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(54) SURGICAL SYSTEM FOR CONVERTING A CLAMP INTO AN ELECTROPHYSIOLOGY DEVICE
OPERATIONSSYSTEM ZUR VERWANDLUNG EINER KLEMME IN EINE 
ELEKTROPHYSIOLOGISCHE VORRICHTUNG
SYSTEME CHIRURGICAL PERMETTANT DE TRANSFORMER UN CLAMP EN DISPOSITIF 
D’ELECTROPHYSIOLOGIE

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(56) References cited: 
US-B1- 6 210 330 US-B1- 6 312 426

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Description

BACKGROUND OF THE INVENTIONS

1. Field of Inventions

[0001] The invention relates to a surgical system according to the preamble of claim 1.  
[0002] The present invention relates generally to structures for positioning diagnostic and therapeutic elements within the body and, more particularly, to devices which are particularly well suited for the treatment of cardiac conditions.

2. Description of the Related Art

[0003] A surgical system of the above-mentioned type is known, for example, from WO 00 24330 A.
[0004] There are many instances where diagnostic and therapeutic elements must be inserted into the body. One instance involves the treatment of cardiac conditions such as atrial fibrillation and atrial flutter which lead to an unpleasant, irregular heart beat, called arrhythmia.
[0005] Normal sinus rhythm of the heart begins with the sinoatrial node (or "SA node") generating an electrical impulse. The impulse usually propagates uniformly across the right and left atria and the atrial septum to the atrioventricular node (or "AV node"). This propagation causes the atria to contract in an organized way to transport blood from the atria to the ventricles, and to provide timed stimulation of the ventricles. The AV node regulates the propagation delay to the atrioventricular bundle (or "HIS" bundle). This coordination of the electrical activity of the heart causes atrial systole during ventricular diastole. This, in turn, improves the mechanical function of the heart. Atrial fibrillation occurs when anatomical obstacles in the heart disrupt the normally uniform propagation of electrical impulses in the atria. These anatomical obstacles (called "conduction blocks") can cause the electrical impulse to degenerate into several circular wavelets that circulate about the obstacles. These wavelets, called "reentry circuits," disrupt the normally uniform activation of the left and right atria. Because of a loss of atrioventricular synchrony, the people who suffer from atrial fibrillation and flutter also suffer the consequences of impaired hemodynamics and loss of cardiac efficiency. They are also at greater risk of stroke and other thromboembolic complications because of loss of effective contraction and atrial stasis.
[0006] One surgical method of treating atrial fibrillation by interrupting pathways for reentry circuits is the so-called "maze procedure" which relies on a prescribed pattern of incisions to anatomically create a convoluted path, or maze, for electrical propagation within the left and right atria. The incisions direct the electrical impulse from the SA node along a specified route through all regions of both atria, causing uniform contraction required for normal atrial transport function. The incisions finally direct the impulse to the AV node to activate the ventricles, restoring normal atrioventricular synchrony. The incisions are also carefully placed to interrupt the conduction routes of the most common reentry circuits. The maze procedure has been found very effective in curing atrial fibrillation. However, the maze procedure is technically difficult to do. It also requires open heart surgery and is very expensive. Thus, despite its considerable clinical success, only a few maze procedures are done each year.
[0007] Maze-like procedures have also been developed utilizing catheters and/or surgical probes (collectively "probes") that form lesions to create a maze for electrical conduction in a predetermined path. Typically, the lesions are formed by ablating tissue with one or more electrodes. Electromagnetic radio frequency ("RF") energy applied by the electrode heats, and eventually kills (i.e., "ablates"), the tissue to form a lesion. During the ablation of soft tissue (i.e. tissue other than blood, bone and connective tissue), tissue coagulation occurs and it is the coagulation that kills the tissue. Thus, references to the ablation of soft tissue are necessarily references to soft tissue coagulation. "Tissue coagulation" is the process of cross-linking proteins in tissue to cause the tissue to jell. In soft tissue, it is the fluid within the tissue cell membranes that jells to kill the cells, thereby killing the tissue.
[0008] Catheters used to create lesions typically include a relatively long and relatively flexible body that has one or more electrodes on its distal portion. The portion of the catheter body that is inserted into the patient is typically from 58.4 to 139.7 cm in length and there may be another 20.3 to 38.1 cm, including a handle, outside the patient. The proximal end of the catheter body is connected to the handle which includes steering controls. The length and flexibility of the catheter body allow the catheter to be inserted into a main vein or artery (typically the femoral artery), directed into the interior of the heart, and then manipulated such that the electrode contacts the tissue that is to be ablated. Fluoroscopic imaging is used to provide the physician with a visual indication of the location of the catheter. Exemplary catheters are disclosed in U.S. Patent No. 5,582,609.
[0009] Surgical probes used to create lesions often include a handle, a relatively short shaft that is from about 10.2 to 45.7 cm in length and either rigid or relatively stiff, and a distal section that is from 2.54 to 25.4 cm in length and either malleable or somewhat flexible. One or more electrodes are carried by the distal section. Surgical probes are used in epicardial and endocardial procedures, including open heart procedures and minimally invasive procedures where access to the heart is obtained via a thoracotomy, thoracostomy or median sternotomy. Exemplary surgical probes are disclosed in U.S. Patent No. 6,142,994.
[0010] Clamps, which have a pair of opposable rigid clamp members that may be used to hold a bodily structure or a portion thereof, are used in many types surgical
procedures. Lesion creating electrodes have also been permanently secured to certain types of clamps. Examples of clamps which carry lesion creating electrodes are disclosed in U.S. Patent No. 6,142,994. Such clamps are particularly useful when the physician intends to position electrodes on opposite sides of a body structure.

[0011] As used herein, the term "clamp" includes, but is not limited to, clamps, clips, forceps, hemostats, and any other surgical device that includes a pair of opposable clamp members that hold tissue, at least one of which is movable relative to the other. In some instances, the rigid clamp members are connected to a scissors-like arrangement including a pair of handle supporting arms that are pivotally connected to one another. The clamp members are secured to one end of the arms and the handles are secured to the other end. The clamp members come together as the handles move toward one another. Certain clamps that are particularly useful in minimally invasive procedures also include a pair of handles and a pair of clamp members. Here, however, the clamp members and handles are not mounted on the opposite ends of the same arm. Instead, the handles are carried by one end of an elongate housing and the clamp members are carried by the other. A suitable mechanical linkage located within the housing causes the clamp members to move relative to one another in response to movement of the handles.

[0012] The rigid clamp members in conventional clamps may be linear or have a predefined curvature that is optimized for a particular surgical procedure or portion thereof. It is, therefore, necessary to have a wide variety of clamps on hand. In the field of electrophysiology, a wide variety of clamps that have electrodes permanently secured thereto must be kept on hand.

[0013] The inventor herein has determined that it would be advantageous to provide physicians with a wide variety of devices, including clamps (both with and without energy transmission devices) and surgical probes that carry energy transmission devices, in a wide variety of shapes, and to do so in a manner that is more cost effective than conventional apparatus.

SUMMARY OF THE INVENTIONS

[0014] The invention provides a surgical system having the features of claim 1. The apparatus provides a number of advantages. For example, such an apparatus may be used to quickly convert a conventional clamp into an electrophysiology device. In those instances where a procedure requires a number of different clamps, the apparatus can be moved from clamp to clamp, thereby eliminating the costs associated with providing a variety of different clamps with energy transmission devices permanently secured thereto. Further embodiments are described in the dependent claims.

[0015] The clamp in accordance with one embodiment includes first and second clamp members, at least one of which is malleable, and a movement apparatus that moves at least one of the first and second clamp members relative to the other. Such a clamp provides a number of advantages. For example, the malleable clamp member allows physicians to readily reconfigure the clamp, thereby reducing the number of clamps that must be provided for a particular surgical procedure.

[0016] For example, the system may be used both as a conventional clamp and an electrophysiology device.

[0017] The above described and many other features and attendant advantages of the present inventions will become apparent as the inventions become better understood by reference to the following detailed description when considered in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

[0018] Detailed description of preferred embodiments of the inventions will be made with reference to the accompanying drawings.

Figure 1 is a plan view of a conventional clamp. Figure 2 is a side view of the clamp illustrated in Figure 1. Figure 3 is an enlarged view of a portion of the clamp illustrated in Figure 1 holding a vein. Figure 4 is plan of a pair of energy transmission assemblies in accordance with a preferred embodiment of a present invention. Figure 5 is plan showing the energy transmission assemblies illustrated in Figure 4 mounted on a clamp. Figure 6 is a front view of an electrosurgical unit. Figure 7a is a section view taken along line 7a-7a in Figure 4. Figure 7b is a section view taken along line 7b-7b in Figure 4. Figure 8 is a section view taken along line 8-8 in Figure 7a. Figure 9a is a section view of an energy transmission assembly in accordance with a preferred embodiment of a present invention. Figure 9b is a section view of an energy transmission assembly in accordance with a preferred embodiment of a present invention. Figure 10 is a plan view of an energy transmission assembly not in accordance with the present invention. Figure 11 is a section view taken along line 11-11 in Figure 10. Figure 12 is a section view of an energy transmission assembly in accordance with a preferred embodiment of a present invention. Figure 13 is a section view of an energy transmission assembly in accordance with a preferred embodiment of a present invention. Figure 14 is a section view taken along line 14-14 in Figure 13.
the general principles of the inventions.

[0020] The detailed description of the preferred embodiments is organized as follows:

I. Energy Transmission Assemblies
II. Energy Transmission Devices, Temperature Sensing and Power Control
III. Tissue Cooling Apparatus
IV. Probe Support Devices
V. Clamp With Malleable Clamp Members

The section titles and overall organization of the present detailed description are for the purpose of convenience only and are not intended to limit the present inventions.

[0021] This specification discloses a number of structures, mainly in the context of cardiac ablation, because the structures are well suited for use with myocardial tissue. Nevertheless, it should be appreciated that the structures are applicable for use in therapies involving other types of soft tissue. For example, various aspects of the present inventions have applications in procedures concerning other regions of the body such as the prostate, liver, brain, gall bladder, uterus and other solid organs.

I. Energy Transmission Assemblies

[0022] Energy transmission assemblies in accordance with a present invention may be used to covert a conventional clamp into a tissue coagulation device. The energy transmission assemblies may also be used to covert a clamp in accordance with the inventions described in Section V below into a tissue coagulation device.

[0023] One example of a conventional clamp that may be used in conjunction with the present inventions is generally represented by reference numeral 10 in Figures 1-3. The clamp 10 includes a pair of rigid arms 12 and 14 that are pivotably connected to one another by a pin 16. The proximal ends of the arms 12 and 14 are respectively connected to a pair handle members 18 and 20, while the distal ends are respectively connected to a pair of rigid clamp members 22 and 24. A locking device 26 locks the clamp in the closed orientation, and prevents the clamp members 22 and 24 from coming any closer to one another than is illustrated in Figure 1, thereby defining a predetermined spacing between the clamp members. The clamp 10 also includes a pair of soft, deformable inserts 28 and 30 that are removably carried by the clamp members 22 and 24. The inserts 28 and 30 allow clamp 10 to firmly grip a bodily structure 32 without damaging the bodily structure. The inserts 28 and 30 include mating structures 34 that extend through corresponding apertures 36 in the clamp members 22 and 24 to hold the inserts in place.

[0024] As illustrated for example in Figures 4 and 5, an apparatus 100 for converting the clamp 10 (which has had the inserts 28 and 30 removed) into a bi-polar tissue coagulation device includes a pair of energy transmission
assemblies 102 and 104. Each of the energy transmission assemblies includes a base member 106 that may be removably secured to one of the clamp members 22 and 24 and an energy transmission device 108. [The energy transmission devices 108 are discussed in greater detail in Section II below.] Although the configuration of the energy transmission assemblies 102 and 104 may vary from application to application, the energy transmission assemblies in the exemplary embodiment are configured such that they will abut one another in the same manner as the inserts 28 and 30 (Figures 1-3) when the clamp 10 is in the closed orientation illustrated in Figure 5. Such an arrangement will allow the energy transmission assemblies 102 and 104 to grip a bodily structure in the same manner as the inserts 28 and 30.

[0025] The exemplary base members 106 are preferably formed from a soft, resilient, low durometer material that is electrically insulating. Suitable materials include polyurethane, silicone and polyurethane/silicone blends having a hardness of between about 20 Shore D and about 72 Shore D. Referring to Figures 7a, 7b and 8, each of the exemplary base members 106 includes a longitudinally extending aperture 110 into which one of the clamp members 22 and 24 may be inserted. The apertures 110 should be sized and shaped such that the base members 106 will be forced to stretch when the clamp members 22 and 24 are inserted. If, for example, the apertures 110 have the same cross-sectional shape as the clamp members 22 and 24 (e.g. both are elliptical), then the apertures should be slightly smaller in their cross-sectional dimensions than the corresponding clamp members. The stretching of the apertures 110 creates a tight interference fit between the base members 106 and clamp members 22 and 24. Additionally, although the apertures 110 have a semi-circular cross-section in the exemplary embodiment, the apertures may have a round, rectangular, square or elliptical cross-section, or define any other cross-sectional shape, depending on the particular application.

[0026] The exemplary base members 106 also include slots 112 (Figure 8) that secure the energy transmission devices 108 in place. The configuration of a slot 112, of course, depend on the configuration of the energy transmission device 108 that it is holding. The illustrated energy transmission device 108 is generally cylindrical in shape and the slot 112 has a corresponding arcuate cross-sectional shape. The arc is preferably greater than 180 degrees so that the base member 106 will deflect when the energy transmission device 108 is inserted into the slot 112 and then snap back to hold the energy transmission device in place. Adhesive may also be used to secure the energy transmission devices 108, especially in those instances where the arc is less than 180 degrees.

[0027] Another exemplary apparatus for converting the clamp 10 (which has had the inserts 28 and 30 removed) into a bipolar tissue coagulation device is illustrated in Figures 9a and 9b. The apparatus includes a pair of energy transmission assemblies 114 and 116 which are substantially similar to the energy transmission assemblies 102 and 104 and similar elements are represented by similar reference numerals. Each of the energy transmission assemblies 114 and 116 includes a base member 106 that may be removably secured to one of the clamp members 22 and 24 and an energy transmission device 108. Here, however, the base members 106 are secured to the clamp members 22 and 24 with mating structures 118 that mechanically engage the clamp members.

[0028] The exemplary mating structures 118, which are preferably integral with the base members 106 and formed from the same resilient material, include a relatively narrow portion 120 and a relatively wide portion 122. The relatively narrow portions are approximately the same size as the clamp member apertures 36 and the relatively wide portions 122 are slightly larger than the clamp member apertures. A removable connection is made by urging the mating structures 118 into one end of the apertures 36, thereby deforming the relatively wide portions 122, and then urging the base members 106 against the clamp members 22 and 24 until the relatively wide portions exit through the other end of the apertures and reassume their original shape.

[0029] The exemplary mating structures 118 may also be reconfigured by eliminating the relatively wide portions 122 and enlarging the relatively narrow portions 120 such that the relatively narrow portions will create an interference fit within the clamp member apertures 36. Alternatively, as discussed below with reference to Figure 12, longitudinally extending mating structures, which also create an interference fit, may be employed when longitudinally extending slots are formed in the clamp members. Another alternative is to provide the clamp members with one or more small mating structures that extend outwardly therefrom. The clamp member mating structures will be received within apertures or slots formed in the base member.

[0030] Turning to Figures 10 and 11 not according to the invention, an energy transmission assembly 124 may be used to convert the clamp 10 (which has had the inserts 28 and 30 removed) into a uni-polar tissue coagulation device. The energy transmission assembly 124 includes a base member 126, which may be removably secured to both of the clamp members 22 and 24, and a plurality of spaced energy transmission devices 108. Although the configuration of the energy transmission assembly 124 may vary from application to application to suit particular situations, the energy transmission assembly in the exemplary embodiment is configured such that it will abut each of the clamp members when the clamp 10 is in the closed orientation illustrated in Figure 10.

[0031] The exemplary base member 126 is preferably formed from a soft, resilient, low durometer material that is electrically insulating. Suitable materials include polyurethane, silicone and polyurethane/silicone blends having a hardness of between about 20 Shore D and about
72 Shore D. A slot 128 secures the energy transmission devices 108 in place. Although the configuration of the slot 128 will depend on the configuration of the energy transmission devices 108, the exemplary slot has an arcuate cross-sectional shape that conforms to the shape of the exemplary cylindrical energy transmission devices. The arc is preferably greater than 180 degrees so that the base member 126 will deflect when the energy transmission devices 108 are inserted into the slot 128 and then snap back to hold the energy transmission devices in place. Adhesive may also be used to secure the energy transmission devices 108 in place, especially in those instances where the arc is less than 180 degrees.

[0032] The base member 126 is removable secured to the clamp members 22 and 24 with two sets of the mating structures 118 that are described above with reference to Figures 9a and 9b (with or without the relatively wide portions 122). Alternatively, and as illustrated for example in Figure 12, in those instances where the clamp members 22" and 24" include longitudinally extending slots 38 instead of the apertures 36, the energy transmission assembly 124 may be provided with longitudinally extending mating structures 130 that extend outwardly from the base member 126". The longitudinally extending mating structures 130, which are preferably integral with the base member 126" and formed from the same resilient material, are sized and shaped to create an interference fit with the slots 38. Still another alternative is to provide the clamp members with one or more small mating structures that are received within apertures or slots formed in the base member.

[0033] Another energy transmission assembly that may be used to convert the clamp 10 into a uni-polar tissue coagulation device is generally represented by reference numeral 132 in Figures 13 and 14. The energy transmission assembly 132 includes a base member 134 that is preferably formed from a soft, resilient, low durometer material and a plurality of energy transmission devices 108. The material which forms the base member 134 should also be electrically insulating. Suitable materials include polyurethane, silicone and polyurethane/silicone blends having a hardness of between about 20 Shore D and about 72 Shore D. A slot 128, which secures the energy transmission devices 108 in place in the manner described above with reference to Figures 10 and 11, is also provided.

[0034] The exemplary base member 134 includes a longitudinally extending aperture 136 into which both of the clamp members 22 and 24 may be inserted. The aperture 136 should be sized and shaped such that the base member 134 will be forced to stretch when the clamp members 22 and 24 are inserted with the clamp 10 in a closed orientation. The stretching creates a tight interference fit between the base member 134 and the clamp members 22 and 24. Additionally, although the apertures 110 have an elliptical cross-section in the exemplary embodiment, the apertures may have a round, rectangular, square or semi-circular cross-section, or de-fine any other cross-sectional shape, depending on the particular application.

[0035] The length of the base members in the exemplary energy transmission assemblies will vary according to the intended application. In the area of cardiovascular treatments, it is anticipated that suitable lengths will range from, but are not limited to, about 2 cm to about 10 cm.

[0036] The exemplary energy transmission assemblies described above may also be modified in a variety of ways. For example, the energy transmission assembly illustrated in Figures 10 and 11 may be converted into a bi-polar device by simply adding a second slot 128 that is preferably spaced apart from and parallel to the existing slot. The second slot 128 could, for example, include a single return energy transmission device 108 or a plurality of spaced return energy transmission devices. Additionally, as illustrated for example in Figures 7a and 13, the base members and energy transmission devices in the illustrated embodiments are configured such that the energy transmission devices are generally linear and parallel to the longitudinal axis of the base members (when the assemblies are in a relaxed state and not being urged against a body structure). The base members and/or energy transmission devices may be reconfigured such that the energy transmission devices, or a portion thereof, are curved and/or non-parallel to the longitudinal axis of the base members when in the relaxed state.

II. Energy Transmission Devices, Temperature Sensing and Power Control

[0037] In the exemplary embodiments illustrated in Figures 4-16b, the energy transmission devices are electrodes. More specifically, the energy transmission devices are preferably in the form of wound, spiral coil electrodes that are relatively flexible. The coils are made of electrically conducting material, like copper alloy, platinum, or stainless steel, or compositions such as draw-filled tubing (e.g. a copper core with a platinum jacket). The electrically conducting material of the coils can be further coated with platinum-iridium or gold to improve its conduction properties and biocompatibility. A preferred coil electrode configuration is disclosed in U.S. Patent No. 5,797,905. Although the diameter of the electrodes will vary from application to application, the diameter preferably ranges from about 1 mm to about 3 mm for cardiovascular applications.

[0038] As an alternative, the electrodes may be in the form of solid rings of conductive material, like platinum, or can comprise a conductive material, like platinum-iridium or gold, coated upon the base member using conventional coating techniques or an ion beam assisted deposition (IBAD) process. For better adherence, an undercoating of nickel or titanium can be applied. The electrodes can also be in the form of helical ribbons. The electrodes can also be formed with a conductive ink compound that is pad printed onto a non-conductive tubular.
body. A preferred conductive ink compound is a silver-based flexible adhesive conductive ink (polyurethane binder), however other metal-based adhesive conductive inks such as platinum-based, gold-based, copper-based, etc., may also be used to form electrodes. Such inks are more flexible than epoxy-based inks.

[0039] When a single flexible coil electrode is carried by a base member (see, for example, Figure 7a), the length will depend on the length of the member and the intended application. When a plurality of spaced flexible coil electrodes are carried by a base member (see, for example, Figure 10), the electrodes will preferably be about 10 mm to about 40 mm in length. Preferably, the electrodes will be 25 mm in length with 1 mm to 2 mm spacing, which will result in the creation of continuous lesion patterns in tissue when coagulation energy is applied simultaneously to adjacent electrodes. For rigid electrodes, the length of each electrode can vary from about 3 mm to about 10 mm. Using multiple rigid electrodes longer than about 10 mm each adversely effects the overall flexibility of the device, while electrodes having lengths of less than about 2 mm do not consistently form the desired continuous lesion patterns.

[0040] It should also be noted that other energy transmission devices, such as laser arrays, ultrasonic transducers, microwave electrodes, and ohmically heated hot wires, may be substituted for the electrodes. Another type of energy transmission device that may be substituted for the electrodes is cryotemperature elements. Here, the energy transmission is the removal of heat from the tissue. Still another type of energy transmission device that may be substituted for the electrodes is needle projections for chemical ablation (which are preferably about 1 to 2 mm in length). Here, the energy transmission is the transmission of chemical energy.

[0041] Referring for example to Figures 5-8, each energy transmission device 108 is individually coupled to a wire 137 (Figure 8) that conducts coagulating energy. The wires 137 pass in conventional fashion through cables 138 to an associated connector (140 or 142). The connectors 140 and 142 are configured to plug into an electrosurgical unit ("ESU") 144 that supplies and controls power, such RF power. A suitable ESU is the Model 4810 ESU sold by EP Technologies, Inc. of San Jose, California. The exemplary ESU 144 illustrated in Figure 6 includes a plurality of displays and buttons that are used to control the level of power supplied to the energy transmission device(s) 108 and the temperature at the energy transmission device(s). When a plurality of spaced energy transmission devices 108 are employed, the ESU 144 may also be used to selectively control which of the energy transmission devices receive power. The amount of power required to coagulate tissue ranges from 5 to 150 W.

[0042] The exemplary ESU 144 illustrated in Figure 6 is operable in a bi-polar mode, where tissue coagulation energy emitted by the energy transmission device(s) 108 on one energy transmission assembly is returned through the energy transmission device(s) on another energy transmission assembly, and a uni-polar mode, where the tissue coagulation energy emitted by the energy transmission device(s) on an energy transmission assembly is returned through one or more indifferent electrodes (not shown) that are externally attached to the skin of the patient with a patch or one or more electrodes (not shown) that are positioned in the blood pool. To that end, the exemplary ESU 144 is provided with a power output connector 141 and a pair of return connectors 143. According to the invention, the ESU output and return connectors 141 and 143 have different shapes to avoid confusion and the connectors 140 and 142 have corresponding shapes. As such, in the exemplary bi-polar arrangement illustrated in Figure 5, the connector 140 associated with energy transmission assembly 102 has a shape corresponding to the ESU output connector 141 and the connector 142 associated with energy transmission assembly 104 has a shape corresponding to the ESU return connector 143.

[0043] The connector (not shown) associated with the energy transmission assembly 124 illustrated in Figure 10, which is intended to be operated in the uni-polar mode, would have a shape corresponding to the ESU output connector 141. In those instances where it is desirable to clamp the indifferent electrode within the patient, as opposed to positioning the indifferent electrode on the patient’s skin, a second energy transmission assembly may be provided with a connector having a shape corresponding to the ESU return connector 143. Additionally, in those instances where the energy transmission assembly 124 has been modified to includes space electrodes (or spaced groups of longitudinally spaced electrodes) that operated in bi-polar fashion, the assembly would be provided with a pair of connectors. One would have a shape corresponding to the ESU output connector 141 and the other would have a shape corresponding to the ESU return connector 143.

[0044] With respect to power and temperature control, one or more temperature sensors 146, such as thermocouples or thermistors, may be located on, under, abutting the longitudinal end edges of, or in between, the energy transmission devices 108. A reference thermocouple (not shown) may also be provided. For temperature control purposes, signals from the temperature sensors 146 are transmitted to the ESU 144 by way of wires 148 (Figure 8) that are connected to the connector 140 and, in some instances, the connector 142. The wires 137 and 148 (which are not shown in all of the Figures for clarity purposes) run through wire apertures 150 and small holes 152, which are formed in the base members 106, 126, 126’, 134 and 134’. Suitable temperature sensors and power control schemes that are based on a sensed temperature are disclosed in U.S. Patent Nos. 5,456,682, 5,582,609 and 5,755,715.

[0045] The actual number of temperature sensors 146 may be varied in order to suit particular applications. In the bi-polar arrangement illustrated in Figures 7a and 7b,
III. Tissue Cooling Apparatus

[0048] Energy transmission devices in accordance with the present inventions may also include apparatus that cools the tissue during tissue coagulation procedures. Examples of suitable cooling apparatus are illustrated in Figures 13-15. Such tissue cooling apparatus may also be used in conjunction with the exemplary devices illustrated in Figures 4, 5, 7a-12, 16a and 16b. The tissue cooling apparatus disclosed herein employ conductive fluid to cool tissue during coagulation procedures.

More specifically, and as described below and in U.S. application Serial No. 09/761,981, heat from the tissue being coagulated is transferred to ionic fluid to cool the tissue while energy is transferred from an electrode or other energy transmission device to the tissue through the fluid by way of ionic transport. The conductive fluid may be pumped through the tissue cooling apparatus (Figures 13 and 14) or the tissue cooling apparatus may be saturated with the fluid prior to use (Figure 15). In either case, cooling tissue during a coagulation procedure facilitates the formation of lesions that are wider and deeper than those that could be realized with an otherwise identical device which lacks tissue cooling apparatus.

[0049] Referring first to Figures 13 and 14, an exemplary tissue cooling apparatus 154 includes a nanoporous outer casing 156 through which ionic fluid (represented by arrows F) is transferred. The ionic fluid preferably flows from one longitudinal end of the tissue cooling apparatus 154 to the other. The outer casing 156 is secured to the base member 134 over the energy transmission devices 108 such that a fluid transmission space 158 is defined therebetween. More specifically, the proximal and distal ends of the outer casing 156 are secured to the base member 134 with anchoring devices (not shown) such as lengths of heat shrink tubing, Nitinol tubing or other mechanical devices that form an interference fit between the casing and the base member. Adhesive bonding is another method of securing the outer casing 156 to the base member 134. The fluid transmission space will typically be about 0.5 mm to about 2.0 mm high and slightly wider than the associated energy transmission device(s) 108.

[0050] The ionic fluid is supplied under pressure from a fluid source (not shown) by way of a supply line 160 and is returned to the source by way of a return line 162. The supply line 160 is connected to a fluid lumen 164 that runs from the proximal end of the base member 134 to the distal region of the outer casing 156. The fluid lumen 164 is connected to the fluid transmission space 158 by an aperture 166.

[0051] The electrically conductive ionic fluid preferably possesses a low resistivity to decrease ohmic losses, and thus ohmic heating effects, within the outer casing 156. The composition of the electrically conductive fluid can vary. In the illustrated embodiment, the fluid is a hypertonic saline solution, having a sodium chloride concentration at or near saturation, which is about 5% to about 25% weight by volume. Hypertonic saline solution has a relatively low resistivity of only about 5 ohm-cm, as compared to blood resistivity of about 150 ohm-cm and myocardial tissue resistivity of about 500 ohm-cm. Alternatively, the ionic fluid can be a hypertonic potassium chloride solution.

[0052] With respect to temperature and flow rate, a suitable inlet temperature for epicardial applications (the temperature will, of course, rise as heat is transferred to the fluid) is about 0 to 25°C with a constant flow rate of...
about 2 to 20 ml/min. The flow rate required for endocar- 
dial applications where blood is present would be about 
three-fold higher (i.e. 6 to 60 ml/min.). Should applica-
tions so require, a flow rate of up to 100 ml/min. may be 
employed. In a closed system where the fluid is stored 
in a flexible bag, such as the Viaflex® bag manufactured 
by Baxter Corporation, and heated fluid is returned to the 
bag, it has been found that a volume of fluid between 
about 200 and 500 ml within the bag will remain at room 
temperature (about 22°C) when the flow rate is between 
about 2 ml/min. and 20 ml/min. Alternatively, in an open 
system, the flexible bag should include enough fluid to 
complete the procedure. 160 ml would, for example, be 
required for a 20 minute procedure where the flow rate 
was 8 ml/min.

[0053] The fluid pressure within the outer casing 156 
should be about 30 mm Hg in order to provide a structure 
that will resiliently conform to the tissue surface in re-
sponse to a relatively small force normal to the tissue. 
Pressures above about 100 mm Hg will cause the outer 
casing 156 to become too stiff to properly conform to the 
tissue surface. For that reason, the flow resistance to and 
from the outer casing 156 should be relatively low.

[0054] The pores in the nanoporous outer casing 156 
allow the transport of ions contained in the fluid through 
the casing and into contact with tissue. Thus, when an 
energy transmission device 108 transmit RF energy into 
the ionic fluid, the ionic fluid establishes an electrically 
conductive path through the outer casing 156 to the tissue 
being coagulated. Regenerated cellulose membrane 
materials, typically used for blood oxygenation, dialysis 
or ultrafiltration, are a suitable nanoporous material for 
the outer casing 156. The thickness of the material should 
be about 0.05 mm to 0.13 mm. Although regenerated 
cellulose is electrically non-conductive, the relatively 
small pores of this material allow effective ionic transport 
in response to the applied RF field. At the same time, the 
relatively small pores prevent transfer of macromole-
cules through the pores, are also more likely to occur.

[0055] Hydro-Fluoro™ material, which is disclosed in 
U.S. Patent No. 6,395,325, is another material that may 
be used. Materials such as nylons (with a softening tem-
perature above 100°C), PTFE, PEI and PEEK that have 
nanopores created through the use of lasers, electrostat-
ic discharge, ion beam bombardment or other processes 
may also be used. Such materials would preferably in-
clude a hydrophilic coating. Nanoporous materials may 
also be fabricated by weaving a material (such as nylon, 
polyester, polyethylene, polypropylene, fluorocarbon, 
fine diameter stainless steel, or other fiber) into a mesh 
having the desired pore size and porosity. These mate-
rials permit effective passage of ions in response to the 
applied RF field. However, as many of these materials 
possess larger pore diameters, pressure driven liquid 
perfusion, and the attendant transport of macromole-
cules through the pores, are also more likely to occur.

[0056] The electrical resistivity of the outer casing 156 
will have a significant influence on lesion geometry and 
controllability. Low-resistivity (below about 500 ohm-cm) 
requires more RF power and results in deeper lesions, 
while high-resistivity (at or above about 500 ohm-cm) 
generates more uniform heating and improves control-
lability. Because of the additional heat generated by the 
increased body resistivity, less RF power is required to 
reach similar tissue temperatures after the same interval 
of time. Consequently, lesions generated with high-re-
sistivity structures usually have smaller depth. The elec-
trical resistivity of the outer casing can be controlled by 
specifying the pore size of the material, the porosity of 
the material, and the water adsorption characteristics 
(hydrophilic versus hydrophobic) of the material. A de-
tailed discussion of these characteristics is found in U.S. 
Patent No. 5,961,513. A suitable electrical resistivity for 
epidermal and endocardial lesion formation is about 1 to 
3000 ohm-cm measured wet.

[0057] Generally speaking, low or essentially no liquid 
perfusion through the nanoporous outer casing 156 is 
preferred. When undisturbed by attendant liquid per-
fusion, ionic transport creates a continuous virtual elec-
trode at the tissue interface. The virtual electrode effi-
ciently transfers RF energy without need for an electric-
cally conductive metal surface.

[0058] Pore diameters smaller than about 0.1 pm re-
tain macromolecules, but allow ionic transport through 
the pores in response to the applied RF field. With smaller 
pore diameters, pressure driven liquid perfusion through 
the pores is less likely to accompany the ionic transport, 
unless relatively high pressure conditions develop within 
the outer casing 156. Larger pore diameters (up to 8 pm) 
can also be used to permit ionic current flow across the 
membrane in response to the applied RF field. With larger 
pore diameters, pressure driven fluid transport across 
the membrane is much higher and macromolecules 
(such as protein) and even small blood cells (such as 
platelets) could cross the membrane and contaminate 
the inside of the probe. Red blood cells would normally 
not cross the membrane barrier, even if fluid perfusion 
across the membrane stops. On balance, a pore diam-
eter of 1 to 5 μm is suitable for epicardial and endocardial 
lesion formation. Where a larger pore diameter is em-
ployed, thereby resulting in significant fluid transfer 
through the porous region, a saline solution having a so-
dium chloride concentration of about 0.9% weight by vol-
ume would be preferred.

[0059] With respect to porosity, which represents the 
volumetric percentage of the outer casing 156 that is 
composed of pores and not occupied by the casing ma-
terial, the magnitude of the porosity affects electrical re-
sistance. Low-porosity materials have high electrical re-
sistivity, whereas high-porosity materials have low elec-
trical resistivity. The porosity of the outer casing 156 
should be at least 1% for epicardial and endocardial ap-
lications employing a 1 to 5 pm pore diameter.
[0060] Turning to water absorption characteristics, hydrophilic materials are generally preferable because they possess a greater capacity to provide ionic transfer of RF energy without significant liquid flow through the material.

[0061] The exemplary tissue cooling apparatus 168 illustrated in Figure 15 consists of a wettable fluid retention element 170 that is simply saturated with ionic fluid (such as saline) prior to use, as opposed to having the fluid pumped through the apparatus in the manner described above with reference to Figures 13 and 14. The energy transmission device(s) 108 are carried within the fluid retention element 170. Suitable materials for the fluid retention element 170 include biocompatible fabrics commonly used for vascular patches (such as woven Dacron®), open cell foam materials, hydrogels, nanoporous balloon materials (with very slow fluid delivery to the surface), and hydrophilic nanoporous materials. The effective electrical resistivity of the fluid retention element 170 when wetted with 0.9% saline (normal saline) should range from about 1 Ω-cm to about 2000 Ω-cm. A preferred resistivity for epicardial and endocardial procedures is about 1000 Ω-cm.

IV. Probe Support Devices

[0062] Probe support devices in accordance with a present invention may be used to covert a conventional clamp, or a clamp in accordance with the inventions described in Section V below, into a tissue coagulation device by securing one or more conventional catheters, surgical probes, or other apparatus that support energy transmission devices, to the clamp. Although the configuration of the probe support devices may vary from application to application to suit particular situations, the exemplary probe support devices are configured such that the probes being supported will abut one another in the same manner as the inserts 28 and 30 (Figures 1-3) when the associated clamp is in the closed orientation. Such an arrangement will allow the energy transmission devices on the probes to face one another in the manner similar to that described in Section I above.

[0063] As illustrated for example in Figures 17 and 18, a probe support device 172 in accordance with one embodiment of a present invention includes a base member 174, a slot 176 configured to receive an electrode support device such as a catheter or surgical probe, and a plurality of mating structures 178 that mechanically engage a clamp member. The exemplary base member 174 is preferably formed from a soft, resilient, low durometer material that is electrically insulating. Suitable materials include polyurethane, silicone and polyurethane/silicone blends having a hardness of between about 20 Shore D and about 72 Shore D.

[0064] The size and shape of the slot 176 will, of course, depend on the size and shape of the probe that it is holding. Many probes are generally cylindrical in shape and, according, the exemplary slot 176 has a corresponding arcuate cross-sectional shape. The arc is preferably greater than 180 degrees so that the base member 174 will deflect when a probe is inserted into the slot 176 and then snap back to hold the probe in place.

[0065] The exemplary mating structures 178, which are preferably integral with the base member 174 and formed from the same resilient material, include a relatively narrow portion 180 and a relatively wide portion 182. The relatively narrow portions 180 are approximately the same size as the clamp member apertures 36 (Figure 3) and the relatively wide portions 182 are slightly larger than the clamp member apertures. A removable connection is made by urging the mating structures 178 into one end of the apertures 36, thereby deforming the relatively wide portions 182, and then urging the base members 174 against the clamp member until the relatively wide portions exit through the other end of the apertures and reassume their original shape.

[0066] Turning to Figures 19 and 20, a pair of the exemplary probe support devices 172 may be used in conjunction with a pair of probes 184 to convert the clamp 10 (which has had the inserts 28 and 30 removed) into a bi-polar tissue coagulation device. Although the present inventions are not limited to use with a particular type of probe, each probe 184 in the exemplary implementation includes a shaft 186, a plurality of spaced electrodes 188, and a plurality of temperature sensors (not shown) respectively associated with the electrodes. Once the probe support devices 172 have been secured to the clamp members 22 and 24, the probes 184 may be snapped into the slots 176 by moving the probes from the dash line positions illustrated in Figure 19 to the solid line positions. One of the probes 184 may be connected to the output connector of an ESU, while the other probe may be connected to the return connector to complete the bi-polar arrangement.

[0067] Another exemplary probe support device 190 is illustrated in Figures 21 and 22. The probe support device 190 is similar to the probe support device 172 illustrated in Figures 17 and 18 and similar structural elements are represented by similar reference numerals. The exemplary probe support device 190 may also be used in the manner described above with reference to Figures 19 and 20. Here, however, the mating structures 178 have been eliminated and the base member 172 is provided with a longitudinally extending aperture 192 into which one of the clamp members 22 and 24 may be inserted.

[0068] The aperture 192 should be sized and shaped such that the base member 174 will be forced to stretch when one of the clamp members 22 and 24 is inserted. If, for example, the apertures 192 have the same cross-sectional shape as the clamp members 22 and 24 (e.g. both are elliptical), then the apertures should be slightly smaller in their cross-sectional dimensions than the corresponding clamp members. The stretching of base member 174 creates a tight interference fit between the base member and the clamp member. Additionally, al-
though the aperture 192 has a semi-circular cross-section in the exemplary embodiment, the apertures may have a round, rectangular, square or elliptical cross-section, or define any other cross-sectional shape, depending on the particular application.

Alternatively, and as illustrated for example in Figure 23, in those instances where the clamp members include longitudinally extending slots instead of apertures (such as the slots 38 described above with reference to Figure 12), the probe support device 172 may be provided with a longitudinally extending mating structure 194 that extends outwardly from the base member 174. The longitudinally extending mating structure 194, which is preferably integral with the base member 174 and formed from the same resilient material, is sized to create an interference fit with a slot. Still another alternative is to provide the clamp members with one or more small mating structures that are received within apertures or slots formed in the base member 174.

An exemplary probe support device 196 that may be used in conjunction with a probe 184 to convert the clamp 10 (which has had the inserts 28 and 30 removed) into a uni-polar tissue coagulation device not according to the invention, is illustrated in Figures 24 and 25. Although the configuration of the probe support device 196 may vary from application to application to suit particular situations, the probe support device in the exemplary embodiment is configured such that it will abut each of the clamp members 22 and 24 when the clamp is in the closed orientation illustrated in Figure 25.

The exemplary probe support device 196 includes a base member 198, a slot 200 configured to receive a probe 184 or other electrode supporting device, and a plurality of mating structures 178 that mechanically engage a clamp members 22 and 24 in the manner described above. The exemplary base member 198 is preferably formed from a soft, resilient, low durometer material that is electrically insulating. Suitable materials include polyurethane, silicone and polyurethane/silicone blends having a hardness of between about 20 Shore D and about 72 Shore D. The size and shape of the slot 200 will depend on the size and shape of the probe that it is intended to hold, as is described above with reference to slot 200. The aperture 208 should be sized and shaped such that the base member 204 will be forced to stretch when the clamp members 22 and 24 are inserted with the clamp 10 in a closed orientation. The stretching creates a tight interference fit between the base member 204 and the clamp members 22 and 24. Additionally, although the aperture 208 has an elliptical cross-section in the exemplary embodiment, the aperture may have a round, rectangular, square or semi-circular cross-section, or define any other cross-sectional shape, depending on the particular application.

The length of the base members in the exemplary probe support devices will vary according to the intended application. In the area of cardiovascular treatments, it is anticipated that suitable lengths will range from, but are not limited to, about 3 cm to about 10 cm.

V. Clamp With Malleable Clamp Members

This portion of the specification refers to rigid and malleable structures. A rigid structure is a structure than cannot be readily bent by a physician. A malleable structure can be readily bent by the physician to a desired shape, without springing back when released, so that it will remain in that shape during the surgical procedure. Thus, the stiffness of a malleable structure must be low enough to allow the structure to be bent, but high enough to resist bending when the forces associated with a surgical procedure are applied to the structure. Rigid structures are preferably formed from stainless steel, while malleable structure are preferably formed from annealed stainless steel or titanium. Additional information concerning malleable structures may be found in U.S. Patent No. 6,142,994.

As illustrated for example in Figure 27, a clamp 210 in accordance with a preferred embodiment of a present invention includes a pair of malleable clamp members 212 and 214. The malleable clamp members 212 and 214 are carried at the distal ends of a pair of arms 216 and 218. The arms 216 and 218 are pivotally secured to one another by a pin 220 to allow the clamp members 212 and 214 to be moved towards and away from one another between opened and closed positions. The arms 216 and 218 are preferably formed from rigid material, but may also be malleable if desired. When rigid, the arms 216 and 218 may be linear or have a preformed curvature.

A pair of handles 222 and 224 are mounted on the proximal ends of the arms 216 and 218. A locking device 226 locks the clamp 210 in the closed orientation
illustrated in Figure 27. The locking device 226 also prevents the clamp members 212 and 214 from coming any closer to one another than is illustrated in Figure 27, thereby defining a predetermined spacing between the clamp members.

[0077] The malleability of the clamp members 212 and 214 allows them to be re-shaped by the physician as needed for particular procedures and body structures. As such, a single clamp 210 is capable of taking the place of a number of conventional clamps with rigid clamp members. In some implementations, the clamp members 212 and 214 will be more malleable (i.e. easier to bend) at their distal end than at their proximal end. This may be accomplished by gradually decreasing the cross-sectional area of each clamp member 212 and 214 from the proximal end to the distal end.

[0078] The clamp members 212 and 214 may also be provided with holes 228 (Figure 31) that allow soft deformable inserts, such as the conventional inserts 28 and 30 described above with reference to Figures 1-3. The exemplary clamp 210 may also be used in conjunction with the energy transmission assemblies, probe support devices, and probes described in Sections I-IV above.

[0079] There will be many instances where it will be important to maintain the predefined spacing between the malleable clamp members 212 and 214 during the bending process in order to insure that the predefined spacing will remain when the bending process is complete. To that end, the exemplary clamp 210 is provided with a malleable insert 230 that is sized and shaped (rectangular in the exemplary implementation) to be held between the malleable clamp members 212 and 214 when the clamp is closed and locked. The friction between the clamp members 212 and 214 and insert 230 will hold the insert in place during bending. Nevertheless, if desired, the insert 230 may be provided with small protrusions that will be received by the holes 228. The malleable insert 230, which is preferably formed from the same material as the malleable clamp members 212 and 214, will bend with the clamp members during the bending process, thereby maintaining the predefined spacing. [Note Figure 32.]

[0080] The exemplary mandrel 232 illustrated in Figures 28 and 29 may be used to bend the malleable clamp members 212 and 214. The exemplary mandrel 232 includes a base 234 and a pair of cylindrical posts 236 and 238. Posts of other shapes, such as elliptical posts, may also be employed to achieve particular bends. The mandrel 232 should also be formed from material that is harder than the malleable clamp members 212 and 214, such as stainless steel or titanium.

[0081] The exemplary mandrel 232 may be used to bend the malleable clamp members 212 and 214 in the manners illustrated in Figures 30 and 31. Referring first to Figure 30, once the malleable clamp members 212 and 214 and malleable insert 230 have been placed between the posts 236 and 238, the clamp 210 may be rotated in the direction of the arrow (or in the opposite direction) until the clamp members 212 and 214 are bent the desired amount. The clamp 210 may then moved longitudinally and the bending process repeated until the desired bend, such as the exemplary bend illustrated in Figure 32, has been achieved. Alternatively, or in addition, the clamp 210 can be rotated about its longitudinal axis and bent in other planes, as is illustrated for example in Figures 31 and 33. It should also be noted that, if desired, the malleable clamp members 212 and 214 may be bent independently of one another and/or into different shapes. Preferably, the physician will simply place the mandrel 232 on a suitable surface and press down the base 234 during a bending procedure. Alternatively, structure may be provided to secure the mandrel 232 to the surface.

[0082] Another example of a clamp in accordance with a preferred embodiment of a present invention is generally represented by reference numeral 240 in Figure 34. Clamp 240 is similar to clamp 210 and similar elements are represented by similar reference numerals. The exemplary clamp 240 includes malleable clamp members 212 and 214, pivotable arms 216 and 218, handles 222 and 224, and a locking device 226. Here, however, the arms 216 and 218 are pivotally carried by one end of an elongate housing 242 and the malleable clamp members 212 and 214 are carried by a pair of supports 244 and 246 that are pivotally carried the other end of the housing. A suitable mechanical linkage (not shown) located within the housing causes the supports 244 and 246 (and clamp members 212 and 214) to move relative to one another in response to movement of the arms 216 and 218. The housing 242 may be rigid or malleable.

[0083] The present clamps with malleable clamp members (such as exemplary clamps 210 and 240) have a wide variety of applications. One example is the formation of transmural epicardial lesions to isolate the sources of focal (or ectopic) atrial fibrillation and, more specifically, the creation of transmural lesions around the pulmonary veins. Energy transmission devices may be permanently affixed to the malleable clamp members. Energy transmission devices may also be added using the structures described in Sections I-IV above and the clamp may be used as a clamp or as a surgical probe, depending on the structure being used in combination with the clamp. Access to the heart may be obtained via a thoracotomy, thoracostomy or median sternotomy. Ports may also be provided for cameras and other instruments.

[0084] Lesions may be created around the pulmonary veins individually or, alternatively, lesions may be created around pairs of pulmonary veins. For example, a first transmural epicardial lesion may be created around the right pulmonary vein pair and a second transmural epicardial lesion may be created around the left pulmonary vein pair. Thereafter, if needed, a linear transmural epicardial lesion may be created between the right and left pulmonary vein pairs. A linear transmural lesion that extends from the lesion between the right and left pulmonary vein pairs to the left atrial appendage may also be
formed. Alternatively, a single lesion may be formed around all four of the pulmonary veins.

Although the present inventions have been described in terms of the preferred embodiments above, numerous modifications and/or additions to the above-described preferred embodiments would be readily apparent to one skilled in the art. It is intended that the scope of the present inventions extend to all such modifications and/or additions and that the scope of the present inventions is limited solely by the claims set forth below.

Claims

1. A surgical system comprising a clamp (10, 210 or 240) including first and second clamp members (22/24, 22'24' or 212/214) defining respective longitudinal axes and movement apparatus (12/14 or 216/218) that moves at least one of the first and second clamp members relative to the other of the first and second clamp members such that the clamp has an open state and a closed state and an apparatus for use with the clamp including a first base member (106, 126', 134 or 134') configured to be removable secured to the first clamp member, at least one energy transmission device (108) carried by the first base member, a second base member (106, 126', 134 or 134') configured to be removable secured to the second clamp member, and at least one energy transmission device (108) carried by the second base member, characterized by a first electrical connector (140) operably connected to the first energy transmission device (108) and defining a first connector configuration; and a second electrical connector (142) operably connected to the second energy transmission device (108) and defining a second connector configuration, the second connector configuration being different than the first connector configuration.

2. A surgical system as claimed in claim 1, wherein at least one of the first and second clamp members (212/214) is malleable.

3. A surgical system as claimed in claim 1, wherein the movement apparatus (12/14 or 216/218) comprises first and second arms pivotably secured to one another.

4. A surgical system as claimed in claim 1, wherein the base member includes a longitudinally extending aperture (110 or 136) configured to receive the at least one of the first and second clamp members (22/24 or 212/214).

5. A surgical system as claimed in claim 4, wherein the at least one of the first and second clamp members (22/24 or 212/214) and the longitudinally extending aperture (110 or 136) are respectively sized and shaped such that the base member will stretch when the at least one of the first and second clamp members is inserted into the longitudinally extending aperture.

6. A surgical system as claimed in claim 1, wherein the base member (126') includes at least one longitudinally extending base member mating structure (130) configured to mate with a longitudinally extending slot (38) in the at least one of the first and second clamp members (22'/24').

7. A surgical system as claimed in claim 1, wherein the at least one energy transmission device (108) comprises an electrode.

8. A surgical system as claimed in claim 1, wherein the base member (106, 126', 134 or 134') is at least one of electrically insulating and resilient.

Patentansprüche


2. Chirurgisches System, wie in Anspruch 1 bean-
1. Chirurgisches System, wie in Anspruch 1 beansprucht, wobei die Bewegungsvorrichtung (12/14 oder 216/218) einen ersten und einen zweiten Arm, die schwenkbar aneinander befestigt sind, umfasst.

2. Chirurgisches System, wie in Anspruch 1 beansprucht, wobei die Basiselemente (126', 126, 134 oder 134') mindestens eines des ersten und des zweiten Klemmelements (212/214) verformbar ist.

3. Chirurgisches System, wie in Anspruch 1 beansprucht, wobei die Bewegungsvorrichtung (12/14 oder 216/218) einen ersten und einen zweiten Arm, die schwenkbar aneinander befestigt sind, umfasst.

4. Chirurgisches System, wie in Anspruch 1 beansprucht, wobei das Basiselement (126') mindestens eine in Längsrichtung erstreckende Öffnung (110 oder 136) umfasst, die dazu konfiguriert ist, das mindestens eine des ersten und des zweiten Klemmelements (22/24 oder 212/214) aufzunehmen.

5. Chirurgisches System, wie in Anspruch 4 beansprucht, wobei das Basiselement (126') mindestens eine in Längsrichtung erstreckende Basiselement-Passtruktur (130) beinhaltet, die dazu konfiguriert ist, mit einem sich in Längsrichtung erstreckenden Schlitz (38) in dem mindestens einen des ersten und des zweiten Klemmelements (22'/24') zusammenzupassen.

6. Chirurgisches System, wie in Anspruch 1 beansprucht, wobei das Basiselement (126') mindestens eine in Längsrichtung erstreckende Basiselement-Passtruktur (130) beinhaltet, die dazu konfiguriert ist, mit einem sich in Längsrichtung erstreckenden Schlitz (38) in dem mindestens einen des ersten und des zweiten Klemmelements (22'/24') zusammenzupassen.

7. Chirurgisches System, wie in Anspruch 1 beansprucht, wobei die mindestens eine Energieübertragungsvorrichtung (108) eine Elektrode umfasst.


Revendications

1. Système chirurgical comprenant un clamp (10, 210 ou 240) englobant un premier et un deuxième membre faisant office de clamp (22/24, 22'/24' ou 212/214) définissant des axes longitu- dinaux respectifs et un appareil de mise en mouvement (12/14 ou 216/218) qui déplace au moins un premier et deuxième membres faisant office de clamp l’un par rapport à l’autre des premier et deuxième membres faisant office de clamp de telle sorte que le clamp possède un état ouvert et un état fermé, et un appareil à utiliser avec le clamp, englobant un premier membre faisant office de base (106, 126', 134 ou 134') configuré pour venir se fixer de manière amovible au premier membre faisant office de clamp, au moins un dispositif de transmission d’énergie (108) supporté par le premier membre faisant office de base, un deuxième membre faisant office de base (106, 126', 134 ou 134') configuré pour venir se fixer de manière amovible au deuxième membre faisant office de clamp, et au moins un dispositif de transmission d’énergie (108) supporté par le deuxième membre faisant office de base, caractérisé par un premier raccord électrique (140) en liaison opérante avec le premier dispositif de transmission d’énergie (108) et définissant une première configuration de raccord ; et un deuxième raccord électrique (142) en liaison opérante avec le deuxième dispositif de transmission d’énergie (108) et définissant une deuxième configuration de raccord, la deuxième configuration de raccord étant différente de la première configuration de raccord.

2. Système chirurgical selon la revendication 1, dans lequel le moins un membre parmi le premier et le deuxième membre faisant office de clamp (212/214) est malléable.

3. Système chirurgical selon la revendication 1, dans lequel l’appareil de mise en mouvement (12/14 ou 216/218) comprend des premier et deuxième bras fixés en pivotement l’un à l’autre.

4. Système chirurgical selon la revendication 1, dans lequel le membre faisant office de base englobe un orifice s’étendant en direction longitudinale (110 ou 136) configuré pour recevoir ledit au moins un membre parmi les premier et deuxième membres faisant office de clamp (22/24 ou 212/214).

5. Système chirurgical selon la revendication 4, dans lequel ledit au moins un membre parmi les premier et deuxième membres faisant office de clamp (22/24 ou 212/214) et l’orifice s’étendant en direction longitudinale (110 ou 136) sont dimensionnés et configurés de manière respective de telle sorte que le membre faisant office de base sera étiré lorsque ledit au moins un membre des premier et deuxième membres de clamp est inséré dans l’orifice s’étendant en direction longitudinale.

6. Système chirurgical selon la revendication 1, dans lequel le membre faisant office de base (126') englobe au moins une structure correspondant au membre faisant office de base s’étendant en direction longitudinale (130), configurée pour correspondre à une fente s’étendant en direction longitudinale (38) pratiquée dans ledit au moins un des premier et deuxième membres faisant office de clamps (22'/24').
7. Système chirurgical selon la revendication 1, dans lequel ledit au moins un dispositif de transmission d’énergie (108) comprend une électrode.

8. Système chirurgical selon la revendication 1, dans lequel le membre faisant office de base (106, 126", 134 ou 134") représente au moins un membre résilient et procurant une isolation électrique.