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Regadas

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(54) **POLYP REMOVAL DEVICE AND METHOD OF USE**

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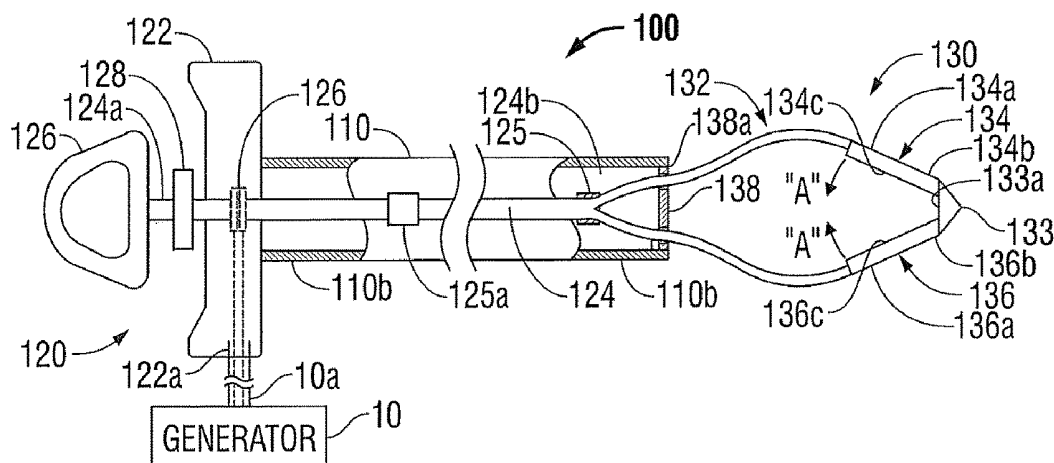
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(57) **ABSTRACT**

A device and system for removing polyps is provided and includes a tubular member having proximal and distal ends, a snare portion operably extending from within the distal end of the tubular member, the snare portion including a cutting member for severing the sealed tissue. The polyp removal device may further include a handle portion operatively extending from within the proximal end of the tubular member. The handle portion may be configured for operable engagement by a user and the cutting member may extend between distal ends of the first and second electrodes.

21 Claims, 4 Drawing Sheets



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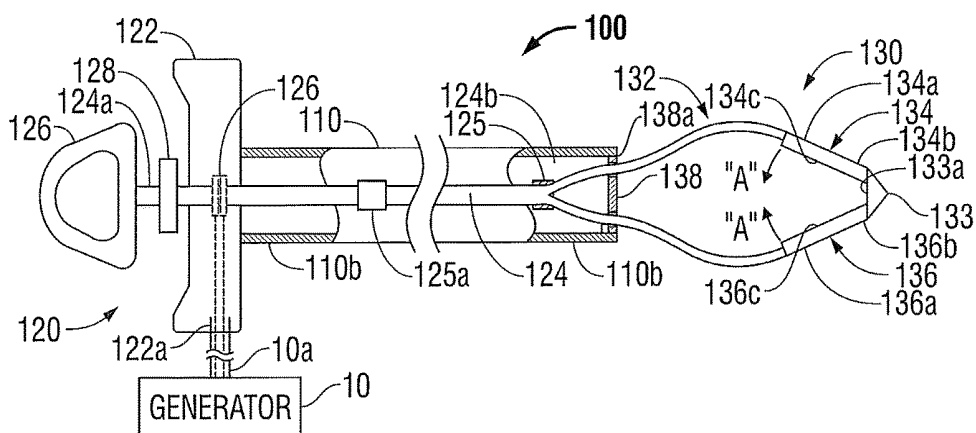


FIG. 1

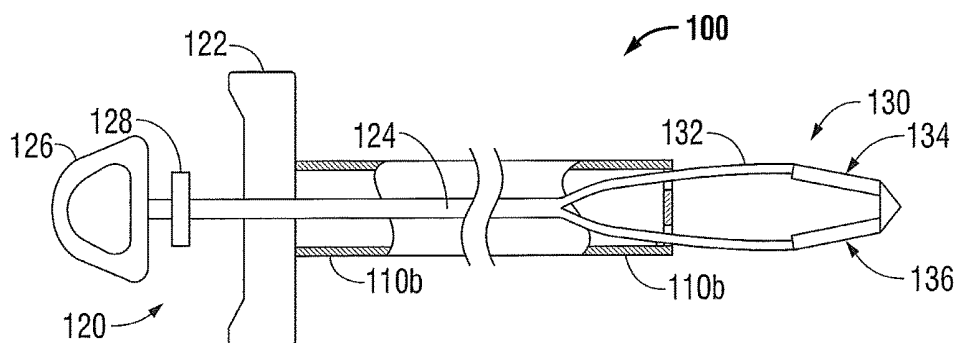


FIG. 2

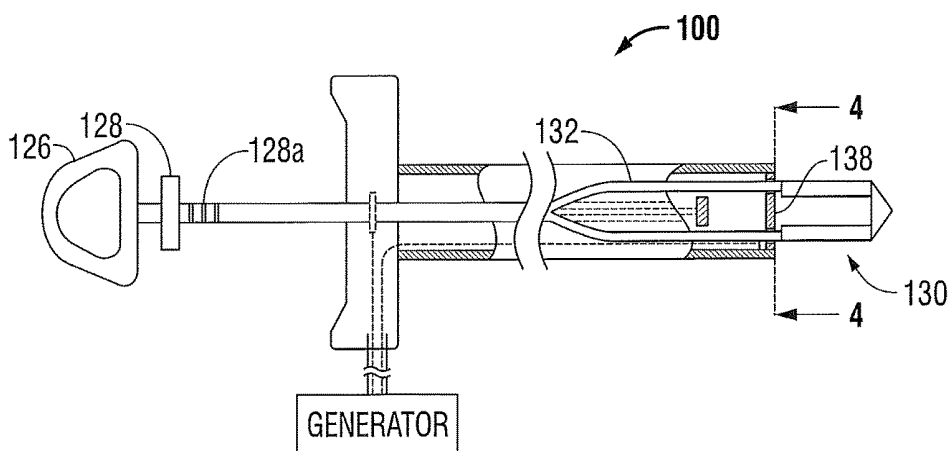


FIG. 3

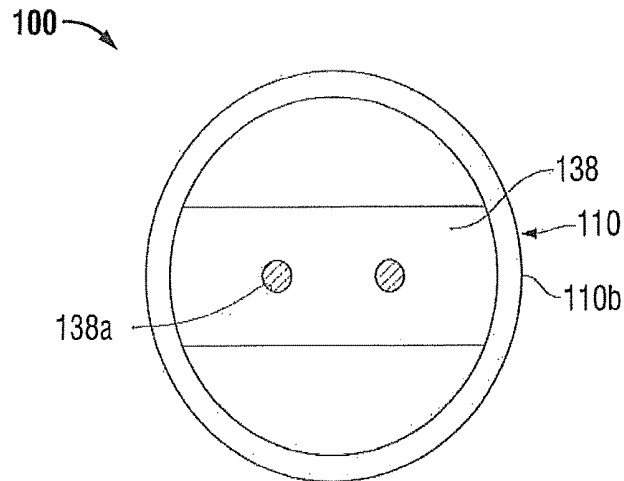


FIG. 4

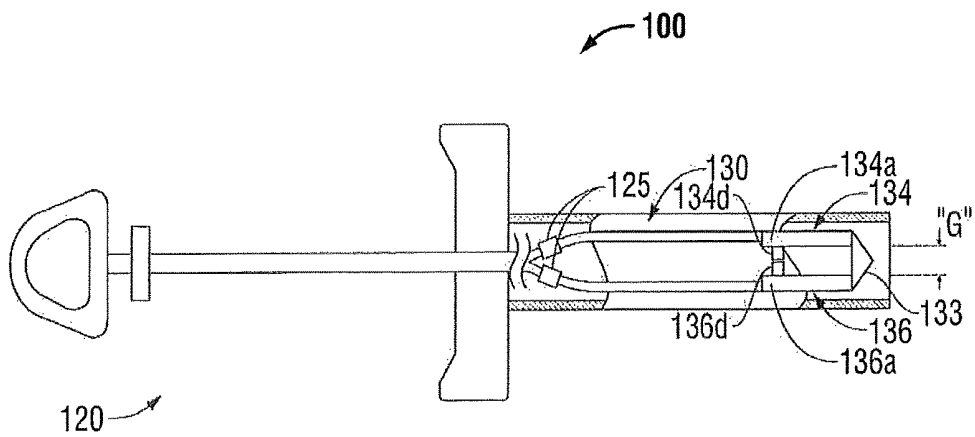


FIG. 5

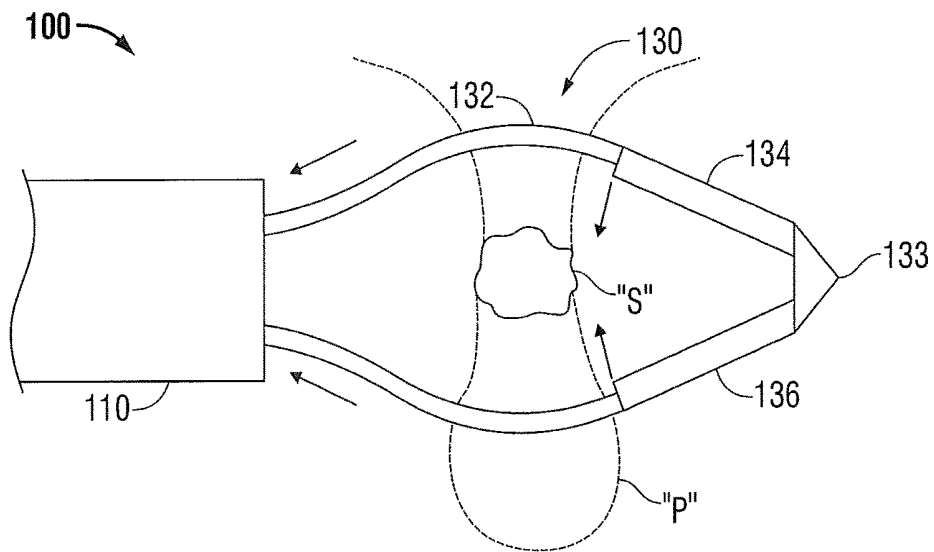


FIG. 6

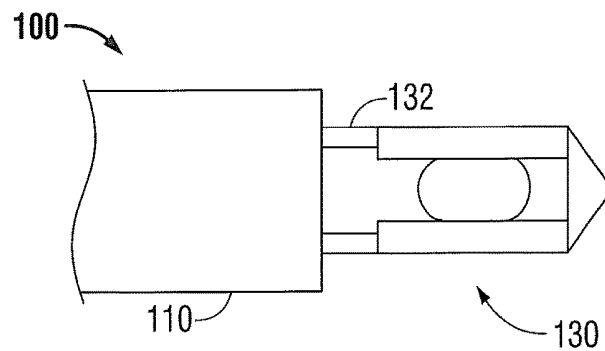


FIG. 7

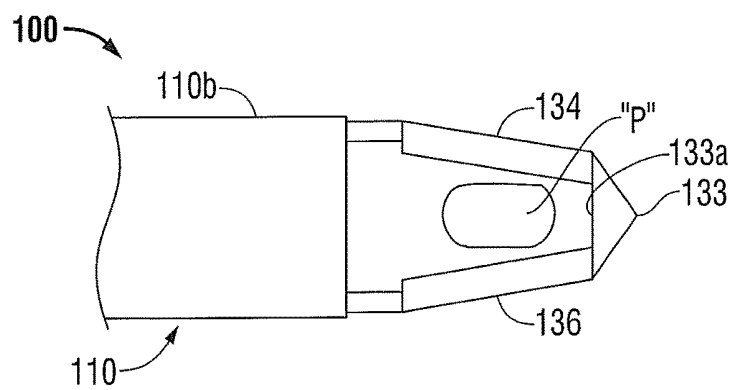


FIG. 8

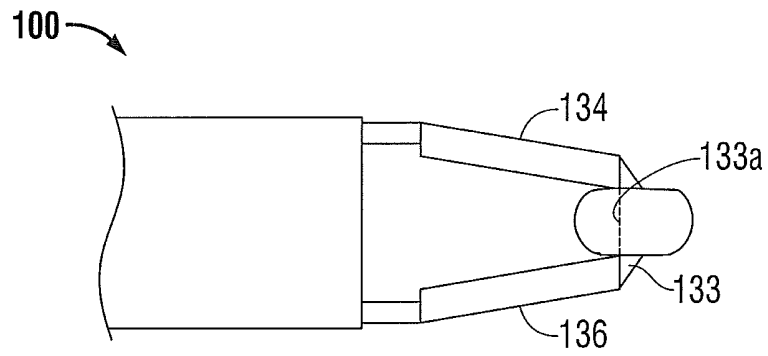


FIG. 9

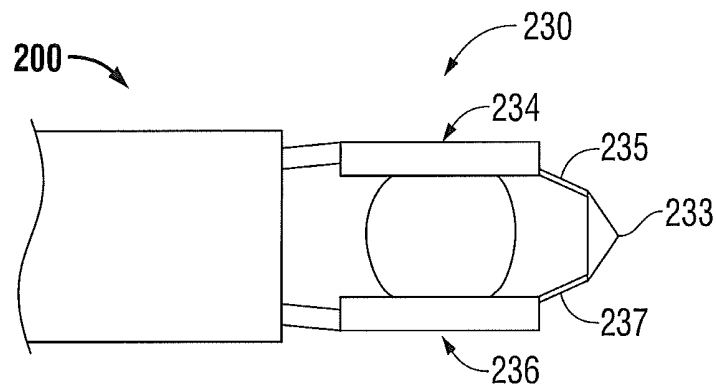


FIG. 10

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POLYP REMOVAL DEVICE AND METHOD OF USE

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation application of U.S. patent application Ser. No. 12/363,086, filed Jan. 30, 2009, which claims the benefit of and priority to U.S. Provisional Patent Application Ser. No. 61/063,158, titled "Endoscopic Flexible Loop for Gastrointestinal Polypectomy and Mucosal Resection Using LigaSure™ Sealing Technology", by Wexner et al., filed Jan. 31, 2008, the entire contents of these applications are incorporated by reference herein.

BACKGROUND

Technical Field

The present disclosure relates to devices and methods for the removal of internal tissue and, more particularly, to snare-type devices including a pair of electrodes for treating tissue prior to excision.

Background of Related Art

A polyp is an abnormal growth of tissue projecting from a mucous membrane. A polyp that is attached to the surface of the mucous membrane by a narrow elongated stalk is said to be pedunculated. If no stalk is present, the polyp is said to be sessile. Polyps are commonly found in the colon, stomach, nose, urinary bladder and uterus. Polyps may also form elsewhere in the body where mucous membranes exist, for example, the cervix and small intestine.

The surgical procedure for removing a polyp is generally referred to as a "polypectomy". Polypectomies are generally endoscopic or laparoscopic procedures performed through the oral or anal cavities. When the location of the polyp permits, the polypectomy may be performed as an open procedure. Conventional polypectomies are completed using various apparatus and techniques known in the art.

As noted above, there are two forms of polyps, sessile and pedunculated. The stalkless or sessile polyps are generally removed using electrical forceps. For example, the excess tissue projecting from the mucous membrane is cauterized and torn from the tissue wall. Pedunculated polyps, or those with stalks, tend to be larger with a greater blood supply. The size and shape of pedunculated polyps typically do not lend themselves to being removed using traditional forceps. Unlike sessile polyps, polyps with a stalk cannot simply be grasped in the jaw members of an electrosurgical forceps and be torn from the tissue wall. Instead, the polypectomy is performed using a surgical snare device.

Conventional snare devices are configured with a snare for looping over the distal end of a hanging polyp and tightening securely around the stalk of the polyp. By constricting the snare, and selectively applying energy to the snare, the device may cauterize or seal the polyp at the stalk as the polyp is severed from the tissue wall. Conventional snare devices may be configured for monopolar or bipolar use. Excising a polyp using a conventional snare device typically involves cutting or otherwise separating the polyp from the tissue wall as the snare device is activated and constricted about the stalk of the polyp. In this manner, the polyp is cauterized as the snare passes through the tissue.

SUMMARY

Disclosed is a device configured for removing polyps. In one embodiment, the polyp removal device includes a

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tubular member having proximal and distal ends, a snare portion operably extending from within the distal end of the tubular member and including first and second electrodes configured to seal tissue therebetween, the snare portion further including a cutting member for severing the sealed tissue. The polyp removal device may further include a handle portion operatively extending from within the proximal end of the tubular member. The handle portion may be configured for operable engagement by a user. The cutting member may extend between distal ends of the first and second electrodes.

The polyp removal device may be configured to grasp the stalk of a polyp between the first and second electrodes. The first and second electrodes may be hingedly attached to the cutting member. The snare portion may be configured to be retracted within the tubular member. The snare portion may be retracted within the tubular member upon retraction of the handle portion relative to the base portion.

A system including the polyp removal device may further include an electrosurgical generator. The electrodes may be electrically connected to the electrosurgical generator. The distal end of the tubular member may include a spacer member for preventing the first and second electrodes from contacting one another. Alternatively, at least one of the first and second electrodes may include a spacer mounted thereon for preventing contact between the first and second electrodes.

Also provided is a polyp removal device including a tubular member having proximal and distal ends, a handle portion operatively extending from the proximal end of the tubular member, and a snare portion slidably supported within the lumen of the tubular member and operably extending from the distal end of the tubular member, the snare portion including first and second electrodes operably mounted thereto for sealing tissue therebetween, wherein at least one of the first and second electrodes includes at least one spacer mounted thereon for preventing contact between the first and second electrodes. The snare portion of the polyp removal device may be retractable with the tubular member.

Additionally provided is a method of removing a polyp. The method includes the steps of providing a polyp removal device including, a tubular member having proximal and distal ends, a snare portion slidably supported within the lumen of the tubular member and operably extending from the distal end of the tubular member, the snare portion including first and second electrodes operably mounted thereto for sealing tissue therebetween, wherein at least one of the first and second electrodes includes at least one spacer mounted thereon for preventing contact between the first and second electrodes, extending the snare portion relative to the tubular member, looping the snare portion about a portion of a polyp, retracting the snare portion relative to the tubular member to capture the portion of the polyp between the first and second electrodes, and activating the first and second electrodes.

The method may further include the steps of partially advancing the snare portion relative to the tubular member, retracting the tubular member and the snare portion relative to the polyp and severing the polyp from surrounding tissue.

BRIEF DESCRIPTION OF THE DRAWINGS

The accompanying drawings, which are incorporated in and constitute a part of this specification, illustrate embodiments of the present disclosure and, together with the

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detailed description of the embodiments given below, serve to explain the principles of the disclosure.

FIG. 1 is a partial cross-sectional, top plan view of a polyp removal device according to an embodiment of the present disclosure, shown in a first or extended position;

FIG. 2 is a partial cross-sectional top plan view of the polyp removal device of FIG. 1, shown in a partially retracted position;

FIG. 3 is a partial cross-sectional top plan view of the polyp removal device of FIG. 1, shown in a retracted position;

FIG. 4 is a cross-section end view of the polyp removal device of FIGS. 1-3 taken along line 4-4 of FIG. 3;

FIG. 5 is a partial cross-section top plan view of an alternative embodiment of a polyp removal device according to the present disclosure in a completely retracted position;

FIG. 6 is a partial top plan view of the distal end of the polyp removal device of FIGS. 1-4, in a first or extending position about a polyp;

FIG. 7 is a partial top plan view of the distal end of the polyp removal device of FIG. 6, in a retracted position about a polyp;

FIG. 8 is a partial top plan view of the distal end of the polyp removal device of FIGS. 6 and 7, in a partially advanced position about a polyp;

FIG. 9 is a partial top plan view of the distal end of the polyp removal device of FIGS. 6-8, as the stalk of the polyp is being severed; and

FIG. 10 is a partial top plan view of another illustrative embodiment of a polyp removal device of the present disclosure, in a retracted position about a polyp.

DETAILED DESCRIPTION OF EMBODIMENTS

The foregoing summary, as well as the following detailed description will be better understood when read in conjunction with the appended figures. For the purpose of illustrating the present disclosure, various embodiments are shown. It is understood, however, that the present disclosure is not limited to the precise arrangement and instrumentalities shown.

As shown in the drawings and described throughout the following description, as is traditional when referring to relative positioning on an object, the term "proximal" refers to the end of the apparatus that is closer to the user and the term "distal" refers to the end of the apparatus that is further from the user.

Referring to FIGS. 1-4, an embodiment of the presently disclosed polyp removal device is shown therein and is generally designated as polyp removal device 100. Polyp removal device 100 includes an elongated tubular member 110, a handle portion 120 extending proximally from tubular member 110, and a snare portion or assembly 130 operably engaged with handle portion 120 and extending distally from within elongated tube 110. Polyp removal device 100 may be any suitable length and size for accessing various locations throughout the body. Device 100 may be configured for removal of suitable types of polyps of any size, or may be configured for general mucosal resection or removal of any suitable tissue mass. Preferably, polyp removal device 100 is configured for endoscopic, laparoscopic or transluminal insertion.

Referring initially to FIGS. 1-3, elongated tubular member 110 includes proximal and distal ends 110a, 110b. Proximal end 110a of tubular member 110 is operably coupled to handle portion 120. As will be described in further detail below, handle portion 120 may be integrally

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formed with tubular member 110. Alternatively, handle portion 120 may be releasably secured to tubular member 110, or may instead be securely affixed to tubular member 110. Tubular member 110 may be flexible, semi-rigid, or rigid and may be constructed of metal, alloy, plastic, polymers, or any other suitable material. Distal end 110b of tubular member 110 is configured to slidably receive a proximal end of snare portion 130.

Although the following description of the polyp removal device 100 will be described in the form of a snare-type device, the aspects of the present disclosure may be modified for use with alternative handle configurations, including a pistol grip style device, mechanically and/or remotely actuated handles, other suitable types of devices or handle arrangements. As one example, polyp removal device 100 may be incorporated as a pistol grip style laparoscopic device like some of the LigaSure™ tissue fusion devices sold by Valleylab of Boulder, Colo.

Still referring to FIGS. 1-3, handle portion 120 of polyp removal device 100 includes a base 122 coupled to proximal end 110a of tubular member 110, a connector shaft 124 extending through base 122, and a handle 126 operably coupled to a proximal end 124a of connector shaft 124. In one embodiment, base 122 of handle portion 120 may be securely affixed to proximal end 110a of tubular member 110 using adhesive, bonding, mechanical fasteners, welding or other suitable methods. Alternatively, base 122 may be releasably connected to tubular member 110 using mechanical fasteners, threaded engagement, friction fitting, bayonet connections, or the like. In this manner, snare portion 130 may be removed and replaced through proximal end 110a of tubular member 110. Base 122 of handle portion 120 may instead be integrally formed with proximal end 110a of tubular member 110. Handle 126 is configured for operable engagement by a user. Handle 126 may be knurled or may include a coating for facilitating engagement by a user.

Base 122 of handle portion 120 is further configured for operable engagement with a generator 10. Base 122 defines a connection port 122a for receiving an electrical cord 10a extending from generator 10. Electrical cord 10a extends from connection port 122a through base 122 and is operably coupled to first and second electrodes 134, 136 of snare portion 130. Electrical cord 10a may be directly coupled to first and second electrodes 134, 136. Alternatively, electrical cord 10a may couple with a coupling member 123 mounted to shaft 124. In this manner, coupling member 123 electrically couples generator 10 with first and second electrodes 134, 136 along shaft 124. Generator 10 may include any suitable generator configured to selectively provide energy to electrodes 134, 136. For example, radiofrequency energy, either monopolar or bipolar may be provided to electrodes 134, 136. In other embodiments, ultrasonic, microwave, or laser energy may be provided. Depending on the energy modality utilized, snare 130 and/or electrodes 134, 136 may have to be reconfigured in order to handle the different types of energy.

In an embodiment where radiofrequency energy is utilized, to increase safety, bipolar radiofrequency energy is preferred because the energy will travel between first and second electrodes 134, 136 instead of between the snare 130 and some remote return pad. In one particular embodiment, generator 10 takes the form of a battery-powered generator that is integral with, or releasably coupled to, polyp removal device 100. In this example, electrical cord 10a is not needed. Generator 10 may also be configured to monitor the electrical properties of the tissue maintained between first

and second electrodes **134**, **136** and signal to the clinician when conditions have been met that are suitable for sealing of tissue.

Still referring to FIGS. 1-3, connector shaft **124** includes an elongated tubular shaft including proximal and distal ends **124a**, **124b**. Proximal end **124a** is configured for operable engagement with handle **126**. Handle **126** may be securely affixed, releasable coupled, or integrally formed with proximal end **124a** of connector shaft **124**. Handle portion **120** may further include a stop member **128** positioned about connector shaft **124** between base **122** and handle **126**. Stop member **128** prevents complete retraction of connector shaft **124** past base **122**. Proximal end **124a** may be configured to securely retain stop member **128**, or instead, may include grooves or indents **128a** for selectively positioning stop member **128** thereabout. In this manner, adjustment of stop member **128** reduces or increases the amount of extension of snare portion **130**. Alternatively, stop member **128** may be configured to selectively engage distal end **124a** of connector shaft **124** through the incorporation of a mechanical fastener or the like (not shown). Distal end **124b** of connector shaft **124** is operably coupled to snare portion **130**. Distal end **124b** may be permanently coupled, releasably coupled or integrally formed with snare portion **130**.

Snare portion **130** includes a snare **132** formed of a loop of rigid or semi-rigid wire or flexible band. Snare **132** may be constructed of metal, polymer or other suitable material. In the illustrated embodiment, snare **132** is non-conductive; however, snare **132** may be conductive or partially conductive. Coupled to snare **132** in any suitable manner near a distal end **130b** of snare portion **130** are first and second electrodes **134**, **136**. As discussed above, first and second electrodes **134**, **136** are electrically coupled to generator **10** to receive electrosurgical energy therefrom. First and second electrodes **134**, **136**, which may be of any suitable construction, each include at least a tissue contacting surface **134c**, **136c**. In some embodiments, first and second electrodes **134**, **136** are each formed from a tissue contacting surface **134c**, **136c** and an insulative body portion. Tissue contacting surfaces **134c**, **136c** may include any configuration suitable for treating tissue, including hatching, grooves and detents (not shown). In one embodiment, generator **10**, first and second electrodes **134**, **136**, and tissue contacting surfaces **134c**, **136c**, are all configured for sealing tissue.

Positioned between first and second electrodes **134**, **136** is a connection member **133**. Distal ends **134b**, **136b** of first and second electrodes **134**, **136**, respectively, are coupled to connection member **133**. First and second electrodes **134**, **136** and connection member **133** are configured such that in a first or extended position, proximal ends **134a**, **136a** of first and second electrodes **134**, **136**, respectively, are substantially spaced from one another. As will be discussed in further detail below, this configuration opens snare **132** to facilitate the looping of snare **132** over a polyp "P" (FIG. 6) and the positioning of first and second electrodes **134**, **136** about stalk "S" of polyp "P".

First and second electrodes **134**, **136** and connection member **133** are further configured such that as proximal end **130a** of snare portion **130** is retracted within distal end **110b** of tubular member **110**, proximal ends **134a**, **136a** of first and second electrodes **134**, **136**, respectively, are approximated towards each other, in the direction of arrows "A" (FIG. 1). In this manner, first and second electrodes **134**, **136** and connection member **133** act like a jaw assembly, thereby compressing stalk "S" of polyp "P" between first and second electrodes **134**, **136**. To prevent first and second

electrodes **134**, **136** from contacting one another, distal end **110b** of tubular member **110** may include a spacer member **138** through which snare **132** may extend. Spacer member **138** spans distal end **110b** of tubular member **110** and includes a pair of apertures **138a** through which snare **132** is received. Apertures **138a** are spaced such that upon retraction of proximal end **130a** of snare portion **130** within tubular member **110**, proximal ends **134a**, **136a** of first and second electrodes **134**, **136** are approximated towards one another. Apertures **138a** are further configured to prevent first and second electrodes **134**, **136** contacting one another. Release of proximal end **130a** of snare portion **130** from within distal end **110b** of tubular member **110** permits proximal ends **134a**, **136a** of first and second electrodes **134**, **136** to approximate away from one another and eventual return to their initial position.

In an alternative embodiment, one or both of first and second electrodes **134**, **136** may include one or more spacers **134d**, **136d** (FIG. 5) formed on respective proximal ends **134a**, **136a**. Spacers **134d**, **136d** are configured to prevent contact between first and second electrodes **134**, **136**.

Connection member **133** extends between first and second electrodes **134**, **136** and is configured to maintain distal ends **134b**, **136b** of first and second electrodes **134**, **136**, in a spaced relationship throughout the tissue excising procedure. As described above, in a first or extended position, proximal ends **134a**, **136a** of first and second electrodes **134**, **136** are maintained in a substantially spaced apart relationship. This configuration facilitates positioning of snare **132**, and first and second electrodes **134**, **136** in particular, about stalk "S" of a polyp "P" (FIG. 6). Distal ends **134b**, **136b** of first and second electrodes **134**, **136** are coupled to connection member **133** in a manner that permits pivoting, hinging, or flexing of first and second electrodes **134**, **136** relative to connection member **133** as snare **132** is constricted about stalk "S" of polyp "P". First and second electrodes **134**, **136** and/or connection member **133** may be formed of flexible material, or may instead include a hinge mechanism or any other suitable configuration capable of permitting first and second electrodes **134**, **136** to pivot, hinge, or flex relative to connection member **133**. In the event that proximal ends **134a**, **136a** of first and second electrodes **134**, **136** are approximated away from one another by the configuration of connection member **133**, snare **132** may provide the spring force to maintain proximal ends **134a**, **136a** substantially spaced apart when polyp removal device **100** is in a first or extended position.

Connection member **133** further includes at least a first sharpened surface **133a**. First sharpened surface **133a** is formed between distal ends **134b**, **136b** of first and second electrodes **134**, **136** and may be configured to sever stalk "S" of polyp "P" after proper sealing of the tissue. In addition, or alternatively, the outer surface of connection member **133** may be sharpened to facilitate excising of polyp "P".

Referring now to FIGS. 1-3 and 6-8, one embodiment of the operation of polyp removal device **100** will be described. Initially, distal end **110b** of tubular member **110** is inserted into a patient. As discussed above, introduction of polyp removal device **100** may be accomplished through an endoscopic or laparoscopic port, or may be inserted transluminally through the mouth or anus. To facilitate insertion of tubular member **110** into the body, in one embodiment (FIG. 5), snare portion **130a** is completely retracted within tubular member **110**.

Once distal end **110b** of tubular member **110** is positioned near a polyp "P" to be removed, snare portion **130** is

extended distally by advancing handle **126** of handle portion **120** relative to base **122**. The spring-like configuration between first and second electrodes **134**, **136** and connection member **133** results in proximal ends **134a**, **136a** of first and second electrodes **134**, **136** approximating away from each other as handle **126** is advanced and proximal end **130a** of snare portion **130** is released from tubular member **110**. The separating of proximal ends **134a**, **136a** of first and second electrodes **134**, **136** opens snare **132** and facilitates positioning of snare **132** about stalk "S" of polyp "P". Once snare **132** is positioned such that stalk "S" is received between first and second electrodes **134**, **136**, handle **126** may be retracted or, in the alternative, tubular member **110** may be advanced to constrict snare **132** about stalk "S". As proximal end **130a** of snare portion **130** is received within distal end **110b** of tubular member **110** through spacer member **138**, proximal ends **134a**, **136a** of first and second electrodes **134**, **136** are approximated towards one another, thereby capturing stalk "S" of polyp "P" therebetween. Other methods of approximating first and second electrodes **134**, **136** towards one another are contemplated by the present invention. For example, another instrument separate from polyp removal device **100** may be utilized.

Continued retraction of snare portion **130** within tubular member **110** causes first and second electrodes **134**, **136** to compress stalk "S" of polyp "P". Once the clinician is satisfied stalk "S" is sufficiently received between first and second electrodes **134**, **136**, generator **10** may be activated to treat the tissue between first and second electrodes **134**, **136**. Alternatively, and as discussed above, generator **10** may include a system for monitoring the electrical properties of the tissue between first and second electrodes **134**, **136**. Once a predetermined condition has been satisfied, preferably, a condition suitable to promote tissue sealing, generator **10** signals to the clinician that the tissue is ready to be sealed, at which point, generator **10** may automatically provide electrosurgical energy to tissue contacting surfaces **134c**, **136c**, or the clinician may activate generator **10** to provide the energy to seal the tissue. Electrosurgical energy is provided to tissue contacting surfaces **134c**, **136c** until the tissue of stalk "S" is properly sealed. Electrosurgical energy may be provided at a range of frequencies, over a variable duration, and may be continuous or intermittent, depending of the type of tissue being treated and the thickness of the tissue. Furthermore, the monitoring function may include for a signal to denote that the sealing is finished and that severing of the polyp "P" may take place.

In some embodiments, in order to effectively seal larger vessels (or tissue) two predominant mechanical parameters should be accurately controlled—the pressure applied to the vessel (tissue) and the gap distance between the electrodes—both of which are affected by the thickness of the sealed vessel. More particularly, accurate application of pressure may be important to oppose the walls of the vessel; to reduce the tissue impedance to a low enough value that allows enough electrosurgical energy through the tissue; to overcome the forces of expansion during tissue heating; and to contribute to the end tissue thickness, which is an indication of a good seal. It has been determined that a typical fused vessel wall is preferably between 0.001 and 0.006 inches. Below this range, the seal may shred or tear and above this range the lumens may not be properly or effectively sealed.

With respect to smaller vessels, the pressure applied to the tissue tends to become less relevant, whereas the gap distance between the electrically conductive surfaces becomes more significant for effective sealing. In other words, the

chances of the two electrically conductive surfaces touching during activation increases as vessels become smaller.

As mentioned above, at least one electrode, e.g., **134**, may include a stop member **134d** that limits the movement of the two opposing electrodes **134**, **136** relative to one another. Stop member **134d** may extend from the tissue contacting surface **134c** a predetermined distance according to the specific material properties (e.g., compressive strength, thermal expansion, etc.) to yield a consistent and accurate gap distance "G" during sealing (FIG. 5). In one embodiment, the gap distance between opposing tissue contacting surfaces **134c**, **136c** during sealing ranges from about 0.001 inches to about 0.006 inches and, preferably, between about 0.002 and about 0.003 inches. The non-conductive stop members **134d**, **136d** may be molded onto electrodes **134**, **136** (e.g., overmolding, injection molding, etc.), stamped onto electrodes **134**, **136** or deposited (e.g., deposition) onto electrodes **134**, **136**. For example, one technique involves thermally spraying a ceramic material onto the surface of the electrodes **134**, **136** to form the stop members **134d**, **136d**, respectively. Several thermal spraying techniques are contemplated that involve depositing a broad range of heat resistant and insulative materials on various surfaces to create stop members **134d**, **136d** for controlling the gap distance between electrically conductive surfaces **134c**, **136c**.

It has also been found that the pressure range for assuring a consistent and effective seal may be between about 3 kg/cm.sup.2 to about 16 kg/cm.sup.2 and, preferably, within a working range of 7 kg/cm.sup.2 to 13 kg/cm.sup.2. Manufacturing an instrument that is capable of providing a closure pressure within this working range has been shown, in some embodiments, to be effective for sealing arteries, tissues and other vascular bundles.

As can be appreciated, controlling the compressive force between with first and second electrodes **134**, **136** as snare portion **130** is retracted may facilitate and assure consistent, uniform and accurate closure pressure about the tissue within the desired working pressure range of about 3 kg/cm.sup.2 to about 16 kg/cm.sup.2 and, preferably, about 7 kg/cm.sup.2 to about 13 kg/cm.sup.2. By controlling the intensity, frequency and duration of the electrosurgical energy applied to the tissue, the user can either cauterize, coagulate/desiccate, seal and/or simply reduce or slow bleeding.

In one embodiment, by controlling the force applied to handle **126**, the resulting tension in snare portion **130** may be adjusted, which, in turn, regulates the overall pressure between electrodes **134**, **136** to within the above-identified desired sealing range. A sensor **125** may be employed to accomplish this purpose and may be mechanically coupled to handle portion **120** and/or snare portion **130**. A visual or audible indicator (not shown) may be operably coupled to handle portion **120**, snare portion **130** and/or sensor **125** to provide visual, audible or tactile feed back to the user to assure that the clamping pressure is within the desired range prior to activation of the instrument.

Alternatively, handle portion **120** may incorporate a spring mechanism **125a** configured to prevent excessive force being applied to snare portion **130**, thereby preventing overcompression of the tissue between first and second electrodes **134**, **136**. Spring mechanism **125a** may include a compression spring (not shown) that deforms once the force applied to handle **126** exceeds the force necessary to accomplish effective sealing between first and second electrodes **134**, **136**. In an alternative embodiment, snare portion **130** may instead be advanced and retracted through the rotation

of handle 126. In this manner, handle portion 120 may include a torque mechanism, e.g., torque wrench or the like, (not shown) which is precisely set to measure the torque (rotational force) applied to snare portion 130 so the closure pressure between electrodes 134, 136 will fall within the desired pressure range.

Once the tissue of stalk "S" is sealed, handle 126 is partially advanced to extend a length of snare portion 130 from tubular member 110, thereby approximating proximal ends 134a, 136a of first and second electrodes, respectively, away from each other and exposing sealed stalk "S" to first sharpened surface 133a of connection member 133. While maintaining snare portion 130 relative to tubular member 110, tubular member 110 is retracted relative to polyp "P", thereby causing the engagement of stalk "S" with first sharpened surface 133a of connection member 133. Continued retraction of tubular member 110 causes the complete severing of stalk "S" (see FIG. 9).

When using polyp removal device 110, the likelihood of excessive bleeding is greatly reduced because the tissue is completely sealed prior to being cut. Using polyp removal device 110 further reduces the likelihood of creating an open wound that is susceptible to infection. In addition, the utilization of bipolar energy to seal a polyp or other tissue is safer than monopolar, e.g., because of less thermal and energy spread, thereby reducing the likelihood of perforations.

With reference now to FIG. 10, an alternative embodiment of the present disclosure is shown generally as polyp removal device 200. Polyp removal device 200 is substantially similar to polyp removal device 100 and will, therefore, only be described as relates to the differences therebetween. Snare portion 230 includes first and second electrodes 234, 236 and a connection member 233. Extending between first and second electrodes 234, 236 and connection member 233 are extension members 235, 237, respectively. Extension member 235, 237 permit first and second electrodes 234, 236 to be spaced further apart from one another in a second or sealing condition. In this manner, polyp removal device 200 may be used to excise larger polyps having a larger stalk.

Various changes in form, detail and operation of the polyp removal devices of the present disclosure may be made without departing from the spirit and scope of the present disclosure.

What is claimed is:

1. A tissue removal system comprising:
a tubular member having proximal and distal ends, the distal end including a spacer member that defines a plurality of apertures therethrough;
a connector shaft supported within the tubular member and having proximal and distal ends; and
a snare portion having first and second ends monolithically formed with the distal end of the connector shaft, the snare portion operably extending through the plurality of apertures of the spacer member and from the distal end of the tubular member, the snare portion movable relative to the spacer member and including a cutting member having a sharpened surface.
2. The tissue removal system of claim 1, further comprising a handle portion secured to the proximal end of the connector shaft and operatively coupled to the proximal end of the tubular member.
3. The tissue removal system of claim 2, wherein the handle portion is configured for operable engagement by a user and movable relative to the tubular member.

4. The tissue removal system of claim 2, wherein the sharpened surface of the cutting member is configured to cut tissue in response to movement of the handle portion in a proximal direction relative to the tubular member.

5. The tissue removal system of claim 1, wherein the snare portion is configured to grasp a stalk of a polyp.

6. The tissue removal system of claim 1, wherein the snare portion is configured to be retracted within the tubular member in response to movement of the connector shaft in a proximal direction.

7. The tissue removal system of claim 1, further including an electrosurgical generator.

8. The tissue removal system of claim 7, wherein the snare portion is electrically coupled to the electrosurgical generator.

9. The tissue removal system of claim 1, wherein first and second apertures of the plurality of apertures of the spacer member are spaced from one another to prevent a first portion of the snare portion from contacting a second portion of the snare portion.

10. The tissue removal system of claim 1, wherein the tubular member is configured for endoscopic insertion, laparoscopic insertion or transluminal insertion.

11. The tissue removal system of claim 2, wherein the handle portion includes handle configurations selected from the group consisting of pistol grip style handles, mechanically-actuated handles and remotely-actuated handles.

12. A tissue removal system comprising:

a tissue removal device comprising:

- a tubular member having proximal and distal ends, the distal end including a spacer member that defines a plurality of apertures therethrough;
 - a handle portion operatively extending from the proximal end of the tubular member;
 - a connector shaft having a proximal end secured to the handle portion and a distal end;
 - a snare portion having first and second ends monolithically formed with the distal end of the connector shaft, the snare portion slidably supported within the tubular member, the snare portion movable through the plurality of apertures of the spacer member and from the distal end of the tubular member; and
 - a cutting member having a sharpened surface positioned on the snare portion; and
- a generator configured to generate and deliver energy to the tissue removal device.

13. The tissue removal system of claim 12, wherein the energy is selected from the group consisting of radiofrequency energy, ultrasonic energy, microwave energy, and laser energy.

14. A method of removing a tissue mass of a body comprising:

- providing a tissue removal device including a snare portion and a cutting member having a sharpened surface positioned on the snare portion, the snare portion including first and second ends monolithically formed with a distal end of a connector shaft;
- moving the snare portion through a plurality of apertures defined through a spacer member to surround the tissue mass;
- retracting the snare portion to capture the tissue mass within the snare portion;
- delivering energy to the snare portion;
- discontinuing delivery of energy to the snare portion; and
- further retracting the snare portion such that the sharpened surface of the cutting member severs the tissue mass from the body.

15. The method of claim 14, wherein delivering energy to the snare portion further includes sealing the tissue mass.

16. The method of claim 15, further comprising:
monitoring electrical properties of the tissue mass during
energy delivery;

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detecting a predetermined condition; and
signaling that the tissue mass is ready to be severed.

17. The method of claim 14, further including preventing contact between a first portion of the snare portion and a second portion of the snare portion with the spacer member.

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18. The tissue removal system of claim 1, wherein the snare portion includes at least one electrode that is hingedly connected to the cutting member.

19. The tissue removal system of claim 12, wherein the snare portion includes a proximal portion and a distal portion, the first and second ends of the snare portion are monolithically formed with the distal end of the connector shaft at the proximal portion of the snare portion.

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20. The tissue removal system of claim 19, wherein a length of the snare portion extends from the proximal portion to the distal portion, the length of the snare portion remaining substantially constant as the snare portion moves between fully approximated and fully unapproximated positions to sever a tissue mass enclosed by the snare portion.

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21. The tissue removal system of claim 1, wherein the spacer member is fixedly secured to the distal end of the tubular member.

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专利名称(译)	息肉清除装置和使用方法		
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[标]申请(专利权)人(译)	柯惠有限合伙公司		
申请(专利权)人(译)	COVIDIEN LP ,		
当前申请(专利权)人(译)	COVIDIEN LP		
[标]发明人	REGADAS F SERGIO P		
发明人	REGADAS, F. SERGIO P.		
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摘要(译)

提供了一种用于移除息肉的装置和系统，其包括具有近端和远端的管状构件，可操作地从管状构件的远端内延伸的圈套部分，圈套部分包括用于切断密封组织的切割构件。息肉去除装置可进一步包括手柄部分，其可操作地从管状部件的近端内延伸。手柄部分可以被配置用于由使用者可操作地接合，并且切割构件可以在第一电极和第二电极的远端之间延伸。

