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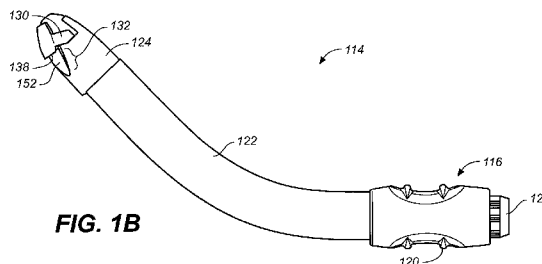
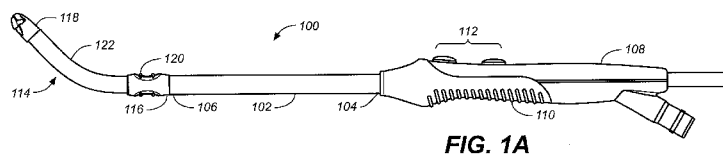
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(54) Title: ELECTROSURGICAL DEVICES FOR TONSILLECTOMY AND ADENOIDECTOMY



(57) Abstract: Monopolar electro surgery devices adapted for resecting tonsil and adenoid tissue. The devices minimize thermal injury by employing a plasma generated by pulsed electrical signals to precisely and effectively cut or coagulate the tissues. Suction may also be applied to the tissues to enhance the cutting, coagulation, and tissue manipulation functions. The devices include an interchangeable tip that may be switched for another tip, depending on which tip may be more suitable for tonsillectomy or adenoidectomy.

ELECTROSURGICAL DEVICES FOR TONSILLECTOMY AND ADENOIDECTOMY

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority to U.S. Patent Application No. 12/348,722 filed January 5, 2009, which is hereby incorporated by reference in its entirety.

FIELD

[0002] The field is electrosurgery, and in particular, tonsillectomy and adenoidectomy.

BACKGROUND

[0003] The tonsils and adenoids are generally located in the back of the human (mammalian) nose and throat, and are part of the lymphatic system that samples bacteria and viruses entering the body. Once sampled, the immune system is activated to produce antibodies that fight infection. When bacteria and viruses become trapped in the tonsils and adenoids, these tissues are able to break down their cell wall and deliver the fragments to areas of the body that produce antibodies modeled against the fragments. However, repeated inflammation of the tonsils and adenoids impedes their ability to destroy the bacteria that become entrapped therein, resulting in bacterial colonization of these tissues. The colonies of bacteria can then serve as a reservoir for repeated infections (e.g., tonsillitis or ear infections). Tonsillectomy and/or adenoidectomy may be indicated when antibiotic treatment fails to remove the bacterial reservoir. Tonsil tissue may also need to be removed if it enlarges to the point of causing airway obstruction, which may manifest as snoring or sleep apnea. Some individuals are also born with larger tonsils that are more prone to cause obstruction. Adenoidectomy may also be required to remove adenoid tissue when ear pain persists, or when nose breathing or function of the Eustachian tubes is impaired.

[0004] The devices and techniques used for tonsillectomy and adenoidectomy will usually depend on such factors as the type and amount of tissue to be removed and surgeon preference. The two procedures are routinely performed together. A common method for

tonsillectomy and adenoidectomy employs cold surgical dissection. Here tissue is removed using a scalpel or other sharp instrument such as a curette or punch device. Sharp dissection oftentimes results in heavy bleeding, which can be stemmed with electrocautery.

[0005] In addition to coagulation, electrosurgery devices (e.g., suction-tipped, blade, or needle tip Bovie) may also be employed to resect tonsil or adenoid tissue. The suction tipped Bovie typically has a hollow center to suction blood, secretions, and smoke from the surgical field, and a rim of metal for cutting and coagulation. A separate aspirator is used when blade and needle tip Bovie are used. Although the use of Bovie reduces blood loss intraoperatively in comparison to cold techniques, it is associated with increased postoperative pain due to spread of the thermal injury from the heat of electrosurgery (average temperatures are above 300°C). Despite their increased thermal injury profile, use of the Bovie remains the most popular method of tonsil removal in the U.S. due to its speed, convenience, universal availability, and surgeon familiarity with the device.

[0006] Other energy-type devices have been commercialized that attempt to minimize thermal injury. These include the Harmonic Scalpel® system (Ethicon Endo-Surgery, Cincinnati, OH) (ultrasonic energy), lasers (e.g., KTP, Nd:YAG, or CO₂ lasers), and Coblation® devices (Arthrocare, Austin, TX) (bipolar radiofrequency ablation). However, the decrease in thermal injury provided by these devices is questionable. Even if they do result in less thermal injury, it is offset by reduced control of bleeding and surrounding tissue trauma, longer operative times, or less precise cutting. Some of the instruments also obscure the surgical field and are difficult to maneuver due to their large size.

[0007] When Coblation® devices are employed, the procedure requires saline delivery in order to establish an electrosurgical effect. Pre-operative time is prolonged due to the inclusion of the saline delivery set-up. Coblation® device aspiration lumens also periodically clog during the procedure, which causes saline to pool in the patient's throat. The pooled saline must then

be suctioned using a separate aspirator. Additionally, the Coblation® device lumen typically has to be manually cleared in order to finish the surgery.

[0008] Accordingly, new devices for resecting tonsil and adenoid tissue would be useful. In particular, devices that precisely cut tonsil and adenoid tissue while effectively controlling bleeding and surrounding tissue trauma would be desirable. Devices that provide easier access to the tonsils and adenoids and manipulation of those tissues would also be desirable.

SUMMARY

[0009] Described here (but not limiting) are electrically “monopolar” electrosurgical devices that optimize many functions required in performing tonsillectomy or adenoidectomy while minimizing side-effects (the return electrode is applied to the patient remote from the active electrode). For example, the precise cutting of tissue is provided by a thin layer of plasma that surrounds the electrode edge. In some instances the plasma is formed with pulsed radiofrequency (RF) energizing waveforms having a lower range of duty cycles so that tissue has time to cool between pulses. This coupled with the mostly insulated tips helps to reduce thermal damage by limiting the extent of thermal diffusion from the electrode into surrounding tissue. Continuous or 100% duty cycle energizing waveforms may also be employed when faster cutting is desired. When hemostasis is needed, suction and plasma may be directly applied to the target area to mechanically and electrically stop the bleeding. Tension of the electrode surface(s) against the tissue is generally optimized with a configuration where suction is applied through the electrode or in very close proximity thereto. The aspiration port being closely associated with the blade allows a “dry field” surgical technique. This is advantageous here because it allows a concave blade to function well unlike the case with a wet field. This is because current flows from the blade to the tissue only via the part of the blade in actual tissue contact with the dry field. Further, the suction decreases the width of the tissue in contact with the blade, further reducing the contact area and also further drying the tissue. Given that it may be difficult to access the portions of the oropharynx and nasopharynx where the tonsils and

adenoids are located, the devices described herein may be designed with one or more malleable portions so that they can be shaped to improve access to these regions and/or accommodate variations in patient anatomy.

[0010] It has been found that for ENT (ear-nose-throat) surgery as described here, a monopolar approach is advantageous because the remote return electrode guarantees good electrical contact. In contrast, given the nature of such energizing, a bipolar device with both electrodes on one shaft makes good electrical contact for the return electrode difficult.

[0011] The electrosurgical devices generally include an elongate body having a proximal end, a distal end, and an aspiration lumen extending therethrough, an interchangeable tip removably attached to the distal end of the elongate body, and a handle at the proximal end of the elongate body where control mechanisms for cutting and coagulation are located. The interchangeable tip typically also has a proximal end, a distal end, and a tip lumen that is fluidly connected to the aspiration lumen of the elongate body. A housing that is secured to the distal end of the interchangeable tip will usually have an aperture and an integrated blade assembly that includes an active electrode. The blade assembly or portion thereof may form an aspiration port with the aperture of the housing.

[0012] The devices generally include a connector at the proximal end of the interchangeable tips for removably attaching them to the elongate body. In some variations, the connector comprises a compressible barrel and one or more tabs that are located circumferentially around the connector. The compressible barrel may be inserted into the distal end of the elongate body to create a friction fit between the components and to thereby removably attach them to one another. The one or more tabs may be used to attach an ergonomic finger grip to the connector that may make insertion and removal of the interchangeable tip easier.

[0013] The interchangeable tips may be specifically configured for adenoidectomy or tonsillectomy. For example, when adenoidectomy is to be performed, the devices may include a

blade assembly that optimizes tissue removal using a raking motion. Here the blade assembly may include an active electrode that has a cutting edge and a flat coagulation surface. A plurality of arms extending from the active electrode may secure the active electrode to the housing. The arms may have an angle that positions the cutting edge against the tissue in a manner that allows it to be easily and precisely scooped or shaved into the aspiration port. The particular arm angle may also be useful in achieving hemostasis by optimizing the position of the coagulation surface against the tissue.

[0014] When tonsillectomy is to be performed, the devices may include a blade assembly having a curved and tapered active electrode. This curvature may result in a more precise tonsil resection because the electrode surface approximately conforms to the contour of tonsil tissue. The curved and tapered electrode may also define an aspiration port therethrough. Here bleeding may be more effectively controlled since suction may be applied directly through the aspiration port of the electrode when the device is in the coagulation mode. This would combine mechanical pressure with plasma coagulation to achieve hemostasis.

[0015] The interchangeable tips may also be configured to be malleable so that their shape can be tailored to navigate access to the tissues or accommodate variations in patient anatomy. In some instances the interchangeable tips may be made for single use. Kits including one or more interchangeable tips are also contemplated.

[0016] Methods for resecting tonsil or adenoid tissue may generally include applying a pulsed electrical signal to the active electrode of the devices described herein to form a plasma on the active electrode, and cutting or coagulating the tissue with the plasma. In some variations, suction is applied simultaneously with proximal movement of the electrode to suction cut the tissue. In other variations, the tissue is coagulated while suction is simultaneously applied. In yet further variations, the methods will include exchanging one interchangeable tip, e.g., a tonsillectomy tip, with another interchangeable tip, e.g., an adenoidectomy tip.

BRIEF DESCRIPTION OF THE DRAWINGS

[0017] FIG. 1A depicts a side view of an exemplary electrosurgical device with an adenoidectomy tip.

[0018] FIG. 1B depicts an expanded side view of the adenoid tip shown in FIG. 1A.

[0019] FIG. 1C is an expanded perspective view of the blade assembly shown in FIGS. 1A and 1B.

[0020] FIG. 1D shows the components of the housing in FIGS. 1A and 1B and how they are joined to integrate the blade assembly of FIG. 1C.

[0021] FIG. 1E is an expanded side view of the blade assembly shown in FIGS. 1A-1D.

[0022] FIG. 1F shows an expanded inferior view of an exemplary housing and blade assembly for adenoidectomy.

[0023] FIG. 1G depicts a variation of the cutting edge that may be used in the blade assembly for adenoidectomy.

[0024] FIG. 2A depicts an exemplary housing and blade assembly for tonsillectomy.

[0025] FIG. 2B shows the components of the housing in FIG. 2A and how they are joined to integrate the blade assembly also shown in FIG. 2A.

DETAILED DESCRIPTION

[0026] Described here are electrosurgical devices that can be customized for either adenoidectomy or tonsillectomy. Customization may be provided by the use of interchangeable tips that can be removably attached to the elongate body of the device. Specifically, the design of the blade assemblies and location of the aspiration port from which suction is applied may be varied to optimize manipulation, cutting, and hemostasis of the tissue being operated on.

[0027] The present electrosurgical devices are generally monopolar instead of bipolar. Both types of devices use high frequency alternating current and a pair of electrodes, one designated the active electrode and the other the return electrode. However, the difference lies in the placement of these electrodes. In a monopolar device, current is passed from the active

electrode through the patient's body to a grounding pad (return electrode) placed on the body, usually the thigh or shoulder. A monopolar device may be particularly useful in tonsil and adenoid surgery because contact with the body is guaranteed. In a bipolar device, the active and return electrodes are both placed at the site of electrosurgery, for example, on the same shaft of the device, and electrosurgery occurs only on the tissue on the active electrode(s) if there is a conductive path back to the return electrode.

[0028] The devices will generally include an elongate body having a proximal end, a distal end, and an aspiration lumen extending therethrough. Depending on the materials used, all or a portion of the elongate body may be formed to be stiff, flexible, or malleable. Exemplary materials for making the elongate body include, without limitation, fluoropolymers; thermoplastics such as polyetheretherketone, polyethylene, polyethylene terephthalate, polyurethane, nylon, and the like; and silicone.

[0029] The elongate body may be between about 5.0 cm to about 20 cm in length. For example, the elongate body may be between about 5.0 cm to about 15 cm, between about 5.0 cm to about 10 cm, or between about 5.0 cm to about 8.0 cm in length. In some instances, the elongate body may be about 8.0 cm in length. In other instances the devices lack an elongate body, and the interchangeable tip is directly connected to the handle.

[0030] In some variations, the elongate body is coated with a lubricious polymer to reduce friction between the mucosa and device during the procedure. Lubricious polymers are well known in the art, and are typically hydrophilic. Exemplary hydrophilic polymers that may be used as lubricious coatings include, without limitation, polyvinyl alcohol, polyethylene glycol, polyvinyl pyrrolidone, cellulosic polymers and polyethylene oxide.

[0031] A handle is usually coupled to the proximal end of the elongate body that allows the surgeon to hold the device. The handle may include a hand grip on the bottom and a pad with at least two control buttons on the top. The control buttons may be depressed to activate electric switches for turning on the cutting or coagulation regimes. The button pad and hand

grip are incorporated into the handle using spring clips or other well-known attachments and conventional techniques such as overmolding. A standard electrical connector and cable may be used with the handle to connect electrical leads to a generator. The handle may be between about 5.0 cm to about 30 cm, about 5.0 cm to about 25 cm, about 5.0 to about 20 cm, or about 5.0 cm to about 15 cm in length. In one variation, the handle is about 20 cm in length.

[0032] The aspiration lumen of the elongate body continues through the handle to a suction connector, which in turn connects tubing to a collection canister. The dimensions of the handle may be varied such that it is comfortably held in a hand, yet able to be manipulated during tonsillectomy and adenoidectomy. In some variations, the handle may be made for single use. As used herein, the term “suction handle” shall refer to the handle when coupled to the elongate body.

[0033] An interchangeable tip, as further described below, may be removably attached to the distal end of the elongate body. The interchangeable tip will typically have a proximal end, a distal end, and a tip lumen that is fluidly connected to the aspiration lumen to allow continuous flow of fluids, tissue, smoke, etc. away from the surgical field. The length of the interchangeable tip varies, but may be from about 1.0 cm to about 15 cm. For example, the adenoidectomy tip may be between about 5.0 cm to about 10 cm, and the tonsillectomy tip may be from about 1.5 cm to about 5.0 cm. In one variation, the adenoidectomy tip is about 9.0 cm. In another variation, the tonsillectomy tip is about 2.5 cm. The materials used to make the elongate body may also be employed in forming the interchangeable tip.

[0034] In one variation, the interchangeable tip is malleable. Malleability may be provided by wires running through the wall of the elongate body that are capable of being shaped by bending the elongate body. For example, the wires may be made from stainless steel or alloys thereof, nickel-titanium alloy, etc. Malleability may also be provided by forming the elongate body or portions thereof with metals or polymers that can be shaped. In some variations, joints or hinges are included. By having the ability to manipulate the shape of the

interchangeable tip, it may be easier and quicker to access the tonsils and adenoids with the devices and easier to accommodate anatomical variations in that area. The interchangeable tips may be bent directly, e.g., by finger manipulation, or remotely through cables or wires using mechanisms commonly employed with steerable catheters. The interchangeable tips may be designed for single use.

[0035] The devices generally include a connector at the proximal end of the interchangeable tips for removably attaching them to the elongate body. An ergonomic finger grip may be used on the connector that may ease insertion and removal of the interchangeable tip. In some variations, the interchangeable tip may be configured to rotate about the connector. The connectors may be configured to removably attach to the elongate body by friction fit mechanisms. For example, they may include a compressible component that has a larger diameter when not connected to the elongate body. Compression of the component typically allows it to be slidably engaged with the elongate body at which point the component is capable of expanding to its uncompressed diameter against the internal wall of the elongate body. In one variation, the connector comprises a compressible barrel and one or more tabs that are located circumferentially around the connector. The compressible barrel may be inserted into the distal end of the elongate body to create a friction fit between the components and to thereby removably attach them to one another. The one or more tabs may be used to attach an ergonomic finger grip to the connector that may make insertion and removal of the interchangeable tip easier.

[0036] Connection between the elongate body and interchangeable tip may also be supplied by locking-unlocking mechanisms. For example, the interchangeable tip may include a groove or channel, e.g., a L-shaped channel, in its wall configured so that upon receipt of a pin or other protrusion on the elongate body, the interchangeable tip may be rotated to be locked in place. Other tongue-and-groove type locks and mechanisms including depressible pins or tabs are also contemplated.

[0037] The electrosurgical devices described here also generally include a housing at the distal end of the interchangeable tip for holding and securing the blade assembly thereto. The housing may be a single unit component or be made from multiple components. For example, the housing may be made by pressing together two components, an upper portion and lower portion. In some variations, the housing is configured with a blunt nose. In other variations, the housing is designed with a taper.

[0038] The housing may include an aperture through which suction can be applied to the tissues. The aperture may be of any shape and size so long as it provides an opening for aspiration of smoke, tissue, fluids, etc. For example, the aperture may be spherical, elliptical, egg-shaped, rectangular, triangular, diamond-shaped, or heart-shaped. These shapes are not meant to be limiting. When tissue is to be aspirated, the aperture may be larger than conventional suction openings. For example, the aperture may have a diameter from about 0.1 cm to about 1.0 cm. In some variations, the aperture diameter is larger than about 0.4 cm or about 0.5 cm.

[0039] In addition, the housing may be made from transparent materials or include a marker to aid visualization or provide an electrode locator in the surgical field. Materials with suitable transparency are typically polymers such as acrylic copolymers, acrylonitrile butadiene styrene (ABS), polycarbonate, polystyrene, polyvinyl chloride (PVC), polyethylene terephthalate glycol (PETG), and styrene acrylonitrile (SAN). Acrylic copolymers that may be particular useful include, but are not limited to, polymethyl methacrylate (PMMA) copolymer and styrene methyl methacrylate (SMMA) copolymer (e.g., Zylar 631® acrylic copolymer).

[0040] A blade assembly may be integrated within the housing, and generally includes the active electrode. The active electrode typically defines a cutting edge. The edge, however, is not like that of an ordinary knife which does the cutting solely via mechanical application to the tissue being cut. Instead here the edge of the electrode focuses the electric field induced by the applied electrical signal. This field that is concentrated at the edge generates a local plasma

discharge. In some instances, the edge is formed by the onlay of metal foil. The tissue cutting may be assisted by mechanical force supplied by the edge of the electrode, as in conventional cutting, or by application of suction.

[0041] The active electrode may be made from any material having suitable electrical properties. For example, the active electrode may include without limitation, metals such as molybdenum, nickel, platinum, stainless steel, tantalum, titanium, tungsten, and alloys thereof. Thickness of the active electrode may be from about 0.1 mm to about 1.0 mm thick with an edge of about 10 μ m to about 70 μ m in thickness. In one variation, the active electrode thickness is about 0.25 mm (0.01 inch). The active electrodes may be etched or stamped or machined out of a large sheet of metal, which has been annealed for hardness.

[0042] An electric insulating layer may be overlaid upon the active electrode to mostly cover it but leaving an exposed edge or surface that would be used for cutting or coagulation. The insulation is typically a coating of glass or ceramic approximately 0.005 mm to 0.5 mm thick, or in some instances approximately 0.01 mm to 0.2 mm thick. When glass insulation is used, it may be applied by a conventional process of dipping each relevant component prior to assembly in liquid (molten) glass and then annealing the glass.

[0043] The configuration of the blade assembly may vary depending on the intended use of the electrode. For example, technical features that enhance removal of adenoid tissue may differ from other features of interest in tonsillectomy. These design features are further elucidated below.

Adenoidectomy Tip

[0044] Adenoid tissue, which lies at the back of the nose/throat, may be removed through the mouth or nose, but typically through the mouth under general anesthesia. Thus, not only are the adenoids difficult to get to, but the presence of an endotracheal tube further cramps an already limited surgical field. The adenoid tips described here are interchangeable tips specifically configured to address many of these difficulties with adenoidectomy.

[0045] In one variation, as illustrated in FIG. 1A, electrosurgical device (100) includes an elongate body (102) having a proximal end (104) and a distal end (106). Proximal end (104) is coupled to a handle (108) where a finger grip (110) and pair of control buttons (112) are located. Adenoid tip (114) also has a proximal end (116) and a distal end (118), and is removably attached to the elongate body distal end (106) by a connector (120) positioned at the elongate body distal end (106). Here connector (120) includes a compressible basket (121) that can be depressed and slid into the distal end of the elongate body where it will then decompress to expand and create a friction fit against the internal wall of the elongate body (FIG. 1B). The elongate body (102) may be permanently or removably attached to the handle (108), but will typically be permanently attached. Adenoid tip (114) is malleable and can be bent, as shown in FIG. 1A. Here the adenoid tube (122) comprises a flexible material such as polyurethane, and includes two lumens (not shown) within its wall where bendable wires (not shown) run longitudinally between the adenoid tip distal end (118) and proximal end (116). Additional lumens and wires may be employed as desired.

[0046] Although shown as bent upwards, the adenoid tip may be bent in other directions, so long as the device materials or functionality is not compromised. For example, the adenoid tip may be bent to form an angle with the elongate body of up to about 45°, up to about 60°, or up to about 90° or more with respect to the horizontal axis of the elongate body. Adenoid tip (114) may also be rotated to any angle about the axis of the connector (120). As previously mentioned, the bendable nature of the adenoid tip allows easier access to the adenoids. It also provides more precise positioning of the active electrode.

[0047] The adenoid distal tip also includes a housing for holding the blade assembly. As shown in FIGS. 1B and 1D, housing (124) is comprised of an upper portion (126) and a lower portion (128) that integrate a blade assembly (130) when both portions are joined together. Upper (126) and lower (128) portions may be joined by snap fitting, adhesives, welding, etc. The lower portion (128) may have an aperture (132) that defines an aspiration port (152) when

the blade assembly (130) is integrated into the housing (124). In some variations, the housing may be formed using a single component instead of a plurality of components.

[0048] As shown in the enlarged housing-blade assembly of FIG. 1F, the housing (124) and integrated blade assembly (130) may define an aspiration port (152), as explained above, through which suction is applied to aspirate fluids and smoke. However, the aspiration port (152) will generally also be dimensioned so that the device is capable of aspirating tissue through the port. In such instances, the port in cross-section may be spherical, ellipsoid, egg-shaped, rectangular, triangular, diamond-shaped, or heart-shaped, and thus wider than conventional suction openings. Here the aspiration port is about 0.1 cm to about 1.5 cm along its major axis and about 0.1 cm to about 1.0 cm along its minor axis. It is understood that these dimensions are exemplary and not limiting.

[0049] Bringing the upper and lower housing portions together with the blade assembly therebetween integrates the blade assembly into the housing. Here the housing is made from a transparent polymer such as Zylar 631® acrylic copolymer (SMMA copolymer) so as not to block visualization at the adenoid tip distal end (118). Although not shown here, a marker can also be included on the surface of upper housing portion (126) to indicate the blade location underneath.

[0050] Turning to FIG. 1C, blade assembly (130) includes an active electrode (136) having a cutting edge (138) and a coagulation surface (140). A plurality of arms (142) having a first portion (144) and a second portion (146) secure the active electrode (134) to the housing (124). The arms may be secured to the housing with medical grade adhesives such as cyanoacrylate or epoxy adhesives, and other adhesives, which are well known. Referring to FIG. 1E, first (144) and second (146) arm portions form an angle (148) of about 110° to about 140°. In one variation, a 120° angle is defined between the first (144) and second (146) portions. This angle may optimize opposition of the coagulation surface (140) onto tissue to effectively achieve hemostasis. Further, the angle (148) may optimize the point of contact of plasma to the

tissue from the cutting edge (138) to effect a more precise cut as well as resection of tissue while minimizing clogging of the aspiration port.

[0051] In another variation, as shown in FIG. 1G, the cutting edge (138) may be shaped to include a leading edge (154). The leading edge may aid initiation of plasma cutting and more effectively remove adenoid tissue. In some variations, the leading edge may extend about 0.2 mm to about 2.0 mm from the cutting edge.

[0052] After the blade assembly is integrated into the housing, cutting edge (138) defines an aspiration port (152) with the aperture (132). With this adenoid tip configuration, cut adenoid tissue is directly and immediately suctioned into the aspiration port. Thus, proximal movement (movement toward the operator) of the adenoid tip while simultaneously maintaining suction may result in shaving strips of tissue.

Tonsillectomy Tip

[0053] The tonsils are located on either side of the throat and removed under general anesthesia. In this procedure, the unattached end of the tonsil is usually grasped with forceps while an incision is made at the other end to remove it from the throat. Thus, similar to adenoidectomy, access is also difficult and the surgical field cramped. The tonsillectomy tips described here are interchangeable tips specifically configured to address many of these difficulties with tonsillectomy.

[0054] The suction handle (handle with the elongate body attached) and connector are generally of the same configuration and made from the same materials as described for the adenoidectomy tips. The tonsillectomy tips may also be malleable and provide the same advantages as those previously described. Further, the housing, blade assembly, and active electrode are typically made with the same materials. However, their configuration will differ.

[0055] In one variation, as depicted in FIG. 2A, the tonsillectomy tip has a housing (200) that is tapered. Blade assembly (202) is integrated within the housing (200). An opening in the active electrode (204) defines an aspiration port (206) through which suction may be applied to a

tissue surface, usually the tonsil bed, as explained above. A rim of active electrode results that has a rim thickness (205) from about 0.5 mm to about 3.0 mm. The rim is insulated except for a about a 5 mm to about a 20 mm portion running along the outer edge (209). About a 2 mm to about a 15 mm exposed (uninsulated) portion may also run along the inner edge (207). Upon application of an energizing electrical signal (waveform), pulsed or continuous, plasma may be formed along the inner (207) and/or outer (209) edges of the rim to cut or coagulate tissue. As previously mentioned, cutting precision is typically not a result of the degree of sharpness of the electrode edge, and is not the case here. However, in some variations, one or more sides of the active electrode may be sharpened to form about a 10° to 40° edge.

[0056] Although shown as a triangular shaped opening in FIGS. 2A and 2B, the aspiration port (206) may take various other shapes. The triangular configuration of the aspiration port may optimize suction at the tip of the device without compromising the preciseness of tissue dissection.

[0057] The active electrode may have a length from about 0.1 cm to about 1.5 cm. In one variation, the active electrode has a length of about 0.4 cm. At its widest portion, the active electrode has a width from about 0.1 cm to about 1.0 cm. In some instances, the active electrode has a width of 0.4 cm. The active electrode may also be configured with an upward curve so that it approximately conforms to the curvature of the tonsil (e.g., the palatine tonsil). This upward angle of the active electrode may be of any degree. For example, the upward angle may be from about 5° to about 60°. In some variations, the upward angle is about 30°.

[0058] The relationship between the housing and blade assembly components of FIG. 2A is further detailed in FIG. 2B. In FIG. 2B, housing (200) is shown as being formed by an upper portion (208) and a lower portion (210). A pin (212) extending inferiorly from the internal surface of the upper portion (208) is threaded through a hole (216) of the blade assembly (214) to integrate the blade assembly (214) within the housing (200). An adhesive may further be used to secure the housing and blade assembly components to one another.

[0059] The housing in FIGS. 2A-2B is shown as tapered, but need not be. In this variation, upper portion (208) is longer than bottom portion (210). The upper portion (208) will generally be from about 0.3 cm to about 3.0 in length. The bottom portion (210) will generally also be from about 0.3 cm to about 3.0 cm in length. The difference in length between the upper and lower portions results in complete coverage of the aspiration port (206) on the top of the active electrode (204) but only partial coverage of the aspiration port (206) on the bottom of the active electrode (204). With this configuration, suction may be provided through the undersurface of the active electrode. The aspiration port (206) may be from about 0.03 cm to about 0.5 cm wide, and about 0.03 cm to about 1.0 cm in length. In one variation, the aspiration port is about 0.17 cm in width and about 0.5 cm in length. A connective piece (218) is also shown in FIG. 2B that allows the integrated housing and blade assembly to be attached to the tip lumen.

[0060] The tonsillectomy tips herein described may provide more precise cutting and coagulation because of the particular suction electrode design. Here not only does the aspiration port suction smoke and fluids from the operative field to improve visualization, but suction and coagulation is capable of being directly and immediately placed on the tonsil bed from which the tonsil tissue has been resected. This is the raw surface where hemostasis is needed most, not on the tonsil itself, which is removed from the body. The active electrode rim also provides a larger surface that could be swept across the tissue to more effectively control bleeding.

[0061] Furthermore, use of the tonsil devices described here may result in less thermal injury. As further detailed below in the single Example, the depth of thermal injury was found to be less when the instant tonsil devices were employed in comparison to Bovie needles (0.26 mm vs. 0.77 mm) on tonsil tissue. Thermal injury was decreased by 66% when cut settings were used and by 73% when coagulation settings were used.

[0062] The electrical signals (radio frequency in this case) for generating the plasma may be provided by generators and electrical circuits of the kind well known in the art, or by those

described in pending co-owned Application Nos. 11/982,734 and 12/126,683, which are incorporated by reference in their entirety. The electrical signals are conventionally applied to the electrode by a conductor(s) extending through the electrosurgical device and not shown here. As previously mentioned, the plasma is formed along the edge of the active electrode by application of the electrical signals to the electrode. To further decrease the heat accumulation and associated collateral tissue damage, low duty cycle waveforms may be used. As well known, duty-cycle refers to the proportion of time that the electrical energy is actually being applied. Low duty-cycle here typically refers to duty-cycles of less than 10% which may be, for instance, 1% or less, or 0.1% or less. In some cases the low duty-cycle refers to the pulse voltage regime that is applied to the active electrode. For instance, a pulsed low duty-cycle signal may include a plurality of pulse bursts that are separated by more than one millisecond (e.g., has a frequency of less than 1 KHz) where each burst is shorter than one millisecond. The burst of pulses may include pulses that are biphasic (e.g., of alternating polarity) and the pulses may have different peak voltages. Again, none of this is limiting. The low duty-cycle is intended to minimize the spread of thermal injury, including tissue charring or burning. However, in some instances the devices described here may employ continuous waveforms or duty cycles of greater than about 50% or greater than about 75%.

[0063] Cutting or coagulation is generally obtained by energizing the active electrode with a suitable electrical signal (typically of different frequency, duty-cycle, etc.) for each surgical function. For example, when cutting is desired, the applied signal is an RF (radio frequency) signal having a frequency in the range of 100 KHz to 10 MHz. This energy may be applied in the form of bursts of pulses. Each burst will typically have a duration in the range of 10 microseconds to 1.0 millisecond. The individual pulses in each burst typically each have a duration of 0.1 to 10 microseconds with an interval therebetween of 0.1 to 10 microseconds. The actual pulses are typically square waves and bi-phasic, that is alternating positive and negative amplitudes. Generally the interval between pulses must be shorter than a lifetime of the

plasma vapor cavity in order to maintain the cavity and the plasma regime during each pulse burst. In one variation the bursts each are separated by a duration of at least one millisecond. Typically the time between the pulse bursts is sufficient so that the duty-cycle is relatively low as explained above. This minimizes the undesirable heating effects. Coagulation may be achieved in the same manner, but by increasing output power of the device.

[0064] The generator associated with the electrosurgical devices may also allow varying degrees of cutting and coagulation. For example, in addition to a default power setting when the device is turned on, other settings to which the device could be adjusted are provided. This would provide in some instances higher hemostasis when cutting.

[0065] The electrosurgical devices described here are intended for single use, and can be employed with tonsillectomy, adenoidectomy, or combined procedures. In use, the tonsils and/or adenoids are accessed and a pulsed electrical signal applied to the active electrode of the interchangeable tip to form a plasma. The tissue is then cut or coagulated with the plasma. Suction may or may not be used when cutting or coagulating. In some instances, suction is applied simultaneously with proximal movement (pulling) of the device to suction cut the tissue. In other instances, suction is applied through the active electrode, as described above.

[0066] The electrosurgical devices described here may also be useful in “dry field” surgical procedures where there is no electrically conductive fluid added to the surgical field to, for example, aid conduction of energy between the active and return electrodes and/or to provide a cooling effect to the electrodes. When a dry field approach is employed, energy is typically transmitted only by the portion(s) of the active electrode that contacts tissue. In this instance the energy can be selectively provided to the areas in which it is desired. The application of suction to the tissue may further remove body fluids that would be naturally present in the field to further facilitate tissue contact.

[0067] Contrastingly, in a wet field (e.g., when the active and/or return electrodes are immersed or submerged in an electrically conductive fluid provided from outside the body), all

parts of the active electrode that are in contact with the electrically conductive fluid provide energy. As a result, more energy than required is generally transmitted to the tissue, and less precisely.

[0068] Moreover it has been found that a concave blade does not operate well in a wet field (conductive medium) because components of the blade's electric field directed in opposite directions cancel each other. This does not occur in a dry field since current (energy) only flows to the tissue from that part of the blade in direct contact with the tissue. Hence a dry field approach is advantageous here, so the use of aspiration as described here is beneficial to create the dry field. Further, the suction (aspiration) here directed from the center of the blade pulls the tissue into the aspiration port thereby further decreasing the width of the tissue touching the blade and rendering the field even drier, thereby further improving the accuracy of current flow and the electric field effects described above.

[0069] The interchangeable tip may be exchanged for another at any time during the procedure. Thus, it may be useful to provide the suction handle and interchangeable tips with blade assemblies in a kit. Given that the device is intended for single use, one or more suction handles could be packaged with one or more interchangeable tips with blade assemblies. The interchangeable tips could be of the same type, e.g., all adenoidectomy tips or all tonsillectomy tips, or a mixture of the two types. In other instances, the kits may include a plurality of interchangeable tips without the suction handle. The interchangeable tips and suction handles may also be individually packaged.

[0070] The kits will also generally contain instructions for use. The instructions may include directions on how to start, operate, and shut down the device, as well as directions on how to adjust power level settings. Steps for changing one interchangeable tip to another, e.g., an adenoidectomy tip for a tonsillectomy tip, may also be provided.

[0071] The invention described herein will be further understood by the following non-limiting example.

[0072] Example: Comparative Data on Thermal Injury

Excised human palatine tonsils were subjected to a series of surgical incisions using conventional “Bovie” needle tips at settings of 30W Cut and 30W Coag. Additional incisions were made using the electrosurgical tonsillectomy devices described here (“Tonsil Blade”) on settings of Cut 5 (20W power output) for cutting and Coag 6 (30W power output) for coagulation. Histology samples were harvested immediately after incision and evaluated for residual thermal injury using microscopy. A comparison of the depth of thermal injury (mm) and percent reduction in thermal injury is provided in Table 1.

Table 1. Comparison of Thermal Injury Depth

	Thermal Injury Depth (mm)	
	Cut	Coag
Tonsil Blade (20W Cut) (30W Coag)	0.26	0.26
Bovie (30W Cut) (30W Coag)	0.77	0.96
Percent Reduction	66%	73%
p value	$p < 0.05$	$p < 0.005$

CLAIMS

1. An electrosurgical device comprising:
 - a) an elongate body having a proximal end, a distal end, and defining a lumen for applying aspiration extending therethrough;
 - b) a tip having a proximal end, a distal end, and defining a tip lumen fluidly connected to the aspiration lumen at the distal end of the elongate body;
 - c) a housing secured to the distal end of the tip and defining an aperture fluidly connected to the tip lumen; and
 - d) a blade assembly in the housing and comprising an electrode,
wherein the blade assembly or portion thereof forms an aspiration port with the aperture of the housing.
2. The electrosurgical device of claim 1, wherein the tip is malleable.
3. The electrosurgical device of claim 1, further comprising a connector at the proximal end of the tip for removably attaching the tip to the distal end of the elongate body.
4. The electrosurgical device of claim 3, wherein the connector comprises a compressible barrel and one or more tabs located about the circumference of the connector.
5. The electrosurgical device of claim 4, wherein the one or more tabs attach a finger grip to the connector.
6. The electrosurgical device of claim 1, wherein the housing comprises an upper portion and a lower portion that define the aperture when joined to each other.

7. The electrosurgical device of claim 1, wherein the aperture is located in a wall of the housing.

8. The electrosurgical device of claim 7, wherein the aperture is spherical or elliptical.

9. The electrosurgical device of claim 7, wherein the aperture is about 0.4 cm to about 1.5 cm along its major axis.

10. The electrosurgical device of claim 7, wherein the aperture is about 0.4 cm to about 1.0 cm along its minor axis.

11. The electrosurgical device of claim 1, wherein the housing comprises a transparent material.

12. The electrosurgical device of claim 11, wherein the transparent material is a polymer.

13. The electrosurgical device of claim 12, wherein the polymer is selected from the group consisting of acrylic copolymers, acrylonitrile butadiene styrene (ABS), polycarbonate, polystyrene, polyvinyl chloride (PVC), polyethylene terephthalate glycol (PETG), and styrene acrylonitrile (SAN).

14. The electrosurgical device of claim 13, wherein the polymer is polymethyl methacrylate (PMMA) copolymer or styrene methyl methacrylate (SMMA) copolymer.

15. The electrosurgical device of claim 1, wherein the blade assembly comprises a plurality of arms for securing it to the housing.

16. The electrosurgical device of claim 15, wherein the plurality of arms each have a first portion and a second portion, and an angle of about 110° to about 140° between the first and second portions.

17. The electrosurgical device of claim 15, wherein the electrode comprises a cutting edge and a coagulation surface.

18. The electrosurgical device of claim 17, wherein the cutting edge and housing aperture define the aspiration port.

19. The electrosurgical device of claim 1, wherein the electrode extends past a remainder of the housing and has a curved portion and an opening defined therethrough.

20. The electrosurgical device of claim 19, wherein the opening is the aspiration port.

21. The electrosurgical device of claim 1, wherein the active electrode comprises a metal selected from the group consisting of molybdenum, nickel, platinum, stainless steel, tantalum, titanium, tungsten, and alloys thereof.

22. The electrosurgical device of claim 1, wherein the device is electrically monopolar.

23. The electrosurgical device of claim 1, wherein the electrode is elongated and defining the aspiration port in a central portion thereof, a cutting portion of the electrode lying

adjacent the aspiration port and being elongated and tapering towards its distal portion, wherein the distal portion lies away from a plane defined by a proximal portion of the cutting portion at an angle of at least 20°.

24. A method for resecting tonsil or adenoid tissue using the device of claim 1 comprising the acts of:

- a) applying a pulsed electrical signal to the electrode to form a plasma thereon;
- and
- b) cutting or coagulating the tissue with the plasma.

25. The method of claim 24, further comprising the act of applying suction to the tissue through the aspiration port.

26. The method of claim 25, wherein suction is applied simultaneously with proximal movement of the electrode to suction cut the tissue.

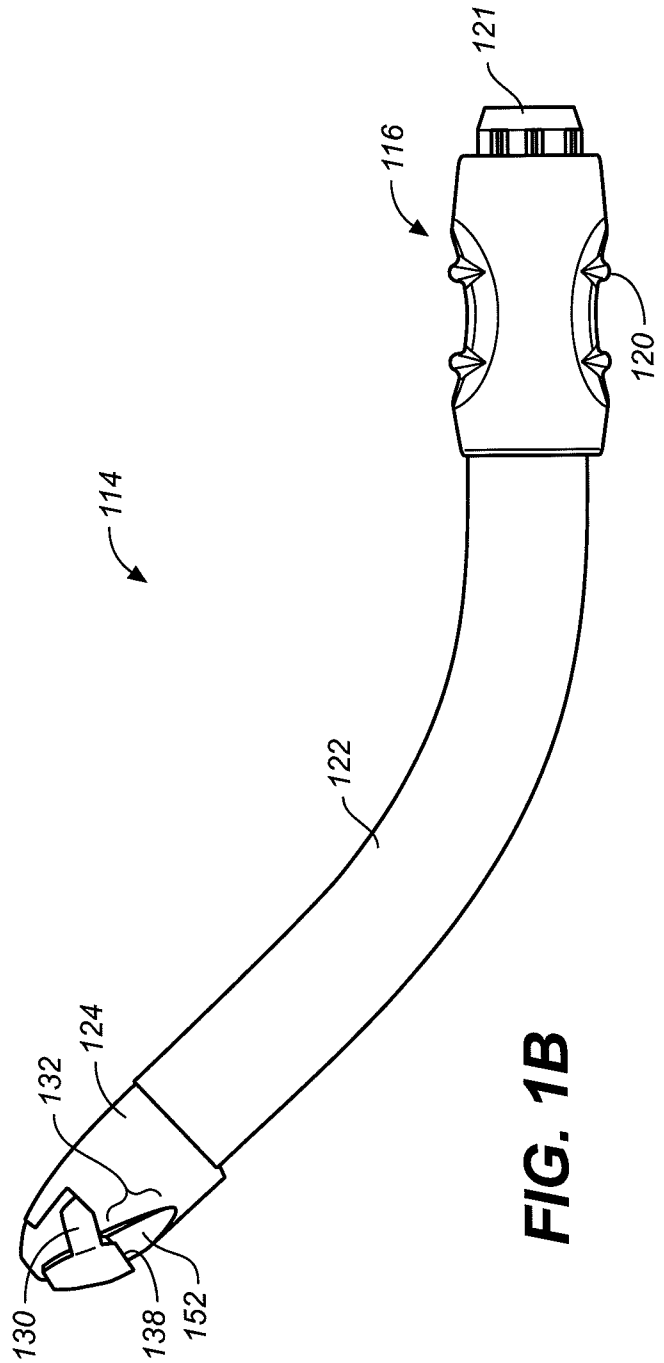
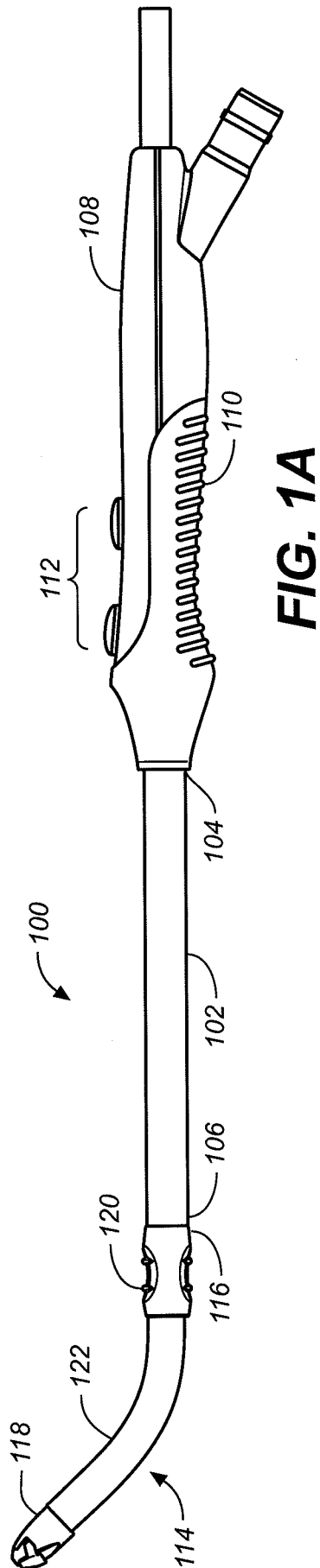
27. The method of claim 24, further comprising removing the tip and replacing it with a second tip during a surgical procedure.

28. A kit comprising:

- a) one or more tips, the tips comprising a housing defining an aperture and having a blade assembly therein comprising an electrode, wherein the blade assembly or portion thereof forms an aspiration port with the aperture of the housing; and
- b) instructions for use.

29. The kit of claim 28 further comprising one or more suction handles couplable to the tips.

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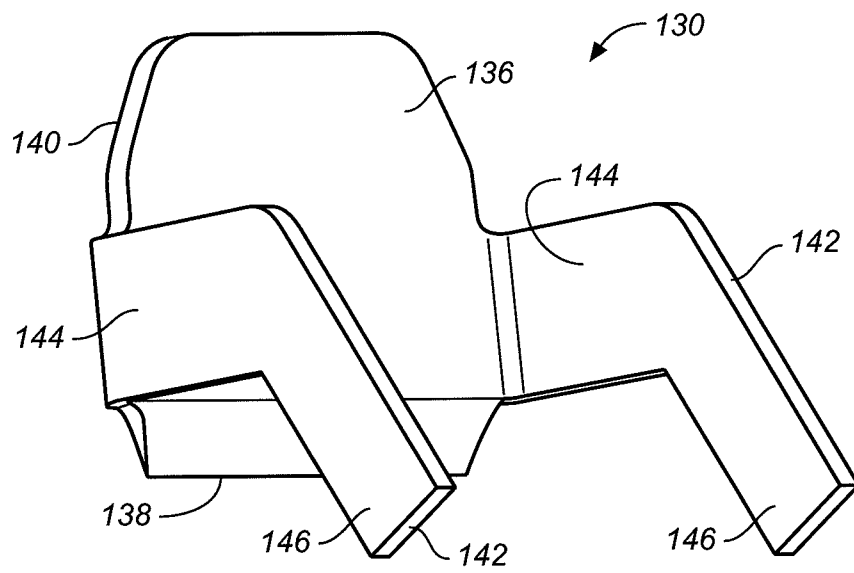


FIG. 1C

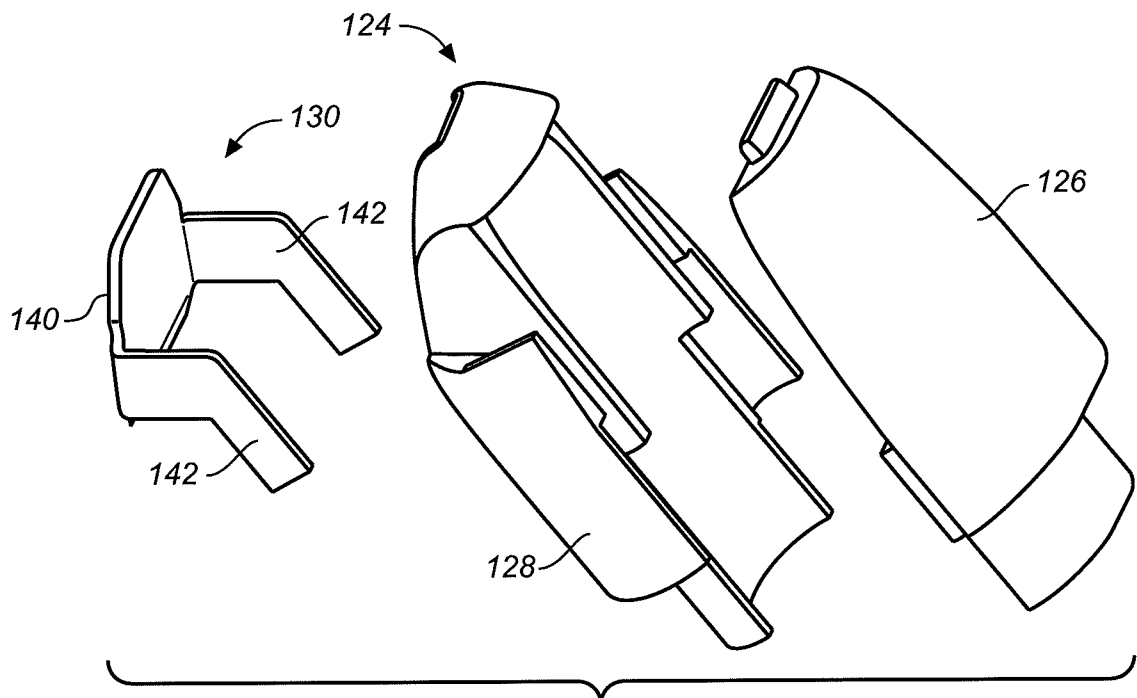


FIG. 1D

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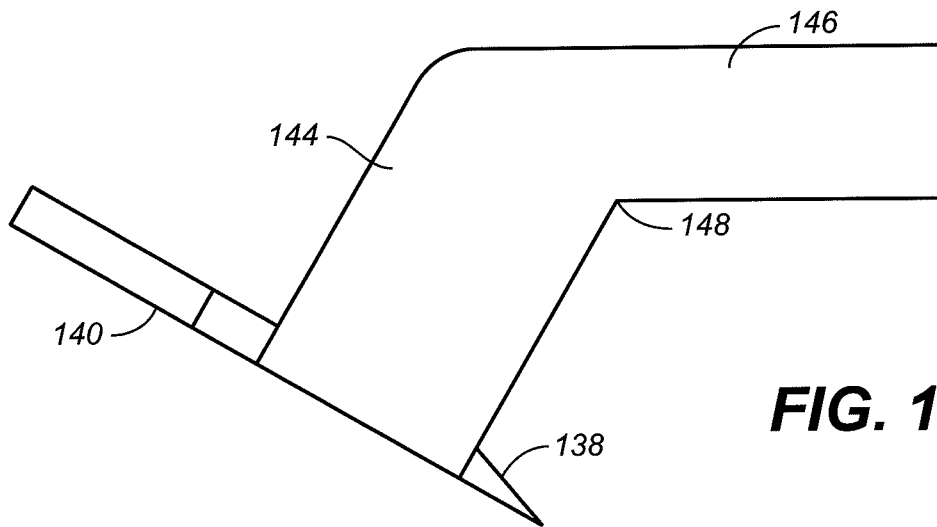


FIG. 1E

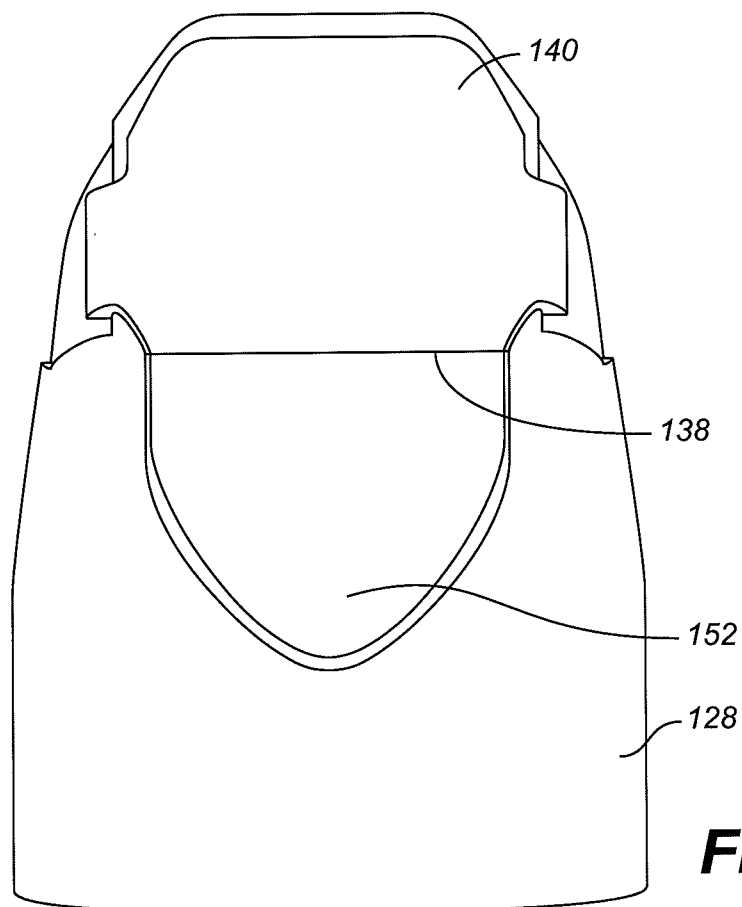


FIG. 1F

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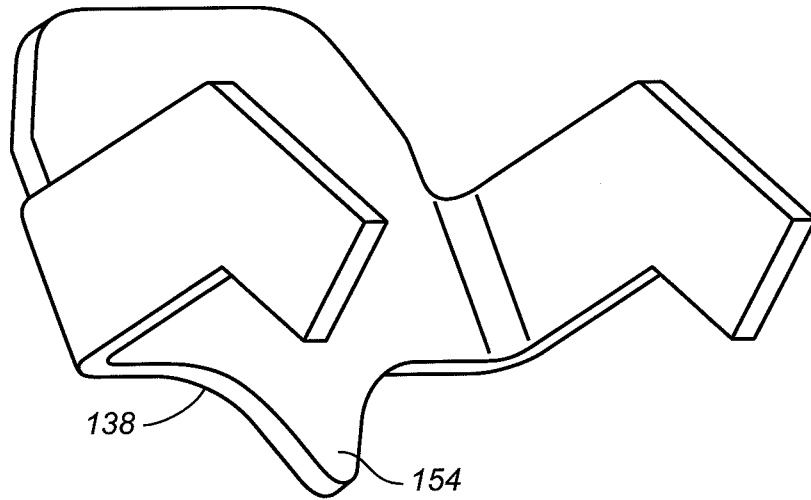


FIG. 1G

5 / 5

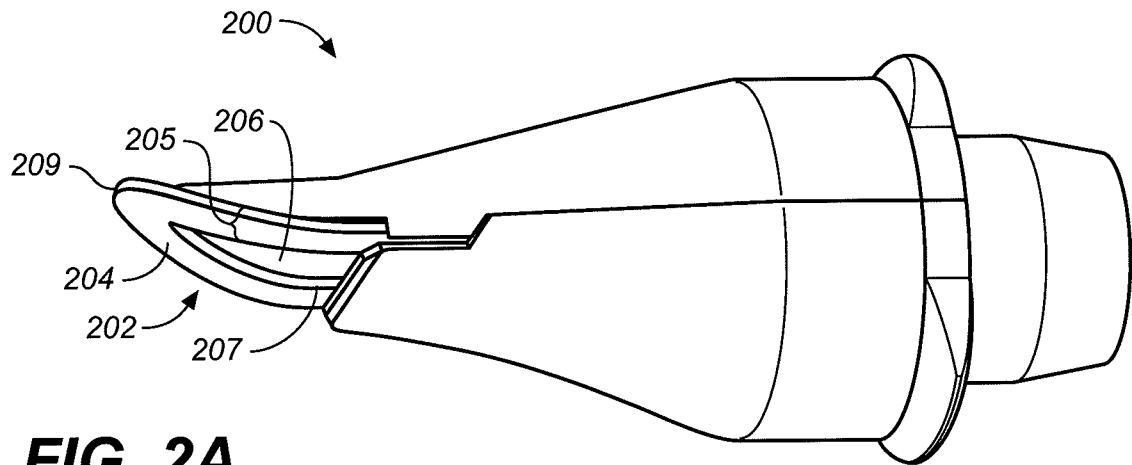


FIG. 2A

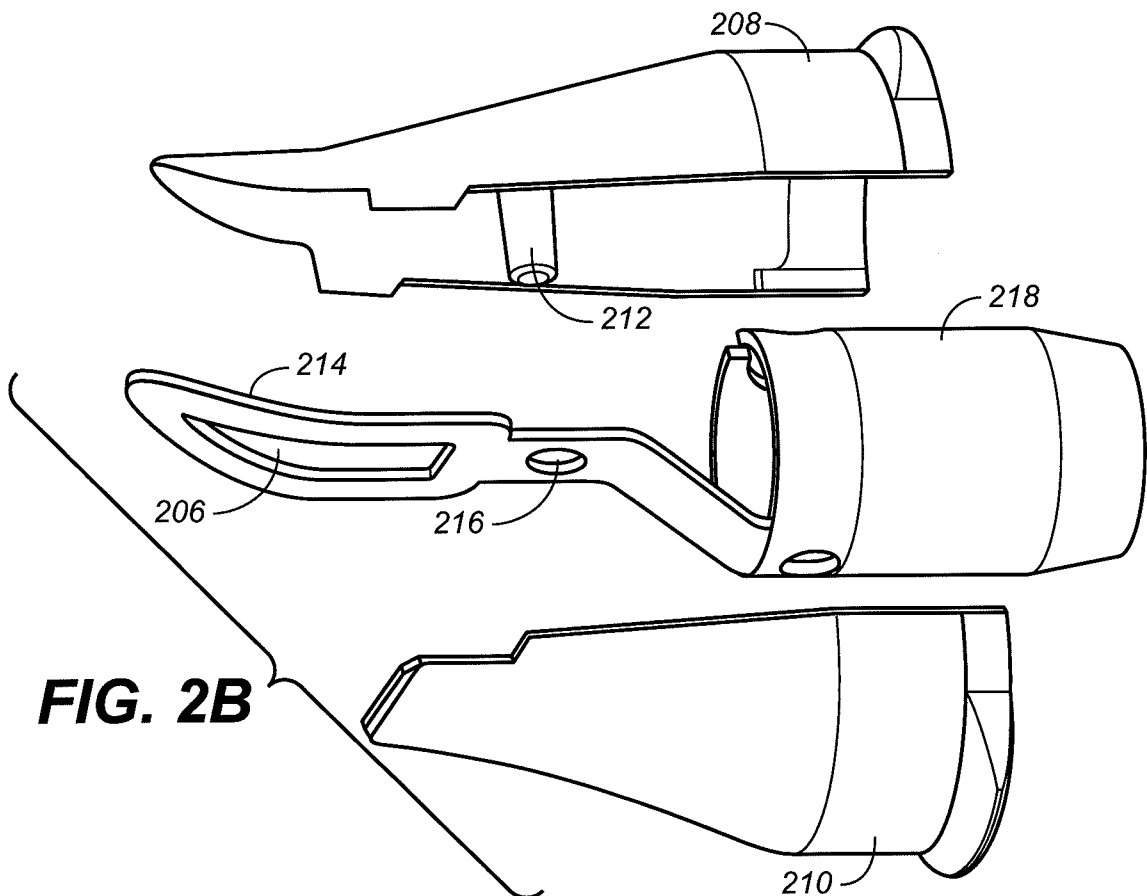


FIG. 2B

INTERNATIONAL SEARCH REPORT

International application No

PCT/US2009/030180

A. CLASSIFICATION OF SUBJECT MATTER

INV. A61B18/14

ADD. A61B18/00

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)

A61B

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 practical, search terms used)

EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X Y	EP 1 602 338 A2 (MEDTRONIC INC [US]) 7 December 2005 (2005-12-07) abstract; figures 1-2 paragraphs [0010] - [0012] -----	1-3, 21-23 17-18
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X A	WO 03/061499 A1 (ETHICON ENDO SURGERY INC [US]) 31 July 2003 (2003-07-31) abstract; figure 1 paragraphs [0032] - [0033] -----	28-29 1-5
	-/--	

☒ 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 document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

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

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

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

& document member of the same patent family

Date of the actual completion of the international search

18 November 2009

Date of mailing of the international search report

30/11/2009

Name and mailing address of the ISA/

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Fax: (+31-70) 340-3016

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Pfeiffer, Uwe

INTERNATIONAL SEARCH REPORT

International application No

PCT/US2009/030180

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 6 530 924 B1 (ELLMAN ALAN G [US] ET AL) 11 March 2003 (2003-03-11) abstract; figures 1,3 column 2, line 42 - column 3, line 48 column 5, lines 11-44 -----	1-3, 19-23, 28-29
X	US 5 234 428 A (KAUFMAN DAVID I [US]) 10 August 1993 (1993-08-10) abstract; figures 1,3 column 5, line 64 - column 6, line 35 -----	1-3, 6-10, 21-22 18
X	US 2007/208335 A1 (WOLOSZKO JEAN [US] ET AL) 6 September 2007 (2007-09-06) A abstract; figures 6,7,11,13,15,16 paragraphs [0007] - [0009], [0045] - [0047], [0 51], [0056], [0063] -----	1-3, 7-10,21 18-20
X	US 2005/080412 A1 (OUCHI TERUO [JP]) 14 April 2005 (2005-04-14) abstract; figures 1,2B,3,4 paragraphs [0001] - [0019], [0036] - [0041] -----	1-5, 7-10, 15-16, 19, 21-22, 28-29 17-18
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A	WO 98/43549 A1 (FEMRX INC [US]) 8 October 1998 (1998-10-08) page 1, line 12 - page 3, line 19 page 4, line 1 - page 6, line 3 figure 1 -----	1,15-18

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2009/030180

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.: 24-27
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

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

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

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of 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.:
1-10, 15-23, 28-29
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.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 1-5, 19-23, 28-29

electrosurgical device with removable tip

2. claims: 6-10

electrosurgical device with a distal housing forming an aperture

3. claims: 11-14

electrosurgical device with a transparent distal housing

4. claims: 15-18

electrosurgical device with an electrode blade secured to the housing

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2009/030180

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