Abstract:

A tissue resector includes an outer cannula with a cutting wire disposed therein. The outer cannula includes an opening that is configured to be associated with a cutting element of the cutting wire in such a way that when tissue is drawn into the opening of the cannula, a cutting element of the cutting wire can be rotated past the opening to engage and cut into and through the tissue. The outer cannula and the cutting wire may also be configured to enable aspiration of the tissue. A tissue resector according to this disclosure may be used with a hand-held, hand-operated rooter.
TISSUE RESECTORS WITH CUTTING WIRES, HAND-OPERATED TISSUE RESECTOR SYSTEMS AND ASSOCIATED METHODS

CROSS-REFERENCE TO RELATED APPLICATION

A claim to the benefit of the April 28, 2014 filing date of U.S. Provisional Patent Application No. 61/985,283, and titled MORCELLATORS WITH CUTTING WIRES, HAND-OPERATED MORCELLATION SYSTEMS AND ASSOCIATED METHODS ("the '283 Provisional Application") is hereby made. The entire disclosure of the '283 Provisional Application is hereby incorporated by reference.

TECHNICAL FIELD

This disclosure relates generally to tissue resectors and, more specifically, to tissue resectors that efficiently aspirate excised tissue. In particular, tissue resectors that employ wires to remove undesired tissue, such as polyps and fibroids, are disclosed. Systems and methods for tissue resecting are also disclosed.

RELATED ART

Morcellation is a process by which undesirable growths, such as benign tumors, polyps and fibroids, are removed from within a subject's body. Physicians have used morcellation in gynecological surgeries. For example, high powered electric morcellators have been used to laparoscopically extract the uterus (i.e., hysterectomy). Specifically, an electric morcellator grinds the entire uterus, including any undiagnosed sarcomas. While the risk is relatively small that these cancerous tissues remain within a woman's body, there is a significant likelihood that any cancerous cells that remain within the body may spread.

In less invasive procedures, smaller morcellators have been used in conjunction with hysteroscopes to remove relatively small uterine polyps (e.g., polyps with diameters of about 3 cm or less, etc.) and fibroids (which typically have diameters of about 1 cm to about 2 cm). Some hysteroscopes are configured to inflate the uterine cavity with fluid or air.

With the uterine cavity inflated, a light source of the hysteroscope may illuminate the interior surfaces of the uterus, and a camera of the hysteroscope and a display associated with the
camera of the hysteroscope may enable a physician to visualize features, such as polyps and fibroids, on interior surfaces of the uterus. While the physician is looking at the interior surface of the uterine wall, he or she may operate a morcellator in conjunction with the hysteroscope to remove any polyps or fibroids that appear on the display. Debris from the morcellation process may be aspirated through the morcellator, and collected for pathology.

The morcellators that are currently used to remove uterine polyps and fibroids are power-driven devices. A typical morcellator includes an outer cannula with an opening located near its distal end and formed in a portion of the circumference of the outer cannula. An inner cannula is positioned within a lumen of the outer cannula, and includes a distal end that defines and blade that communicates with the opening near the distal end of the outer cannula. Depending upon the configurations of the opening and the blade, the inner cannula may rotate within the lumen of the outer cannula, or the inner cannula may move longitudinally back and forth within the outer cannula. In any configuration, a polyp or fibroid may be drawn into the opening in the outer cannula, and then cut with the blade of the inner cannula. Once the polyp or fibroid, or a portion thereof, has been cut from the inner surface of the uterus, it may be drawn, by way of a vacuum, through a lumen in the inner cannula.

One example of an existing morcellator is the TRUCLEAR® morcellator offered by Smith & Nephew. That morcellator includes an outer cannula that has an outer diameter of 0.14 inch (2.9 mm) \( (i.e., \) a cross-sectional area of 0.0102 in\(^2\) or 6.6 mm\(^2\) \) and an inner cannula with an inner diameter of about 0.070 inch (1.8 mm) \( (i.e., \) a cross-sectional area of 0.00385 in\(^2\) or 2.5 mm\(^2\) \). Thus, the available, open cross-sectional area through that device only comprises 37.7% of the overall cross-sectional area occupied by the cannula of that device.

Hysteroscopy and morcellation can be painful. The relatively small inner diameters of the inner cannulas of existing morcellators limit the rate at which excised tissues \( (e.g., \) polyps, fibroids, etc.) may be collected, which unfortunately and undesirably prolongs the morcellation procedure and the pain caused by that procedure.

SUMMARY
A tissue resector according to this disclosure includes a cannula and a cutting wire. The cannula has a lumen within which the cutting wire may be positioned. The cannula includes an opening located at or adjacent to its distal end. The opening may be configured to receive uterine polyps and fibroids. A cutting feature at or near a distal end of the cutting wire cooperates with the opening of the cannula in such a way that the cutting feature will cut, slice, shear, chew or tear tissue that is introduced (e.g., by suction, etc.) into the opening. For the sake of simplicity, the acts of cutting, slicing, shearing, chewing, tearing and similar actions are individually and collectively referred to herein as "cutting." The use of a cutting wire within a single cannula provides for a relatively large cross-sectional area through which tissue may be aspirated through the cannula.

In various embodiments, of the entire cross-sectional area occupied by the cannula, at least 50% may be available for aspiration. In some embodiments (e.g., embodiments where smaller cutting wires are used, etc.), the open area may comprise 60% or even 70% of the entire cross-sectional area occupied by the cannula.

In another aspect of this disclosure, a tissue resector according to this disclosure may be part of a tissue resector system. In some embodiments, the tissue resector may be used with a rooter that operates under manual power, such as that disclosed by U.S. Patent Application Publication No. US 2012/0239008 of Fojtik, the entire disclosure of which is hereby incorporated herein. Such a rooter is also referred to herein as a "hand-powered rooter" and as a "manual spinning instrument." In other embodiments, the tissue resector may be used with a power-driven instrument, such as those used to drive the inner cannulas of existing morcellators, and an appropriate adapter, which translates actions of the power-driven instrument to the features of a tissue resector with a single cannula and a cutting wire.

Another embodiment of tissue resector system includes an existing tissue resector blade (with an outer cannula and an inner cannula on which a blade is defined), a hand-operated rooter and an adapter for converting actions of the rooter to the features of existing tissue resector blade.

As an example of use, a tissue resector or tissue resector system according to this disclosure may be used to remove undesired growths from a woman's uterus, including, without limitation, polyps, fibroids and other undesirable growths. While viewing such a
growth, it may be drawn into the opening of a cannula under suction (i.e., a vacuum) applied to the lumen of the cannula. With the tissue in the opening, the cutting wire may be rotated, and its cutting feature may cut tissue from the growth. This process may continue until the growth and immediately adjacent tissues have been removed. The tissues, which are aspirated, may then be collected and evaluated by a pathologist.

Other aspects, as well as features and advantages of various aspects, of the disclosed subject matter will become apparent to those of ordinary skill in the art through consideration of the ensuing description, the accompanying drawings and the appended claims.

BRIEF DESCRIPTION OF THE DRAWINGS

In the drawings:

FIGs. 1A and 1B depict an embodiment of a tissue resector that includes a cannula and a cutting wire within a lumen of the cannula;

FIGs. 2A through 2H depict various embodiments of distal ends of and openings of the cannulas of various embodiments of tissue resectors that incorporate the features shown in FIGs. 1A and 1B;

FIGs. 3A and 3B illustrate some embodiments of cannulas that include two or more openings;

FIGs. 4A through 4G show some embodiments of cutting wires that may be used with a tissue resector, such as that depicted by FIGs. 1A and 1B;

FIGs. 5A through 5C illustrate an embodiment of a tissue resector system that includes a tissue resector of the type shown in FIGs. 1A and 1B, as well as a hand-powered rooter, with FIG. 5A illustrating the tissue resector system in an assembled state, FIG. 5B providing an exploded view of the elements of the hand-powered rooter and the tissue resector and FIG. 5C providing a cross-sectional representation of the tissue resector system;

FIGs. 6A and 6B show an embodiment of an adapter for enabling use a hand-powered rooter with an existing morcellator, as well as an embodiment of a tissue resector system that includes the hand-powered rooter, the adapter and the morcellator; and

FIG. 7 depicts an embodiment of an adapter for enabling use of a tissue resector of the type illustrated by FIGs. 1A and 1B with a power-drive instrument of a conventional
morcellation device, as well as an embodiment of a tissue resector system including the power-drive instrument, the adapter and the tissue resector.

DETAILED DESCRIPTION

As shown in FIGs. 1A and IB, a tissue resector 10 according to this disclosure includes a cannula 20 and a cutting wire 30. The cannula 20 is an elongated tubular element with a wall 22 that defines a lumen 24 along its length. In the illustrated embodiment, the cannula 20 is substantially straight; however, a tissue resector 10 according to this disclosure may include a curved cannula 20 or even a bent cannula 20.

The cannula 20 of a tissue resector 10 may have any of a variety of different dimensions. Without limitation, a cannula 20 may have an outer diameter of about 5 French (i.e., 0.066 inch; 1.67 mm), about 7 French (i.e., 0.092 inch; 2.33 mm) or about 9 French (i.e., 0.118 inch; 3 mm), which may correspond to the size of a hysteroscope (e.g., to the size of an access lumen through the hysteroscope, etc.) with which the tissue resector 10 is to be used. In a specific embodiment, the cannula 20 may comprise a hypotube with an outer diameter of 0.115 inch (2.9 mm) and an inner diameter of 0.095 inch (2.4 mm).

At or near its distal end 26, the cannula 20 includes an opening 27. The opening 27 is configured to receive undesirable growths, such as uterine polyps and fibroids. In some embodiments, the edges 23 of the outer wall 22 that define the opening 27 may be configured to facilitate separation of unwanted tissue from adjacent (e.g., healthy) tissue.

FIGs. 2A through 2H illustrate various embodiments of openings 27 in the cannula 20 or a tissue resector 10.

The cannula 20' depicted by FIG. 2A includes an opening 27' through its wall 22' and positioned proximal to the distal end 26' of the cannula 20'. This configuration retains the tubularity of distal-most portion of the cannula 20', which may hold a distal end of a cutting wire 30 (FIGs. 1A and IB) in place as the cutting wire 30 is rotated within the lumen 24' of the cannula 20'. The opening 27' has a width that is about the same as a diameter of the cannula 20', and is at least partially defined by edges 23' that are oriented longitudinally relative to the length of the cannula 20'. The opening 27' is elongated, with its length being oriented along the length of the cannula 20'.
FIG. 2B depicts an embodiment of a cannula 20" that includes an opening 27" with a position and a configuration similar to those of the opening 27' of the embodiment of cannula 20' depicted by FIG. 2A. However, the edges 23" that define the periphery of the opening 27" are configured as blades, with sharper edges adjacent to an interior surface 25" of the lumen 24" than the corresponding edges 23' that define the opening 27' of cannula 20'. Thus, the edges 23" that define the opening 27" may cut into tissue as a cutting wire 30 (FIGs. 1A and IB) rotates within the lumen 24" that extends through the cannula 20".

Another similarly positioned and configured opening 27'" is depicted by FIG. 2C, but with edges 23'" that include teeth 23τ. The teeth 23τ may be configured to cut into tissue as a cutting wire 30 (FIGs. 1A and IB) rotates within the lumen 24" that extends through the cannula 20".

The cannula 20"" depicted by FIG. 2D includes an opening 27"" with edges 23"" that comprise serrations 23s. Accordingly, each edge 23"" may include a series of short, curved sections that are adjoined at teeth 23τ"".

The embodiment of tissue resector 10"" illustrated by FIG. 2E includes a cannula 20"" with an opening 27"" that is at least partially defined by edges 23"" that are oriented oblique to the length, or longitudinal axis, of the cannula 20"".

In FIG. 2F, an embodiment of tissue resector 110 is shown with an opening 127 that extends to the distal end 126 of the cannula 120. In that embodiment, the portion of the distal end 126 that remains is closed. Such a configuration may be used with a cutting wire 30' that has a cutting loop 37' at its distal end 36', and may at least partially hold the distal end 36' in place as the cutting wire 30' rotates. The edges 123 that define the opening 127 may have any configuration, including, but not limited to, a substantially square configuration, or any of the configurations shown in and described in reference to FIGs. 2B through 2E.

The embodiment of tissue resector 110' shown in FIG. 2G includes a cannula 120' with an opening 127' similar in configuration to the opening 127 of the embodiment of cannula 120 depicted by FIG. 2E; however, the entire distal end 126' of the cannula 120' is open. The edges 123' that define the opening 127' may have any configuration, including, but not limited to, a substantially square configuration, or any of the configurations shown in and described in reference to FIGs. 2B through 2E.
In the embodiment of tissue resector 210 depicted by FIG. 2H, the cannula 220 includes an open distal end 226, but no opening that extends into any portion of the circumference (i.e., the wall 222) of the cannula 220. The distal end 36' of a cutting wire 30', which may be configured (e.g., with a loop, etc.) to cut or otherwise separate tissues that are drawn into or otherwise come into contact with the distal end 226 of the cannula 220, may protrude distally beyond the distal end 226 of the cannula 220.

In some embodiments, such as those depicted by FIGs. 3A and 3B, the cannula 320, 320' of a tissue resector 310, 310' may include at least one additional opening 328, 328'. Such an opening 328, 328' may tailor the manner in which debris is aspirated through the lumen 324, 324' of the cannula 320, 320' (e.g., by balancing suction, etc.), or it may provide another location (i.e., in addition to the opening 327, 327' located closer to the distal end 326, 326' of the cannula 320, 320') at which an undesired growth, such as a uterine polyp or fibroid, may be engaged, severed and aspirated by the tissue resector 310, 310'. In the embodiment depicted by FIG. 3A, the opening 328 is longitudinally aligned with and located proximal to the opening 327, but is smaller than the opening 327. FIG. 3B shows an embodiment in which the openings 327 and 328 are located on different sides, or at different locations around the circumference, of the cannula 320.

While FIGs. 2A through 2F respectively show embodiments of cannulas with openings that are at least partially defined by edges that are linear or substantially linear, and, with the exception of the embodiment depicted by FIG. 2D, are oriented substantially parallel to the lengths of their respective cannulas, the edges that define the openings through the illustrated cannulas or any other embodiments of cannulas may be curved, may include curved sections and/or may be oriented at oblique angles to their respective cannulas.

With returned reference to FIGs. 1A and IB, the cutting wire 30 of a tissue resector 10 may comprise a guide wire of a known type and configuration. Without limitation, the cutting wire 30 may comprise or consist of a solid filament. Alternatively, the cutting wire 30 may include a coiled filament, which may surround a solid filament. As another alternative, the cutting wire 30 may include features that facilitate engagement and/or cutting of tissue, such as grooves or teeth that engage tissue or teeth or a sharpened edge that cuts into tissue. In still another alternative, the cutting wire 30 may include proximal features
(e.g., helical grooves, teeth, a helical thread, etc.) that facilitate the proximal movement of tissues through the lumen 24 of the cannula 20 of a tissue resector 10, for example, by breaking down tissues and other materials as they move proximally through the lumen 24, by forcing larger pieces proximally through the lumen 24 or by any other suitable mechanism.

FIGs. 4A through 4F depict a few non-limiting embodiments of cutting wires 30 that may be used in a tissue resector 10 (FIGs. 1A and IB) according to this disclosure.

In FIG. 4A, an embodiment of a cutting wire 30' is illustrated that is substantially linear along a majority of its length, with a cutting loop 37' at its distal end 36'. The cutting loop 37' of the cutting wire 30' may be elongated. The length of the cutting loop 37' may be about the same as or exceed the length of an opening 27 (FIGs. 1A and IB) of a cannula 20 (FIGs. 1A and IB) with which the cutting wire 30' is used. The width of the cutting loop 37' may be the same as, substantially the same as or less than the inner diameter of the lumen 24 (FIGs. 1A and IB) of a cannula 20 (FIGs. 1A and IB) with which the cutting wire 30' is configured for use.

In use, both sides of an open cutting loop, such as the embodiment of cutting loop 37' depicted by FIG. 4A, may cut into and through tissue. Thus, a cutting wire 30 with an open cutting loop (e.g., cutting loop 37', etc.) may cut tissue twice with each rotation.

FIG. 4B illustrates an embodiment of cutting wire 30" with a twisted cutting loop 37". As shown, a twisted cutting loop 37" may have a single twist, imparting it with a configuration that resembles a three-dimensional figure eight. Of course, a cutting loop 37" may have fewer than one twist or it may include more than one twist. The diameter of a twisted loop 37" may be the same as, substantially the same as or less than the inner diameter of the lumen 24 (FIGs. 1A and IB) of a cannula 20 (FIGs. 1A and IB) with which the cutting wire 30" is configured for use. A twisted cutting loop 37" may comprise an open loop, or it may comprise a solid, flat element having a helical configuration.

As another option, a cutting wire 30‴ may have a cutting feature 37‴ formed in its distal end 36‴, as illustrated by FIG. 4C. The cutting feature 37‴ may comprise one or more sharp points, edges, indentations or other features that may enable it to engage and cut tissue.

Another embodiment of cutting wire 130 is shown in FIG. 4D. That cutting wire 130 includes a cutting element 137 located near its distal end 136. The cutting element 137 comprises an elongated region of the cutting wire 130 that is offset from a remainder of the
length of the cutting wire 130. The distance of the offset may be the same as, substantially
the same as, or less than the radius of a lumen 24 (FIGs. 1A and IB) of a cannula 20 (FIGs.
1A and IB) within which the cutting wire 130 is configured to be positioned. The distal
end 136 of such a cutting wire 130 may be configured to facilitate smooth rotation of the
cutting element 137 relative to an opening 27 (FIGs. 1A and IB) through a wall 22 of the
cannula 20 with which the cutting wire 130 is configured to be used. In the depicted
embodiment, the distal end 136 of the cutting wire 130 is configured complementary to the
distal end 26 (FIGs. 1A and IB) of the cannula 20.

FIG. 4E illustrates an embodiment of cutting wire 230 with a centering feature 239
along at least a portion of its length, at a location proximal to a cutting element (not shown)
of the cutting wire 230. The illustrated embodiment of centering feature 239, or any
equivalently configured centering feature 239, may align an axis of rotation (not shown) of
the cutting feature (not shown) with a longitudinal axis through the center of the lumen 24
(FIGs. 1A and IB) of a cannula 20 (FIGs. 1A and IB) with which the cutting wire 230 is to be used. Such a configuration may optimize the stability with which the cutting wire 230 rotates, providing for smooth rotation of the cutting wire 230 within the lumen 24 of the cannula 20.

Another embodiment of cutting wire 330 is shown in FIG. 4F. That cutting wire 330 includes augers 339 along at least a portion of its length. The augers 339 may be configured
and oriented to facilitate the flow of excised tissue proximally through the lumen 24 (FIGs.
1A and IB) of a cannula 20 (FIGs. 1A and IB) with which the cutting wire 330 is used. In some embodiments, at least some of the augers 339 may be configured and/or oriented to break the excised tissue into smaller pieces, further facilitating the rate at which they may flow proximally through the lumen 24 of the cannula 20. In some embodiments may comprise discrete elements. In other embodiments, the augers 339 may have helical configurations, which may comprise one or more Archimedes screws positioned along the length of the cutting wire 330.

FIG. 4G illustrates an embodiment of cutting wire 430 that includes a solid, elongated
element 431 with a flattened distal end 432 and a cutter 435 secured to the flattened distal
end 431. The cutter 435 may comprise a cylindrical element with an open proximal end 436,
into which the flattened distal end 431 of the elongated element extends. A distal end 437 of
the cutter 435 may also be open (e.g., to enable samples and/or debris to be aspirated therein, etc.). The cutter 435 may also include one or more features that may cooperate with an opening 27 through a cannula 20 (see, e.g., FIG. IB). In the illustrated embodiment, a slot 438 extends along a portion of a length of the cutter 435, with opposed edges 439 of the slot 438 being configured to cut into body tissue or other materials as the cutting wire 430 is rotated within a lumen 21 of a cannula 20. In some embodiments, the edges 439 may comprise blades, teeth or serrations or other features that enable them to cut readily into and through body tissue or other materials.

Returning reference again to FIGs. 1A and IB, the cutting wire 30 of a tissue resector 10 may be of any suitable size (e.g., have an outer diameter) that will cut tissue in the desired manner while enabling the tissue to be aspirated through the lumen 24 of the cannula 20 at an acceptable rate (e.g., at a rate that will minimize the duration of a tissue resection procedure and, thus, the pain suffered by a patient, etc.). As an example, acceptable rates of aspiration may be achieved with a tissue resector 10 that has a lumen 24 with an open cross-sectional area (i.e., the cross-sectional area of the lumen 24 minus the cross-sectional area of the cutting wire 30) that is at least 50% of the cross-sectional area of the cannula 20. In a specific embodiment, a cannula 20 with an outer diameter of 0.115 inch (2.9 mm) (i.e., a cross-sectional area of 0.0104 in² or 6.7 mm²) and an inner diameter of 0.095 inch (2.4 mm) (i.e., a cross-sectional area of 0.00709 in² or 4.6 mm²), when used with a 0.040 inch (1 mm) (i.e., 0.00126 in² or 0.8 mm² in cross-sectional area) cutting wire 30, will have an open cross-sectional area of 0.00583 in² or 3.8 mm², which accounts for 56.1% of the entire cross-sectional area occupied by the cannula 20. Of course, the use of smaller cutting wires 30 would provide an even larger percentage of open area (e.g., at least 60%, at least 65%, at least 70%, etc.) and enable even greater rates of aspiration.

As illustrated by FIGs. 5A through 5C, a tissue resector 10 according to this disclosure may be operated under manual power (i.e., by hand) with a rooter 50, such as that described by U.S. Patent Application Publication No. US 2012/0239008 of Fojtik. Together, the tissue resector 10 and the rooter 50 may provide a tissue resector system 1. Such a rooter 50 may easily be used to rotate a cutting wire 30 at about 500 revolutions per minute (rpm), and can generate up to about 3,000 rpm, which may result in about 1,000 cuts per minute and 6,000 cuts per minute, respectively.
The rooter 50 of a tissue resector system 1 may enable rotation of a cutting wire 30 of the tissue resector 10 as the cannula 20 of the tissue resector 10 is held substantially stationary. As the rooter 50 is manually operated, it may spin the cutting wire 30 in a repetitious back-and-forth (i.e., clockwise and counterclockwise) manner, which may provide for two sets of cutting wire 30 rotations with each pull (or push) on an actuator of the rooter 50.

When an embodiment of cutting wire 30' (FIG. 4A) having an open cutting loop 37' (FIG. 4A) is used in the tissue resector system 1, since the cutting loop 37' cuts tissue twice with each rotation of the cutting wire 30', manual operation of the rooter 50 and, thus, hand-operated rotation of the cutting wire 30' may efficiently cut tissue and, thus remove the same from an individual's body.

Turning now to FIGs. 6A and 6B, with a proper adapter 60, a rooter 50 that is configured for hand-powered operation may also be used with the limited use (e.g., disposable, etc.) portion 70 of a conventional morcellator, providing another embodiment of tissue resector system 1'.

The depicted embodiment of adapter 60 includes a series of elements that translate the action generated by the rooter 50 to an action that rotates the inner cannula, or blade 72, of a conventional morcellator. More specifically, the adapter 60 may include elements 61 and 62 that are respectively configured to engage a rotatable element of the rooter 50 and a proximal end 73 of the blade 72. Element 62, which is configured to engage the proximal end 73 of the blade 72, may be configured to rotate the blade 72 about an eccentric axis (e.g., an axis that enables the blade 72 to move rotational over an inner circumference of a lumen 74 of an outer cannula 76, etc.).

In addition, the adapter 60 may include one or more stationary elements 63 and/or a distal end cap 66 that may be configured to hold the outer cannula 76 that surrounds the blade 72 stationary, even while the blade 72 is rotated within a lumen 74 of the outer cannula 76.

In addition, the adapter 60 may include seals 67, 68, which enable movement of the blade 72 within the lumen 74 of the outer cannula 76, while simultaneously enabling the aspiration of body tissues, fluids or the like through the lumen 74 of the outer cannula 76, through the housing 65 of the adapter 60 and through an aspiration port 64 that
communicates with an interior of the housing 65 of the adapter 60. Thus, the seals 67, 68 enable a suction (i.e., a vacuum) to be applied to the aspiration port 64 and communicated through the housing 65 of the adapter to the lumen 74 of the outer cannula 76 to drawn tissue, fluid or other materials proximally therethrough.

As depicted by FIG. 7, in another embodiment of tissue resector system 1", a tissue resector 10 may also be used with a power-drive instrument 90 of a conventional morcellator when coupled to the power-drive instrument 90 with an appropriately configured adapter 80. The adapter 80 may be configured to translate rotating action by the power-drive instrument 90 to the cutting wire 30 (FIGs. 1A and IB) of the tissue resector 10 and convert a system that aspirates through the power-drive instrument 90 for communication with the lumen 24 (FIGs. 1A and IB) that extends through the outer cannula 20 (FIGs. 1A and IB) of the tissue resector 10.

Although the foregoing disclosure provides many specifics, these should not be construed as limiting the scope of any of the ensuing claims. Other embodiments may be devised which do not depart from the scopes of the claims. Features from different embodiments may be employed in combination. The scope of each claim is, therefore, indicated and limited only by its plain language and the full scope of available legal equivalents to its elements.
What is claimed:

1. A tissue resector, comprising:
   a cannula including:
   a wall;
   a lumen defined through the wall and extending along a length of the cannula;
   a distal end; and
   an opening at or adjacent to the distal end and establishing communication between
   an exterior of the wall and the lumen; and
   a cutting wire within the lumen, the cutting wire including:
   a cutting feature that cooperates with the opening through the cannula.

2. The tissue resector of claim 1, wherein an open cross-sectional area including
   a cross-sectional area of the lumen less a cross-sectional area of the cutting wire comprises at
   least 50% of the cross-sectional area of the cannula.

3. The tissue resector of claim 2, wherein an outer diameter of the cannula is
   about 2.9 mm and a diameter of the lumen is about 2.4 mm.

4. The tissue resector of claim 3, wherein the cutting wire has an outer diameter
   of about 1 mm or less.

5. The tissue resector of any of claims 1-4, wherein the opening of the cannula is
   defined through a circumference of the cannula.

6. The tissue resector of claim 5, wherein the opening of the cannula is spaced
   proximally from the distal end of the cannula.

7. The tissue resector of claim 5, wherein the opening of the cannula extends to
   the distal end of the cannula.
8. The tissue resector of any of claims 1-4, wherein the opening of the cannula is defined through the distal end of the cannula.

9. The tissue resector of any of claims 1-8, wherein the opening of the cannula includes at least one edge configured to cut into tissue.

10. The tissue resector of any of claims 1-9, wherein the cutting feature of the cutting wire includes at least one edge configured to cut into tissue.

11. The tissue resector of any of claims 1-10, wherein the cutting wire is configured to rotate eccentrically within the lumen of the cannula.

12. The tissue resector of any of claims 1-11, wherein the cannula further includes:

at least one secondary opening through the wall of the cannula and located proximal to the opening.

13. A tissue resector system, comprising:
a tissue resector according to any of claims 1-12; and

a hand-powered rooter for rotating the cutting wire of the tissue resector.

14. A method for removing tissue from a body of a subject, comprising:
drawing tissue into an opening of a tissue resector of any of claims 1-12;
rotating a cutting wire of the tissue resector to cut the tissue; and

aspirating cut tissue.

15. The method of claim 14, further comprising:
repeating the acts of drawing, rotating and aspirating.
**INTERNATIONAL SEARCH REPORT**

**Application No.**
PCT/US 15/28084

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC (8) - A61B 17/3205 (2015.01)

CPC - A61B 2017/320024

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 (8): A61B 17/3205 (2015.01)

CPC: A61B 2017/320024

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)
PatBase, Google Patent, Google Scholar

Search terms used: Tissue resector cannula resection cutting wire morcellator device morcellation side window resect cut

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

<table>
<thead>
<tr>
<th>Category</th>
<th>Citation of document, with indication, where appropriate, of the relevant passages</th>
<th>Relevant to claim No.</th>
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<tr>
<td>X</td>
<td>US 2008/0103504 A1 (Schmitz et al.) 01 May 2008 (01.05.2008), Entire document, especially figs 3B-4, para [0074])</td>
<td>1-6, 8</td>
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<td>Y</td>
<td>US 8,574,253 B2 (Gruber et al.) 05 November 2013 (05.1.2013), abstract, fig 34</td>
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<td>US 6,066,153 A (Lev) 23 May 2000 (23.05.2000), Entire document</td>
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Further documents are listed in the continuation of Box C.

* 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

**Date of the actual completion of the international search**

08 July 2015 (08.07.2015)

**Date of mailing of the international search report**

27 JUL 2015

**Name and mailing address of the ISA/US**

Mail Stop PCT, Attn: ISA/US, Commissioner for Patents
P.O. Box 1450, Alexandria, Virginia 22313-1450
Facsimile No. 571-273-8300

**Authorized officer**

Lee W. Young

PCT Handling: 571-272-4300
PCT OSP: 571-272-7774

Form PCT/ISA/2 10 (second sheet) (January 2015)
INTERNATIONAL SEARCH REPORT

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.: 9-15
   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:

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 additional fees, this Authority did not invite payment of 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.

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