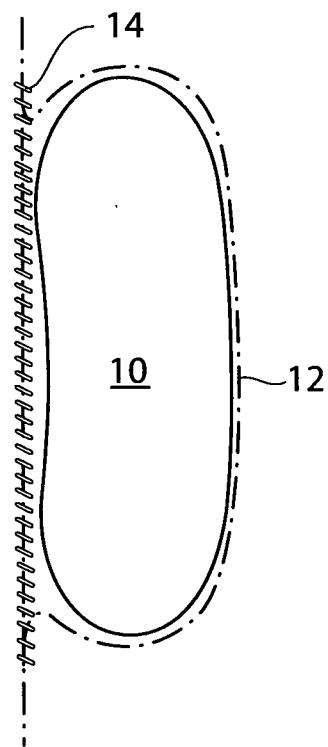


Fig. 20a

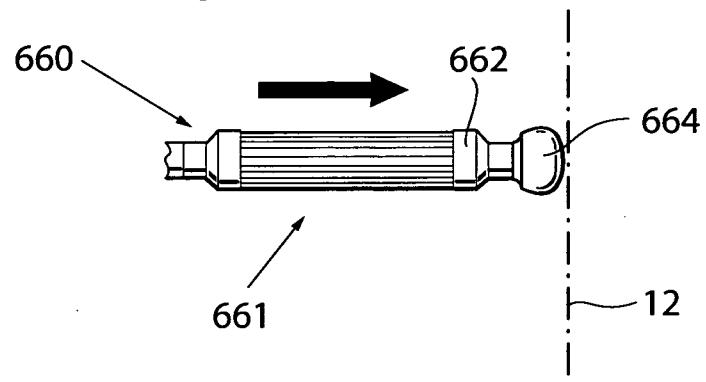


Fig. 20b

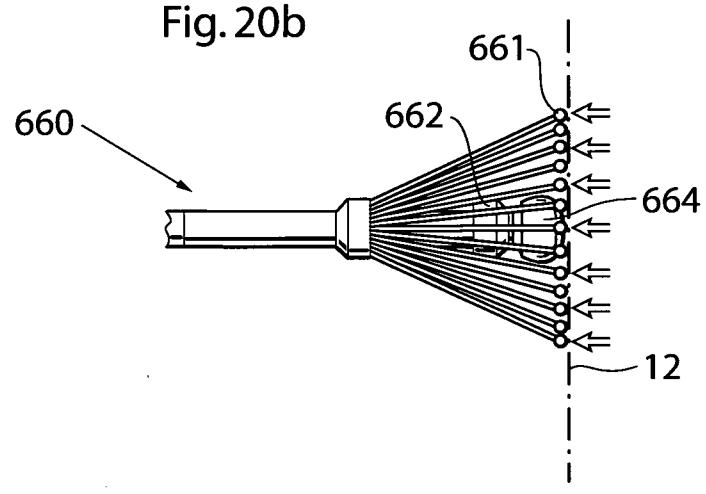


Fig. 20c

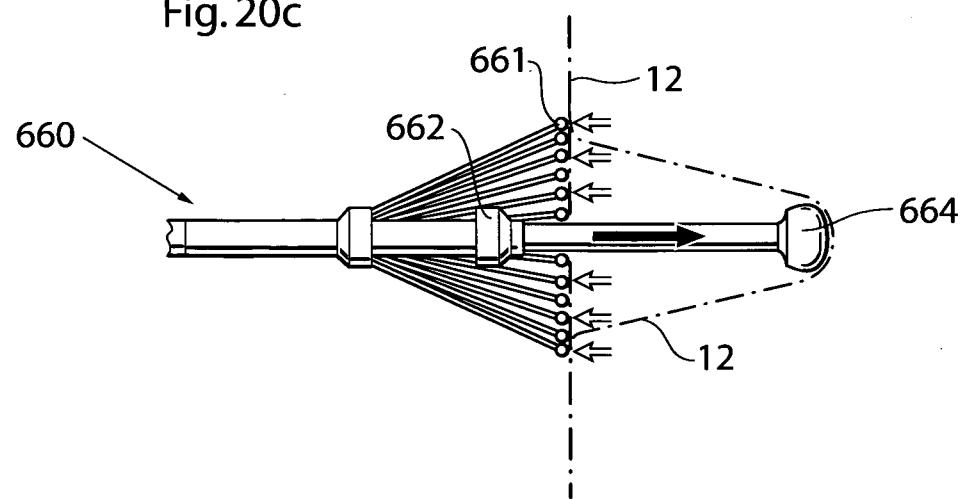


Fig. 20d

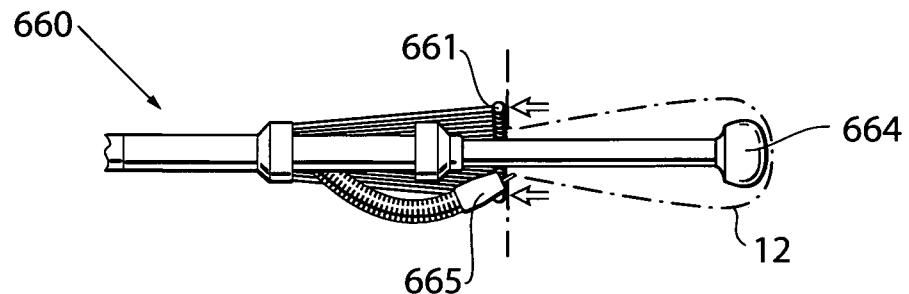


Fig. 20e

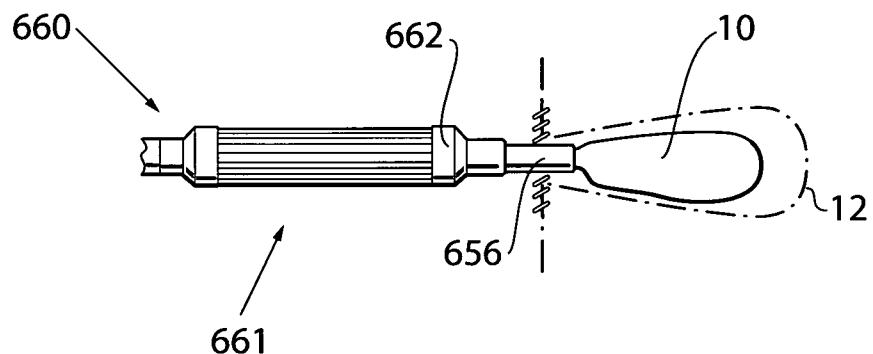


Fig. 20f

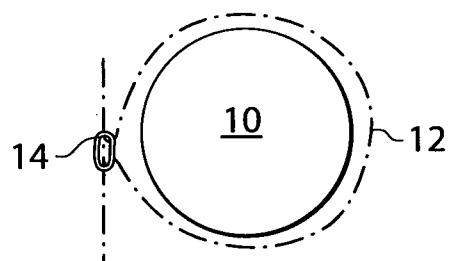


Fig. 21a

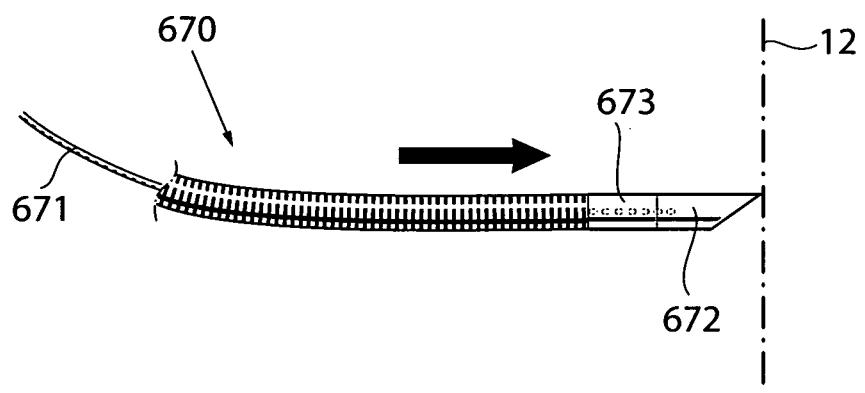


Fig. 21b

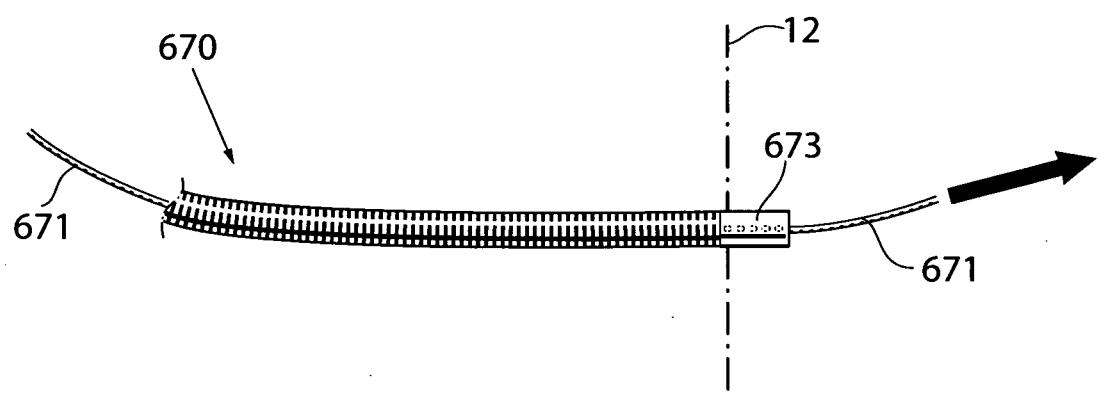
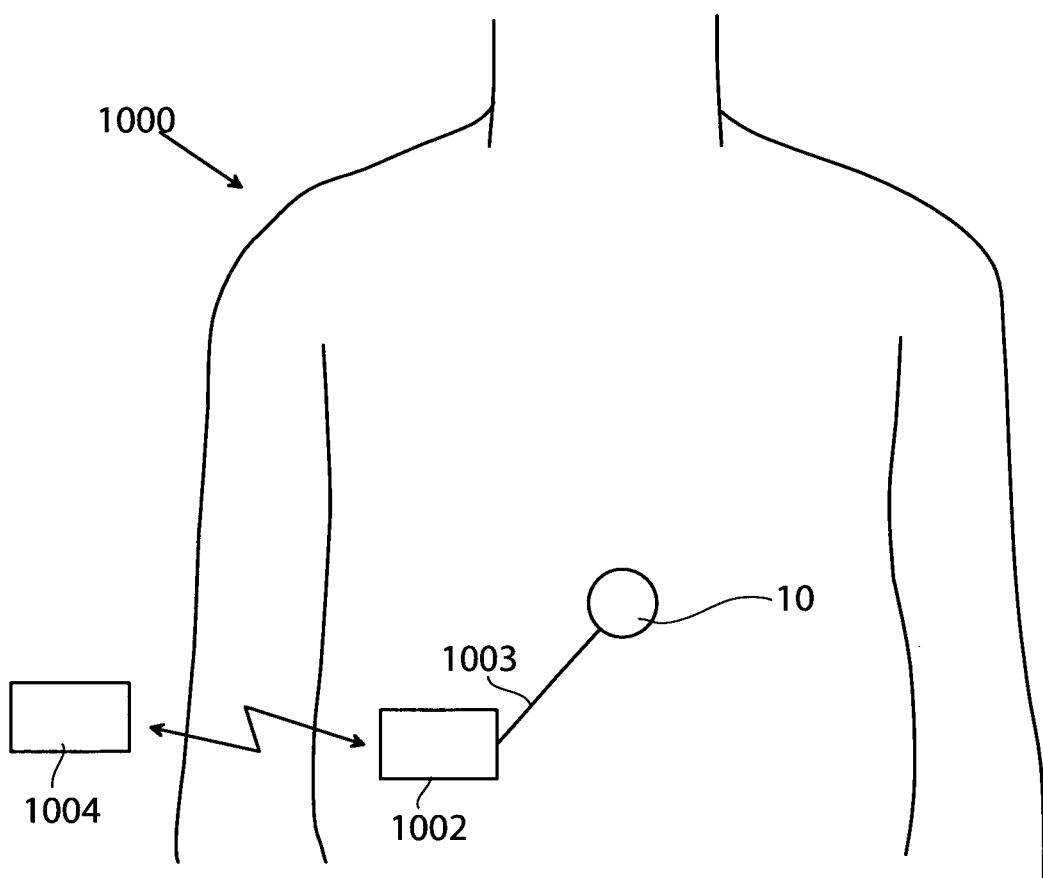


Fig. 22



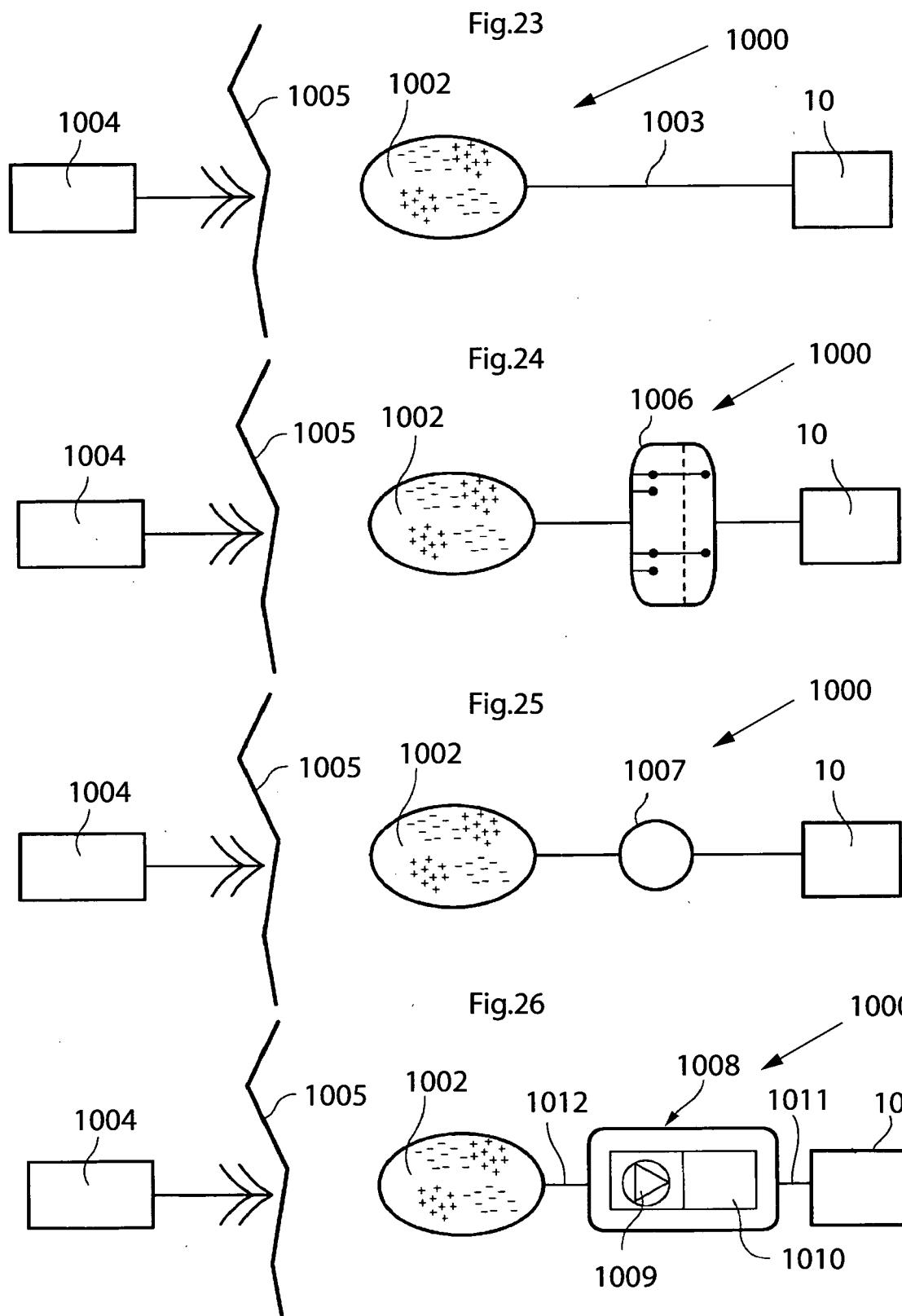


Fig.27

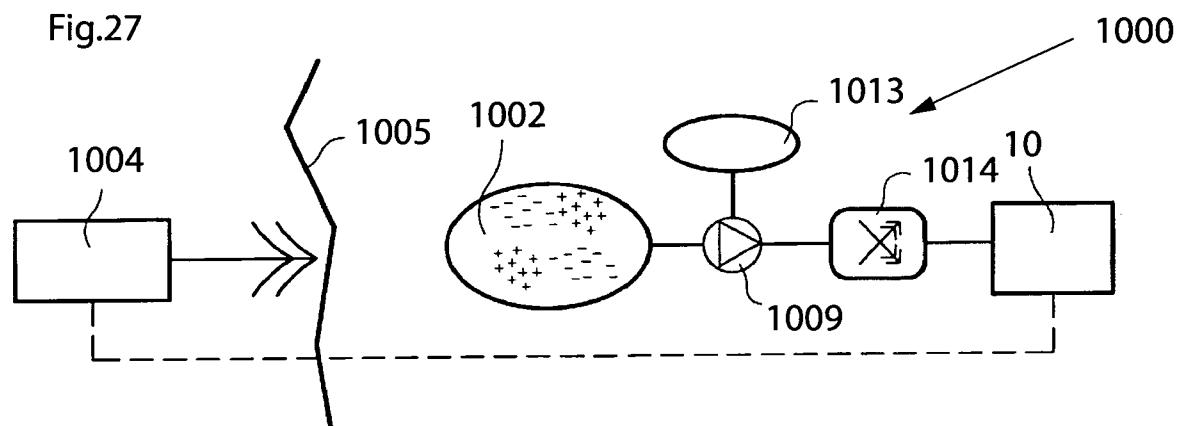


Fig.28

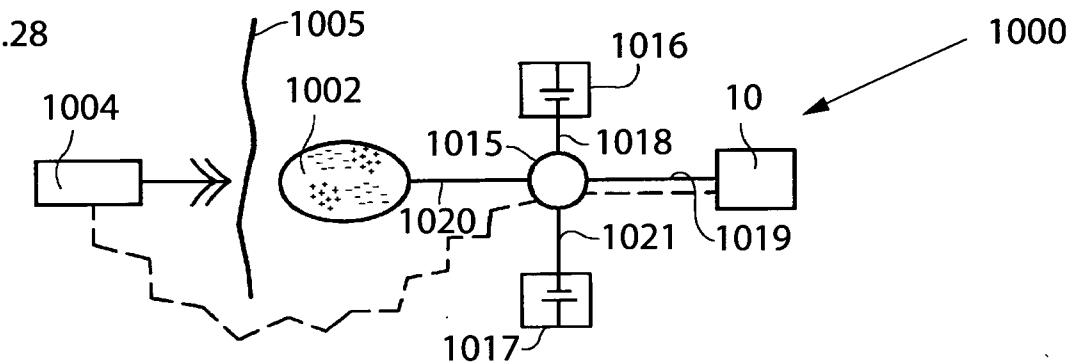


Fig.29

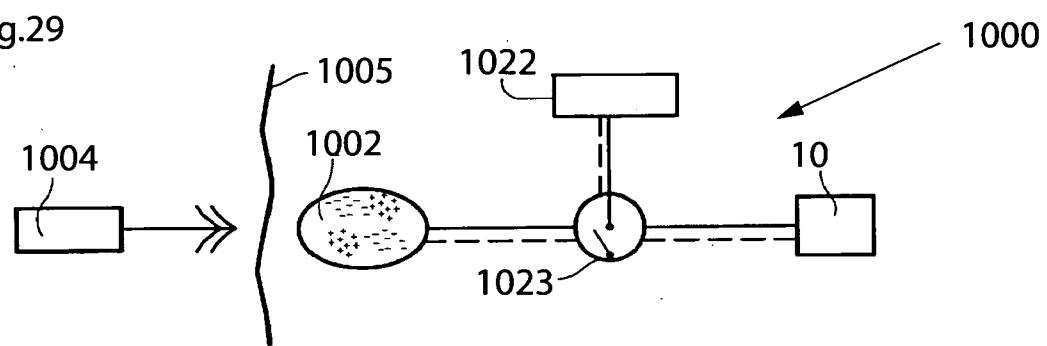


Fig.30

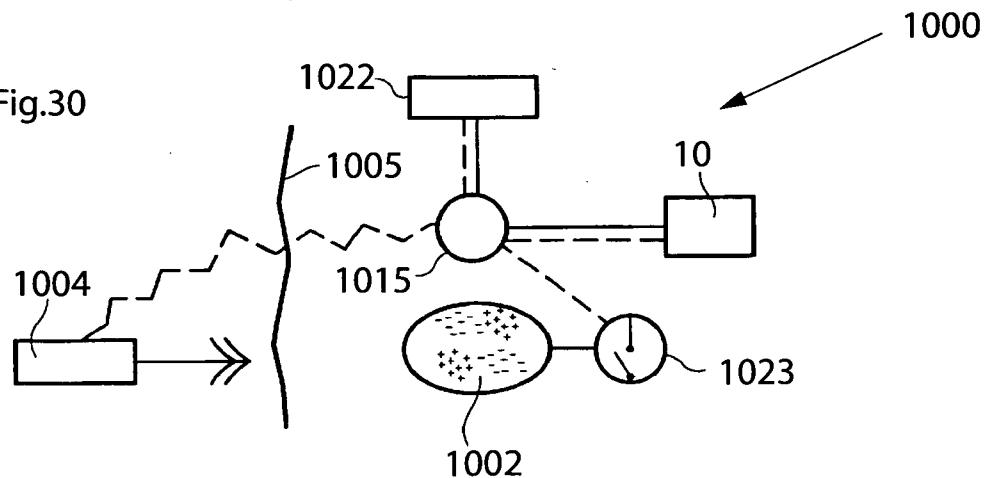


Fig.31

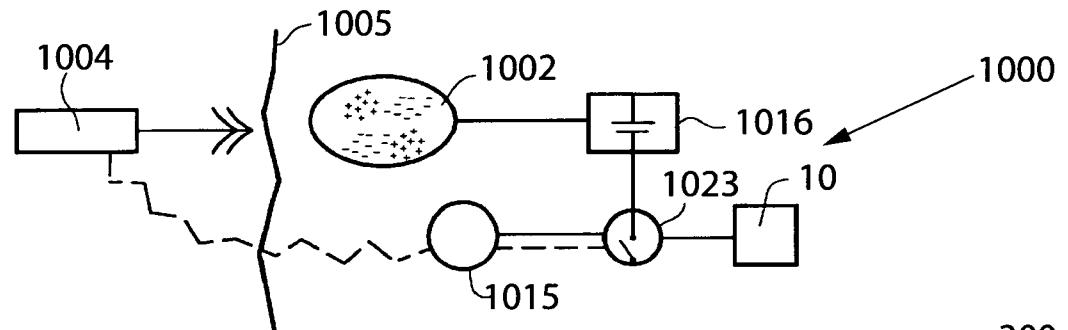


Fig.32

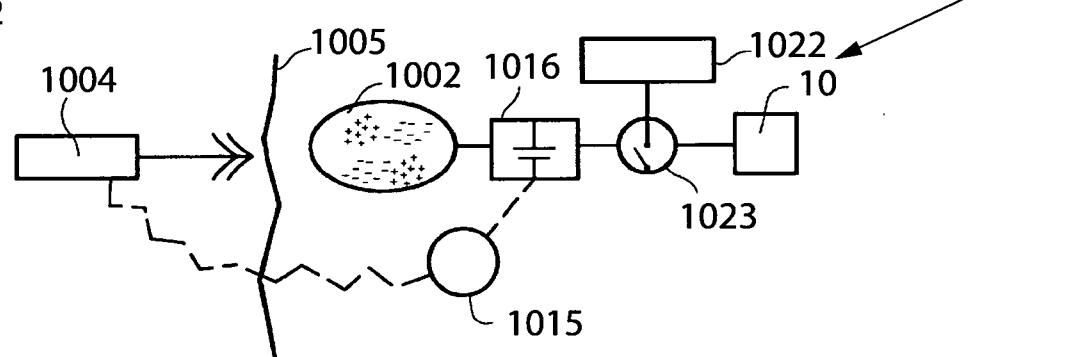


Fig.33

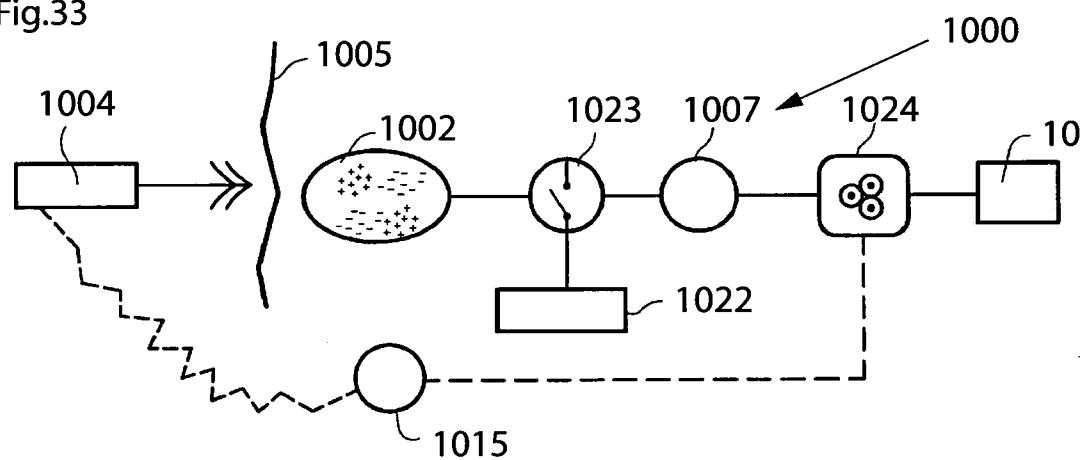


Fig.34

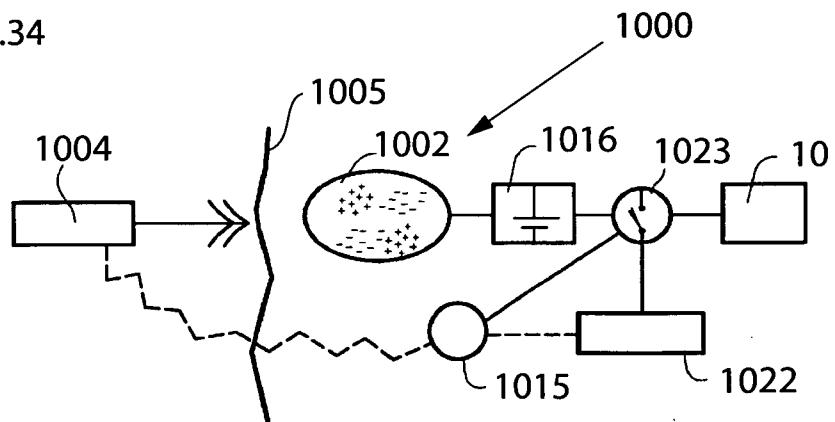


Fig.35

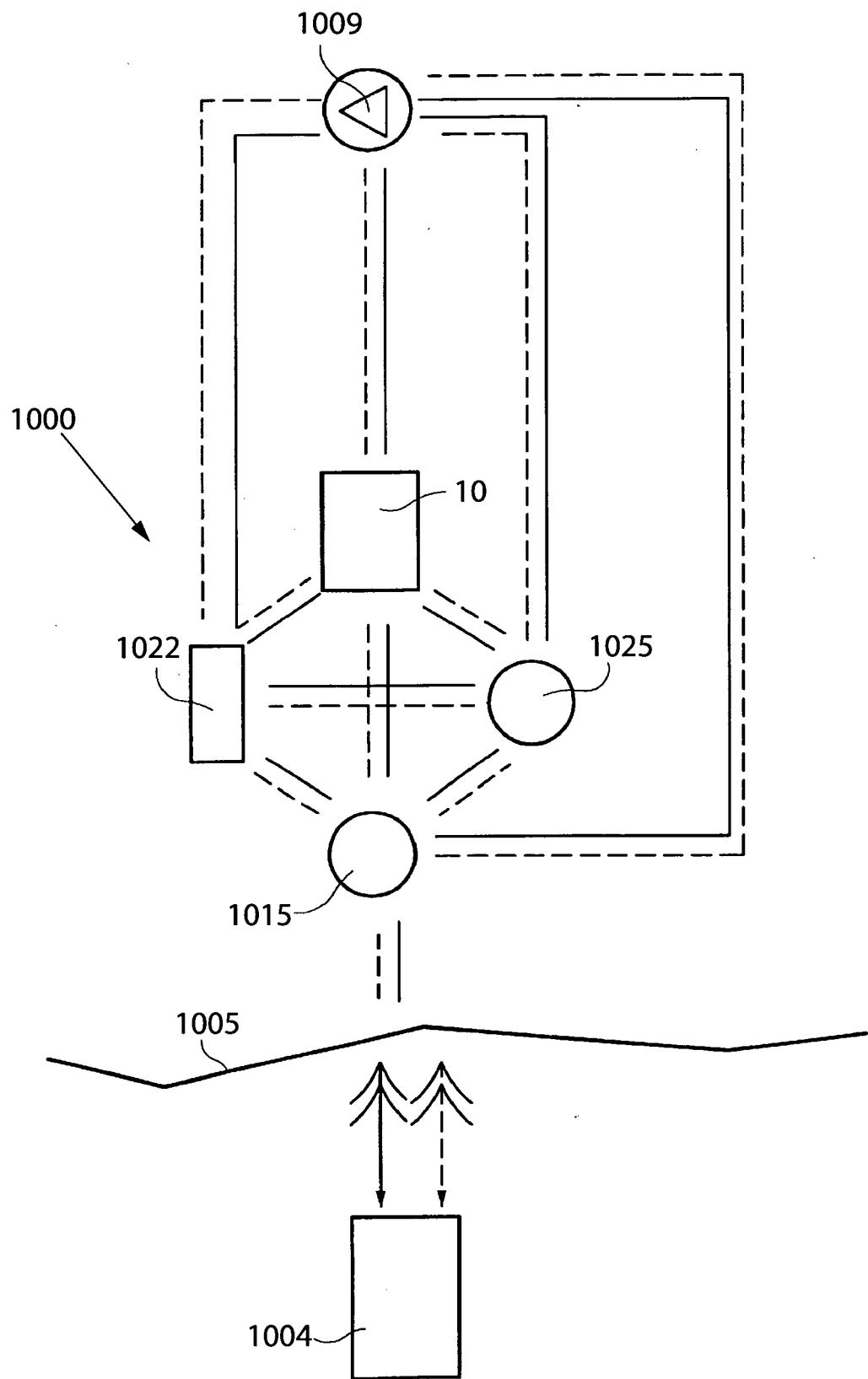


Fig.36

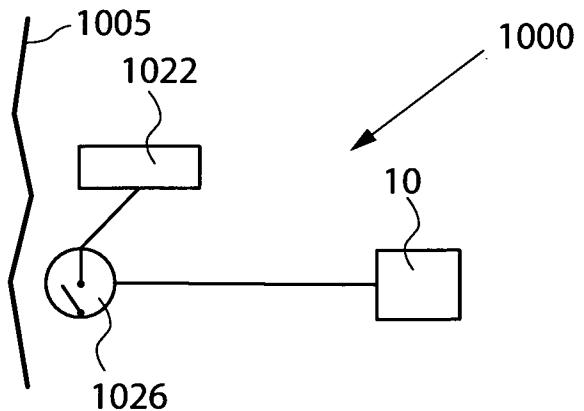


Fig.37

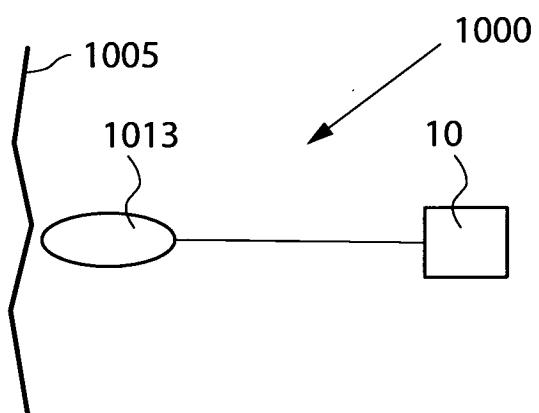


Fig.38

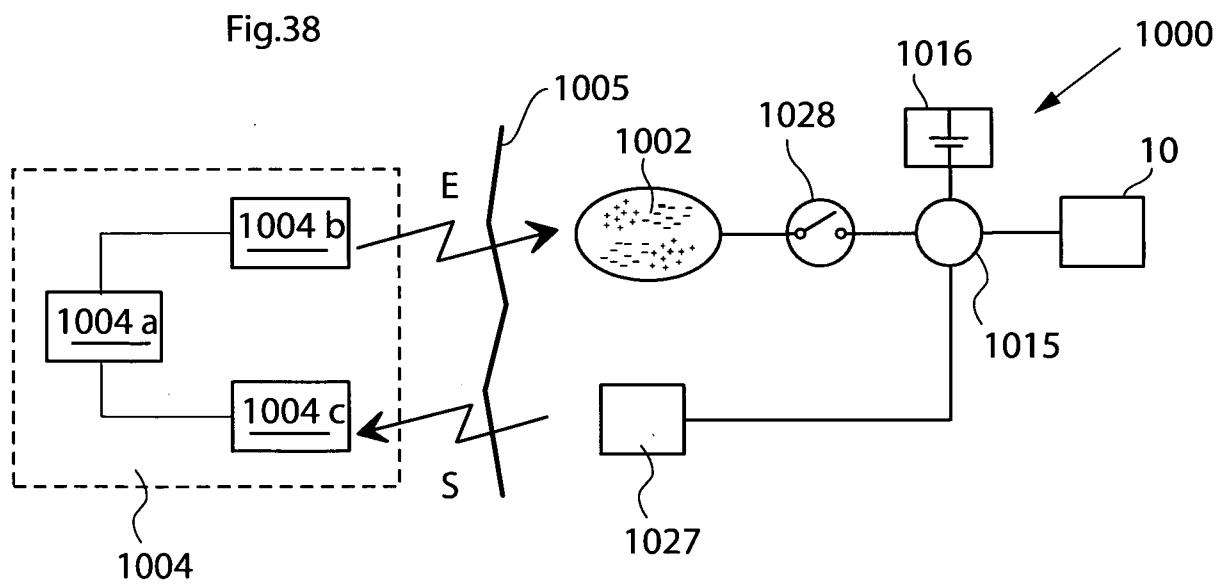


Fig.39

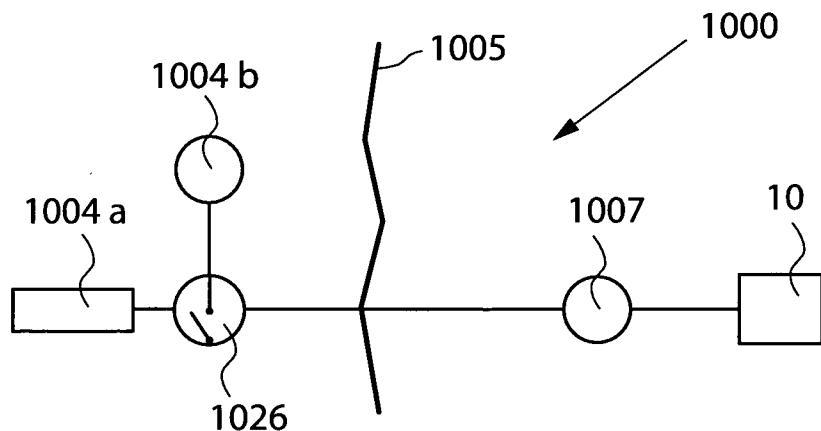
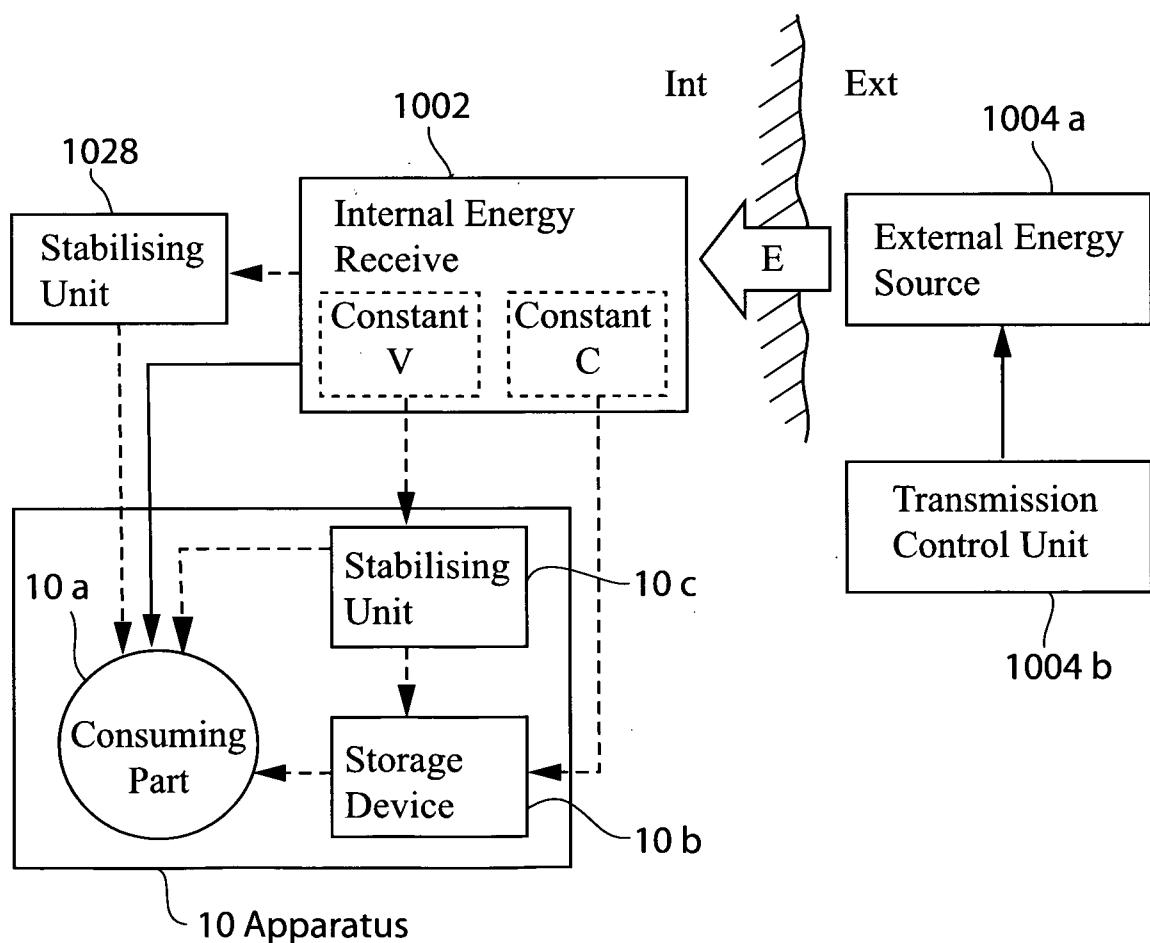


Fig.40



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Fig.41

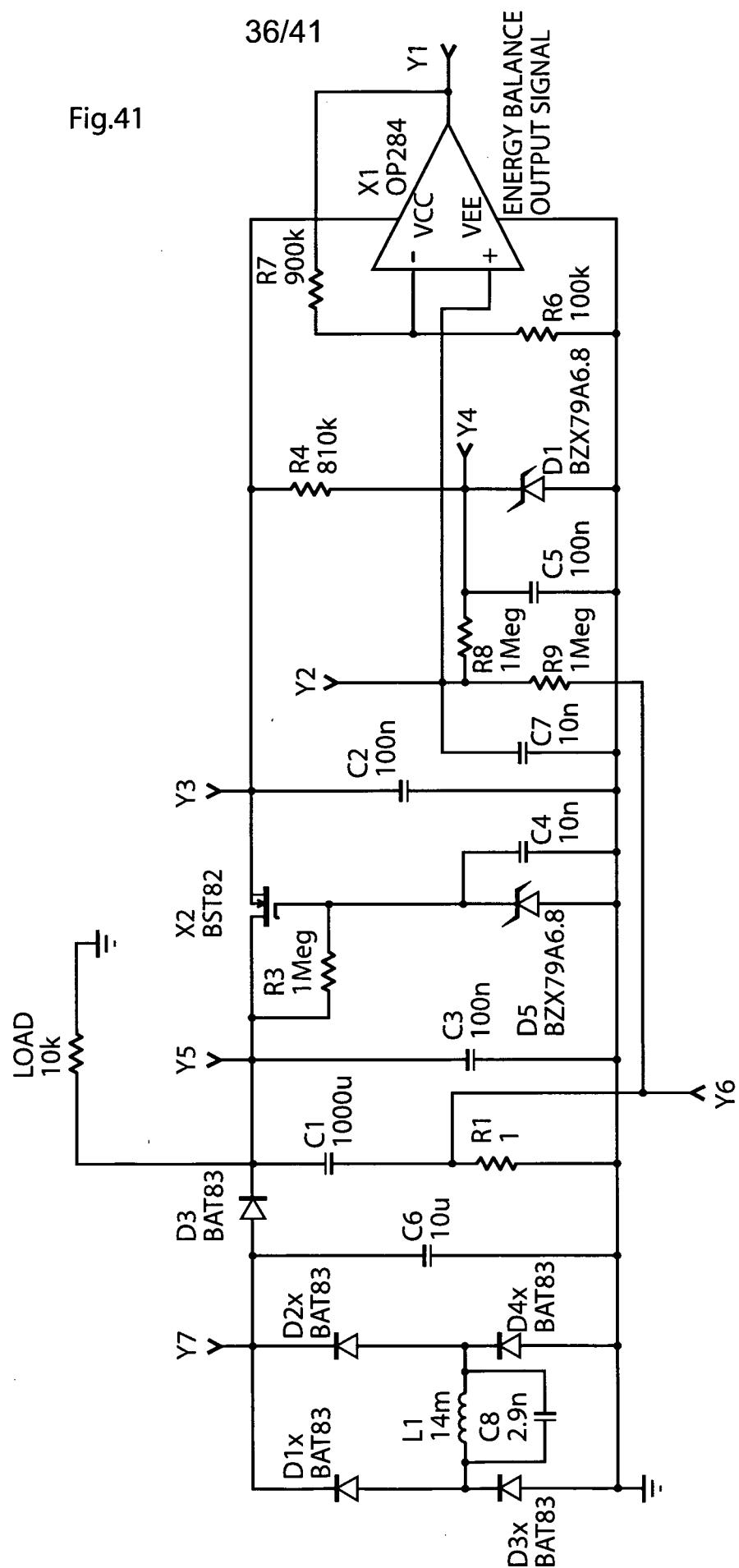


Fig.42

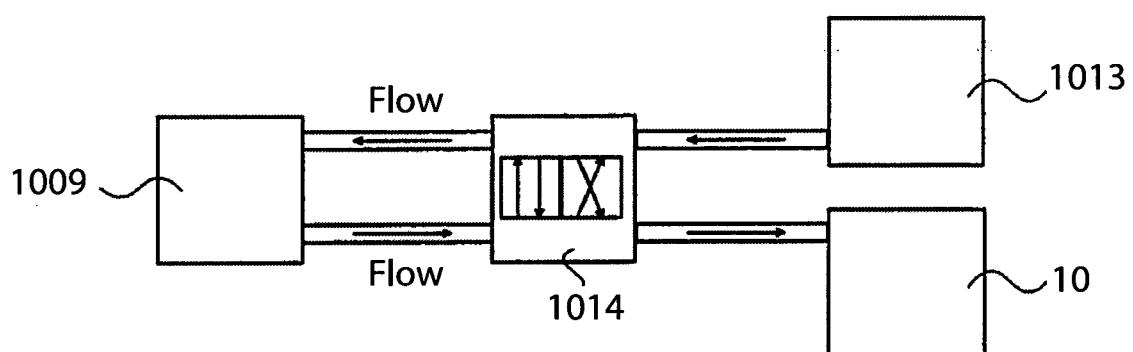


Fig.43

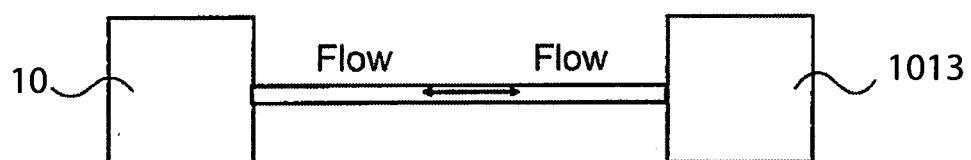


Fig.44

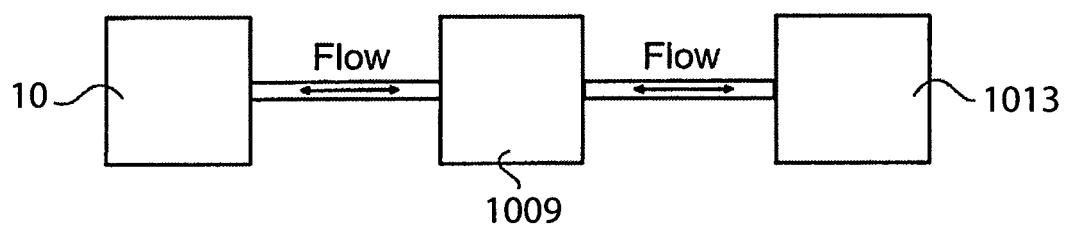


Fig.45

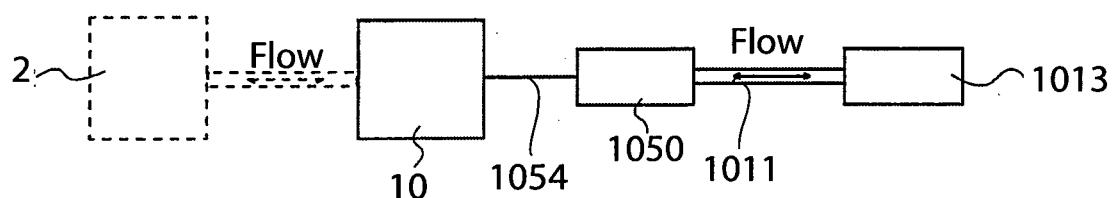


Fig.45 a

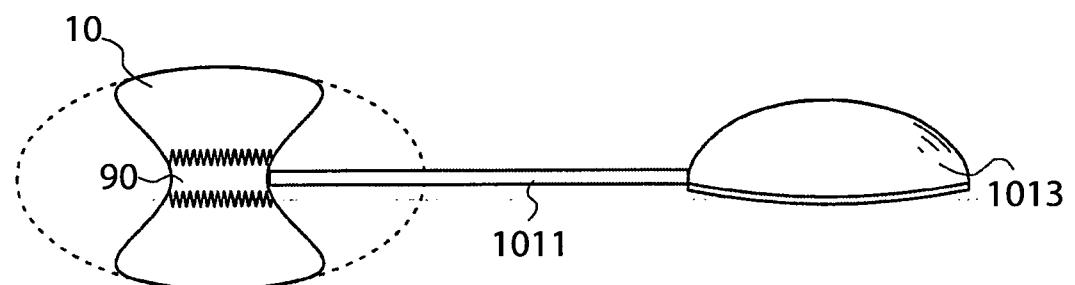


Fig.45 b

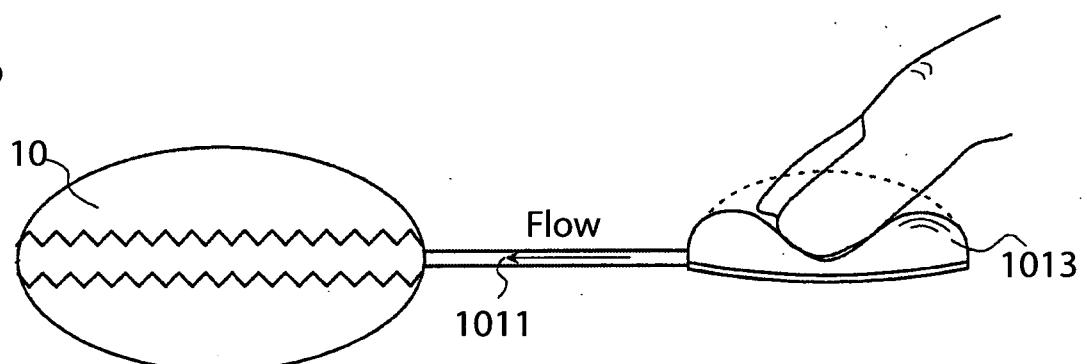


Fig.45 c

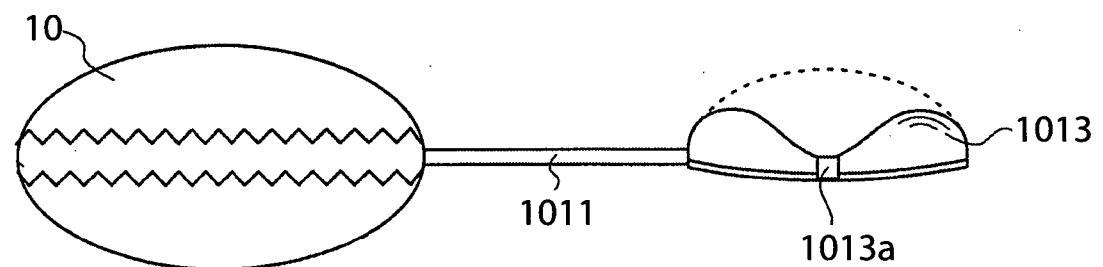


Fig.46

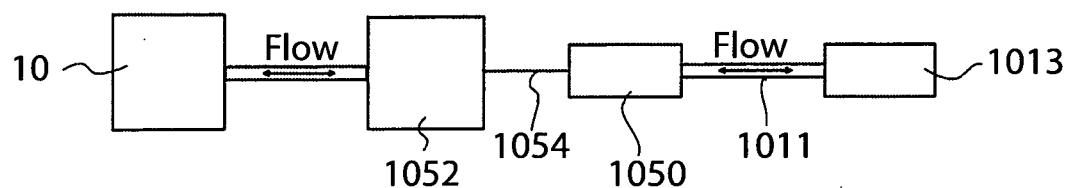


Fig.47 a

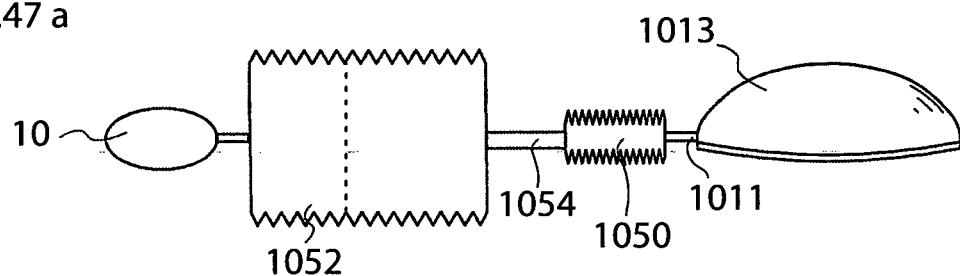


Fig.47 b

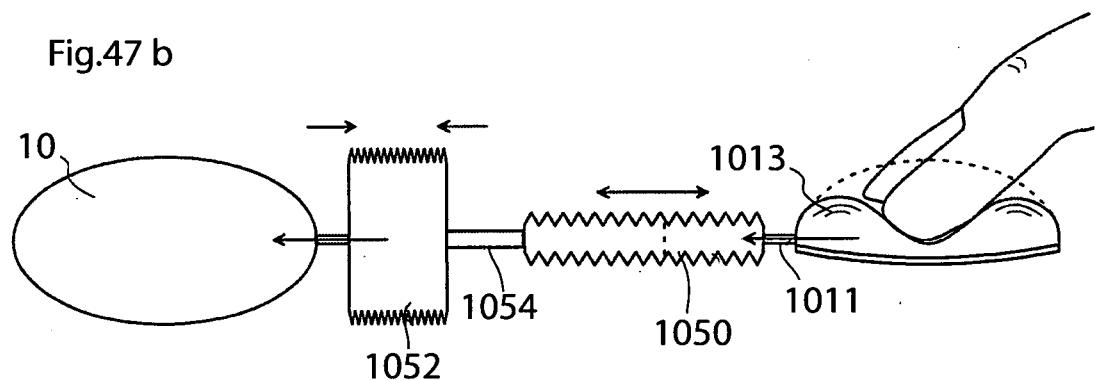


Fig.47 c

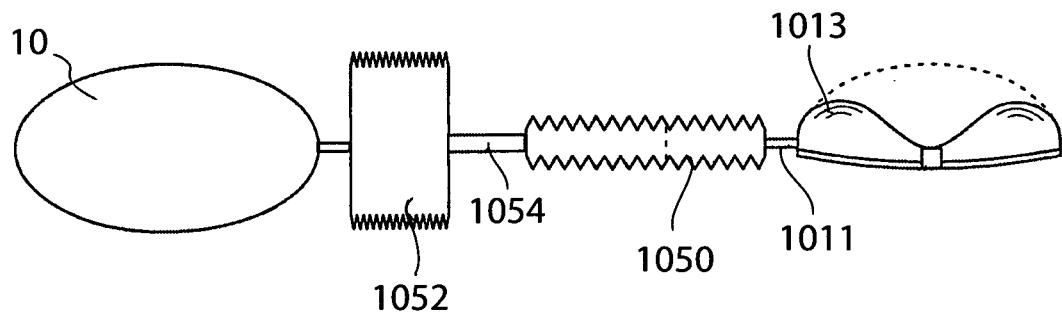
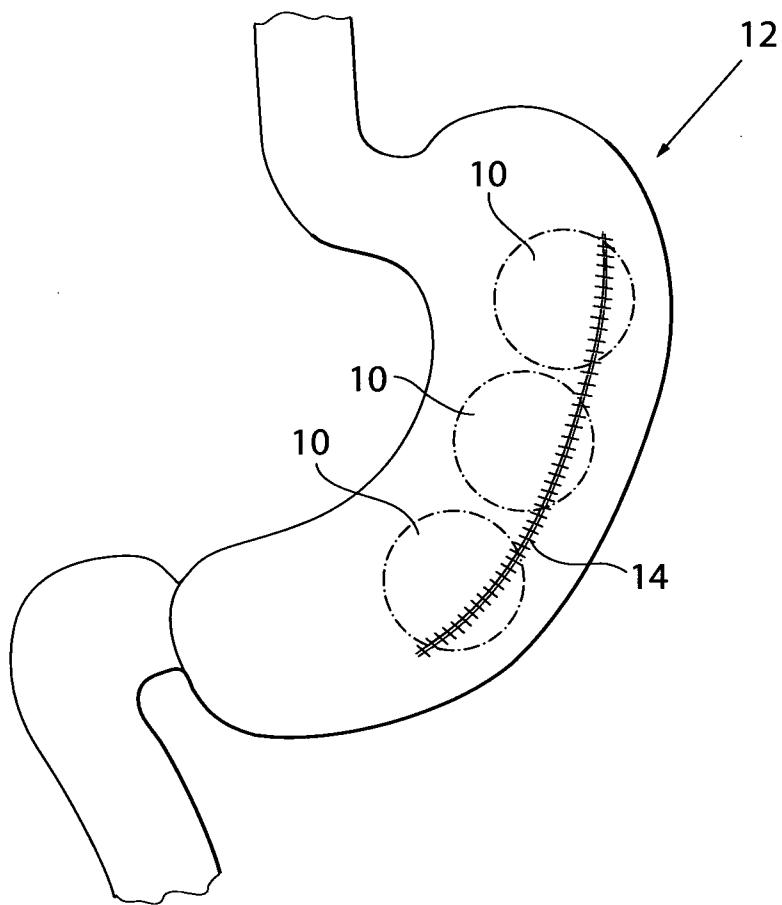


Fig.48



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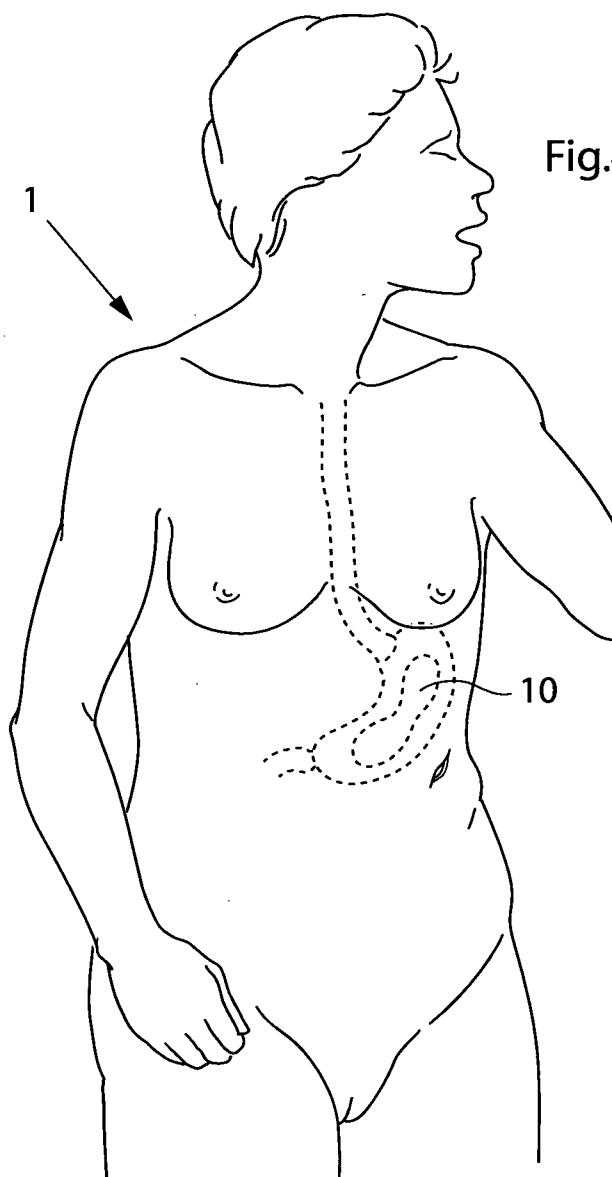
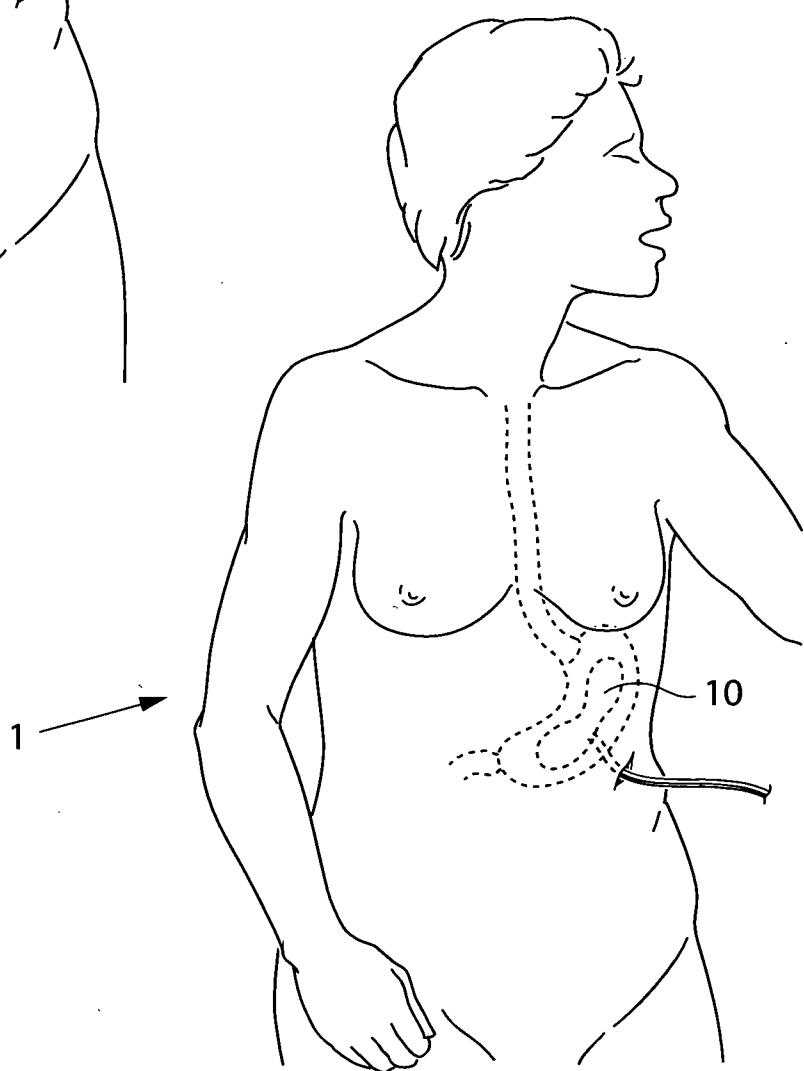


Fig.49a

Fig.49b



SUBSTITUTE SHEET (RULE 26)

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE2009/000047

A. CLASSIFICATION OF SUBJECT MATTER

IPC: **see extra sheet**

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC: **A61F**

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-INTERNAL, WPI DATA, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 20050261712 A1 (D.J. BALBIERZ ET AL), 24 November 2005 (24.11.2005), figures 10,18,19, claims 3,23, paragraphs (0045),(0046),(0048)	1-37,39-68, 81-83,85,102
Y		69,84,86-89, 98-101

Y	WO 0158391 A1 (OBTECH MEDICAL AG), 16 August 2001 (16.08.2001), figures 1-17	69,84,86-89, 98-101

X	WO 2005105003 A1 (SYNECOR, LLC), 10 November 2005 (10.11.2005), page 8, line 26 - page 9, line 22, figures 10,18,19, claims 3,23	1-37,39-68, 81-83,85,102

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

29 May 2009

Date of mailing of the international search report

29-05-2009

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Authorized officer

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Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE2009/000047

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 2004019765 A2 (SATIETY, INC.), 11 March 2004 (11.03.2004), figure 5, paragraphs (0010)-(0011), (0042), (0052) --	1-102
A	WO 2007017880 A2 (STIMPLANT LTD.), 15 February 2007 (15.02.2007), figure 5, abstract, paragraph (0055) --	1-102
A	US 20050267405 A1 (T.M. SHAH), 1 December 2005 (01.12.2005), figures 1-2, abstract --	1-102
A	US 20060257446 A1 (Y.G. TROPSHA ET AL), 16 November 2006 (16.11.2006), figure 7, abstract -- -----	1-102

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE2009/000047

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

See extra sheet.

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of any additional fees.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE2009/000047

Box III

The following separate inventions were identified:

- 1: Claims 1-7, 17-21, 23-37, 39-40, 45-54, 56-63, 68, 81-82, 85 and 102 directed to arrangements and design of a volume filling device.
- 2: Claims 8-16, 22, 41-44, 55 and 64-67 directed to details regarding material choice for the volume filling device.
- 3: Claims 38 and 70-80 directed to arrangements related to a volume filling device with fluid connection with a stretching device adapted to stretch the stomach.
- 4: Claims 69, 83-84, 86-89 and 98-101 directed to arrangements for controlling an implanted device.
- 5: Claims 90-93 directed to arrangements for additional treatment of gastric reflux disease (GERD).
- 6: Claims 94-97 directed to arrangements where the stomach wall is stretched between separate stretching devices.

The present application has been considered to contain six inventions which are not linked such that they form a single general inventive concept, as required by Rule 13 PCT for the following reasons:

The single general concept of the present application is the teaching that the volume of the stomach cavity can be reduced by invagination of a device in the stomach wall, in order to treat obesity.

Document D1 discloses a system for treating obesity. The system comprises an inflatable implant (10) invaginated in the stomach wall (W) and arranged to reduce the effective volume of the stomach (S).

Thus, the single general concept is known and cannot be considered as a single general inventive concept in the sense of Rule 13.1 PCT.

.../...

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE2009/000047

Box III

Invention 1 differs mainly from D1 in details regarding the arrangement of a volume filling device.

Invention 2 differs from D1 in aspects related to materials of the volume filling device.

Invention 3 differs from D1 in arrangements regarding a volume filling device and associated stretching devices.

Invention 4 differs from D1 in arrangements for controlling the device.

Invention 5 differs from D1 in that it is also directed to the treatment of gastric reflux disease.

Invention 6 differs from D1 in that the stretching is performed between separate stretching devices.

No other features can be distinguished which can be considered as the same or corresponding special technical features in the sense of Rule 13.2 PCT.

Thus, the application lacks unity of invention.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE2009/000047

International patent classification (IPC)

A61F 5/00 (2006.01)

A61F 2/04 (2006.01)

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Information on patent family members

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PCT/SE2009/000047

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专利名称(译)	用于治疗肥胖症的装置		
公开(公告)号	EP2249752A4	公开(公告)日	2012-10-31
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IPC分类号	A61F5/00 A61F2/04 A61B17/04		
CPC分类号	A61F5/0036 A61F5/0063 A61B17/00234 A61B17/0469 A61B17/068 A61B17/0682 A61B2017/00278 A61B2017/00561 A61B2017/00818 A61B2017/00827 A61B2017/306 A61B2017/308 A61F2/04 A61F5 /0003 A61F5/0013 A61F5/0026 A61F5/003 A61F5/0033 A61F5/004 A61F5/0043 A61F5/0046 A61F5 /005 A61F5/0069 A61F5/0073 A61F5/0076 A61F5/0079 A61F5/0086 A61F5/0089 A61F2005/0016 A61F2005/002 A61F2005/0023 A61F2250/0001 A61F2250/0004 A61N1/36007 A61B1/04 A61B1/2736 A61B1/3132 A61B17/00 A61B17/064 A61B17/08 A61B17/30 A61B17/320016 A61B17/3423 A61B17 /3474 A61B2017/081 A61F2002/044 A61F2002/045		
优先权	61/006719 2008-01-29 US 0802138 2008-10-10 SE		
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摘要(译)

一种用于治疗人类或动物哺乳动物患者的反流疾病的设备。该装置包括适于与患者的心脏括约肌接合的可植入刺激装置(26)和用于控制刺激装置的控制装置(56)，该控制装置(56)通过提供电能脉冲以增加括约肌张力来刺激stimulate门括约肌，从而使ia门关闭。所述控制装置适于在操作状态下使所述刺激装置以至少两个脉冲序列刺激所述心脏括约肌。至少一个脉冲序列内的至少一些脉冲通过第一时间间隔彼此分开。至少两个脉冲序列彼此相隔第二时间间隔，并且第二时间间隔比第一时间间隔长。