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

(19) World Intellectual Property Organization
International Bureau

(43) International Publication Date
27 December 2001 (27.12.2001)

PCT

(10) International Publication Number
WO 01/97897 A1

(51) International Patent Classification[7]: A61M 29/00, A61F 7/00

(21) International Application Number: PCT/US01/19470

(22) International Filing Date: 19 June 2001 (19.06.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
09/598,852 20 June 2000 (20.06.2000) US

(71) Applicant: **STARION INSTRUMENTS, INC.** [US/US]; 20665 4th Street, Saratoga, CA 95070 (US).

(72) Inventor: **MOLLENAUER, Kenneth, H.**; 20665 4th Street, Saratoga, CA 95070 (US).

(74) Agent: **CROCKETT, K., David**; Crockett & Crockett, 24012 Calle de la Plata, Suite 400, Laguna Hills, CA 92653 (US).

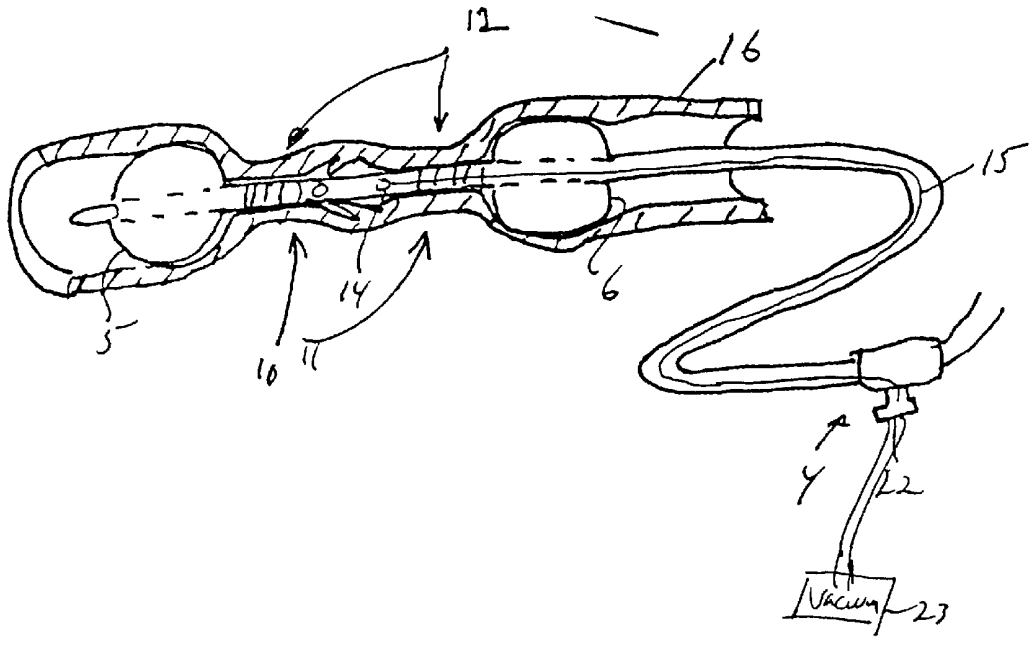
(81) Designated States *(national)*: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.

(84) Designated States *(regional)*: ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

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

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: DEVICES AND METHODS FOR REPAIR OF VALVES IN THE HUMAN BODY



(57) Abstract: Devices and method for treating various incompetent anatomical valves by thermally damaging the nearby supporting tissue of the body vessel controlled by the valve with the use of electrical wires (12, 13). The device having two balloons (5, 6) and suction port (14) to draw the tissue of the vessel near the heating element (10).

WO 01/97897 A1

Devices And Methods For Repair Of Valves In The Human Body

Field of the Inventions

The inventions described below relate to the fields of minimally invasive surgery, vascular surgery, urology and general surgery.

Background of the Inventions

Several diseases of the human body result from poor performance of natural valves within the body. Venous insufficiency, gastrointestinal reflux, and urinary incontinence are examples of maladies which are the result of the failure of valves to perform normally.

Venous insufficiency generally refers to conditions which cause the veins in the leg to become incapable of functioning properly due to failure of venous valves. Generally, the return of blood from the veins in the leg to the heart is caused by the interaction of the leg muscles with venous valves that function as check valves. Muscular activity in the legs forces venous blood upward toward the heart, through the venous valves which are generally found in the perforating veins between the superficial veins and the deep veins in the leg.

Venous insufficiency is often caused by failure of the valves located in small communicating veins which connect large superficial veins such as the posterior arch vein or saphenous vein, with large deep veins such as the peroneal or tibial veins. The communicating veins have valves which look like duckbill valves or leaflet valves that act as check valves, allowing blood to flow from the superficial veins into the deep veins, but blocking flow in the reverse direction.

Exercise and movement of the calf muscles around the communicating veins squeezes blood through the communicating veins. This mechanical pumping action, combined with the function of the valves, is responsible for returning blood flow to the heart.

When the valves fail, venous blood in the superficial veins cannot be pumped into the deeper veins, resulting in blood pooling in the legs. The condition causes very poor circulation in the legs and can lead to varicose veins and skin ulcers. Large varicose veins in the lower leg, skin ulcers just above the ankle bone on the inside of the calf, and discolored skin on the lower leg, are common symptoms. Similar symptoms are seen in other areas of the body, particularly the thighs and arms, when perforating veins in those areas become incompetent.

Venous insufficiency is generally attributed to the failure of certain groups of perforating veins. There are about 150 perforating veins in the leg, but there are several major perforating veins which are important contributors to the problem of venous insufficiency. An important group of perforators is found high on the inside of the calf, over the calf muscle. Another important group of perforators is found low on the inside of the calf, just above the ankle and toward the back of the leg. Another set of perforators is found on the lateral or outside of the leg run though the muscles on the outside of the calf.

An effective surgical treatment of this condition was developed by Linton circa 1938. In the Linton procedure, also referred to as the Medial subfascial approach, the calf is cut open along the Linton line, extending from just above the ankle bone (or medial malleolus) on the inside or medial side of the foot, up the inside of the calf almost to the knee.

The incision is deep enough to cut the skin and fat, and also the deep fascia which is a filmy fibrous layer of tissue which covers the muscles of the calf. Upon peeling away the skin, fat and fascia, some of the communicating veins can be seen, and these are cut and tied off. A minimally invasive version of this procedure is described in U.S. Patent 5,979,452.

An alternative to the Linton procedure is repair of the veins. One method for repairing the venous valves is disclosed in Farley, et al., Catheter Having Expandable Electrodes And Adjustable Stent, U.S. Patent 6,014,589 (Jan. 11, 2000). Farley proposes a catheter having expandable electrodes for applying energy to a vein and having expandable stent members for limiting vein shrinkage to a final desired vein diameter. The catheter includes a set of expandable arms that are pre-formed into an outwardly bowed configuration. An electrode is mounted on each arm. The catheter is delivered percutaneously into the veins of a patient, and positioned near a failed valve. The stent arms are expanded outward to the desired final diameter of the vein. The electrode arms are then expanded into apposition with the vein wall and energy is applied to shrink the vein into contact with the stent arms.

Urinary incontinence is another condition caused, at least in part, by failure of an anatomical valve. Stress Urinary incontinence, or SUI, effects females and results from the inoperability of the bladder neck sphincter, which controls flow or urine from the bladder. Various treatments have been tried, and the bladder neck suspension is the predominant surgical cure. Bladder neck suspension methods include various procedures for lifting the bladder neck and urging it anteriorly, toward the front of the body. The bladder neck is tied to another structure in the body, and is literally suspended from these structures. This suspension

alters the forces countering the closure of the bladder neck sphincter, and cures the condition. Bladder neck suspension may be accomplished with open surgical techniques or minimally invasive techniques. Minimally invasive techniques, such as those indicated in Benderev, U.S. Patent 5,860,425, require penetration of the skin and insertion of various suturing and knot tying devices into the body.

Gastroesophageal reflux disease, or GERD, is another condition caused, at least in part, by failure of an anatomical valve. In GERD, the stomach contents are regurgitated from the stomach into the lower esophagus due to a failure of the lower esophageal sphincter. Since the stomach contents are highly acidic, the condition is uncomfortable and, if left untreated, potentially dangerous. Symptoms range from mere heartburn to pulmonary disorders, ulcer formation, or esophagitis esophageal obstruction and perforation of the esophagus. One surgical treatment of GERD is the Nissen fundoplication, in which a surgeon constructs a new valve to support LES by pulling the gastric fundis upward and wrapping it around the lower esophagus. The procedure is accomplished through open surgery, but a version of the Nissen fundoplication can now be accomplished with minimally invasive techniques. Recently, RF ablation of the lower esophageal sphincter has been proposed, under the assumption that aberrant electrical activity in the lower esophageal sphincter causes the reflux, and the ablation of aberrant electrical tissue will reduce lower esophageal sphincter relaxations.

Summary

The devices and methods described below provide for the treatment of various incompetent valves and sphincters throughout the body. The catheters provide for location of a heating element or other tissue necrosing tool in the lumen of

the vessel controlled by the valve, at or near the base of the valve (but not on the valve itself). Additionally, the catheters include balloons for locating and anchoring the distal section of the catheter within the lumen, such that the heating element is positioned near the base of the valve, in contact with lumenal tissue at the base of the valve. The catheters also include suction ports on the distal end of the catheter which can be operated to size or draw down the vessel to the diameter of the catheter, so that the vessel walls are in contact with the heating elements. The catheters may be used for treatment of venous insufficiency, urinary incontinence, and gastroesophageal reflux disease.

Brief Description of The Drawings

Figure 1 illustrates a venous repair catheter positioned with a vein, in the proximity of a venous valve.

Figure 2 illustrates a venous repair catheter in one step of operation, drawing the vein segment to be treated toward the catheter body.

Figure 3 illustrates the segment of diseased vein after treatment and withdrawal of the venous repair catheter.

Figure 4 illustrates a venous repair catheter positioned within a vein, where the catheter is adapted to treat a vein segment including several valves in a single application .

Figure 5 illustrates the anatomy involved in stress urinary incontinence in a female patient.

Figure 6 illustrates the repair catheter adapted for use in treatment of stress urinary incontinence in a female patient.

Figure 7 illustrates the anatomy involved in gastroesophageal reflux disease.

Figure 8 illustrates the repair catheter of Figure 7 in use in the treatment or gastroesophageal reflux disease in a patient.

Detailed Description of the Inventions

Figure 1 illustrates a venous repair catheter positioned within a vein, in the proximity of a venous valve that is to be repaired. The catheter **1** includes a tubular catheter body **2** characterized by a distal section **3** and a proximal section **4**. The distal section is adapted for percutaneous insertion into the blood vessel, and the proximal end, which remains outside the body, is adapted for connection to various external systems, such as to vacuum source, inflation reservoirs or pumps, and a heating power source. In the distal section, a pair of inflatable balloons including a distal balloon **5** and a proximal balloon **6** are disposed on the catheter body 2 and bound and define a heating segment **7**. The catheter body in the heating segment has an outer diameter of about 2 to 5 mm, depending on the initial size and desired post-treatment size of the vein in which the catheter is to be used. The catheter body is preferably a relatively flexible catheter body, to permit percutaneous insertion and navigation through the veins of the patient, although it may be provided as a stiff and inflexible catheter body for treatment of valves readily accessible from nearby percutaneous access sited. The balloons are supplied with fluid through lumens **8** and **9** (they may also be supplied by a single shared lumen). Within the heating segment, a pair of heating elements, including the distal heating element **10** and a proximal heating element **11** are mounted on the outside of the catheter body 2, and electrical wires **12** and **13** running through the catheter body connect the heating element to a source of power through the proximal end of the catheter. The heating elements are comprised of a resistive heating elements, electrodes, RF

electrodes, ultrasonic heat sources, LED's and other light or laser sources, or other suitable heating mechanisms. Where the heating elements are resistive heating elements, the electrical wires will comprise a ground wire and a hot wire, and while a minor amount of current may pass through the body to ground, the bulk heating of the venous tissue will be caused by conductive heating from the heating elements which are in turn heated due to resistance of the elements and the passage of current through the elements. Appropriate materials for the resistive heating elements include nichrome and nickel-titanium alloys such as nitinol. One or more suction ports **14** connected to a suction lumen **15** (shown in phantom) within the catheter body communicates with the exterior of the catheter body in the vicinity of the heating segment. The suction port is located relative to the heating elements such that suction applied to the vessel through the suction port will draw the tissue of the vessel near the valve toward the heating elements. When placed within the body as illustrated, the catheter distal segment 3 is inserted into a section of the patient's vein **16** which has a venous valve **17** which is incompetent and requires treatment. The balloons reside on the distal side **18** of the venous valve and the proximal side **19**, and when inflated create a segment of isolated vein which includes the venous valve. The distal heating element 10 is located on the distal side 18 of the valve, while the proximal heating element 11 is located on the proximal side 19 of the valve. The proximal and distal sides of the valve are defined here in relation to blood flow within the vein, with the distal side being downstream from the proximal side, or, in other words, in relation to the origin of blood flow. In the illustration, access is gained through a superficial vein upstream of the valve. The proximal and distal components of the catheter would of course be placed in the distal and proximal sides, respectively, if access is

gained from a point downstream of the valve and the approach into the vein is retrograde, or upstream, in the blood flow. Proper placement of the catheter relative to the venous valve can be confirmed with ultrasound imaging, endoscopic viewing, palpation by the operating surgeon, or any other means.

Referring now to the proximal section 4 of the catheter body, this section includes a hub **21**. The hub includes a luer fitting **22** which is connected to the suction lumen 15. The luer fitting is adapted for air-tight connection to a source of vacuum **23**. Shutoff valves **24** and throttle valves **25** may be placed in line between the vacuum and the luer fitting, so that suction can be controlled or terminated as desired by the operator. The hub also includes luer fittings **26** and **27** connected in fluid communication with the inflation lumens 8 and 9, respectively. These luer fittings are adapted for connection to the inflation fluid source **28** which provides pressurized fluid to the system. A shutoff valve **29** and throttle valve **30** may be placed in line between the inflation fluid source and the luer fittings, so that inflation pressure can be controlled or terminated as desired by the operator. The hub also includes the proximal end of the heating element electrical wires 12 and 13, and these are conveniently terminated in an electrical connector **31** which may be integral or separate from the hub. The electrical connector provides for connection of the electrical wires to the direct current power supply **32**. The direct current power supply, should be capable of delivering direct current power in the range of 1 to 72 watts, with voltage in the range of 1-12 volts and amperage in the range of 1-6 amps.

While Figure 1 illustrates the device components and the initial placement of the catheter within a segment of diseased vein, Figure 2 illustrates the catheter in operation, drawing the vein segment to be treated toward the catheter body. With

the balloon inflation reservoir connected to the inflation luer fittings, the distal and proximal balloons have been inflated to create a cylindrical lumenal space between the balloons having the vein as the wall of the cylinder. Optionally, saline or other solution (such as an anti-coagulant) has been flushed through the vacuum lumen 15 and suction port 14 to flush out the blood within the cylinder or dilute or treat the blood within the cylinder prior to application of suction. The vacuum source 23 has been connected to the vacuum line luer fitting 22, and the valves have been opened to create a vacuum in the cylindrical space between the balloons. Under the vacuum, the vein between the balloons has collapsed upon the catheter body and heating segment 7. In the next step of the procedure, electrical current is applied to the collapsed vein segment through the heating elements, supplied with direct current from the DC source. Preferably, the direct current is applied at power levels adequate to shrink the vein walls on either side of the venous valve, but not high enough to thermally damage the valve itself. Should thermal damage to the valve itself be medically indicated, the power levels may be increased and the heating elements may be located on the catheter body to correspond with the location of the venous valve. After treatment, the catheter may be withdrawn, leaving the vein in the condition illustrated in Figure 3, where the valve 17 now spans a segment **33** of vein which is constricted vis-à-vis its original state.

Figure 4 illustrates a venous repair catheter positioned with a vein, where the catheter is adapted to treat a vein segment including several valves in a single application of heating power. In the distal section, a pair of inflatable balloons including a distal balloon 5 and a proximal balloon 6 are disposed on the catheter body 2 and bound and define a heating segment. The balloons are supplied with fluid through

lumens 8 and 9. Within the heating segment 7, a first pair of heating elements, including the distal heating element 10a and a proximal heating element 11a are mounted on the outside of the catheter body 2, and electrical wires 12a and 13a running through the catheter body connect the first heating elements to a source of power through the proximal end of the catheter. These heating elements are located relative to the proximal balloon so that they will be located with on either side of a venous valve 17a upon positioning of the balloon near the venous valve. A second pair of heating elements, including the distal heating element 10b and a proximal heating element 11b are mounted on the outside of the catheter body 2, and electrical wires 12b and 13b running through the catheter body connect the second pair of heating elements to a source of power through the proximal end of the catheter. This second pair of heating elements are located relative to the proximal balloon so that they will be located with on either side of a second venous valve 17b upon positioning of the balloon near the first venous valve. A third pair of heating elements, including the distal heating element 10c and a proximal heating element 11c are mounted on the outside of the catheter body 2, and electrical wires 12c and 13c running through the catheter body connect the third pair of heating elements to a source of power through the proximal end of the catheter. This third pair of heating elements are located relative to the proximal balloon so that they will be located with on either side of a third venous valve 17c upon positioning of the balloon near the first venous valve. As in Figure 1, one or more suction ports 14 are connected to a suction lumen 15 (shown in phantom) within the catheter body, and provides suction to the exterior of the catheter body in the vicinity of the heating segment. Using this catheter, several incompetent valves and surrounding areas may be treated at the same time.

Figure 5 illustrates the anatomy involved in stress urinary incontinence in a female patient. The typical anatomy of a female patient **40** is illustrated in this sagittal cross section of the pelvic area, and includes anatomical structures such as the uterus **41**, the vagina **42** the urethra **43**, the bladder **44**, and the pubis symphasis **45**. The bladder neck sphincter **46** is located at the junction between the bladder and the urethra, at the proximal end of the urethra. In a bladder neck suspension procedure, sutures are used to tie the tissue posterior to the proximal urethra to anterior structures such as the Coopers ligament **47** located above the pubis symphasis 45.

Figure 6 illustrates the bladder neck sphincter repair catheter **48** adapted for use in treatment or stress urinary incontinence in a female patient. The repair catheter includes the tubular body 2 characterized by a distal section 3, a proximal section 4. The distal section in adapted for insertion into the urethra, and preferably has a distal tip adapted for insertion into the bladder. The proximal end, which remains outside the body, is adapted for connection various external systems, such as to source of vacuum, inflation reservoirs or pumps, and a heating power source, as described in reference to the venous valve repair catheter. In this embodiment, a single balloon **49** is mounted at the distal tip of the catheter, and a heating element **50** is mounted a short distance proximal to the balloon, spaced from the balloon a distance **51** chosen to facilitate and ensure placement of the heating element distal (relative to the bladder and urethra) to the bladder neck sphincter, but still fairly close to the bladder neck sphincter and in the proximal portion of the urethra. Also on the distal end of the catheter, proximal to the balloon and located in the vicinity of the heating element, one or more suction ports **52** in communication with the exterior of the catheter may be

provided. The catheter body in the heating segment has an outer diameter of about 2 to 5 mm, depending on the initial size and desired post-treatment size of the urethra in which the catheter is to be used. The catheter body is preferably a relatively stiff catheter body, to permit insertion through the urethra. The proximal end of the catheter is provided with a hub **53** fitted inflation luer **54** and suction luer connection **55**, should suction be required to draw the urethra to the catheter body, and an electrical connector **56** for providing power to the heating element. Within the catheter body 2, the requisite inflation lumen and suction lumen connect the balloon and suction ports to their respective luer connections, and electrical leads connect the heating element to the electrical connector on the proximal hub.

In use, the operator inserts the catheter of Figure 6 into the urethra, until the balloon enters the bladder. The operator then inflates the balloon so that it impedes pullout of the catheter from the urethra, and then seats the proximal surface of the balloon on the urethral opening into the bladder. This should locate the heating element just distal to the bladder neck sphincter, and the operator should confirm this with ultrasound imaging, palpation, fluoroscopy, MRI or other imaging means. With the heating element properly placed, the operator applies direct current energy through the heating element to the urethral wall distal to the bladder neck sphincter. This heating will then result in shrinkage of the urethral section distal to the bladder neck sphincter. The effect of the bladder neck suspension will thus be achieved without sutures or surgery. If the diameter of the urethra exceeds the diameter of the catheter body in the heating segment, the operator may apply suction to the urethra through the suction ports on the catheter and thereby draw down the proximal urethra onto the catheter body. After drawing down the urethra to the diameter of the catheter body,

the direct current energy is applied to the heating elements. The power supply, when used to treat the proximal urethra, should be capable of delivering direct current power in the range of 1 to 72 watts, with voltage in the range of 1-12 volts and amperage in the range of 1-6 amps.

Figure 7 illustrates the anatomy involved in gastroesophageal reflux disease, and its relationship to the parts of the repair catheter. The patient **60** is shown with the repair catheter 1 extending from outside the body, entering the mouth and extending down the esophagus **61** until the distal tip extends into the stomach **62**. When properly positioned, the heating segment will be positioned above (proximal to) the lower esophageal valve **63**. The distal balloon **64** is located at the tip of the catheter so that it may be inflated, as shown, within the stomach and below the lower esophageal sphincter. The proximal balloon **65** is located proximal to the heating segment, so that it can be inflated as shown within the esophagus to seal off a cylindrical space between the proximal balloon and the distal balloon. One or more heating elements **66** are mounted on the heating segment. Also on the distal end of the catheter, proximal to the distal balloon and located in the vicinity of the heating element, one or more suction ports **67** in communication with the exterior of the catheter may be provided. The catheter body in the heating segment has an outer diameter of about 15 to 25 mm, depending on the initial size and desired post-treatment size of the esophagus in which the catheter is to be used. The catheter body is preferably a sufficiently flexible to permit negotiation of through the mouth and into the esophagus. The proximal end of the lower esophageal sphincter repair catheter is provided with a hub **68** fitted inflation luer **69** and suction luer connection **70**, should suction be required to draw the urethra to the catheter body, and an electrical connector **71** for providing power to

the heating element. Within the catheter body 2, the requisite inflation lumens and suction lumen connect the balloon and suction ports to their respective luer connections, and electrical leads connect the heating element to the electrical connector on the proximal hub.

Figure 8 illustrates the repair catheter in use in the treatment or gastroesophageal reflux disease in a patient. The distal balloon 64 and the proximal balloon 65 have been inflated to seal off a cylindrical segment of the lower esophagus. Vacuum has been applied through the suction ports 67, drawing the esophagus surrounding the heating segment into close proximity of the heating element(s) 66. With the esophageal tissue immediately above the lower esophageal sphincter drawn down to the heating segment, the operator can energize the heating element, with direct current passed through the connector in the hub, to cause ablation of the esophagus just above the lower esophageal sphincter. The direct current power supply, when used to treat the lower esophagus, should be capable of delivering direct current power in the range of 1 to 72 watts, with voltage in the range of 1-12 volts and amperage in the range of 1-6 amps.

Determination of the endpoint of the treatment may be accomplished in several ways. Temperature sensors may be placed on the outer surface of the heating segment, and application of the heating power may be limited to maintain a temperature, based on feedback from the temperature sensors, at the surface of the probe in the range of about 45-50°C. Total power applied to the tissue supporting the valve can then be controlled by limiting the duration or time period in which power is applied. Additionally, suction and heating may be interrupted, and the vessel observed through ultrasound or other imaging technique. When the operator observes that the

vessel has shrunk to the desired size, treatment may be halted and deemed complete.

Direct current, applied to the resistive heating elements, has been discussed as the preferred power source for applying thermal energy to the structures surrounding and supporting anatomical valves. Other power source may be used, such as alternating current and radiofrequency current (RF). RF power may be applied in bipolar mode or monopolar modes. For bipolar application, RF energy will flow from one electrode on the catheter to another, such as from electrode 66d and 66p shown in Figures 7 and 8. For monopolar RF application, a ground electrical on the surface of the patient's body must be provided, and RF energy will flow from each electrode 66d and 66p to the surface ground electrode. Various other sources of ablative or injurious power may be used, including lower frequency AC electrical power, ultrasound energy, radiation, cryosurgical devices and chemical ablating agents. The energy is applied to damage or injure tissue in the body of the vessel that supports the valve which controls flow of fluids through the vessel. This tissue in the body of the vessel may be distal to the valve, proximal to the valve, or both. Preferably, the valve itself is not injured unless injury is indicated for additional treatment of the incompetence.

Thus, while the preferred embodiments of the devices and methods have been described in reference to the environment in which they were developed, they are merely illustrative of the principles of the inventions. Other embodiments and configurations may be devised without departing from the spirit of the inventions and the scope of the appended claims.

I claim:

1. A device for treating an incompetent anatomical valve or sphincter within the body of a patient, wherein said valve or sphincter controls flow of fluid through a vessel of the body and is supported by tissue of the vessel near the valve, said device comprising:

a catheter body having a distal end and a proximal end, said distal end being adapted for insertion into the body;

a first balloon located at the distal end of the catheter, said first balloon being inflatable to a diameter greater than the catheter body distal end, and a first inflation lumen communicating from the proximal end of the catheter body to the distal end of the catheter body;

a heating element mounted on the distal end of the catheter, proximal to the first balloon;

a suction lumen communicating from the proximal end of the catheter body to the distal end of the catheter body, and a suction port located on the distal end of the catheter communicating from suction lumen to the exterior of the catheter body, said suction port being located proximate the heating element; whereby suction applied to the vessel through the suction port will draw the tissue of the vessel near the valve toward the heating element.

2. The device of claim 1 further comprising:

a second balloon located at the distal end of the catheter, proximal to the first balloon, the heating

element and suction port, said second balloon being inflatable to a diameter greater than the catheter body distal end.

3. The device of claim 2 further comprising:

a second inflation lumen communicating from the proximal end of the catheter body to the second balloon on the distal end of the catheter body.

4. The device of claim 1 further comprising:

a pair of wires running from the heating element to the proximal end of the catheter, said wires adapted to electrically connect the heating element to direct current power supply; and

wherein the heating element is a resistive heating element.

5. The device of claim 1 further comprising:

a wire running from the heating element to the proximal end of the catheter, said wire adapted to electrically connect the heating element to a radiofrequency power supply; and

wherein the heating element is a radiofrequency electrode adapted for transmission of radiofrequency energy into the tissue of the vessel.

6. A method of treating a vessel of the human body, wherein the vessel includes an anatomical valve which controls the flow of fluids through the vessel, said method comprising;

providing a catheter having a distal end adapted for insertion into the vessel, a first balloon disposed on the distal end of the catheter, a heating element

disposed on the distal end of the catheter a short distance proximal to the balloon, a suction port located near the heating element;

inserting the distal end of the catheter into the vessel so that the heating element is located near the valve;

inflating the first balloon;

applying suction to the vessel through the suction port to draw down the vessel wall until the vessel is in contact with the heating element;

heating the heating element to cause thermal injury to the vessel; and

withdrawing the catheter from the vessel.

7. The method of claim 6 further comprising the steps of:

providing a second balloon on the distal end of the catheter, said second balloon located proximal to the first balloon, the heating element and the suction port;

after inserting the distal end of the catheter into the vessel, inflating the second balloon to isolate the section of the vessel including the valve between the first and second balloons.

8. The method of claim 6 further comprising:

flushing a fluid through the suction port into the vessel prior to applying suction to the vessel through the suction port.

Abstract

Devices and method for treating various incompetent anatomical valves by thermally damaging the nearby supporting tissue of the body vessel controlled by the valve.

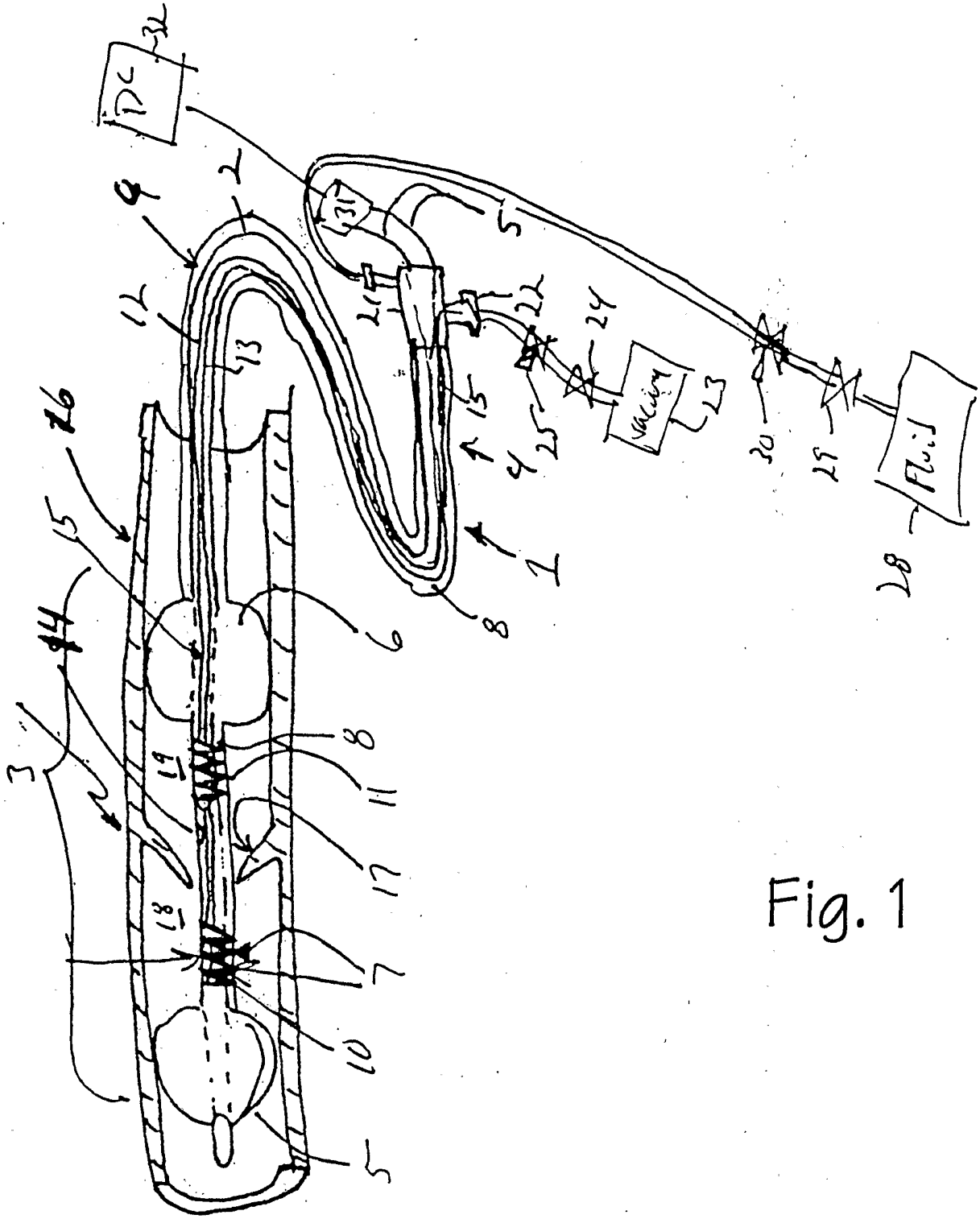

IDC
32
31
2
9
12
13
16
15
14
3
19
18
6
8
11
17
7
10
5
1
4
25
22
24
23
Vacuum
30
29
Fluid
28

Fig. 1

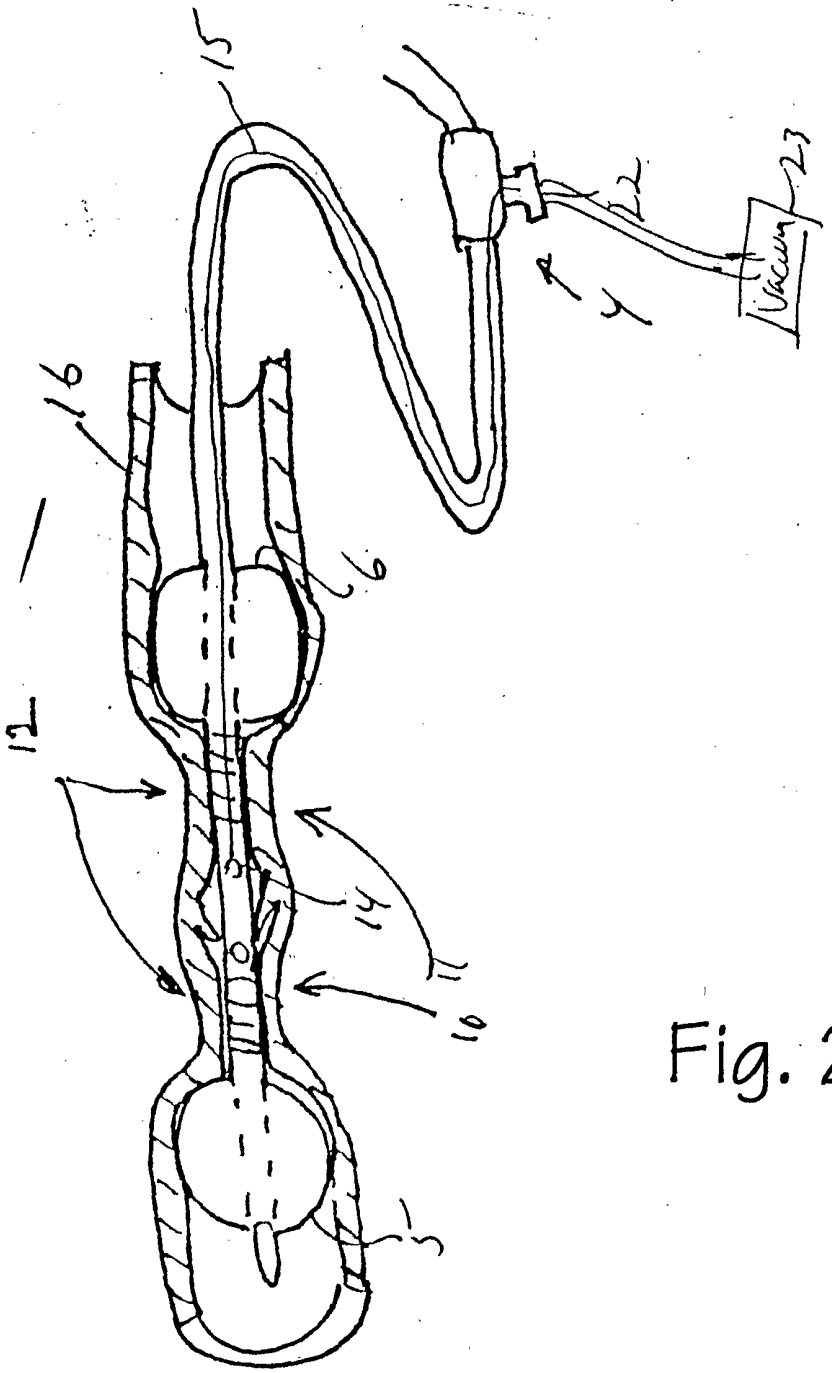


Fig. 2

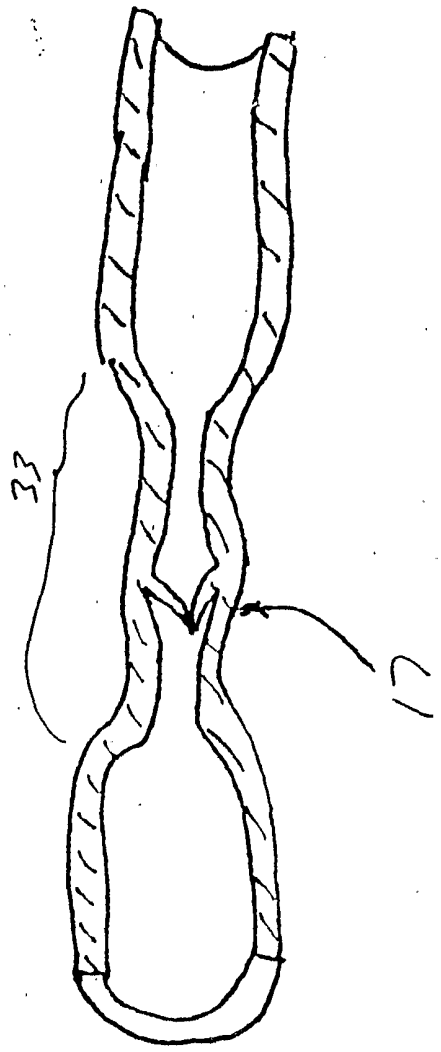


Fig. 3

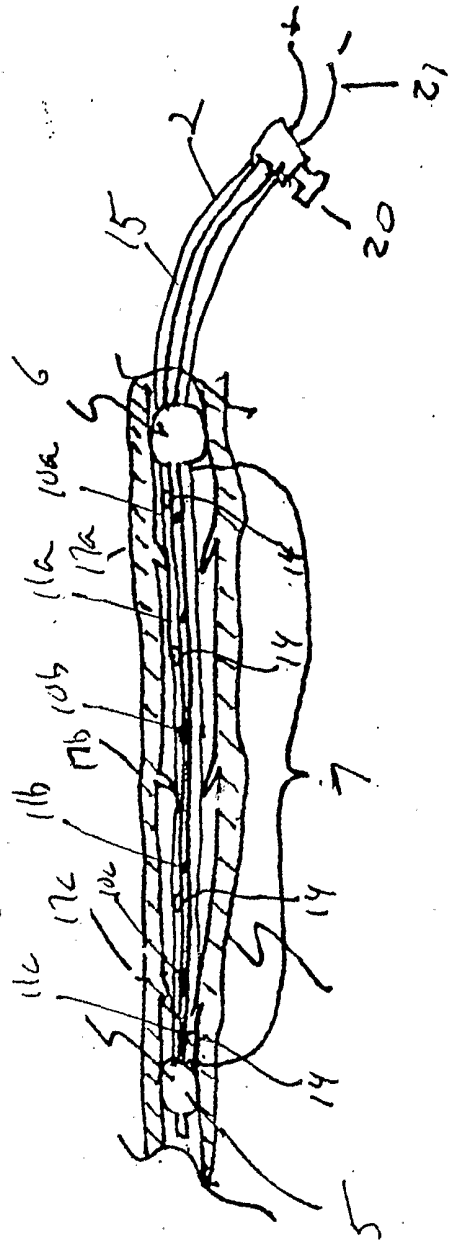


Fig. 4

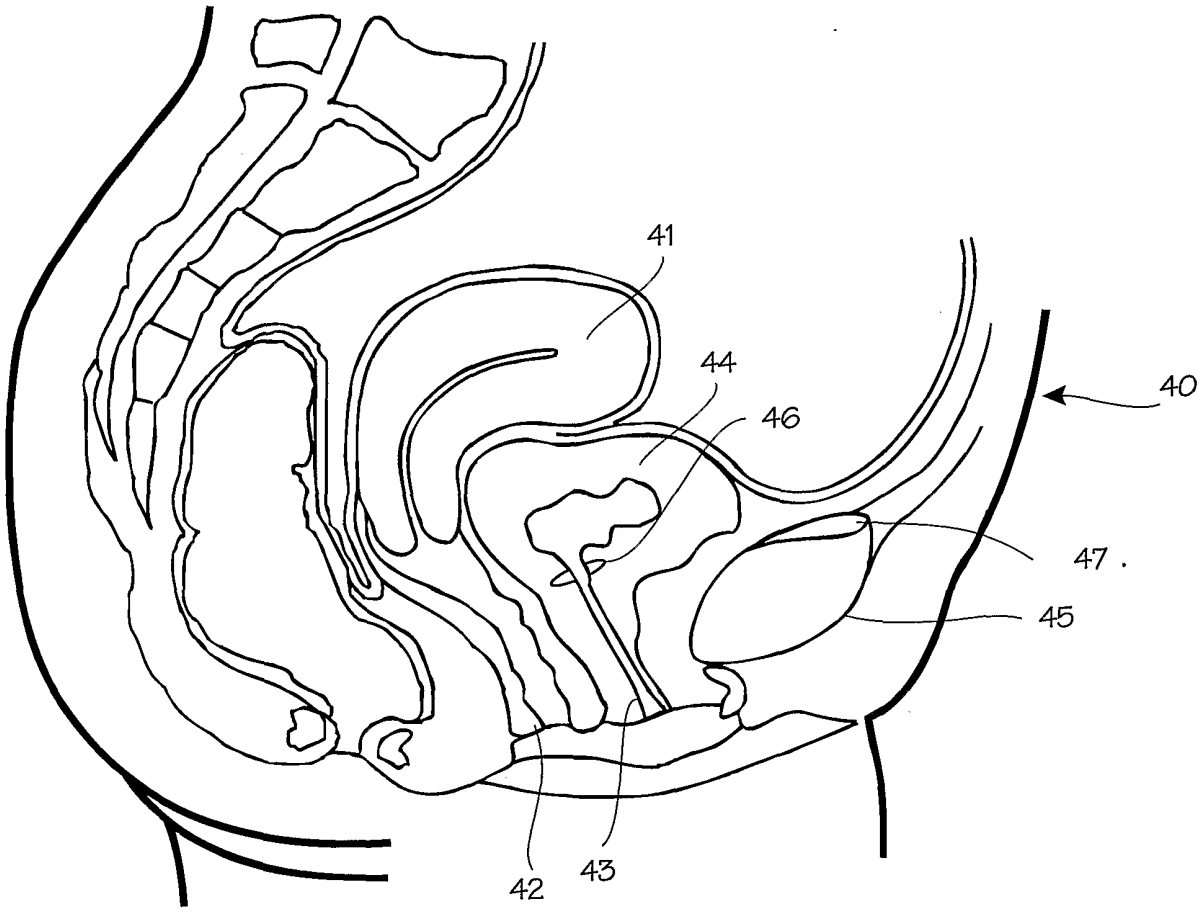


Fig. 5

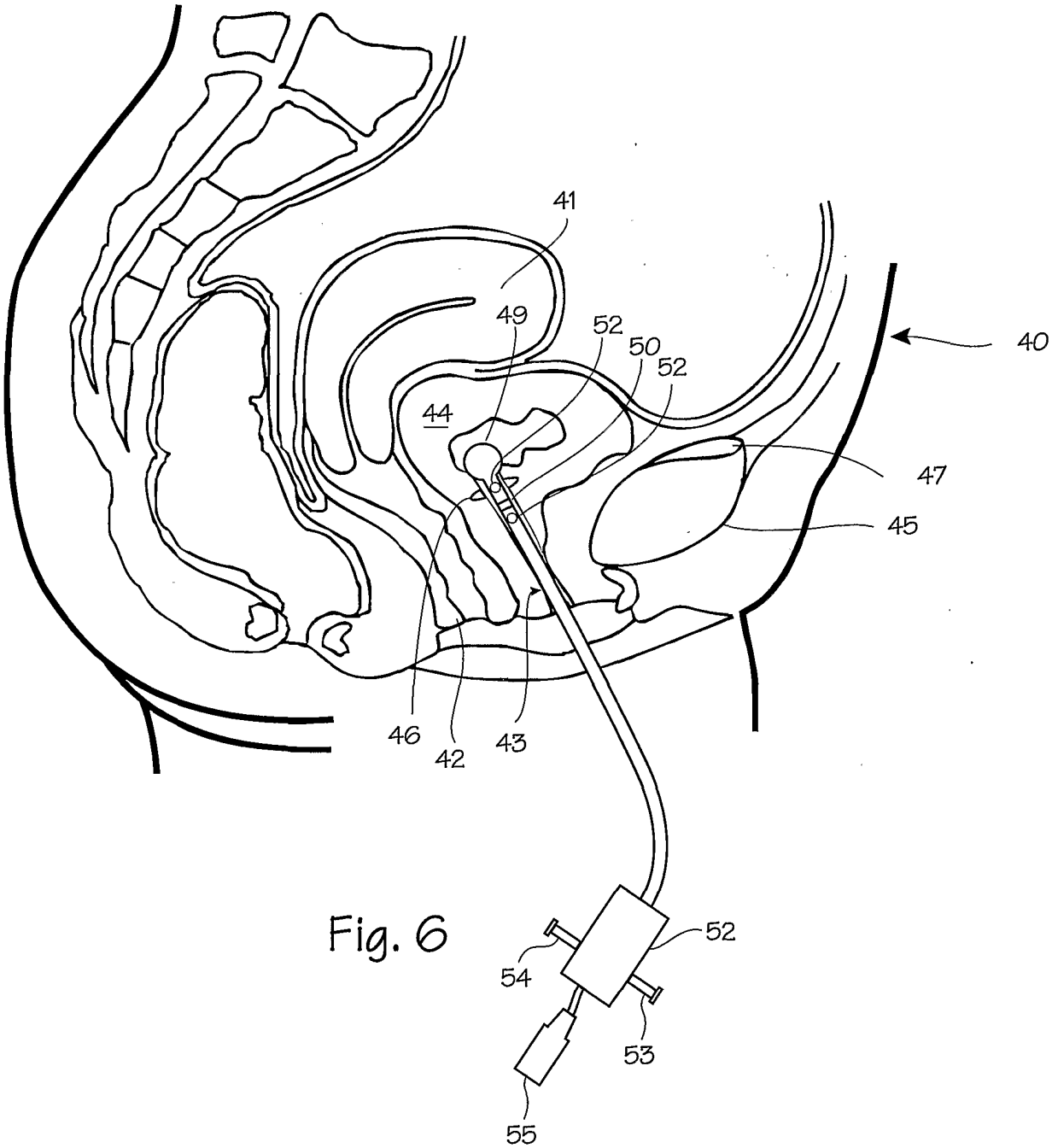


Fig. 6

Fig. 7

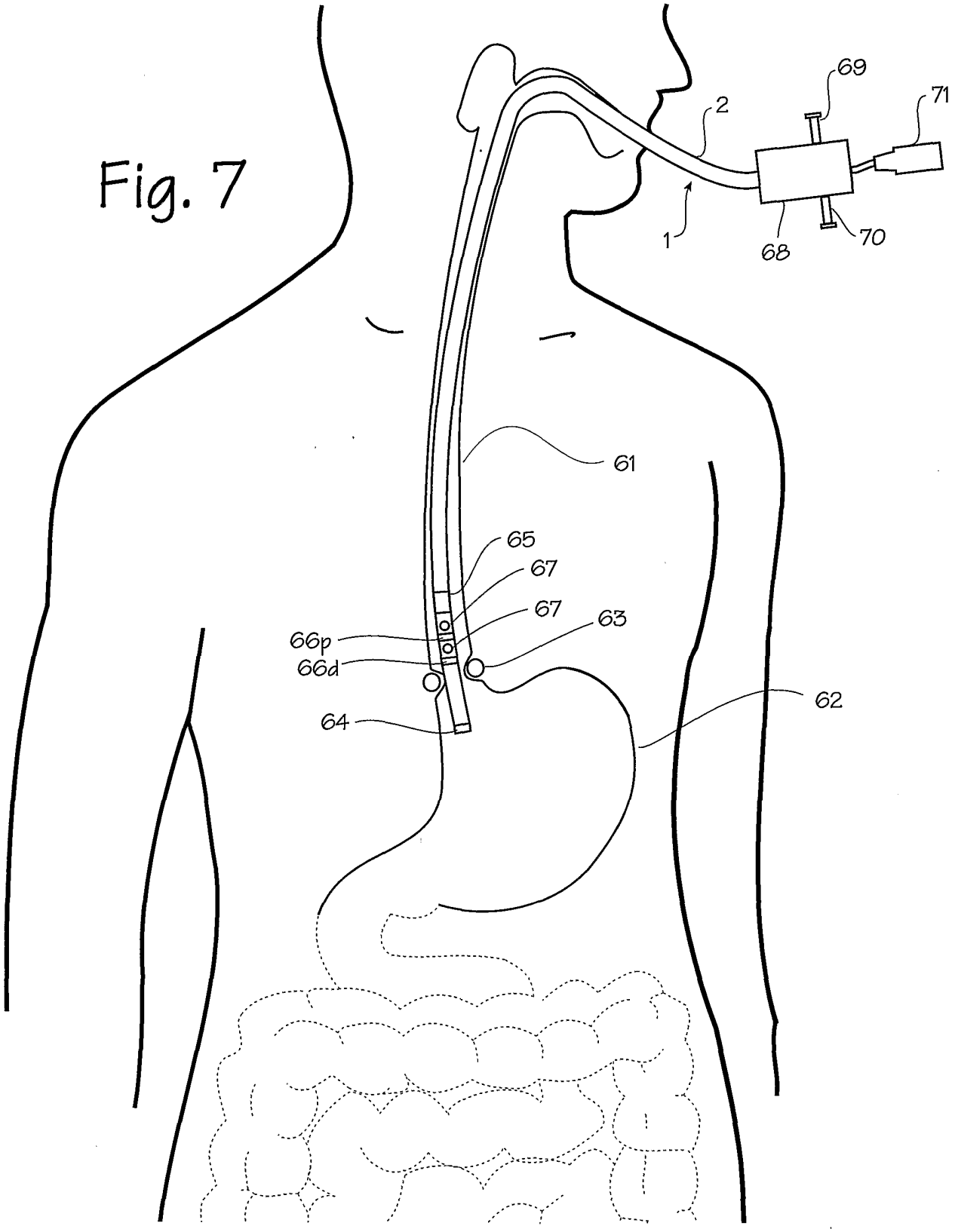
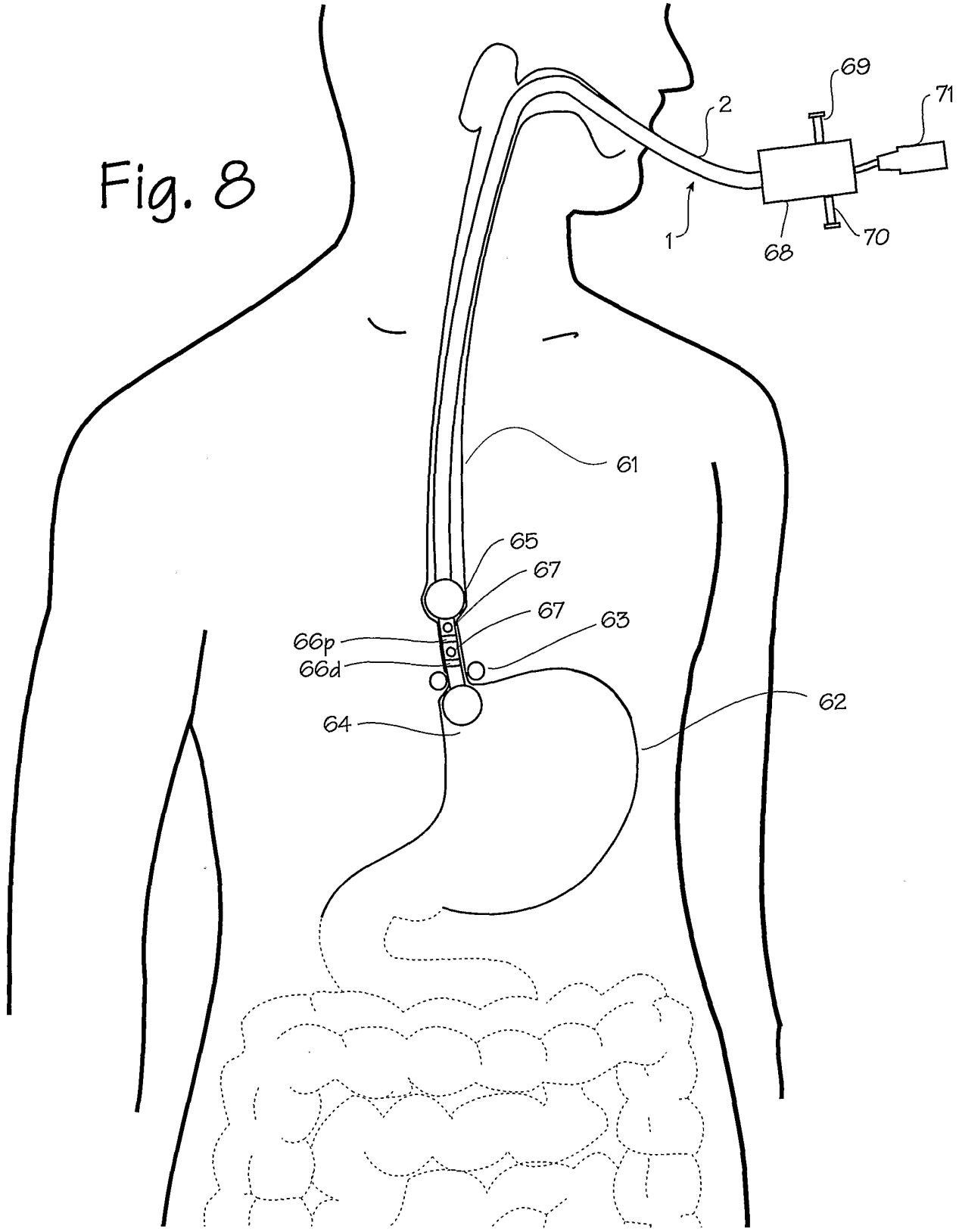


Fig. 8



INTERNATIONAL SEARCH REPORT

International application No.
PCT/US01/19470

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61M 29/00 ; A61F 7/00
US CL : 604/101.01; 606/194 ;607/96

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)
U.S. : 604/96.01, 101.01-101.05, 102.01-102.03, 103.01; 606/182-197, 26, 32 ;607/96, 100-102, 104-105

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

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

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,749,922 A (SLEPIAN et al.) 12 May 1998, see abstract and figs. and cols. 8-10	1-3,5
---		---
Y		4,6-8
X	US 5,405,322 A (LENNOX et al.) 11 April 1995, see abstract and figs. and cols. 3-4.	1-3,5
---		---
Y		6-8
X	US 4,776,349 A (NASHEF et al.) 11 Octeober 1988, see abstract and fig. 2 and cols. 1-3.	1-3,5
---		---
Y		4,6-8

☐ 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 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
26 July 2001 (26.07.2001)

Date of mailing of the international search report
16 NOV 2001

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231
Facsimile No. (703)305-3230

Authorized officer
LoAn H Thanh
Telephone No. 703-308-0858

Form PCT/ISA/210 (second sheet) (July 1998)