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(54) **SURGICAL SYSTEM WITH USER ADAPTABLE TECHNIQUES BASED ON TISSUE TYPE**

CHIRURGISCHES SYSTEM MIT BENUTZERSEITIG ANPASSBAREN VERFAHREN AUSGEHEND  
VOM GEWEBETYP

SYSTÈME CHIRURGICAL À TECHNIQUES ADAPTABLES PAR L'UTILISATEUR SUR LA BASE DU  
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**Description****TECHNICAL FIELD**

**[0001]** The present disclosure generally relates to ultrasonic surgical systems and, more particularly, to ultrasonic and electrosurgical systems that allows surgeons to perform cutting and coagulation and adapt and customize techniques for performing such procedures based on the type of tissue being treated.

**BACKGROUND**

**[0002]** Ultrasonic surgical instruments are finding increasingly widespread applications in surgical procedures by virtue of the unique performance characteristics of such instruments. Depending upon specific instrument configurations and operational parameters, ultrasonic surgical instruments can provide substantially simultaneous cutting of tissue and hemostasis by coagulation, desirably minimizing patient trauma. The cutting action is typically realized by an end effector, or blade tip, at the distal end of the instrument, which transmits ultrasonic energy to tissue brought into contact with the end effector. Ultrasonic instruments of this nature can be configured for open surgical use, laparoscopic, or endoscopic surgical procedures including robotic-assisted procedures.

**[0003]** Some surgical instruments utilize ultrasonic energy for both precise cutting and controlled coagulation. Ultrasonic energy cuts and coagulates by vibrating a blade in contact with tissue. Vibrating at high frequencies (e.g., 55,500 times per second), the ultrasonic blade denatures protein in the tissue to form a sticky coagulum. Pressure exerted on tissue with the blade surface collapses blood vessels and allows the coagulum to form a hemostatic seal. The precision of cutting and coagulation is controlled by the surgeon's technique and adjusting the power level, blade edge, tissue traction, and blade pressure.

**[0004]** Electrosurgical devices for applying electrical energy to tissue in order to treat and/or destroy the tissue are also finding increasingly widespread applications in surgical procedures. An electrosurgical device typically includes a hand piece, an instrument having a distally-mounted end effector (e.g., one or more electrodes). The end effector can be positioned against the tissue such that electrical current is introduced into the tissue. Electrosurgical devices can be configured for bipolar or monopolar operation. During bipolar operation, current is introduced into and returned from the tissue by active and return electrodes, respectively, of the end effector. During monopolar operation, current is introduced into the tissue by an active electrode of the end effector and returned through a return electrode (e.g., a grounding pad) separately located on a patient's body. Heat generated by the current flowing through the tissue may form hemostatic seals within the tissue and/or between tissues and thus may be particularly useful for sealing blood vessels, for example. The end effector of an electrosurgical device also may include a cutting member that is movable relative to the tissue and the electrodes to transect the tissue.

**[0005]** Electrical energy applied by an electrosurgical device can be transmitted to the instrument by a generator in communication with the hand piece. The electrical energy may be in the form of radio frequency ("RF") energy. RF energy is a form of electrical energy that may be in the frequency range of 200 kilohertz (kHz) to 1 megahertz (MHz). In application, an electrosurgical device can transmit low frequency RF energy through tissue, which causes ionic agitation, or friction, in effect resistive heating, thereby increasing the temperature of the tissue. Because a sharp boundary is created between the affected tissue and the surrounding tissue, surgeons can operate with a high level of precision and control, without sacrificing un-targeted adjacent tissue. The low operating temperatures of RF energy is useful for removing, shrinking, or sculpting soft tissue while simultaneously sealing blood vessels, RF energy works particularly well on connective tissue, which is primarily comprised of collagen and shrinks when contacted by heat.

**[0006]** The RF energy may be in a frequency range described in EN 60601-2-2:2009+A11:2011, Definition 201.3.218 - HIGH FREQUENCY. For example, the frequency in monopolar RF applications may be typically restricted to less than 5MHz. However, in bipolar RF applications, the frequency can be almost anything. Frequencies above 200 kHz can be typically used for monopolar applications in order to avoid the unwanted stimulation of nerves and muscles that would result from the use of low frequency current. Lower frequencies may be used for bipolar applications if the risk analysis shows the possibility of neuromuscular stimulation has been mitigated to an acceptable level. Normally, frequencies above 5 MHz are not used in order to minimize the problems associated with high frequency leakage currents. Higher frequencies may, however, be used in the case of bipolar applications. It is generally recognized that 10 mA is the lower threshold of thermal effects on tissue.

**[0007]** A challenge of using these medical devices is the inability to control and customize the power output depending on the type of tissue being treated by the devices. It would be desirable to provide a surgical instrument that overcomes some of the deficiencies of current instruments. The surgical system described herein overcomes those deficiencies.

**[0008]** US2009254080 (A1) relates to a surgical operation apparatus which includes a handpiece to be held by an operator, and an apparatus main body to be connected with the handpiece, wherein the handpiece includes a high-frequency treatment unit to treat a living tissue by a high-frequency current, an ultrasonic treatment unit to treat a living

tissue by an ultrasonic vibration, and a sensor unit to perform detection for calculation of an impedance of a living tissue  
**[0009]** US2015141981 (A1) relates to a surgical instrument system including an end effector, a generator, and a controller. The end effector includes an ultrasonic blade and at least one electrode surface. The generator provides power to the end effector.

**[0010]** EP2866698 (A1) relates to surgical instruments comprising an end effector, a shaft and a jaw assembly. The end effector may comprise an ultrasonic blade extending distally substantially parallel to a longitudinal axis.

**[0011]** US2013282038 (A1) relates to an apparatus which comprises an end effector, a body assembly, a power source, and a control module. The end effector is operable for use in a surgical procedure and can deliver energy to a surgical site. The end effector comprises at least one sensor.

**[0012]** US2012136386 (A1) relates to a heat-generating device which includes a heat-generating unit 5a, having a resin member 10a which generates heat upon application of vibration and a vibration unit 11a which imparts vibration to the resin member 10a, and a heat generation control unit 6a which, by controlling the vibration generated by the vibration unit 11a, controls the heat generation of the heat-generating unit 5a in such a manner that the heat-generating unit 5a has a prescribed temperature.

## SUMMARY

**[0013]** The present invention is defined by the independent claims. Further embodiments of the invention are described in the dependent claims.

**[0014]** In addition to the foregoing, various other method and/or system and/or program product aspects of the disclosure are set forth and described in the teachings such as text (e.g., claims and/or detailed description) and/or drawings of the present disclosure.

**[0015]** The foregoing is a summary and thus may contain simplifications, generalizations, inclusions, and/or omissions of detail; consequently, those skilled in the art will appreciate that the summary is illustrative only. Other aspects, features, and advantages of the devices and/or processes and/or other subject matter described herein will become apparent in the teachings set forth herein.

**[0016]** In one or more various aspects, related systems include but are not limited to circuitry and/or programming for effecting herein-referenced method aspects; the circuitry and/or programming can be virtually any combination of hardware, software, and/or firmware configured to affect the herein-referenced method aspects depending upon the design choices of the system designer. In addition to the foregoing, various other method and/or system aspects are set forth and described in the teachings such as text (e.g., claims and/or detailed description) and/or drawings of the present disclosure.

**[0017]** Further, it is understood that any one or more of the following-described forms, expressions of forms, examples, can be combined with any one or more of the other following-described forms, expressions of forms, and examples.

**[0018]** The foregoing summary is illustrative only. In addition to the illustrative aspects, embodiments, and features described above, further aspects, embodiments, and features will become apparent by reference to the drawings and the following detailed description. The present invention is defined by the independent claims. Further embodiments of the invention are described in the dependent claims.

## FIGURES

**[0019]** The novel features of the described forms are set forth with particularity in the appended claims. The described forms, however, both as to organization and methods of operation, may be best understood by reference to the following description, taken in conjunction with the accompanying drawings in which:

FIG. 1 illustrates one aspect of a surgical system comprising a generator and various surgical instruments usable therewith;

FIG. 2 is a diagram of one aspect of the ultrasonic surgical instrument of FIG. 16;

FIG. 3 is a diagram of one aspect of the surgical system of FIG. 16;

FIG. 4 is a model illustrating one aspect of a motional branch current;

FIG. 5 is a structural view of one aspect of a generator architecture;

FIG. 6 illustrates one aspect of a drive system of a generator, which creates the ultrasonic electrical signal for driving an ultrasonic transducer;

FIG. 7 illustrates one aspect of a drive system of a generator comprising a tissue impedance module;

FIG. 8 illustrates one aspect of a generator for delivering multiple energy modalities to a surgical instrument;

FIG. 9 is an example graph of two waveforms of energy from one aspect of a generator;

FIG. 10 is an example graph of the sum of the waveforms of FIG. 9;

FIG. 11 is an example graph of sum of the waveforms of FIG. 9 with the RF waveform dependent on the ultrasonic

waveform;

FIG. 12 is an example graph of the sum of the waveforms of FIG. 9 with the RF waveform being a function of the ultrasonic waveform;

FIG. 13 is an example graph of a complex RF waveform;

FIG. 14 illustrates one aspect of an end effector comprising RF data sensors located on the clamp arm;

FIG. 15 illustrates one aspect of the flexible circuit shown in FIG. 14 in which the sensors may be mounted to or formed integrally therewith;

FIG. 16 is a cross-sectional view of the flexible circuit shown in FIG. 15;

FIG. 17 illustrates one aspect of a segmented flexible circuit configured to fixedly attach to a clamp arm of an end effector;

FIG. 18 illustrates one aspect of a segmented flexible circuit configured to mount to a clamp arm of an end effector;

FIG. 19 illustrates one aspect of an end effector configured to measure a tissue gap  $G_T$ ;

FIG. 20 illustrates one aspect of a left-right segmented flexible circuit;

FIG. 21 illustrates one aspect of an end effector comprising segmented flexible circuit as shown in FIG. 20;

FIG. 22 illustrates the end effector shown in FIG. 21 with the clamp arm clamping tissue between the clamp arm and the ultrasonic blade;

FIG. 23 illustrates graphs of energy applied by the right and left side of an end effector based on locally sensed tissue parameters;

FIG. 24 illustrates a graph depicting one aspect of adjustment of threshold due to the measurement of a secondary tissue parameter such as continuity, temperature, pressure, and the like;

FIG. 25 is a cross-sectional view of one aspect of a flexible circuit comprising RF electrodes and data sensors embedded therein;

FIG. 26 is a cross-sectional view of one aspect of an end effector configured to sense force or pressure applied to tissue located between a clamp arm and an ultrasonic blade;

FIG. 27 is a schematic diagram of one aspect of a signal layer of a flexible circuit;

FIG. 28 is a schematic diagram of sensor wiring for the flexible circuit shown in FIG. 27;

FIG. 29 is a schematic diagram of one aspect of an RF energy drive circuit;

FIG. 30 is a graphical representation of measuring tissue gap at a preset time;

FIG. 31 is a time to preset force versus time graph for thin, medium, and thick tissue types;

FIG. 32 is a graphical depiction of a graph of three curves, where the first curve represents power (P), voltage ( $V_{RF}$ ), and current ( $I_{RF}$ ) versus tissue impedance (Z), the second curve and third curve represent tissue impedance (Z) versus time (t);

FIG. 33 is a plan view of one aspect of an end effector;

FIG. 34 is a side view of the end effector shown in FIG. 33 with a partial cut away view to expose the underlying structure of the clamp arm and an ultrasonic blade;

FIG. 35 is partial sectional view of the end effector shown in FIGS. 33, 34 to expose the ultrasonic blade and right and left electrodes, respectively;

FIG. 36 is a cross-sectional view taken at section 36--36 of the end effector shown in FIG. 33;

FIG. 37 is cross-sectional view taken at section 37--37 of the end effector shown in FIG. 33;

FIG. 38 is a cross-sectional view taken at section 36--36 of the end effector shown in FIG. 33, except that the ultrasonic blade has a different geometric configuration;

FIG. 39 is cross-sectional view taken at section 37--37 of the end effector shown in FIG. 33, except that the ultrasonic blade has a different geometric configuration;

FIG. 40 is a cross-sectional view taken at section 36--36 of the end effector shown in FIG. 33, except that the ultrasonic blade has a different geometric configuration;

FIG. 41 is cross-sectional view taken at section 37--37 of the end effector shown in FIG. 33, except that the ultrasonic blade has a different geometric configuration;

FIG. 42A is a graphical representation of one aspect of a medical device surrounding tissue;

FIG. 42B is a graphical representation of one aspect of a medical device compressing tissue;

FIG. 43A is a graphical representation of one aspect of a medical device compressing tissue;

FIG. 43B also depicts example forces exerted by one aspect of an end-effector of a medical device compressing tissue;

FIG. 44 illustrates a logic diagram of one aspect of a feedback system;

FIG. 45 is a graph of power versus force as measured with a plurality of plot points for various tissue types;

FIG. 46 is another graph of power versus force as measured with a plurality of plot points for various tissue types;

FIG. 47 is a logic flow diagram of one aspect of dynamically changing the energy delivered to a surgical instrument based on a determination of tissue type being treated by the instrument;

FIG. 48 is a logic flow diagram of one aspect of dynamically changing the energy delivered to a surgical instrument

based on a determination of tissue type being treated by the instrument;

FIG. 49 is a logic flow diagram of one aspect of a method of dynamically changing the energy delivered to a surgical instrument based on a determination of the hydration level of tissue being treated by a surgical instrument;

FIG. 50 is a logic flow diagram of one aspect of a method of dynamically changing energy being delivered from a generator based on the type of tissue being treated by a surgical instrument and various characteristics of the tissue;

FIG. 51 is a logic flow diagram of one aspect of a technique for dynamically changing the energy delivered from a generator based on aperture defined by the end effector and energy parameters;

FIG. 52 is a logic flow diagram of one aspect of a technique for dynamically changing the energy delivered from a generator based on aperture defined by the end effector and energy parameters; and

FIG. 53 is a logic flow diagram of one aspect of a dynamic tissue sensing technique.

## DESCRIPTION

**[0020]** Before explaining various forms of surgical instruments in detail, it should be noted that the illustrative forms are not limited in application or use to the details of construction and arrangement of parts illustrated in the accompanying drawings and description. The illustrative forms may be implemented or incorporated in other forms, variations and modifications, and may be practiced or carried out in various ways. Further, unless otherwise indicated, the terms and expressions employed herein have been chosen for the purpose of describing the illustrative forms for the convenience of the reader and are not for the purpose of limitation thereof.

**[0021]** Further, it is understood that any one or more of the following-described forms, expressions of forms, examples, can be combined with any one or more of the other following-described forms, expressions of forms, and examples.

**[0022]** Various forms are directed to improved ultrasonic and/or electrosurgical (RF) instruments configured for effecting tissue dissecting, cutting, and/or coagulation during surgical procedures. In one form, a combined ultrasonic and electrosurgical instrument may be configured for use in open surgical procedures, but has applications in other types of surgery, such as laparoscopic, endoscopic, and robotic-assisted procedures. Versatile use is facilitated by selective use of ultrasonic and RF energy.

**[0023]** The various forms will be described in combination with an ultrasonic instrument as described herein. Such description is provided by way of example, and not limitation, and is not intended to limit the scope and applications thereof. For example, any one of the described forms is useful in combination with a multitude of ultrasonic instruments including those described in, for example, U.S. Patent Nos. 5,938,633; 5,935,144; 5,944,737; 5,322,055; 5,630,420; and 5,449,370.

**[0024]** As will become apparent from the following description, it is contemplated that forms of the surgical instruments described herein may be used in association with an oscillator unit of a surgical system, whereby ultrasonic energy from the oscillator unit provides the desired ultrasonic actuation for the present surgical instrument. It is also contemplated that forms of the surgical instrument described herein may be used in association with a signal generator unit of a surgical system, whereby electrical energy in the form of radio frequencies (RF), for example, is used to provide feedback to the user regarding the surgical instrument. The ultrasonic oscillator and/or the signal generator unit may be non-detachably integrated with the surgical instrument or may be provided as separate components, which can be electrically attachable to the surgical instrument.

**[0025]** One form of the present surgical apparatus is particularly configured for disposable use by virtue of its straight-forward construction. However, it is also contemplated that other forms of the present surgical instrument can be configured for non-disposable or multiple uses. Detachable connection of the present surgical instrument with an associated oscillator and signal generator unit is presently disclosed for single-patient use for illustrative purposes only. However, non-detachable integrated connection of the present surgical instrument with an associated oscillator and/or signal generator unit is also contemplated. Accordingly, various forms of the presently described surgical instruments may be configured for single use and/or multiple use with either detachable and/or non-detachable integral oscillator and/or signal generator unit, without limitation, and all combinations of such configurations are contemplated to be within the scope of the present disclosure.

**[0026]** With reference to FIGS. 1-5, one form of a surgical system 10 including an ultrasonic surgical instrument is illustrated. FIG. 1 illustrates one form of a surgical system 100 comprising a generator 102 and various surgical instruments 104, 106, 108 usable therewith. FIG. 2 is a diagram of the ultrasonic surgical instrument 104 of FIG. 1.

**[0027]** FIG. 1 illustrates a generator 102 configured to drive multiple surgical instruments 104, 106, 108. The first surgical instrument 104 comprises a handpiece 105, an ultrasonic transducer 120, a shaft 126, and an end effector 122. The end effector 122 comprises an ultrasonic blade 128 acoustically coupled to the transducer 120 and a clamp arm 140. The handpiece 105 comprises a trigger 143 to operate the clamp arm 140 and a combination of the toggle buttons 134a, 134b, 134c to energize and drive the ultrasonic blade 128 or other function. The toggle buttons 134a, 134b, 134c can be configured to energize the ultrasonic transducer 120 with the generator 102.

**[0028]** Still with reference to FIG. 1, the generator 102 also is configured to drive a second surgical instrument 106.

The second surgical instrument 106 is an RF electrosurgical instrument and comprises a handpiece 107, a shaft 127, and an end effector 124. The end effector 124 comprises electrodes in the clamp arms 143 and return through the ultrasonic blade 149. The electrodes are coupled to and energized by a bipolar energy source within the generator 102. The handpiece 107 comprises a trigger 147 to operate the clamp arm 145 and an energy button 135 to actuate an energy switch to energize the electrodes in the end effector 124.

**[0029]** Still with reference to FIG. 1, the generator 102 also is configured to drive a combination electrosurgical and ultrasonic instrument 108. The combination electrosurgical and ultrasonic multifunction surgical instrument 108 comprises a handpiece 109, a shaft 129, and an end effector 125. The end effector comprises an ultrasonic blade 149 and a clamp arm 145. The ultrasonic blade 149 is acoustically coupled to the ultrasonic transducer 120. The handpiece 109 comprises a trigger 147 to operate the clamp arm 145 and a combination of the toggle buttons 137a, 137b, 137c to energize and drive the ultrasonic blade 149 or other function. The toggle buttons 137a, 137b, 137c can be configured to energize the ultrasonic transducer 120 with the generator 102 and energize the ultrasonic blade 149 with a bipolar energy source also contained within the generator 102.

**[0030]** With reference to both FIGS. 1 and 2, the generator 102 is configurable for use with a variety of surgical devices. According to various forms, the generator 102 may be configurable for use with different surgical devices of different types including, for example, the ultrasonic surgical instrument 104, the electrosurgical or RF surgical devices, such as, the RF electrosurgical instrument 106, and the multifunction surgical instrument 108 that integrate electrosurgical RF and ultrasonic energies delivered simultaneously from the generator 102. Although in the form of FIG. 1, the generator 102 is shown separate from the surgical instruments 104, 106, 108, in one form, the generator 102 may be formed integrally with either of the surgical instrument 104, 106, 108 to form a unitary surgical system. The generator 102 comprises an input device 110 located on a front panel of the generator 102 console. The input device 110 may comprise any suitable device that generates signals suitable for programming the operation of the generator 102. The generator 102 also may comprise one or more output devices 112.

**[0031]** The generator 102 is coupled to an ultrasonic transducer 120 via a cable 144. The ultrasonic transducer 120 and a waveguide extending through a shaft 126 (waveguide not shown in FIG. 2) may collectively form an ultrasonic drive system driving an ultrasonic blade 128 of an end effector 122. The end effector 122 further may comprise a clamp arm 140 to clamp tissue between the clamp arm 140 and the ultrasonic blade 128. In one form, the generator 102 may be configured to produce a drive signal of a particular voltage, current, and/or frequency output signal that can be stepped or otherwise modified with high resolution, accuracy, and repeatability.

**[0032]** Still with reference to FIG. 2, it will be appreciated that a surgical instrument 104 may comprise any combination of the toggle buttons 134a, 134b, 134c. For example, the surgical instrument 104 could be configured to have only two toggle buttons: a toggle button 134a for producing maximum ultrasonic energy output and a toggle button 134c for producing a pulsed output at either the maximum or less than maximum power level. In this way, the drive signal output configuration of the generator 102 could be 5 continuous signals and 5 or 4 or 3 or 2 or 1 pulsed signals. In certain forms, the specific drive signal configuration may be controlled based upon, for example, EEPROM settings in the generator 102 and/or user power level selection(s).

**[0033]** In certain forms, a two-position switch may be provided as an alternative to a toggle button 134c. For example, a surgical instrument 104 may include a toggle button 134a for producing a continuous output at a maximum power level and a two-position toggle button 134b. In a first detented position, toggle button 134b may produce a continuous output at a less than maximum power level, and in a second detented position the toggle button 134b may produce a pulsed output (e.g., at either a maximum or less than maximum power level, depending upon the EEPROM settings).

**[0034]** Still with reference to FIG. 2, forms of the generator 102 may enable communication with instrument-based data circuits. For example, the generator 102 may be configured to communicate with a first data circuit 136 and/or a second data circuit 138. For example, the first data circuit 136 may indicate a burn-in frequency slope, as described herein. Additionally or alternatively, any type of information may be communicated to second data circuit for storage therein via a data circuit interface (e.g., using a logic device). Such information may comprise, for example, an updated number of operations in which the instrument has been used and/or dates and/or times of its usage. In certain forms, the second data circuit may transmit data acquired by one or more sensors (e.g., an instrument-based temperature sensor). In certain forms, the second data circuit may receive data from the generator 102 and provide an indication to a user (e.g., an LED indication or other visible indication) based on the received data. The second data circuit 138 contained in the multifunction surgical instrument 108 of a surgical device. In some forms, the second data circuit 138 may be implemented in a many similar to that of the first data circuit 136 described herein. An instrument interface circuit may comprise a second data circuit interface to enable this communication. In one form, the second data circuit interface may comprise a tri-state digital interface, although other interfaces also may be used. In certain forms, the second data circuit may generally be any circuit for transmitting and/or receiving data. In one form, for example, the second data circuit may store information pertaining to the particular surgical instrument with which it is associated. Such information may include, for example, a model number, a serial number, a number of operations in which the surgical instrument has been used, and/or any other type of information. In some forms, the second data circuit 138 may store information

about the electrical and/or ultrasonic properties of an associated transducer 120, end effector 122, or ultrasonic drive system. Various processes and techniques described herein may be executed by a generator. It will be appreciated, however, that in certain example forms, all or a part of these processes and techniques may be performed by internal logic 139 of the multifunction surgical instrument 108.

**[0035]** FIG. 3 is a diagram of the surgical system 100 of FIG. 1. In various forms, the generator 102 may comprise several separate functional elements, such as modules and/or blocks. Different functional elements or modules may be configured for driving the different kinds of surgical instruments 104, 106, 108. For example, an ultrasonic generator drive circuit 114 may drive ultrasonic devices such as the ultrasonic surgical instrument 104 via a cable 142. An electrosurgery/RF generator drive circuit 116 may drive the electrosurgical instrument 106 via a cable 144. For example, the respective drive circuits 114, 116 may generate respective drive signals for driving the surgical instruments 104, 106, 108. In various forms, the ultrasonic generator drive circuit 114 (e.g., ultrasonic drive circuit) and/or the electrosurgery/RF generator drive circuit 116 (e.g., RF drive circuit) each may be formed integrally with the generator 102. Alternatively, one or more of the drive circuits 114, 116 may be provided as a separate circuit module electrically coupled to the generator 102. (The drive circuits 114 and 116 are shown in phantom to illustrate this option.) Also, in some forms, the electrosurgery/RF generator drive circuit 116 may be formed integrally with the ultrasonic generator drive circuit 114, or vice versa. Also, in some forms, the generator 102 may be omitted entirely and the drive circuits 114, 116 may be executed by processors or other hardware within the respective surgical instruments 104, 106, 108.

**[0036]** In other forms, the electrical outputs of the ultrasonic generator drive circuit 114 and the electrosurgery/RF generator drive circuit 116 may be combined into a single drive circuit to provide a single electrical signal capable of driving the multifunction surgical instrument 108 simultaneously with electrosurgical RF and ultrasonic energies via a cable 146. The multifunction surgical instrument 108 comprises an ultrasonic transducer 120 coupled to an ultrasonic blade 149 and one or more electrodes in the end effector 124 to receive electrosurgical RF energy. In such implementations, the combined RF/ultrasonic signal is coupled to the multifunction surgical instrument 108. The multifunction surgical instrument 108 comprises signal processing components to split the combined RF/ultrasonic signal such that the RF signal can be delivered to the electrodes in the end effector 124 and the ultrasonic signal can be delivered to the ultrasonic transducer 120.

**[0037]** In accordance with the described forms, the ultrasonic generator drive circuit 114 may produce a drive signal or signals of particular voltages, currents, and frequencies, e.g., 55,500 cycles per second (Hz). The drive signal or signals may be provided to the ultrasonic surgical instrument 104, and specifically to the transducer 120, which may operate, for example, as described herein. The transducer 120 and a waveguide extending through the shaft 126 (waveguide not shown in FIG. 2) may collectively form an ultrasonic drive system driving an ultrasonic blade 128 of an end effector 122. In one form, the generator 102 may be configured to produce a drive signal of a particular voltage, current, and/or frequency output signal that can be stepped or otherwise modified with high resolution, accuracy, and repeatability.

**[0038]** The generator 102 may be activated to provide the drive signal to the transducer 120 in any suitable manner. For example, the generator 102 may comprise a foot switch 130 coupled to the generator 102 via a foot switch cable 132. A clinician may activate the transducer 120 by depressing the foot switch 130. In addition, or instead of the foot switch 130 some forms of the ultrasonic surgical instrument 104 may utilize one or more switches positioned on the hand piece that, when activated, may cause the generator 102 to activate the transducer 120. In one form, for example, the one or more switches may comprise a pair of toggle buttons 134a, 134b (FIG. 2), for example, to determine an operating mode of the surgical instrument 104. When the toggle button 134a is depressed, for example, the ultrasonic generator 102 may provide a maximum drive signal to the transducer 120, causing it to produce maximum ultrasonic energy output. Depressing toggle button 134b may cause the ultrasonic generator 102 to provide a user-selectable drive signal to the transducer 120, causing it to produce less than the maximum ultrasonic energy output. The surgical instrument 104 additionally or alternatively may comprise a second switch (not shown) to, for example, indicate a position of a jaw closure trigger for operating jaws of the end effector 122. Also, in some forms, the ultrasonic generator 102 may be activated based on the position of the jaw closure trigger, (e.g., as the clinician depresses the jaw closure trigger to close the jaws, ultrasonic energy may be applied).

**[0039]** Additionally or alternatively, the one or more switches may comprises a toggle button 134c that, when depressed, causes the generator 102 to provide a pulsed output. The pulses may be provided at any suitable frequency and grouping, for example. In certain forms, the power level of the pulses may be the power levels associated with toggle buttons 134a, 134b (maximum, less than maximum), for example.

**[0040]** In accordance with the described forms, the electrosurgery/RF generator drive circuit 116 may generate a drive signal or signals with output power sufficient to perform bipolar electrosurgery using radio frequency (RF) energy. In bipolar electrosurgery applications, the drive signal may be provided, for example, to electrodes of the electrosurgical instrument 106, for example. Accordingly, the generator 102 may be configured for therapeutic purposes by applying electrical energy to the tissue sufficient for treating the tissue (e.g., coagulation, cauterization, tissue welding).

**[0041]** The generator 102 may comprise an input device 110 (FIG. 1) located, for example, on a front panel of the

generator 102 console. The input device 110 may comprise any suitable device that generates signals suitable for programming the operation of the generator 102. In operation, the user can program or otherwise control operation of the generator 102 using the input device 110. The input device 110 may comprise any suitable device that generates signals that can be used by the generator (e.g., by one or more processors contained in the generator) to control the operation of the generator 102 (e.g., operation of the ultrasonic generator drive circuit 114 and/or electrosurgery/RF generator drive circuit 116). In various forms, the input device 110 includes one or more of buttons, switches, thumbwheels, keyboard, keypad, touch screen monitor, pointing device, remote connection to a general purpose or dedicated computer. In other forms, the input device 110 may comprise a suitable user interface, such as one or more user interface screens displayed on a touch screen monitor, for example. Accordingly, by way of the input device 110, the user can set or program various operating parameters of the generator, such as, for example, current (I), voltage (V), frequency (f), and/or period (T) of a drive signal or signals generated by the ultrasonic generator drive circuit 114 and/or electrosurgery/RF generator drive circuit 116.

**[0042]** The generator 102 also may comprise an output device 112 (FIGS. 1, 3), such as an output indicator, located, for example, on a front panel of the generator 102 console. The output device 112 includes one or more devices for providing a sensory feedback to a user. Such devices may comprise, for example, visual feedback devices (e.g., a visual feedback device may comprise incandescent lamps, light emitting diodes (LEDs), graphical user interface, display, analog indicator, digital indicator, bar graph display, digital alphanumeric display, LCD display screen, LED indicators), audio feedback devices (e.g., an audio feedback device may comprise speaker, buzzer, audible, computer generated tone, computerized speech, voice user interface (VUI) to interact with computers through a voice/speech platform), or tactile feedback devices (e.g., a tactile feedback device comprises any type of vibratory feedback, haptic actuator).

**[0043]** Although certain modules, circuits, and/or blocks of the generator 102 may be described by way of example, it can be appreciated that a greater or lesser number of modules, circuits, and/or blocks may be used and still fall within the scope of the forms. Further, although various forms may be described in terms of modules, circuits, and/or blocks to facilitate description, such modules, circuits, and/or blocks may be implemented by one or more hardware components, e.g., processors, Digital Signal Processors (DSPs), Programmable Logic Devices (PLDs), Application Specific Integrated Circuits (ASICs), circuits, registers and/or software components, e.g., programs, subroutines, logic and/or combinations of hardware and software components. Also, in some forms, the various modules described herein may be implemented utilizing similar hardware positioned within the surgical instruments 104, 106, 108 (i.e., the generator 102 may be omitted).

**[0044]** In one form, the ultrasonic generator drive circuit 114 and electrosurgery/RF drive circuit 116 may comprise one or more embedded applications implemented as firmware, software, hardware, or any combination thereof. The drive circuits 114, 116 may comprise various executable modules such as software, programs, data, drivers, application program interfaces (APIs), and so forth. The firmware may be stored in nonvolatile memory (NVM), such as in bit-masked read-only memory (ROM) or flash memory. In various implementations, storing the firmware in ROM may preserve flash memory. The NVM may comprise other types of memory including, for example, programmable ROM (PROM), erasable programmable ROM (EPROM), electrically erasable programmable ROM (EEPROM), or battery backed random-access memory (RAM) such as dynamic RAM (DRAM), Double-Data-Rate DRAM (DDR), and/or synchronous DRAM (SDRAM).

**[0045]** In one form, the drive circuits 114, 116 comprise a hardware component implemented as a processor for executing program instructions for monitoring various measurable characteristics of the surgical instruments 104, 106, 108 and generating a corresponding output control signals for operating the surgical instruments 104, 106, 108. In forms in which the generator 102 is used in conjunction with the surgical instrument 104, the output control signal may drive the ultrasonic transducer 120 in cutting and/or coagulation operating modes. Electrical characteristics of the surgical instrument 104 and/or tissue may be measured and used to control operational aspects of the generator 102 and/or provided as feedback to the user. In forms in which the generator 102 is used in conjunction with the electrosurgical instrument 106, the output control signal may supply electrical energy (e.g., RF energy) to the end effector 124 in cutting, coagulation and/or desiccation modes. Electrical characteristics of the electrosurgical instrument 106 and/or tissue may be measured and used to control operational aspects of the generator 102 and/or provide feedback to the user. In various forms, as previously discussed, the hardware component may be implemented as a DSP, PLD, ASIC, circuits, and/or registers. In one form, the processor may be configured to store and execute computer software program instructions to generate the output signal functions for driving various components of the surgical instruments 104, 106, 108, such as the ultrasonic transducer 120 and the end effectors 122, 124.

**[0046]** FIG. 4 illustrates an equivalent circuit 150 of an ultrasonic transducer, such as the ultrasonic transducer 120 shown in FIGS. 1-3, according to one form. The circuit 150 comprises a first "motional" branch having a serially connected inductance  $L_s$ , resistance  $R_s$  and capacitance  $C_s$  that define the electromechanical properties of the resonator, and a second capacitive branch having a static capacitance  $C_0$ . Drive current  $I_g$  may be received from a generator at a drive voltage  $V_g$ , with motional current  $I_m$  flowing through the first branch and current  $I_g - I_m$  flowing through the capacitive branch. Control of the electromechanical properties of the ultrasonic transducer may be achieved by suitably controlling  $I_g$  and  $V_g$ . As explained above, conventional generator architectures may include a tuning inductor  $L_t$  (shown in phantom



in FIG. 4) for tuning out in a parallel resonance circuit the static capacitance  $C_o$  at a resonant frequency so that substantially all of generator's current output  $I_g$  flows through the motional branch. In this way, control of the motional branch current  $I_m$  is achieved by controlling the generator current output  $I_g$ . The tuning inductor  $L_t$  is specific to the static capacitance  $C_o$  of an ultrasonic transducer, however, and a different ultrasonic transducer having a different static capacitance requires a different tuning inductor  $L_t$ . Moreover, because the tuning inductor  $L_t$  is matched to the nominal value of the static capacitance  $C_o$  at a resonant frequency, accurate control of the motional branch current  $I_m$  is assured only at that frequency, and as frequency shifts down with transducer temperature, accurate control of the motional branch current is compromised.

**[0047]** Forms of the generator 102 shown in FIGS. 1-3 do not rely on a tuning inductor  $L_t$  to monitor the motional branch current  $I_m$ . Instead, the generator 102 may use the measured value of the static capacitance  $C_o$  in between applications of power for a specific ultrasonic surgical instrument 104 (along with drive signal voltage and current feedback data) to determine values of the motional branch current  $I_m$  on a dynamic and ongoing basis (e.g., in real-time). Such forms of the generator 102 are therefore able to provide virtual tuning to simulate a system that is tuned or resonant with any value of static capacitance  $C_o$  at any frequency, and not just at the resonant frequency dictated by a nominal value of the static capacitance  $C_o$ .

**[0048]** FIG. 5 is a simplified block diagram of a generator 200 which is one form of the generator 102 shown in FIGS. 1-3 for providing inductorless tuning as described herein, among other benefits. Additional details of the generator 102 are described in US2011087256. With reference to FIG. 5, the generator 200 may comprise a patient isolated stage 202 in communication with a non-isolated stage 204 via a power transformer 206. A secondary winding 208 of the power transformer 206 is contained in the isolated stage 202 and may comprise a tapped configuration (e.g., a center-tapped or a non-center-tapped configuration) to define drive signal outputs 210a, 210b, 210c for outputting drive signals to different surgical devices, such as, for example, an ultrasonic surgical instrument 104 and an electrosurgical instrument 106 (as shown in FIGS. 1-3). In particular, the drive signal outputs 210a, 210c may output an ultrasonic drive signal (e.g., a 420V RMS drive signal) to an ultrasonic surgical instrument 104, and the drive signal outputs 210b, 210c may output an electrosurgical RF drive signal (e.g., a 100V RMS drive signal) to an electrosurgical instrument 106, with the output 210b corresponding to the center tap of the power transformer 206.

**[0049]** In certain forms, the ultrasonic and electrosurgical drive signals may be provided simultaneously to distinct surgical instruments and/or to a single surgical instrument having the capability to deliver both ultrasonic and electrosurgical energy to tissue, such as multifunction surgical instrument 108 (FIGS. 1 and 3). It will be appreciated that the electrosurgical signal, provided either to a dedicated electrosurgical instrument and/or to a combined multifunction ultrasonic/electrosurgical instrument may be either a therapeutic or sub-therapeutic level signal. For example, the ultrasonic and radio frequency signals can be delivered separately or simultaneously from a generator with a single output port in order to provide the desired output signal to the surgical instrument, as will be discussed in more detail below. Accordingly, the generator can combine the ultrasonic and electrosurgical RF energies and deliver the combined energies to the multifunction ultrasonic/electrosurgical instrument. Bipolar electrodes can be placed on one or both jaws of the end effector. One jaw may be driven by ultrasonic energy in addition to electrosurgical RF energy, working simultaneously. The ultrasonic energy may be employed to dissect tissue while the electrosurgical RF energy may be employed for vessel sealing.

**[0050]** The non-isolated stage 204 may comprise a power amplifier 212 having an output connected to a primary winding 214 of the power transformer 206. In certain forms the power amplifier 212 may comprise a push-pull amplifier. For example, the non-isolated stage 204 may further comprise a logic device 216 for supplying a digital output to a digital-to-analog converter (DAC) 218, which in turn supplies a corresponding analog signal to an input of the power amplifier 212. In certain forms the logic device 216 may comprise a programmable gate array (PGA), a field-programmable gate array (FPGA), programmable logic device (PLD), among other logic circuits, for example. The logic device 216, by virtue of controlling the input of the power amplifier 212 via the DAC 218, may therefore control any of a number of parameters (e.g., frequency, waveform shape, waveform amplitude) of drive signals appearing at the drive signal outputs 210a, 210b, 210c. In certain forms and as discussed below, the logic device 216, in conjunction with a processor (e.g., a digital signal processor discussed below), may implement a number of digital signal processing (DSP)-based and/or other control techniques to control parameters of the drive signals output by the generator 200.

**[0051]** Power may be supplied to a power rail of the power amplifier 212 by a switch-mode regulator 220. In certain forms the switch-mode regulator 220 may comprise an adjustable buck regulator, for example. The non-isolated stage 204 may further comprise a first processor such as DSP processor 222, which in one form may comprise a DSP processor such as an Analog Devices ADSP-21469 SHARC DSP, available from Analog Devices, Norwood, MA, for example, although in various forms any suitable processor may be employed. In certain forms the DSP processor 222 may control operation of the switch-mode power converter 220 responsive to voltage feedback data received from the power amplifier 212 by the DSP processor 222 via an analog-to-digital converter (ADC) 224. In one form, for example, the DSP processor 222 may receive as input, via the ADC 224, the waveform envelope of a signal (e.g., an RF signal) being amplified by the power amplifier 212. The DSP processor 222 may then control the switch-mode regulator 220 (e.g., via a pulse-

width modulated (PWM) output) such that the rail voltage supplied to the power amplifier 212 tracks the waveform envelope of the amplified signal. By dynamically modulating the rail voltage of the power amplifier 212 based on the waveform envelope, the efficiency of the power amplifier 212 may be significantly improved relative to a fixed rail voltage amplifier schemes.

**[0052]** In certain forms, the logic device 216, in conjunction with the DSP processor 222, may implement a direct digital synthesizer (DDS) control scheme to control the waveform shape, frequency and/or amplitude of drive signals output by the generator 200. In one form, for example, the logic device 216 may implement a DDS control technique by recalling waveform samples stored in a dynamically-updated look-up table (LUT), such as a RAM LUT, which may be embedded in an FPGA. This control technique is particularly useful for ultrasonic applications in which an ultrasonic transducer, such as the ultrasonic transducer 120 (FIGS. 1-3), may be driven by a clean sinusoidal current at its resonant frequency. Because other frequencies may excite parasitic resonances, minimizing or reducing the total distortion of the motional branch current may correspondingly minimize or reduce undesirable resonance effects. Because the waveform shape of a drive signal output by the generator 200 is impacted by various sources of distortion present in the output drive circuit (e.g., the power transformer 206, the power amplifier 212), voltage and current feedback data based on the drive signal may be input into a technique, such as an error control technique implemented by the DSP processor 222, which compensates for distortion by suitably pre-distorting or modifying the waveform samples stored in the LUT on a dynamic, ongoing basis (e.g., in real-time). In one form, the amount or degree of pre-distortion applied to the LUT samples may be based on the error between a computed motional branch current and a desired current waveform shape, with the error being determined on a sample-by-sample basis. In this way, the pre-distorted LUT samples, when processed through the drive circuit, may result in a motional branch drive signal having the desired waveform shape (e.g., sinusoidal) for optimally driving the ultrasonic transducer. In such forms, the LUT waveform samples will therefore not represent the desired waveform shape of the drive signal, but rather the waveform shape that is required to ultimately produce the desired waveform shape of the motional branch drive signal when distortion effects are taken into account.

**[0053]** The non-isolated stage 204 may further comprise an ADC 226 and an ADC 228 coupled to the output of the power transformer 206 via respective isolation transformers 230, 232 for respectively sampling the voltage and current of drive signals output by the generator 200. In certain forms, the ADCs 226, 228 may be configured to sample at high speeds (e.g., 80 MSPS) to enable oversampling of the drive signals. In one form, for example, the sampling speed of the ADCs 226, 228 may enable approximately 200x (depending on frequency) oversampling of the drive signals. In certain forms, the sampling operations of the ADC 226, 228 may be performed by a single ADC receiving input voltage and current signals via a two-way multiplexer. The use of high-speed sampling in forms of the generator 200 may enable, among other things, calculation of the complex current flowing through the motional branch (which may be used in certain forms to implement DDS-based waveform shape control described herein), accurate digital filtering of the sampled signals, and calculation of real power consumption with a high degree of precision. Voltage and current feedback data output by the ADCs 226, 228 may be received and processed (e.g., FIFO buffering, multiplexing) by the logic device 216 and stored in data memory for subsequent retrieval by, for example, the DSP processor 222. As noted above, voltage and current feedback data may be used as input to a technique for pre-distorting or modifying LUT waveform samples on a dynamic and ongoing basis. In certain forms, this may require each stored voltage and current feedback data pair to be indexed based on, or otherwise associated with, a corresponding LUT sample that was output by the logic device 216 when the voltage and current feedback data pair was acquired. Synchronization of the LUT samples and the voltage and current feedback data in this manner contributes to the correct timing and stability of the pre-distortion technique.

**[0054]** In certain forms, the voltage and current feedback data may be used to control the frequency and/or amplitude (e.g., current amplitude) of the drive signals. In one form, for example, voltage and current feedback data may be used to determine impedance phase. The frequency of the drive signal may then be controlled to minimize or reduce the difference between the determined impedance phase and an impedance phase setpoint (e.g., 0°), thereby minimizing or reducing the effects of ultrasonic distortion and correspondingly enhancing impedance phase measurement accuracy. The determination of phase impedance and a frequency control signal may be implemented in the DSP processor 222, for example, with the frequency control signal being supplied as input to a DDS control technique implemented by the logic device 216.

**[0055]** In another form, for example, the current feedback data may be monitored in order to maintain the current amplitude of the drive signal at a current amplitude setpoint. The current amplitude setpoint may be specified directly or determined indirectly based on specified voltage amplitude and power setpoints. In certain forms, control of the current amplitude may be implemented by control technique, such as, for example, a PID control technique, in the DSP processor 222. Variables controlled by the control technique to suitably control the current amplitude of the drive signal may include, for example, the scaling of the LUT waveform samples stored in the logic device 216 and/or the full-scale output voltage of the DAC 218 (which supplies the input to the power amplifier 212) via a DAC 234.

**[0056]** The non-isolated stage 204 may further comprise a second processor such as UI processor 236 for providing, among other things user interface (UI) functionality. In one form, the UI processor 236 may comprise an Atmel

AT91SAM9263 processor having an ARM 926EJ-S core, available from Atmel Corporation, San Jose, CA, for example. Examples of UI functionality supported by the UI processor 236 may include audible and visual user feedback, communication with peripheral devices (e.g., via a Universal Serial Bus (USB) interface), communication with the foot switch 130, communication with an input device 118 (e.g., a touch screen display) and communication with an output device 112 (e.g., a speaker), as shown in FIG. 3, for example. The UI processor 236 may communicate with the DSP processor 222 and the logic device 216 (e.g., via serial peripheral interface (SPI) buses). Although the UI processor 236 may primarily support UI functionality, it also may coordinate with the DSP processor 222 to implement hazard mitigation in certain forms. For example, the UI processor 236 may be programmed to monitor various aspects of user input and/or other inputs (e.g., touch screen inputs, foot switch 130 inputs (FIG. 3), temperature sensor inputs) and may disable the drive output of the generator 200 when an erroneous condition is detected.

**[0057]** In certain forms, both the DSP processor 222 and the UI processor 236, for example, may determine and monitor the operating state of the generator 200. For the DSP processor 222, the operating state of the generator 200 may dictate, for example, which control and/or diagnostic processes are implemented by the DSP processor 222. For the UI processor 236, the operating state of the generator 200 may dictate, for example, which elements of a user interface (e.g., display screens, sounds) are presented to a user. The respective DSP and UI processors 222, 236 may independently maintain the current operating state of the generator 200 and recognize and evaluate possible transitions out of the current operating state. The DSP processor 222 may function as the master in this relationship and determine when transitions between operating states are to occur. The UI processor 236 may be aware of valid transitions between operating states and may confirm if a particular transition is appropriate. For example, when the DSP processor 222 instructs the UI processor 236 to transition to a specific state, the UI processor 236 may verify that requested transition is valid. In the event that a requested transition between states is determined to be invalid by the UI processor 236, the UI processor 236 may cause the generator 200 to enter a failure mode.

**[0058]** The non-isolated stage 204 may further comprise a controller 238 for monitoring input devices 110 (e.g., a capacitive touch sensor used for turning the generator 200 on and off, a capacitive touch screen, e.g., as shown in FIGS. 1 and 3). In certain forms, the controller 238 may comprise at least one processor and/or other controller device in communication with the UI processor 236. In one form, for example, the controller 238 may comprise a processor (e.g., a Mega168 8-bit controller available from Atmel) configured to monitor user input provided via one or more capacitive touch sensors. In one form, the controller 238 may comprise a touch screen controller (e.g., a QT5480 touch screen controller available from Atmel) to control and manage the acquisition of touch data from a capacitive touch screen.

**[0059]** In certain forms, when the generator 200 is in a "power off" state, the controller 238 may continue to receive operating power (e.g., via a line from a power supply of the generator 200. In this way, the controller 238 may continue to monitor an input device 110 (e.g., a capacitive touch sensor located on a front panel of the generator 200) for turning the generator 200 on and off. When the generator 200 is in the power off state, the controller 238 may wake the power supply (e.g., enable operation of one or more DC/DC voltage converters of the power supply) if activation of the "on/off" input device 110 by a user is detected. The controller 238 may therefore initiate a sequence for transitioning the generator 200 to a "power on" state. Conversely, the controller 238 may initiate a sequence for transitioning the generator 200 to the power off state if activation of the "on/off" input device 110 is detected when the generator 200 is in the power on state. In certain forms, for example, the controller 238 may report activation of the "on/off" input device 110 to the UI processor 236, which in turn implements the necessary process sequence for transitioning the generator 200 to the power off state. In such forms, the controller 238 may have no independent ability for causing the removal of power from the generator 200 after its power on state has been established.

**[0060]** In certain forms, the controller 238 may cause the generator 200 to provide audible or other sensory feedback for alerting the user that a power on or power off sequence has been initiated. Such an alert may be provided at the beginning of a power on or power off sequence and prior to the commencement of other processes associated with the sequence.

**[0061]** In certain forms, the isolated stage 202 may comprise an instrument interface circuit 240 to, for example, provide a communication interface between a control circuit of a surgical device (e.g., a control circuit comprising hand piece switches) and components of the non-isolated stage 204, such as, for example, the programmable logic device 216, the DSP processor 222 and/or the UI processor 236. The instrument interface circuit 240 may exchange information with components of the non-isolated stage 204 via a communication link that maintains a suitable degree of electrical isolation between the stages 202, 204, such as, for example, an infrared (IR)-based communication link. Power may be supplied to the instrument interface circuit 240 using, for example, a low-dropout voltage regulator powered by an isolation transformer driven from the non-isolated stage 204.

**[0062]** In one form, the instrument interface circuit 240 may comprise a logic device 242 (e.g., logic circuit, programmable logic circuit, PGA, FPGA, PLD) in communication with a signal conditioning circuit 242. The signal conditioning circuit 244 may be configured to receive a periodic signal from the logic circuit 242 (e.g., a 2 kHz square wave) to generate a bipolar interrogation signal having an identical frequency. The interrogation signal may be generated, for example, using a bipolar current source fed by a differential amplifier. The interrogation signal may be communicated to a surgical

device control circuit (e.g., by using a conductive pair in a cable that connects the generator 200 to the surgical device) and monitored to determine a state or configuration of the control circuit. The control circuit may comprise a number of switches, resistors and/or diodes to modify one or more characteristics (e.g., amplitude, rectification) of the interrogation signal such that a state or configuration of the control circuit is uniquely discernable based on the one or more characteristics. In one form, for example, the signal conditioning circuit 244 may comprises an ADC for generating samples of a voltage signal appearing across inputs of the control circuit resulting from passage of interrogation signal therethrough. The logic device 242 (or a component of the non-isolated stage 204) may then determine the state or configuration of the control circuit based on the ADC samples.

**[0063]** In one form, the instrument interface circuit 240 may comprise a first data circuit interface 246 to enable information exchange between the logic circuit 242 (or other element of the instrument interface circuit 240) and a first data circuit disposed in or otherwise associated with a surgical device. In certain forms, for example, a first data circuit 136 (FIG. 2) may be disposed in a cable integrally attached to a surgical device hand piece, or in an adaptor for interfacing a specific surgical device type or model with the generator 200. The first data circuit 136 may be implemented in any suitable manner and may communicate with the generator according to any suitable protocol including, for example, as described herein with respect to the first circuit 136. In certain forms, the first data circuit may comprise a non-volatile storage device, such as an electrically erasable programmable read-only memory (EEPROM) device. In certain forms and referring again to FIG. 5, the first data circuit interface 246 may be implemented separately from the logic device 242 and comprise suitable circuitry (e.g., discrete logic devices, a processor) to enable communication between the programmable logic device 242 and the first data circuit. In other forms, the first data circuit interface 246 may be integral with the logic device 242.

**[0064]** In certain forms, the first data circuit 136 (FIG. 2) may store information pertaining to the particular surgical device with which it is associated. Such information may include, for example, a model number, a serial number, a number of operations in which the surgical device has been used, and/or any other type of information. This information may be read by the instrument interface circuit 1098 (e.g., by the logic device 242), transferred to a component of the non-isolated stage 204 (e.g., to logic device 216, DSP processor 222 and/or UI processor 236) for presentation to a user via an output device 112 (FIGS. 1 and 3) and/or for controlling a function or operation of the generator 200. Additionally, any type of information may be communicated to first data circuit 136 for storage therein via the first data circuit interface 246 (e.g., using the logic device 242). Such information may comprise, for example, an updated number of operations in which the surgical device has been used and/or dates and/or times of its usage.

**[0065]** As discussed previously, a surgical instrument may be detachable from a hand piece (e.g., as shown in FIGS. 1 and 2, the transducer 120 and the shaft 126 is detachable from the handpiece 105 of the ultrasonic surgical instrument 104) to promote instrument interchangeability and/or disposability. In such cases, conventional generators may be limited in their ability to recognize particular instrument configurations being used and to optimize control and diagnostic processes accordingly. The addition of readable data circuits to surgical device instruments to address this issue is problematic from a compatibility standpoint, however. For example, designing a surgical device to remain backwardly compatible with generators that lack the requisite data reading functionality may be impractical due to, for example, differing signal schemes, design complexity, and cost. Forms of instruments discussed herein address these concerns by using data circuits that may be implemented in existing surgical instruments economically and with minimal design changes to preserve compatibility of the surgical devices with current generator platforms.

**[0066]** With reference to FIGS. 1-3 and 5, additionally, forms of the generator 200 may enable communication with instrument-based data circuits. For example, the generator 200 may be configured to communicate with a second data circuit 138 contained in the ultrasonic surgical instrument 104 (e.g., and/or the other surgical instruments 106, 108). In some forms, the second data circuit 138 may be implemented in a many similar to that of the first data circuit 136 described herein. The instrument interface circuit 240 may comprise a second data circuit interface 248 to enable this communication. In one form, the second data circuit interface 248 may comprise a tri-state digital interface, although other interfaces also may be used. In certain forms, the second data circuit may generally be any circuit for transmitting and/or receiving data. In one form, for example, the second data circuit may store information pertaining to the particular surgical instrument with which it is associated. Such information may include, for example, a model number, a serial number, a number of operations in which the surgical instrument has been used, and/or any other type of information. In some forms, the second data circuit 138 may store information about the electrical and/or ultrasonic properties of an associated transducer 120, end effector 122, or ultrasonic drive system. For example, the first data circuit 136 may indicate a burn-in frequency slope, as described herein. Additionally or alternatively, any type of information may be communicated to second data circuit for storage therein via the second data circuit interface 248 (e.g., using the logic device 242). Such information may comprise, for example, an updated number of operations in which the instrument has been used and/or dates and/or times of its usage. In certain forms, the second data circuit may transmit data acquired by one or more sensors (e.g., an instrument-based temperature sensor). In certain forms, the second data circuit may receive data from the generator 200 and provide an indication to a user (e.g., an LED indication or other visible indication) based on the received data.

**[0067]** In certain forms, the second data circuit and the second data circuit interface 248 may be configured such that communication between the logic device 242 and the second data circuit can be effected without the need to provide additional conductors for this purpose (e.g., dedicated conductors of a cable connecting a hand piece to the generator 200). In one form, for example, information may be communicated to and from the second data circuit using a 1-wire bus communication scheme implemented on existing cabling, such as one of the conductors used transmit interrogation signals from the signal conditioning circuit 244 to a control circuit in a hand piece. In this way, design changes or modifications to the surgical device that might otherwise be necessary are minimized or reduced. Moreover, because different types of communications implemented over a common physical channel can be frequency-band separated, the presence of a second data circuit may be "invisible" to generators that do not have the requisite data reading functionality, thus enabling backward compatibility of the surgical device instrument.

**[0068]** In certain forms, the isolated stage 202 may comprise at least one blocking capacitor 250-1 connected to the drive signal output 210b to prevent passage of DC current to a patient. A single blocking capacitor may be required to comply with medical regulations or standards, for example. While failure in single-capacitor designs is relatively uncommon, such failure may nonetheless have negative consequences. In one form, a second blocking capacitor 250-2 may be provided in series with the blocking capacitor 250-1, with current leakage from a point between the blocking capacitors 250-1, 250-2 being monitored by, for example, an ADC 252 for sampling a voltage induced by leakage current. The samples may be received by the logic device 242, for example. Based changes in the leakage current (as indicated by the voltage samples in the form of FIG. 5), the generator 200 may determine when at least one of the blocking capacitors 250-1, 250-2 has failed. Accordingly, the form of FIG. 5 provides a benefit over single-capacitor designs having a single point of failure.

**[0069]** In certain forms, the non-isolated stage 204 may comprise a power supply 254 for outputting DC power at a suitable voltage and current. The power supply may comprise, for example, a 400 W power supply for outputting a 48 VDC system voltage. The power supply 254 may further comprise one or more DC/DC voltage converters 256 for receiving the output of the power supply to generate DC outputs at the voltages and currents required by the various components of the generator 200. As discussed above in connection with the controller 238, one or more of the DC/DC voltage converters 256 may receive an input from the controller 238 when activation of the "on/off" input device 110 (FIG. 3) by a user is detected by the controller 238 to enable operation of, or wake, the DC/DC voltage converters 256.

**[0070]** With reference back to FIG. 1, having described operational details of various forms of the surgical system 100 operations for the above surgical system 100 may be further described generally in terms of a process for cutting and coagulating tissue employing a surgical instrument comprising an input device 110 and the generator 102. Although a particular process is described in connection with the operational details, it can be appreciated that the process merely provides an example of how the general functionality described herein can be implemented by the surgical system 100. Further, the given process does not necessarily have to be executed in the order presented herein unless otherwise indicated. As previously discussed, the input devices 110 may be employed to program the output (e.g., impedance, current, voltage, frequency) of the surgical instruments 104, 106, 108.

**[0071]** FIG. 6 illustrates a generator 300 comprising one form of drive system 302, according to one aspect of the present disclosure. The generator 300 is similar to the generators 102, 200 described in connection with in FIGS. 1 and 5. The generator 300 produces an ultrasonic electrical signal for driving an ultrasonic transducer, also referred to as a drive signal. The drive system 302 is flexible and can create an ultrasonic electrical output drive signal 304 at a desired frequency and power level setting for driving an ultrasonic transducer 306. In various forms, the generator 300 may comprise several separate functional elements, such as modules and/or blocks. Although certain modules, circuits, and/or blocks may be described by way of example, it can be appreciated that a greater or lesser number of modules, circuits, and/or blocks may be used and still fall within the scope of the forms. Further, although various forms may be described in terms of modules, circuits, and/or blocks to facilitate description, such modules, circuits, and/or blocks may be implemented by one or more hardware components, e.g., processors, Digital Signal Processors (DSPs), Programmable Logic Devices (PLDs), Application Specific Integrated Circuits (ASICs), circuits, registers and/or software components, e.g., programs, subroutines, logic and/or combinations of hardware and software components.

**[0072]** In one form, the drive system 302 of the generator 300 may comprise one or more embedded applications implemented as firmware, software, hardware, or any combination thereof. The drive system 302 may comprise various executable modules such as software, programs, data, drivers, application program interfaces (APIs), and so forth. The firmware may be stored in nonvolatile memory (NVM), such as in bit-masked read-only memory (ROM) or flash memory. In various implementations, storing the firmware in ROM may preserve flash memory. The NVM may comprise other types of memory including, for example, programmable ROM (PROM), erasable programmable ROM (EPROM), electrically erasable programmable ROM (EEPROM), or battery backed random-access memory (RAM) such as dynamic RAM (DRAM), Double-Data-Rate DRAM (DDRAM), and/or synchronous DRAM (SDRAM).

**[0073]** In one form, the drive system 302 comprises a hardware component implemented as a processor 308 for executing program instructions for monitoring various measurable characteristics of the ultrasonic surgical instrument 104 (FIG. 1) and generating various functions as an output signal for driving the ultrasonic transducer 306 in cutting

and/or coagulation operating modes. It will be appreciated by those skilled in the art that the generator 300 and the drive system 302 may comprise additional or fewer components and only a simplified version of the generator 300 and the drive system 302 are described herein for conciseness and clarity. In various forms, as previously discussed, the hardware component may be implemented as a DSP, PLD, ASIC, circuits, and/or registers. In one form, the processor 308 may be configured to store and execute computer software program instructions to generate the output signal functions for driving various components of the ultrasonic surgical instrument 104 (FIG. 1), such as an ultrasonic transducer 306, an end effector, and/or a blade 340.

**[0074]** In one form, under control of one or more software program routines, the processor 308 executes the methods in accordance with the described forms to generate a function formed by a stepwise waveform of drive signals comprising current (I), voltage (V), and/or frequency (f) for various time intervals or periods (T). The stepwise waveforms of the drive signals may be generated by forming a piecewise linear combination of constant functions over a plurality of time intervals created by varying the generator 300 drive signals, e.g., output drive current (I), voltage (V), and/or frequency (f). The time intervals or periods (T) may be predetermined (e.g., fixed and/or programmed by the user) or may be variable. Variable time intervals may be defined by setting the drive signal to a first value and maintaining the drive signal at that value until a change is detected in a monitored characteristic. Examples of monitored characteristics may comprise, for example, transducer impedance, tissue impedance, tissue heating, tissue transection, tissue coagulation, and the like. The ultrasonic drive signals generated by the generator 300 include, without limitation, ultrasonic drive signals capable of exciting the ultrasonic transducer 306 in various vibratory modes such as, for example, the primary longitudinal mode and harmonics thereof as well flexural and torsional vibratory modes.

**[0075]** In one form, the executable modules comprise one or more technique(s) 310 stored in memory that when executed causes the processor 308 to generate a function formed by a stepwise waveform of drive signals comprising current (I), voltage (V), and/or frequency (f) for various time intervals or periods (T). The stepwise waveforms of the drive signals may be generated by forming a piecewise linear combination of constant functions over two or more time intervals created by varying the generator 300 output drive current (I), voltage (V), and/or frequency (f). The drive signals may be generated either for predetermined fixed time intervals or periods (T) of time or variable time intervals or periods of time in accordance with the one or more technique(s) 310. Under control of the processor 308, the generator 300 varies (e.g., increment or decrement over time) the current (I), voltage (V), and/or frequency (f) up or down at a particular resolution for a predetermined period (T) or until a predetermined condition is detected, such as a change in a monitored characteristic (e.g., transducer impedance, tissue impedance). The steps can change in programmed increments or decrements. If other steps are desired, the generator 300 can increase or decrease the step adaptively based on measured system characteristics.

**[0076]** In operation, the user can program the operation of the generator 300 using the input device 312 located on the front panel of the generator 300 console. The input device 312 may comprise any suitable device that generates signals 314 that can be applied to the processor 308 to control the operation of the generator 300. In various forms, the input device 312 includes buttons, switches, thumbwheels, keyboard, keypad, touch screen monitor, pointing device, remote connection to a general purpose or dedicated computer. In other forms, the input device 312 may comprise a suitable user interface. Accordingly, by way of the input device 312, the user can set or program the current (I), voltage (V), frequency (f), and/or period (T) for programming the output function of the generator 300. The processor 308 then displays the selected power level by sending a signal on line 316 to an output indicator 318.

**[0077]** In various forms, the output indicator 318 may provide visual, audible, and/or tactile feedback to the surgeon to indicate the status of a surgical procedure, such as, for example, when tissue cutting and coagulating is complete based on a measured characteristic of the ultrasonic surgical instrument 104 (FIG. 1), e.g., transducer impedance, tissue impedance, or other measurements as subsequently described. By way of example, and not limitation, visual feedback comprises any type of visual indication device including incandescent lamps or light emitting diodes (LEDs), graphical user interface, display, analog indicator, digital indicator, bar graph display, digital alphanumeric display. By way of example, and not limitation, audible feedback comprises any type of buzzer, computer generated tone, computerized speech, voice user interface (VUI) to interact with computers through a voice/speech platform. By way of example, and not limitation, tactile feedback comprises any type of vibratory feedback provided through an instrument housing handle assembly.

**[0078]** In one form, the processor 308 may be configured or programmed to generate a digital current drive signal 320 and a digital frequency signal 322. These drive signals 320, 322 are applied to a direct digital synthesizer (DDS) circuit 324 to adjust the amplitude and the frequency (f) of the output drive signal 304 to the ultrasonic transducer 306. The output of the DDS circuit 324 is applied to an amplifier 326 whose output is applied to a transformer 328. The output of the transformer 328 is the output drive signal 304 applied to the ultrasonic transducer 306, which is coupled to the blade 340 by way of a waveguide.

**[0079]** In one form, the generator 300 comprises one or more measurement modules or components that may be configured to monitor measurable characteristics of the ultrasonic surgical instrument 104 (FIG. 1). In the illustrated form, the processor 308 may be employed to monitor and calculate system characteristics. As shown, the processor

308 measures the impedance  $Z$  of the ultrasonic transducer 306 by monitoring the current supplied to the transducer 306 and the voltage applied to the ultrasonic transducer 306. In one form, a current sense circuit 330 is employed to sense the current supplied to the ultrasonic transducer 306 and a voltage sense circuit 332 is employed to sense the output voltage applied to the ultrasonic transducer 306. These signals may be applied to the analog-to-digital converter 336 (ADC) via an analog multiplexer 334 circuit or switching circuit arrangement. The analog multiplexer 334 routes the appropriate analog signal to the ADC 336 for conversion. In other forms, multiple ADCs 336 may be employed for each measured characteristic instead of the multiplexer 334 circuit. The processor 308 receives the digital output 338 of the ADC 336 and calculates the transducer impedance  $Z$  based on the measured values of current and voltage. The processor 308 adjusts the output drive signal 304 such that it can generate a desired power versus load curve. In accordance with programmed techniques 310, the processor 308 can vary the drive signal 320, e.g., the current or frequency, in any suitable increment or decrement in response to the transducer impedance  $Z$ .

**[0080]** Having described operational details of various forms of the surgical system 100 shown in FIG. 1, operations for the above surgical system 100 may be further described in terms of a process for cutting and coagulating a blood vessel employing a surgical instrument comprising the input device 110 and the transducer impedance measurement capabilities of the drive system 302 described with reference to FIG. 6. Although a particular process is described in connection with the operational details, it can be appreciated that the process merely provides an example of how the general functionality described herein can be implemented by the surgical system 100. Further, the given process does not necessarily have to be executed in the order presented herein unless otherwise indicated.

**[0081]** FIG. 7 illustrates one aspect of a drive system of a generator 400 comprising a tissue impedance module 442. The drive system 402 generates the ultrasonic electrical drive signal 404 to drive the ultrasonic transducer 406. In one aspect, the tissue impedance module 442 may be configured to measure the impedance  $Z_t$  of tissue grasped between the blade 440 and the clamp arm assembly 444. The tissue impedance module 442 comprises an RF oscillator 446, a voltage sensing circuit 448, and a current sensing circuit 450. The voltage and current sensing circuits 448, 450 respond to the RF voltage  $V_{rf}$  applied to the blade 440 electrode and the RF current  $I_{rf}$  flowing through the blade 440 electrode, the tissue, and the conductive portion of the clamp arm assembly 444. The sensed current  $I_{rf}$  and the sensed voltage  $V_{rf}$  from the current sense circuit 430 and the voltage sense circuit 432 are converted to digital form by the ADC 436 via the analog multiplexer 434. The processor 408 receives the digitized output 438 of the ADC 436 and determines the tissue impedance  $Z_t$  by calculating the ratio of the RF voltage  $V_{rf}$  to current  $I_{rf}$  measured by the voltage sensing circuit 448 and the current sense circuit 450.

**[0082]** In one form, the processor 408 may be configured or programmed to generate a digital current signal 420 and a digital frequency signal 422. These signals 420, 422 are applied to a direct digital synthesizer (DDS) circuit 424 to adjust the amplitude and the frequency ( $f$ ) of the current output signal 404 to the transducer 406. The output of the DDS circuit 424 is applied to an amplifier 426 whose output is applied to a transformer 428. The output of the transformer 428 is the signal 404 applied to the ultrasonic transducer 406, which is coupled to the blade 440 by way of a waveguide.

**[0083]** In one aspect, the transection of the inner muscle layer and the tissue may be detected by sensing the tissue impedance  $Z_t$ . Accordingly, detection of the tissue impedance  $Z_t$  may be integrated with an automated process for separating the inner muscle layer from the outer adventitia layer prior to transecting the tissue without causing a significant amount of heating, which normally occurs at resonance.

**[0084]** In one form, the RF voltage  $V_{rf}$  applied to the blade 440 electrode and the RF current  $I_{rf}$  flowing through the blade 440 electrode, the tissue, and the conductive portion of the clamp arm assembly 444 are suitable for vessel sealing and/or dissecting. Thus, the RF power output of the generator 400 can be selected for non-therapeutic functions such as tissue impedance measurements as well as therapeutic functions such as vessel sealing and/or dissection. It will be appreciated, that in the context of the present disclosure, the ultrasonic and the RF electrosurgical energies can be supplied by the generator either individually or simultaneously.

**[0085]** In operation, the user can program the operation of the generator 400 using the input device 412 located on the front panel of the generator 400 console. The input device 412 may comprise any suitable device that generates signals 414 that can be applied to the processor 408 to control the operation of the generator 400. In various forms, the input device 412 includes buttons, switches, thumbwheels, keyboard, keypad, touch screen monitor, pointing device, remote connection to a general purpose or dedicated computer. In other forms, the input device 412 may comprise a suitable user interface. Accordingly, by way of the input device 412, the user can set or program the current ( $I$ ), voltage ( $V$ ), frequency ( $f$ ), and/or period ( $T$ ) for programming the function output of the generator 400. The processor 408 then displays the selected power level by sending a signal on line 416 to an output indicator 418.

**[0086]** In various forms, feedback is provided by the output indicator 418. The output indicator 418 is particularly useful in applications where the tissue being manipulated by the end effector is out of the user's field of view and the user cannot see when a change of state occurs in the tissue. The output indicator 418 communicates to the user that a change in tissue state has occurred. As previously discussed, the output indicator 418 may be configured to provide various types of feedback to the user including, without limitation, visual, audible, and/or tactile feedback to indicate to the user (e.g., surgeon, clinician) that the tissue has undergone a change of state or condition of the tissue. By way of example,

and not limitation, as previously discussed, visual feedback comprises any type of visual indication device including incandescent lamps or LEDs, graphical user interface, display, analog indicator, digital indicator, bar graph display, digital alphanumeric display. By way of example, and not limitation, audible feedback comprises any type of buzzer, computer generated tone, computerized speech, VUI to interact with computers through a voice/speech platform. By way of example, and not limitation, tactile feedback comprises any type of vibratory feedback provided through the instrument housing handle assembly. The change of state of the tissue may be determined based on transducer and tissue impedance measurements as previously described, or based on voltage, current, and frequency measurements.

**[0087]** In one form, the various executable modules (e.g., algorithms 410) comprising computer readable instructions can be executed by the processor 408 portion of the generator 400. In various forms, the operations described with respect to the techniques may be implemented as one or more software components, e.g., programs, subroutines, logic; one or more hardware components, e.g., processors, DSPs, PLDs, ASICs, circuits, registers; and/or combinations of software and hardware. In one form, the executable instructions to perform the techniques may be stored in memory. When executed, the instructions cause the processor 408 to determine a change in tissue state provide feedback to the user by way of the output indicator 418. In accordance with such executable instructions, the processor 408 monitors and evaluates the voltage, current, and/or frequency signal samples available from the generator 400 and according to the evaluation of such signal samples determines whether a change in tissue state has occurred. As further described below, a change in tissue state may be determined based on the type of ultrasonic instrument and the power level that the instrument is energized at. In response to the feedback, the operational mode of the ultrasonic surgical instrument may be controlled by the user or may be automatically or semi-automatically controlled.

**[0088]** As noted above, a single output generator can deliver both RF and ultrasonic energy through a single port and these signals can be delivered separately or simultaneously to the end effector to treat tissue. A single output port generator can include a single output transformer with multiple taps to provide power, either RF or ultrasonic energy, to the end effector depending on the type of treatment of tissue being performed. For example, the generator can deliver energy with higher voltage and lower current to drive an ultrasonic transducer, with lower voltage and higher current as required to drive electrodes for sealing tissue, or with a coagulation waveform for spot coagulation using either monopolar or bipolar electrosurgical electrodes. The output waveform from the generator can be steered, switched, or filtered to provide the desired frequency to the end effector of the surgical instrument.

**[0089]** FIG. 8 illustrates an example of a generator 500 for delivering multiple energy modalities to a surgical instrument. The generator 500 is similar to the generator 102 described in connection with FIG. 1 and includes functionalities of the generators 200, 300, 400 shown in FIGS. 5-7. For conciseness and clarity of disclosure, hereinbelow, the various logic flow diagrams are described in connection with the generator 500, which is a high level block diagram representation. Accordingly, the reader is directed to the description of the functional blocks of the generators 200, 300, 400 in FIGS. 5-7 for additional details that may be necessary to understand and practice the logic flow diagrams described hereinbelow in connection with the generator 500.

**[0090]** Turning back to FIG. 8, the generator 500 provides radio frequency and ultrasonic signals for delivering energy to a surgical instrument. The radio frequency and ultrasonic signals may be provided alone or in combination and may be provided simultaneously. As noted above, at least one generator output can deliver multiple energy modalities (e.g., ultrasonic, bipolar or monopolar RF, irreversible and/or reversible electroporation, and/or microwave energy, among others) through a single port and these signals can be delivered separately or simultaneously to the end effector to treat tissue. The generator 500 comprises a processor 502 coupled to a waveform generator 504. The processor 502 and waveform generator 504 are configured to generate a variety of signal waveforms based on information stored in a memory coupled to the processor 502, not shown for clarity of disclosure. The digital information associated with a waveform is provided to the waveform generator 504 which includes one or more digital-to-analog (DAC) converters to convert the digital input into an analog output. The analog output is fed to an amplifier 506 for signal conditioning and amplification. The conditioned and amplified output of the amplifier 506 is coupled to a power transformer 508. The signals are coupled across the power transformer 508 to the secondary side, which is in the patient isolation side. A first signal of a first energy modality is provided to the surgical instrument between the terminals labeled ENERGY1 and RETURN. A second signal of a second energy modality is coupled across a capacitor 510 and is provided to the surgical instrument between the terminals labeled ENERGY2 and RETURN. It will be appreciated that more than two energy modalities may be output and thus the subscript "n" may be used to designate that up to n ENERGYn terminals may be provided, where n is a positive integer greater than 1. It also will be appreciated that up to "n" return paths RETURNn may be provided without departing from the scope of the present disclosure.

**[0091]** A first voltage sensing circuit 512 is coupled across the terminals labeled ENERGY1 and the RETURN path to measure the output voltage therebetween. A second voltage sensing circuit 524 is coupled across the terminals labeled ENERGY2 and the RETURN path to measure the output voltage therebetween. A current sensing circuit 514 is disposed in series with the RETURN leg of the secondary side of the power transformer 508 as shown to measure the output current for either energy modality. If different return paths are provided for each energy modality, then a separate current sensing circuit should be provided in each return leg. The outputs of the first and second voltage sensing circuits 512,



524 are provided to respective isolation transformers 516, 522 and the output of the current sensing circuit 514 is provided to another isolation transformer 518. The outputs of the isolation transformers 516, 518, 522 in the on the primary side of the power transformer 508 (non-patient-isolated side) are provided to a one or more analog-to-digital converters 526 (ADC). The digitized output of the ADC 526 is provided to the processor 502 for further processing and computation. The output voltages and output current feedback information can be employed to adjust the output voltage and current provided to the surgical instrument and to compute output impedance, among other parameters. Input/output communications between the processor 502 and patient isolated circuits is provided through an interface circuit 520. Sensors also may be in electrical communication with the processor 502 by way of the interface 520.

**[0092]** In one aspect, the impedance may be determined by the processor 502 by dividing the output of either the first voltage sensing circuit 512 coupled across the terminals labeled ENERGY1/RETURN or the second voltage sensing circuit 524 coupled across the terminals labeled ENERGY2/RETURN by the output of the current sensing circuit 514 disposed in series with the RETURN leg of the secondary side of the power transformer 508. The outputs of the first and second voltage sensing circuits 512, 524 are provided to separate isolations transformers 516, 522 and the output of the current sensing circuit 514 is provided to another isolation transformer 516. The digitized voltage and current sensing measurements from the ADC 526 are provided the processor 502 for computing impedance. As an example, the first energy modality ENERGY1 may be ultrasonic energy and the second energy modality ENERGY2 may be RF energy. Nevertheless, in addition to ultrasonic and bipolar or monopolar RF energy modalities, other energy modalities include irreversible and/or reversible electroporation and/or microwave energy, among others. Also, although the example illustrated in FIG. 8 shows a single return path RETURN may be provided for two or more energy modalities, in other aspects multiple return paths RETURNn may be provided for each energy modality ENERGYn. Thus, as described herein, the ultrasonic transducer impedance may be measured by dividing the output of the first voltage sensing circuit 512 by the current sensing circuit 514 and the tissue impedance may be measured by dividing the output of the second voltage sensing circuit 524 by the current sensing circuit 514.

**[0093]** As shown in FIG. 8, the generator 500 comprising at least one output port can include a power transformer 508 with a single output and with multiple taps to provide power in the form of one or more energy modalities, such as ultrasonic, bipolar or monopolar RF, irreversible and/or reversible electroporation, and/or microwave energy, among others, for example, to the end effector depending on the type of treatment of tissue being performed. For example, the generator 500 can deliver energy with higher voltage and lower current to drive an ultrasonic transducer, with lower voltage and higher current to drive RF electrodes for sealing tissue, or with a coagulation waveform for spot coagulation using either monopolar or bipolar RF electrosurgical electrodes. The output waveform from the generator 500 can be steered, switched, or filtered to provide the frequency to the end effector of the surgical instrument. The connection of an ultrasonic transducer to the generator 500 output would be preferably located between the output labeled ENERGY1 and RETURN as shown in FIG. 8. An In one example, a connection of RF bipolar electrodes to the generator 500 output would be preferably located between the output labeled ENERGY2 and RETURN. In the case of monopolar output, the preferred connections would be active electrode (e.g., pencil or other probe) to the ENERGY2 output and a suitable return pad connected to the RETURN output.

**[0094]** In other aspects, the generators 102, 200, 300, 400, 500 described in connection with FIGS. 1-3 and 5-8, the ultrasonic generator drive circuit 114, and/or electrosurgery/RF drive circuit 116 as described in connection with FIG. 3 may be formed integrally with any one of the surgical instruments 104, 106, 108 described in connection with FIGS. 1 and 2. Accordingly, any of the processors, digital signal processors, circuits, controllers, logic devices, ADCs, DACs, amplifiers, converters, transformers, signal conditioners, data interface circuits, current and voltage sensing circuits, direct digital synthesis circuits, multiplexer (analog or digital), waveform generators, RF generators, memory, and the like, described in connection with any one of the generators 102, 200, 300, 400, 500 can be located within the surgical instruments 104, 106, 108 or may be located remotely from the surgical instruments 104, 106, 108 and coupled to the surgical instruments via wired and/or wireless electrical connections.

**[0095]** Examples of waveforms representing energy for delivery from a generator are illustrated in FIGS. 9-13. FIG. 9 illustrates an example graph 600 showing first and second individual waveforms representing an RF output signal 602 and an ultrasonic output signal 604 superimposed on the same time and voltage scale for comparison purposes. These output signals 602, 604 are provided at the ENERGY output of the generator 500 shown in FIG. 8. Time (t) is shown along the horizontal axis and voltage (V) is shown along the vertical axis. The RF output signal 602 has a frequency of about 330kHz RF and a peak-to-peak voltage of  $\pm 1V$ . The ultrasonic output signal 604 has a frequency of about 55kHz and a peak-to-peak voltage of  $\pm 1V$ . It will be appreciated that the time (t) scale along the horizontal axis and the voltage (V) scale along the vertical axis are normalized for comparison purposes and may be different actual implementations, or represent other electrical parameters such as current.

**[0096]** FIG. 10 illustrates an example graph 610 showing the sum of the two output signals 602, 604 shown in FIG. 9. Time (t) is shown along the horizontal axis and voltage (V) is shown along the vertical axis. The sum of the RF output signal 602 and the ultrasonic output signal 604 shown in FIG. 9 produces a combined output signal 612 having a 2V peak-to-peak voltage, which is twice the amplitude of the original RF and ultrasonic signals shown (1V peak-to-peak)

shown in FIG. 9. An amplitude of twice the original amplitude can cause problems with the output section of the generator, such as distortion, saturation, clipping of the output, or stresses on the output components. Thus, the management of a single combined output signal 612 that has multiple treatment components is an important aspect of the generator 500 shown in FIG. 8. There are a variety of ways to achieve this management. In one form, one of the two RF or ultrasonic output signals 602, 604 can be dependent on the peaks of the other output signal.

**[0097]** For example, FIG. 11 illustrates an example graph 620 showing a combined output signal 622 representative of a dependent sum of the output signals 602, 604 shown in FIG. 9. Time (t) is shown along the horizontal axis and voltage (V) is shown along the vertical axis. As shown in FIG. 11, the RF output signal 602 component of FIG. 9 depends on the peaks of the ultrasonic output signal 604 component of FIG. 9 such that the amplitude of the RF output signal component of the dependent sum combined output signal 622 is reduced when an ultrasonic peak is anticipated. As shown in the example graph 620 in FIG. 11, the peaks have been reduced from 2 to 1.5. In another form, one of the output signals is a function of the other output signal.

**[0098]** For example, FIG. 11 illustrates an example graph 630 showing an output signal 632 representative of a dependent sum of the output signals 602, 604 shown in FIG. 9. Time (t) is shown along the horizontal axis and voltage (V) is shown along the vertical axis. As shown in FIG. 12, the RF output signal is a function of the ultrasonic output signal. This provides a hard limit on the amplitude of the output. As shown in FIG. 12, the ultrasonic output signal is extractable as a sine wave while the RF output signal has distortion but not in a way to affect the coagulation performance of the RF output signal.

**[0099]** A variety of other techniques can be used for compressing and/or limiting the waveforms of the output signals. It should be noted that the integrity of the ultrasonic output signal 604 (FIG. 9) can be more important than the integrity of the RF output signal 602 (FIG. 9) as long as the RF output signal 602 has low frequency components for safe patient levels so as to avoid neuro-muscular stimulation. In another form, the frequency of an RF waveform can be changed on a continuous basis in order to manage the peaks of the waveform. Waveform control is important as more complex RF waveforms, such as a coagulation-type waveform 644, as illustrated in the graph 640 shown in FIG. 13, are implemented with the system. Again, time (t) is shown along the horizontal axis and voltage (V) is shown along the vertical axis.

**[0100]** FIGS. 14-42 (26-54) illustrate various configurations of sensors, circuits, and techniques for measuring tissue parameters to facilitate executing the various adaptive tissue identification and treatment technique described herein. FIG. 14 illustrates one aspect of an end effector 700 comprising RF data sensors 706, 708a, 708b located on the clamp arm 702. The end effector 700 comprises a clamp arm 702 and an ultrasonic blade 704. The clamp arm 702 is shown clamping tissue 710 located between the clamp arm 702 and the ultrasonic blade 704. A first sensor 706 is located in a center portion of the clamp arm 702. Second and third sensors 708a, 708b are located on lateral portions of the clamp arm 702. The sensors 706, 708a, 708b are mounted or formed integrally with on a flexible circuit 712 (shown more particularly in FIG. 15 and more particularly segmented flexible circuits 800, 900 shown in FIGS. 17 and 18) configured to be fixedly mounted to the clamp arm 702.

**[0101]** The end effector 700 is an example end effector for the multifunction surgical instrument 108 shown in FIGS. 1 and 2. The sensors 706, 708a, 708b are electrically connected to an energy source, such as for example, the generator 500 shown in FIG. 8. The sensors 706, 708a, 708b are powered by suitable sources within the generator and the signals generated by the sensors 706, 708a, 708b are provided to analog and/or digital processing circuits of the generator 500.

**[0102]** In one aspect, the first sensor 706 is a force sensor to measure a normal force  $F_3$  applied to the tissue 710 by the clamp arm 702. The second and third sensors 708a, 708b include one or more elements to apply RF energy to the tissue 710, measure tissue impedance, down force  $F_1$ , transverse forces  $F_2$ , and temperature, among other parameters. Electrodes 709a, 709b are electrically coupled to the generator and apply RF energy to the tissue 710. In one aspect, the first sensor 706 and the second and third sensors 708a, 708b are strain gauges to measure force or force per unit area. It will be appreciated that the measurements of the down force  $F_1$ , the lateral forces  $F_2$ , and the normal force  $F_3$  may be readily converted to pressure by determining the surface area upon which the force sensors 706, 708a, 708b are acting upon. Additionally, as described with particularity herein, the flexible circuit 712 may comprise temperature sensors embedded in one or more layers of the flexible circuit 712. The one or more temperature sensors may be arranged symmetrically or asymmetrically and provide tissue 710 temperature feedback to control circuits of the generator.

**[0103]** FIG. 15 illustrates one aspect of the flexible circuit 712 shown in FIG. 14 in which the sensors 706, 708a, 708b may be mounted to or formed integrally therewith. The flexible circuit 712 is configured to fixedly attach to the clamp arm 702. As shown particularly in FIG. 15, asymmetric temperature sensors 714a, 714b are mounted to the flexible circuit 712 to enable measuring the temperature of the tissue 710 (FIG. 14).

**[0104]** FIG. 16 is a cross-sectional view of the flexible circuit 712 shown in FIG. 15. The flexible circuit 712 comprises multiple layers and is fixedly attached to the clamp arm 702. A top layer of the flexible circuit 712 is an electrode 709a, which is electrically coupled to an energy source, such as the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), to apply RF energy to the tissue 710 (FIG. 14). A layer of electrical insulation 718 is provided below the electrode 709a layer to electrically isolate the sensors 714a, 706, 708a from the electrode 709a. The temperature sensors 714a are

disposed below the layer of electrical insulation 718. The first force (pressure) sensor 706 is located below the layer containing the temperature sensors 714a and above a compressive layer 720. The second force (pressure) sensor 708a is located below the compressive layer 720 and above the clamp arm 702 frame.

**[0105]** FIG. 17 illustrates one aspect of a segmented flexible circuit 800 configured to fixedly attach to a clamp arm 804 of an end effector. The segmented flexible circuit 800 comprises a distal segment 802a and lateral segments 802b, 802c that include individually addressable sensors to provide local tissue control, as described herein in connection with FIGS. 14-16, for example. The segments 802a, 802b, 802c are individually addressable to treat tissue and to measure tissue parameters based on individual sensors located within each of the segments 802a, 802b, 802c. The segments 802a, 802b, 802c of the segmented flexible circuit 800 are mounted to the clamp arm 804 and are electrically coupled to an energy source, such as the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), via electrical conductive elements 806. A Hall effect sensor 808, or any suitable magnetic sensor, is located on a distal end of the clamp arm 804. The Hall effect sensor 808 operates in conjunction with a magnet to provide a measurement of an aperture defined by the clamp arm 804, which otherwise may be referred to as a tissue gap, as shown with particularity in FIG. 19.

**[0106]** FIG. 18 illustrates one aspect of a segmented flexible circuit 900 configured to mount to a clamp arm 904 of an end effector. The segmented flexible circuit 900 comprises a distal segment 902a and lateral segments 902b, 902c that include individually addressable sensors for tissue control, as described herein in connection with FIGS. 14-17, for example. The segments 902a, 902b, 902c are individually addressable to treat tissue and to read individual sensors located within each of the segments 902a, 902b, 902c. The segments 902a, 902b, 902c of the segmented flexible circuit 900 are mounted to the clamp arm 904 and are electrically coupled to an energy source, such as the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), via electrical conductive elements 906. A Hall effect sensor 908, or other suitable magnetic sensor, is provided on a distal end of the clamp arm 904. The Hall effect sensor 908 operates in conjunction with a magnet to provide a measurement of an aperture defined by the clamp arm 904 of the end effector or tissue gap as shown with particularity in FIG. 19. In addition, a plurality of lateral asymmetric temperature sensors 910a, 910b are mounted on or formally integrally with the segmented flexible circuit 900 to provide tissue temperature feedback to control circuits in the generator.

**[0107]** FIG. 19 illustrates one aspect of an end effector 1000 configured to measure a tissue gap  $G_T$ . The end effector 1000 comprises a jaw member 1002 and a clamp arm 904. The flexible circuit 900 as described in FIG. 18, is mounted to the clamp arm 904. The flexible circuit 900 comprises a Hall effect sensor 908 that operates with a magnet 1004 mounted to the jaw member 1002 to measure the tissue gap  $G_T$ . This technique can be employed to measure the aperture defined between the clamp arm 904 and the jaw member 1002. The jaw member 1002 may be an ultrasonic blade.

**[0108]** FIG. 20 illustrates one aspect of a left-right segmented flexible circuit 1100. The left-right segmented flexible circuit 1100 comprises a plurality of segments L1-L5 on the left side of the left-right segmented flexible circuit 1100 and a plurality of segments R1-R5 on the right side of the left-right segmented flexible circuit 1100. Each of the segments L1-L5 and R1-R5 comprise temperature sensors and force sensors to sense tissue parameters locally within each segment L1-L5 and R1-R5. The left-right segmented flexible circuit 1100 are configured to influence the RF treatment energy based on tissue parameters sensed locally within each of the segments L1-L5 and R1-R5.

**[0109]** FIG. 21 illustrates one aspect of an end effector 1200 comprising segmented flexible circuit 1100 as shown in FIG. 20. The end effector 1200 comprises a clamp arm 1202 and an ultrasonic blade 1204. The segmented flexible circuit 1100 is mounted to the clamp arm 1202. Each of the sensors disposed within the segments 1-5 are configured to detect the presence of tissue positioned between the clamp arm 1202 and the ultrasonic blade 1204 and represent tissue zones 1-5. In the configuration shown in FIG. 21, the end effector 1200 is shown in an open position ready to receive or grasp tissue between the clamp arm 1202 and the ultrasonic blade 1204.

**[0110]** FIG. 22 illustrates the end effector 1200 shown in FIG. 21 with the clamp arm 1202 clamping tissue 1206 between the clamp arm 1202 and the ultrasonic blade 1204. As shown in FIG. 22, the tissue 1206 is positioned between segments 1-3 and represents tissue zones 1-3. Accordingly, tissue 1206 is detected by the sensors in segments 1-3 and the absence of tissue (empty) is detected in section 1208 by segments 4-5. The information regarding the presence and absence of tissue 1206 positioned within certain segments 1-3 and 4-5, respectively, is communicated to a control circuit of the generator, such as the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8). The generator 500 is configured to energize only the segments 1-3 where tissue 1206 is detected and does not energize the segments 4-5 where tissue is not detected. It will be appreciated that the segments 1-5 may contain any suitable temperature, force/pressure, and/or Hall effect magnetic sensors to measure tissue parameters of tissue located within certain segments 1-5 and electrodes to deliver RF energy to tissue located in certain segments 1-5.

**[0111]** FIG. 23 illustrates graphs 1300 of energy applied by the right and left side of an end effector based on locally sensed tissue parameters. As discussed herein, the clamp arm of an end effector may comprise temperature sensors, force/pressure sensors, Hall effector sensors, among others, along the right and left sides of the clamp arm as shown, for example, in FIGS. 14-22. Thus, RF energy can be selectively applied to tissue positioned between the clam jaw and the ultrasonic blade. The top graph 1302 depicts power  $P_R$  applied to a right side segment of the clamp arm versus time (t) based on locally sensed tissue parameters. Thus, the generator, such as the generators 102, 200, 300, 400, 500

(FIGS. 1-3 and 4-8), is configured to measure the sensed tissue parameters and to apply power  $P_R$  to a right side segment of the clamp arm. The generator 500 delivers an initial power level  $P_1$  to the tissue via the right side segment and then decreases the power level to  $P_2$  based on local sensing of tissue parameters (e.g., temperature, force/pressure, thickness) in one or more segments. The bottom graph 1304 depicts power  $P_L$  applied to a left side segment of the clamp arm versus time (t) based on locally sensed tissue parameters. The generator 500 delivers an initial power level of  $P_1$  to the tissue via the left side segment and then increases the power level to  $P_3$  based on local sensing of tissue parameters (e.g., temperature, force/pressure, thickness). As depicted in the bottom graph 1304, the generator is configured to re-adjust the energy delivered  $P_3$  based on sensing of tissue parameters (e.g., temperature, force/pressure, thickness).

**[0112]** FIG. 24 illustrates a graph 1400 depicting one aspect of adjustment of threshold due to the measurement of a secondary tissue parameter such as continuity, temperature, pressure, and the like. The horizontal axis of the graph 1400 is time (t) and the vertical axis is tissue impedance (Z). The curve 1412 represents the change of tissue impedance (Z) over time (t) as different energy modalities are applied to the tissue. With reference also to FIGS. 20-22, the original threshold 1402 is applied when tissue is detected in all five segments 1-5 (tissue zones 1-5) and the adjusted threshold 1404 is applied when the tissue is detected in tissue segments 1-3 (tissue zones 1-3). Accordingly, once the tissue is located in particular segments (zones) the control circuit in the generator adjusts the threshold accordingly.

**[0113]** As shown in FIG. 24, the curve 1412 includes three separate sections 1406, 1408, 1410. The first section 1406 of the curve 1412 represents the time when RF energy is applied to the tissue in tissue zones 1-3 until the tissue impedance drops below the adjusted threshold 1404. At that point 1414, which may indicate that a tissue seal is completed, the energy modality applied to tissue zones 1-3 is changed from RF energy to ultrasonic energy. The ultrasonic energy is then applied in the second and third sections 1408, 1410 and the impedance rises exponentially until the tissue is severed or cut.

**[0114]** FIG. 25 is a cross-sectional view of one aspect of a flexible circuit 1500 comprising RF electrodes and data sensors embedded therein. The flexible circuit 1500 can be mounted to the right or left portion of an RF clamp arm 1502, which is made of electrically conductive material such as metal. Below the RF clamp arm 1502 down force/pressure sensors 1506a, 1506b are embedded below a laminate layer 1504. A transverse force/pressure sensor 1508 is located below the down force/pressure sensor 1506a, 1506b layer and a temperature sensor 1510 is located below the transverse force/pressure sensor 1508. An electrode 1512 electrically coupled to the generator and configured to apply RF energy to the tissue 1514 is located below the temperature sensor 1510.

**[0115]** FIG. 26 is a cross-sectional view of one aspect of an end effector 1600 configured to sense force or pressure applied to tissue located between a clamp arm and an ultrasonic blade. The end effector 1600 comprises a clamp jaw 1602 and a flexible circuit 1604 fixedly mounted to the clamp arm 1602. The clamp arm 1602 applies forces  $F_1$  and  $F_2$  to the tissue 1606 of variable density and thickness, which can be measure by first and second force/pressure sensors 1608, 1610 located in different layers of the flexible circuit 1604. A compressive layer 1612 is sandwiched between the first and second force/pressure sensors 1608, 1610. An electrode 1614 is located on outer portion of the flexible circuit 1604 which contacts the tissue. As described herein, other layers of the flexible circuit 1604 may comprise additional sensors such temperature sensors, thickness sensors, and the like.

**[0116]** FIGS. 27-29 illustrate various schematic diagrams of flexible circuits of the signal layer, sensor wiring, and an RF energy drive circuit. FIG. 27 is a schematic diagram of one aspect of a signal layer of a flexible circuit 1700. The flexible circuit 1700 comprises multiple layers (~4 to ~6, for example). One layer will supply the integrated circuits with power and another layer with ground. Two additional layers will carry the RF power RF1 and RF2 separately. An analog multiplexer switch 1702 has eight bidirectional translating switches that can be controlled through the I<sup>2</sup>C bus. The SCL/SDA upstream pair fans out to eight downstream pairs, or channels. Any individual SCn/SDn channel or combination of channels can be selected, determined by the contents of a programmable control register. The upstream pairs SCL/SDA are connected to a control circuit in the generator. There are six down stream sensors, three on each side of the clamp arm. A first side 1704a comprises a first thermocouple 1706a, a first pressure sensor 1708a, and a first Hall effect sensor 1710a. A second side 1704b comprises a second thermocouple 1706b, a second pressure sensor 1708b, and a second Hall effect sensor 1710b. FIG. 28 is a schematic diagram 1750 of sensor wiring for the flexible circuit 1700 shown in FIG. 27.

**[0117]** FIG. 29 is a schematic diagram of one aspect of an RF energy drive circuit 1800. The RF energy drive circuit 1800 comprises an analog multiplexer 1702 described in connection with FIG. 27. The analog multiplexer multiplexes various signals from the upstream channels SCL/SDA. A current sensor 1802 is coupled in series with the return or ground leg of the power supply circuit to measure the current supplied by the power supply. An FET temperature sensor 1804 provided the ambient temperature. A pulse width modulation (PWM) watchdog timer 1808 automatically generates a system reset if the main program neglects to periodically service it. It is provided to automatically reset the RF energy drive circuit 1800 when it hangs because of a software or hardware fault.

**[0118]** A drive circuit 1806 provides left and right RF energy outputs. The digital signal is provided to the SCL/SDA inputs of the analog multiplexer 1702 from a control circuit of the generator. A digital-to-analog converter (DAC) converts the digital input to an analog output to drive a pulse width modulation (PWM) circuit 1812 coupled to an oscillator 1814.

The PWM circuit 1812 provides a first gate drive signal 1816a to a first transistor output stage 1818a to drive a first RF (Left) energy output. The PWM circuit 1812 also provides a second gate drive signal 1816b to a second transistor output stage 1818 to drive a second RF (Right) energy output.

**[0119]** The circuits 1700, 1750, 1800 described in connection with FIGS. 27-29 are electrically coupled to the generators 200, 300, 400, 500 shown in FIGS. 5-7. For example, the circuits 1700, 1750, 1800 may be coupled to the generator 200 via the signal conditioning circuit 244 and may be coupled to the generator 500 through the interface circuit 520.

**[0120]** FIG. 30 is a graphical representation 1900 of measuring tissue gap at a preset time. A first graph 1902 represents tissue impedance  $Z$  versus time ( $t$ ) where the horizontal axis represents time ( $t$ ) and the vertical axis represents tissue impedance  $Z$ . A second graph 1904 represents change in tissue gap  $\Delta_{\text{gap}}$  versus time ( $t$ ) where the horizontal axis represents time ( $t$ ) and the vertical axis represents change in tissue gap  $\Delta_{\text{gap}}$ . A third graph 1906 represents force  $F$  versus time ( $t$ ) where the horizontal axis represents time ( $t$ ) and the vertical axis represents force  $F$ . With a constant force  $F$  applied to tissue and impedance  $Z$  interrogation to define a wait period, energy modality (e.g., RF and ultrasonic) and motor control parameters, displacement at a time provides velocity. With reference to the three graphs 1902, 1904, 1906, impedance sensing energy is applied during a first period 1908 to determine the tissue type such as thin mesentery tissue (solid line), intermediate thickness vessel tissue (dashed line), or thick uterus/bowel tissue (dash-dot line).

**[0121]** As shown in the third graph 1906, the clamp arm initially applies a force which ramps up from zero exponentially until it reaches a constant force 1924. The preset time  $t_1$  is selected such that it occurs some time after the clamp arm force reaches a constant force 1924. As shown in the first and second graphs 1902, 1904, from the time the clamp force is applied to the mesentery tissue until the preset time  $t_1$  is reached, the change in tissue gap  $\Delta_{\text{gap}}$  curve 1912 decreases exponentially and the tissue impedance curve 1918 also decreases until the preset time  $t_1$  is reached. From the preset time  $t_1$ , a short delay 1928 is applied before treatment energy is applied to the mesentery tissue at  $t_{E1}$ .

**[0122]** As shown in the first and second graphs 1902, 1904, from the time the clamp force is applied to the vessel tissue until the preset time  $t_1$  is reached, the change in tissue gap  $\Delta_{\text{gap}}$  curve 1916 also decrease exponentially and the tissue impedance curve 1920 also decreases until the preset time  $t_1$  is reached. From the preset time  $t_1$ , a medium delay 1930 is applied before treatment energy is applied to the vessel tissue at  $t_{E2}$ .

**[0123]** As shown in the first and second graphs 1902, 1904, from the time the clamp force is applied to the uterus/bowel tissue until the preset time  $t_1$  is reached, the change in tissue gap  $\Delta_{\text{gap}}$  curve 1914 drops exponentially and the tissue impedance curve 1914 also drops until the preset time  $t_1$  is reached. From the preset time  $t_1$ , a short delay 1928 is applied before treatment energy is applied to the mesentery tissue at  $t_{E1}$ .

**[0124]** FIG. 31 is a time to preset force 2008 versus time graph 2000 for thin, medium, and thick tissue types. The horizontal axis represents time ( $t$ ) and the vertical axis represents force ( $F$ ) applied by the clamp arm to the tissue. The graph 2000 depicts three curves, one for thin tissue 2002 shown in solid line, one for medium thickness tissue 2004 shown in dash-dot line, and thick tissue 2006 in dashed line. The graph 2000 depicts measuring time at a preset force as an alternative to tissue gap to control delayed energy mode and other control parameters. Accordingly, the time to preset force 2008 for thick tissue 2006 is  $t_{1a}$ , the time to preset force 2008 for medium thickness tissue 2004 is  $t_{1b}$ , and the time to preset force 2008 for thin tissue 2002 is  $t_{1c}$ .

**[0125]** Once the force reaches the preset force 2008, energy is applied to the tissue. For thin tissue 2002 the time to preset  $t_{1c} > 0.5$  seconds and then RF energy is applied for an energizing period  $t_e$  of about 1-3 seconds. For thick tissue 2006 the time to preset  $t_{1a} < 0.5$  seconds and then RF energy is applied for an energizing period  $t_e$  of about 5-9 seconds. For medium thickness tissue 2004 the time to preset  $t_{1b}$  is about 0.5 seconds and then RF energy is applied for an energizing period  $t_e$  of about 3 to 5 seconds.

**[0126]** FIG. 32 is a graphical depiction of a graph 2100 of three curves 2102, 2104, 2106, where the first curve 2102 represents power ( $P$ ), voltage ( $V_{\text{RF}}$ ), and current ( $I_{\text{RF}}$ ) versus tissue impedance ( $Z$ ), the second curve 2104 and third curve 2106 represent tissue impedance ( $Z$ ) versus time ( $t$ ). The first curve 2102 illustrates the application of power ( $P$ ) for thick tissue impedance range 2110 and thin tissue impedance range 2112. As the tissue impedance  $Z$  increases, the current  $I_{\text{RF}}$  decrease and the voltage  $V_{\text{RF}}$  increases. The power curve  $P$  increases until it reaches a maximum power output 2108 which coincides with the intersection 2114 of the current  $I_{\text{RF}}$  and voltage  $V_{\text{RF}}$  curves.

**[0127]** The second curve 2104 represents the measured tissue impedance  $Z$  versus time ( $t$ ). The tissue impedance threshold limit 2120 is the cross over limit for switching between the RF and ultrasonic energy modalities. For example, as shown in FIG. 32, RF energy is applied while the tissue impedance is above the tissue impedance threshold limit 2120 and ultrasonic energy 2124 is applied while the tissue impedance is below the tissue impedance threshold limit 2120. Accordingly, with reference back to the second curve 2104, the tissue impedance of the thin tissue curve 2116 remains above the tissue impedance threshold limit 2120, thus only RF energy modality is applied to the tissue. On the other hand, RF energy modality is applied to the thick tissue while the impedance is above the tissue impedance threshold limit 2120 and ultrasonic energy is applied to the tissue when the impedance is below the tissue impedance threshold limit 2120. Accordingly, the energy modality switches from RF to ultrasonic when the tissue impedance falls below the tissue impedance threshold limit 2120 and the energy modality switches from ultrasonic to RF when the tissue impedance rises above the tissue impedance threshold limit 2120.

**[0128]** FIG. 33 is a plan view of one aspect of an end effector 2200. The end effector 2200 comprises a clamp arm 2202 and a shaft 2204. The clamp arm 2202 pivots about pivot point 2206 and defines a pivot angle. FIG. 34 is a side view of the end effector 2200 shown in FIG. 33 with a partial cut away view to expose the underlying structure of the clamp arm 2202 and an ultrasonic blade 2208. An electrode 2210 is fixedly mounted to the clamp arm 2202. The electrode 2210 is electrically coupled to the generator and is configured to apply RF energy to tissue located between the clamp arm 2202 and the ultrasonic blade 2208. FIG. 35 is partial sectional view of the end effector shown in FIGS. 33, 34 to expose the ultrasonic blade and right and left electrodes 2210a, 2210b, respectively.

**[0129]** FIG. 36 is a cross-sectional view taken at section 36--36 of the end effector 2200 shown in FIG. 33. The end effector 2200 comprises an ultrasonic blade 2208 acoustically coupled to an ultrasonic transducer which is electrically driven by the generator. The clamp arm 2202 comprises an electrode 2210a on the right side and an electrode 2210b on the left side (from the perspective of the operator). The right side electrode 2210a defines a first width  $W_1$  and defines a first gap  $G_1$  between the electrode 2210a and the ultrasonic blade 2208. The left side electrode 2210b defines a second width  $W_2$  and defines a second gap  $G_2$  between the electrode 2210b and the ultrasonic blade 2208. In one aspect the first width  $W_1$  is less than the second width  $W_2$  and the first gap  $G_1$  is less than the second gap  $G_2$ . With reference also to FIG. 35, a soft polymeric pad 2212 is located between the ultrasonic blade 2208 and the clamp arm 2202. A high density polymeric pad 2214 is located adjacent the soft polymeric pad 2212 to prevent the ultrasonic blade 2208 from shorting the electrodes 2210a, 2210b. In one aspect, the soft polymeric pads 2212, 2214 can be made of polymers known under the tradename TEFLON (polytetrafluoroethylene polymers and copolymers), for example.

**[0130]** FIG. 37 is cross-sectional view taken at section 37--37 of the end effector 2200 shown in FIG. 33. At the plane where section 37--37 the end effector 2200 is thinner and has more curvature than section 36--36. The right side electrode 2210a defines a third width  $W_3$  and defines a third gap  $G_3$  between the electrode 2210a and the ultrasonic blade 2208. The left side electrode 2210b defines a fourth width  $W_4$  and defines a fourth gap  $G_4$  between the electrode 2210b and the ultrasonic blade 2208. In one aspect the third width  $W_3$  is less than the fourth width  $W_4$  and the third gap  $G_3$  is less than the fourth gap  $G_4$ .

**[0131]** FIG. 38 is a cross-sectional view taken at section 36--36 of the end effector 2200 shown in FIG. 33, except that the ultrasonic blade 2208' has a different geometric configuration. The end effector 2200' comprises an ultrasonic blade 2208' acoustically coupled to an ultrasonic transducer which is electrically driven by the generator. The clamp arm 2202' comprises an electrode 2210a' on the right side and an electrode 2210b' on the left side (from the perspective of the operator). The right side electrode 2210a' defines a first width  $W_1$  and defines a first gap  $G_1$  between the electrode 2210a' and the ultrasonic blade 2208'. The left side electrode 2210b' defines a second width  $W_2$  and defines a second gap  $G_2$  between the electrode 2210b' and the ultrasonic blade 2208'. In one aspect the first width  $W_1$  is less than the second width  $W_2$  and the first gap  $G_1$  is less than the second gap  $G_2$ . A high density polymeric pad 2214' is located adjacent the soft polymeric pad 2212' to prevent the ultrasonic blade 2208' from shorting the electrodes 2210a', 2210b'. In one aspect, the soft polymeric pads 2212', 2214' can be made of polymers known under the tradename TEFLON (polytetrafluoroethylene polymers and copolymers), for example.

**[0132]** FIG. 39 is cross-sectional view taken at section 37--37 of the end effector 2200 shown in FIG. 33, except that the ultrasonic blade 2208' has a different geometric configuration. At the plane where section 37--37 the end effector 2200' is thinner and has more curvature than the end effector 2200' at section 36--36. The right side electrode 2210a' defines a third width  $W_3$  and defines a third gap  $G_3$  between the electrode 2210a' and the ultrasonic blade 2208'. The left side electrode 2210b' defines a fourth width  $W_4$  and defines a fourth gap  $G_4$  between the electrode 2210b' and the ultrasonic blade 2208'. In one aspect the third width  $W_3$  is less than the fourth width  $W_4$  and the third gap  $G_3$  is less than the fourth gap  $G_4$ .

**[0133]** FIG. 40 is a cross-sectional view taken at section 36--36 of the end effector 2200 shown in FIG. 33, except that the ultrasonic blade 2208" has a different geometric configuration. The end effector 2200" comprises an ultrasonic blade 2208" acoustically coupled to an ultrasonic transducer which is electrically driven by the generator. The clamp arm 2202" comprises an electrode 2210a" on the right side and an electrode 2210b" on the left side (from the perspective of the operator). The right side electrode 2210a" defines a first width  $W_1$  and defines a first gap  $G_1$  between the electrode 2210a" and the ultrasonic blade 2208". The left side electrode 2210b" defines a second width  $W_2$  and defines a second gap  $G_2$  between the electrode 2210b" and the ultrasonic blade 2208". In one aspect the first width  $W_1$  is less than the second width  $W_2$  and the first gap  $G_1$  is less than the second gap  $G_2$ . A high density polymeric pad 2214" is located adjacent the soft polymeric pad 2212" to prevent the ultrasonic blade 2208" from shorting the electrodes 2210a", 2210b". In one aspect, the polymeric pads 2212", 2214" can be made of polymers known under the tradename TEFLON (polytetrafluoroethylene polymers and copolymers), for example.

**[0134]** FIG. 41 is cross-sectional view taken at section 37--37 of the end effector 2200 shown in FIG. 33, except that the ultrasonic blade 2208" has a different geometric configuration. At the plane where section 37--37 the end effector 2200" is thinner and has more curvature than the end effector 2200" at section 36--36. The right side electrode 2210a" defines a third width  $W_3$  and defines a third gap  $G_3$  between the electrode 2210a" and the ultrasonic blade 2208". The left side electrode 2210b" defines a fourth width  $W_4$  and defines a fourth gap  $G_4$  between the electrode 2210b" and the

ultrasonic blade 2208". In one aspect the third width  $W_3$  is less than the fourth width  $W_4$  and the third gap  $G_3$  is less than the fourth gap  $G_4$ .

**[0135]** The surgical instruments described herein also can include features to allow the energy being delivered by the generator to be dynamically changed based on the type of tissue being treated by an end effector of a surgical instrument and various characteristics of the tissue. In one aspect, a technique for controlling the power output from a generator, such as the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), that is delivered to the end effector of the surgical instrument can include an input that represents the tissue type to allow the energy profile from the generator to be dynamically changed during the procedure based on the type of tissue being effected by the end effector of the surgical instrument.

**[0136]** As disclosed herein, techniques for controlling a generator based on the tissue type may be provided. Various techniques can be used to select a power profile to allow the energy being delivered from the generator to dynamically change based on the tissue type being treated by the surgical instrument.

**[0137]** FIG. 42A illustrates an end effector 2300 comprising a clamp arm 2302 and an ultrasonic blade 2304, where the clamp arm 2302 includes electrodes 2306. The end effector 2300 can be employed in one of the surgical instruments 104, 106, 108 referred to in FIGS. 1-3. In addition to the end effector 122, 124, 125, the surgical instruments 104, 106, 108 include a handpiece 105, 107, 109 and a shaft 126, 127, 129, respectively. The end effectors 122, 124, 125 may be used to compress, cut, and/or seal tissue. Referring to FIG. 42A, the end effector 2300, similar to the end effectors 122, 124, 125 shown in FIGS. 1-3, may be positioned by a physician to surround tissue 2308 prior to compression, cutting, or stapling. As shown in FIG. 42A, no compression may be applied to the tissue while preparing to use the end effector 2300. As shown in FIG. 42A, the tissue 2308 is not under compression between the clamp arm 2302 and the ultrasonic blade 2304.

**[0138]** Referring now to FIG. 42B, by engaging the trigger on the handle of a surgical instrument, the physician may use the end effector 2300 to compress the tissue 2308. In one aspect, the tissue 2308 may be compressed to its maximum threshold, as shown in FIG. 42B. As shown in FIG. 42A, the tissue 2308 is under maximum compression between the clamp arm 2302 and the ultrasonic blade 2304.

**[0139]** Referring to FIG. 43A, various forces may be applied to the tissue 2308 by the end effector 2300. For example, vertical forces  $F_1$  and  $F_2$  may be applied by the clamp arm 2302 and the ultrasonic blade 2304 of the end effector 2300 as tissue 2308 is compressed between the two. Referring now to FIG. 43B, various diagonal and/or lateral forces also may be applied to the tissue 2308 when compressed by the end effector 2300. For example, a force  $F_3$  may be applied. For the purposes of operating a medical device such as the surgical instruments 104, 106, 108 it may be desirable to sense or calculate the various forms of compression being applied to the tissue by the end effector. For example, knowledge of vertical or lateral compression may allow the end effector to more precisely or accurately apply a staple operation or may inform the operator of the surgical instrument such that the surgical instrument can be used more properly or safely.

**[0140]** In one form, a strain gauge can be used to measure the force applied to the tissue 2308 by the end effector shown in FIGS. 42A-B and 43A-B. A strain gauge can be coupled to the end effector 2300 to measure the force on the tissue 2308 being treated by the end effector 2300. With reference now also to FIG. 44, in the aspect illustrated in FIG. 44, a system 2400 for measuring forces applied to the tissue 2308 comprises a strain gauge sensor 2402, such as, for example, a micro-strain gauge, is configured to measure one or more parameters of the end effector 2300 such as, for example, the amplitude of the strain exerted on a clamp arm of an end effector, such as the clamp arm 2302 of FIGS. 43A-B, during a clamping operation, which can be indicative of the tissue compression. The measured strain is converted to a digital signal and provided to a processor 2410 of a microcontroller 2408. A load sensor 2404 can measure the force to operate the ultrasonic blade 2304 to cut the tissue 2308 captured between the clamp arm 2302 and the ultrasonic blade 2304 of the end effector 2300. A magnetic field sensor 2406 can be employed to measure the thickness of the captured tissue 2308. The measurement of the magnetic field sensor 2406 also may be converted to a digital signal and provided to the processor 2410.

**[0141]** Further to the above, a feedback indicator 2414 also can be configured to communicate with the microcontroller 2408. In one aspect, the feedback indicator 2414 can be disposed in the handle of a surgical instrument, such as those shown in FIGS. 1-3. Alternatively, the feedback indicator 2414 can be disposed in a shaft assembly of a surgical instrument, for example. In any event, the microcontroller 2408 may employ the feedback indicator 2414 to provide feedback to an operator of the surgical instrument with regard to the adequacy of a manual input such as, for example, a selected position of a firing trigger that is used to cause the end effector to clamp down on tissue. To do so, the microcontroller 2408 may assess the selected position of the clamp arm 2302 and/or firing trigger. The measurements of the tissue 2308 compression, the tissue 2308 thickness, and/or the force required to close the end effector 2300 on the tissue, as respectively measured by the sensors 2402, 2404, 2406, can be used by the microcontroller 2408 to characterize the selected position of the firing trigger and/or the corresponding value of the speed of end effector. In one instance, a memory 2412 may store a technique, an equation, and/or a look-up table which can be employed by the microcontroller 2408 in the assessment.

[0142] The generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), surgical instruments 104, 106, 108 (FIGS. 1-3), and end effectors 122, 124, 125, 700, 800, 900, 1000, 1100, 1200, 2200, 2200', 2200", 2300 (FIGS. 1-3, 14-22, 33-43B) described herein may be employed alone or in combination to perform surgical procedures in accordance with the techniques and processes described hereinbelow. Nevertheless, for conciseness and clarity, the surgical procedures are described with reference to the multifunction surgical instrument 108 and the generator 500. The multifunction surgical instrument 108 comprises an end effector 125 which includes a clamp arm 145 and an ultrasonic blade 149. The end effector 125 may be configured with any of the structural or functional features of any one of the end effectors 122, 124, 125, 700, 800, 900, 1000, 1100, 1200, 2200, 2200', 2200", 2300 to provide electrodes to apply RF energy to tissue, temperature sensors, force/pressure sensors, and gap measurement sensors, as described hereinabove.

#### **Techniques For Determining Tissue Coefficient Of Friction/Tissue Coefficient of Coagulation**

[0143] In one aspect, the present disclosure provides a technique for determining a tissue coefficient of friction/tissue coefficient of coagulation to control the power output from a generator, such as any one of the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), or a surgical instrument, such as the surgical instrument 108 (FIGS. 1-3). The power delivered to an end effector of a surgical instrument can vary based on a tissue coefficient of friction. The energy profile from the generator can be adjusted based on the tissue coefficient of friction and dynamically switched during the procedure between RF and ultrasonic energy modalities based on the tissue impedance to treat the tissue clamped between a clamp arm and an ultrasonic blade of the end effector of the surgical instrument. For conciseness and clarity of disclosure, the techniques for determining a tissue coefficient of friction will be described with reference to the multifunction surgical instrument 108 of FIG. 2 coupled to the generator 500 of FIG. 8, although it will be appreciated that other configurations of instruments, generators, and end effectors described herein may be readily substituted without departing from the scope of the present disclosure.

[0144] The following description provides techniques for determining a tissue coefficient of friction  $\mu$ . In order to accurately calculate the tissue coefficient of friction  $\mu$  of the tissue 2308, the force applied to the tissue 2308 must be in a certain range to ensure that there is sufficient contact between the tissue 2308 and the end effector 2300 and that therapeutic amounts of energy are being delivered to the end effector 2300. For example, in one aspect, a minimum load on the tissue 2308 can be 1.5 lbs. to ensure that there is enough contact between the tissue 2308 and the end effector 2300, and a maximum load on the tissue 2308 can be 2.2 lbs. to ensure that a therapeutic amount of energy is being used. These values can be used with a power level 1. In addition, the force is measured using any of the above measurement components and techniques at various loads on the tissue 2308. At least two measurements can be taken at two different loads. For example, power can be measured at 1.76 lbs. (800 grams) and 2.2 lbs. (1000 grams) in order to graph power versus force. It can be more accurate, however, to take measurements at a plurality of loads and store the values in a buffer. In one form, a buffer can be filled with values for each incremental gram between 2400 grams and 1000 grams and can use various rules, such as the first in first out rule, to store the values until the slope value is maximized. A new regression can be performed each time a new value is added to the buffer and a value is dropped from the buffer. Various other methods can be used as well to calculate the slope value.

[0145] A functional model for heating of tissue can be represented in a simple frictional model:

$$\dot{Q} = \mu \cdot v \cdot N \quad \text{Equation 1}$$

where  $\dot{Q}$  is the rate of heat generation,  $v$  is the rms velocity of the ultrasonic motion at the tip of the blade, and  $N$  is the normal force driving tissue against the ultrasonic blade. The tissue coefficient of friction,  $\mu$ , is the proportionality constant that makes the statement true and it relates to tissue properties. It is therefore referred to as the tissue coefficient of friction  $\mu$  and may be referred to interchangeably as the tissue coefficient of coagulation. As is known in the art, friction is the force resisting the relative motion of solid surfaces, fluid layers, and material elements sliding against each other.

[0146] In practice,  $\dot{Q}$  is the measured amount power equal to the product of the rms values of current and voltage when driven at zero phase. The velocity is the product of the rms displacement,  $d$ , times the radian frequency or  $v = 2\pi f \cdot d$ . In the case of an ultrasonic surgical system this is the frequency at resonance which equals 55.5 kHz for the ultrasonic system. Furthermore  $d$  is proportional to the current driving the system, which is set by the level on the generator. The factor  $v$  is readily calculated.  $N$  is the normal force which is either set by the instrument design at full trigger closure or can be measured with a force gage/strain gage. Since  $\dot{Q}$ ,  $v$ , and  $N$  are calculable from known parameters, then can be estimated by rearranging Equation 1:

$$\mu = \frac{\dot{Q}}{v \cdot N}$$



[0147] Note that removing the quiescent power from  $\dot{Q}$  and updating the frequency  $f$  as the resonance drifts lower, will increase the repeatability and accuracy of the estimated tissue coefficient of friction  $\mu$ . In order to determine the type of tissue being treated by the end effector of the surgical instrument, a tissue coefficient of friction  $\mu$  can be calculated. The calculated tissue coefficient of friction is compared to a database of tissue coefficients of friction that correlates each tissue coefficient of friction with a tissue type, as will be discussed in more detail below. The calculated tissue coefficient of friction and its related tissue type are used by a technique to control the energy being delivered from the generator to the surgical instrument. In one form, the tissue coefficient of friction  $\mu$  is described by the above where  $\dot{Q}$  is the rate of heat generation,  $v$  is the velocity of the ultrasonic motion of the end effector, and  $N$  is the force applied to the tissue by the end effector. The velocity  $v$  of the ultrasonic motion is a known value from the settings of the generator. Since the value  $v$  is a known value, the tissue coefficient of friction  $\mu$  can be calculated using the slope of a graph of heat generation versus force on the tissue.

[0148] The force applied to the tissue by the end effector can be measured in a variety of ways using different type of components to measure force. This force measurement can be used, for example in the equation above, to determine the tissue coefficient of friction of the tissue being treated to determine its tissue type.

[0149] Looking at the equation  $\dot{Q} = \mu \cdot v \cdot N$  the product of the tissue coefficient of friction and velocity,  $\mu \cdot v$ , is the slope of the relation of heat generation and normal load. Because  $v$  is known,  $\mu$  can be determined by the slope of a graph of heat generation versus normal load. An example of a graph 2500 of power (Watts) shown along the vertical axis versus force (g) shown along the horizontal axis as measured with a plurality of plotted data points is illustrated in FIGS. 45-46. FIG. 45 illustrates graphs 2500 for a porcine carotid curves 2502, 2504 at two different power levels, porcine bowel curve 2506, and dry chamois curve 2508. The data for a carotid artery at Level 3 is also shown. These are single measurements and there are no means and standard deviations to test for differences. While ideally the values of the tissue coefficient of friction  $\mu$  at Levels 1 and 3 should be the same, the values at 0.30 and 0.35 are sufficiently close to believe that  $\mu$  is an intrinsic property of the tissue.

[0150] FIG. 46 illustrates the same graphs 2500' as FIG. 45, but only the sections of each graph that are substantially linear. The linear sections 2502', 2504', 2506', 2508' of each curve 2502, 2504, 2506, 2508 (FIG. 45) are located in the region of the curve where the force was greater than the initial low level of force used. Each of the linear sections 2502', 2504', 2506', 2508' can be modeled as a regression line in the form of:

$$y = mx + b \quad \text{Equation 2}$$

where  $y$  is the dependent variable (Power [Watts]),  $x$  is the independent variable (Force [g]),  $m$  is the slope of the line, and  $b$  is the  $y$ -intercept.

[0151] Each of the linear sections 2502', 2504', 2506', 2508' also are characterized by R-squared ( $R^2$ ), where  $R^2$  is a statistical measure of how close the data are to the fitted regression line  $y$ . It is also known as the coefficient of determination, or the coefficient of multiple determination for multiple regression. The definition of  $R^2$  is the percentage of the response variable variation that is explained by the linear model  $y$ . In other words:

$$R^2 = \text{Explained variation} / \text{Total variation}$$

[0152]  $R^2$  is always between 0 and 100% where 0% indicates that the model explains none of the variability of the response data around its mean and 100% indicates that the model explains all the variability of the response data around its mean. In general, the higher the  $R^2$ , the better the model fits the data.

[0153] In the example illustrated in FIG. 46, the regression line and  $R^2$  for the linear section 2502' representing porcine carotid tissue at power level 1 are:

$$y = 0.0137x + 0.2768$$

$$R^2 = 0.9902$$

[0154] The regression line and  $R^2$  for the linear section 2504' representing porcine carotid tissue at power level 3 are:

$$y = 0.0237x + 8.9847$$

$$R^2 = 0.978$$

[0155] The regression line and  $R^2$  for the linear section 906' representing bowel tissue at power level 1 are:

$$y = 0.0085x + 10.587$$

$$R^2 = 0.9147$$

[0156] The regression line and  $R^2$  for the linear section 2508' representing chamois tissue at power level 1 are:

$$y = 0.034x + 0.0735$$

$$R^2 = 0.9949$$

[0157] The calculated tissue coefficient of friction  $\mu$  is compared to a database of tissue coefficients of friction that correlated each coefficient with a tissue type. For example, Table 1 includes example tissue coefficients of friction  $\mu$  as calculated from the plots of power and force illustrated in FIGS. 45-46 for each of the porcine carotid curves 2502, 2504 at two different power levels, porcine bowel curve 2506, and dry chamois curve 2508.

Table 1

Tissue Type	Power Level	Coefficient of Friction $\mu$
Porcine carotid	1	0.30
Porcine carotid	3	0.35
Porcine bowel	1	0.19
Dry chamois	1	0.75

Assumptions: Ultrasonic bade amplitude is nominally 75 $\mu$ m p-p at level 5 and the frequency  $f$  is 55.5kHz. It will be appreciated that the power levels referenced above are specific to the LCS product and level 5 of the GEN11 generator available from Ethicon Endo-Surgery, Inc. that is compatible with devices known under the tradename HARMONIC and ENSEAL, also available from Ethicon Endo-Surgery, Inc.

[0158] It should be noted that the tissue coefficient of friction  $\mu$  for the porcine carotid tissue at various power levels should be the same as the tissue coefficient of friction  $\mu$  is a value intrinsic to the tissue. The values of 0.30 and 0.35 for the calculated tissue coefficients of friction  $\mu$  are substantially close enough to show that the tissue coefficient of friction  $\mu$  is intrinsic to the tissue itself.

[0159] One technique for determining the slope is to measure power delivered at two loads. Assuming Level 1 is used for this calculation, then the power could be measured at nominally 800 grams (1.76 lbs) and 1000 grams (2.2 lbs). However just using the two points can be inaccurate. It is better to fill a buffer of approximately 201 points for every gram in the 800 to 1000 gram range inclusive. The buffer could be slide advanced using a first-in-first-out (FIFO) rule until the  $R^2$  value is maximized. A new regression would be performed each time a new point is added and an old point is dropped. Other schemes can be envisioned as well. The calculated slope is then compared with known values for specific tissue types. For example if the calculated slope is in the region of 0.30 to 0.35, then an artery is determined to be in the jaws. If the values are in the range of 0.15 to 0.20 then a section of bowel is determined to be in the jaws. In one technique, the  $R^2$  value may be used as an indication of specificity. For example if  $R^2$  is less than 0.90, for example, then an indication is given that a determination cannot be made.  $R^2$  values in the range of 0.60 to 1.00 may be used to indicate that a determination of tissue type can be made and preferably tis threshold may be about 0.90.

[0160] In practice, sampling would be done on a time basis and both power and force would be sampled. These configurations are captured in the logic flow diagram of FIG. 47. The technique can be implemented in the generator software and/or the surgical instrument software.

[0161] It will be appreciated that using the averages for power and force at two positions can improve the accuracy of a two-point slope calculation. For accuracy it may be necessary to measure closure force (and moment) in addition

to power.

**[0162]** The benefits of using the tissue coefficient of friction  $\mu$  as a means to characterize tissue is discussed hereinbelow in connection with FIG. 48. Accordingly, the tissue coefficient of friction  $\mu$  can be used to discriminate tissues and can be calculated using a technique for including steps to determine that a selection cannot be made.

**[0163]** FIG. 47 illustrates a logic flow diagram 2600 of one form of a method for dynamically changing the energy delivered to a surgical instrument based on a determination of tissue type being treated by the instrument. As described herein, the logic flow diagram 2600 may be implemented in the generator 500, the multifunction surgical instrument 108, or a combination thereof. With reference now to the logic flow diagram 2600 shown in FIG. 47, a characterization mode of the system is started by the processor 502 in which the type of tissue being treated by the surgical instrument is determined 2602 by the processor 502. The end effector 125 is positioned such that tissue is positioned within the clamp arm 145 and the ultrasonic blade 149 of the end effector 125. The clamp arm 145 is used to apply a force to the tissue. The force of the end effector 125 on the tissue is measured and compared 2604 by the processor 502 to a threshold minimum force. If the force applied to the tissue is below a minimum threshold force, the force applied to the tissue is increased 2606 and is again measured and compared 2604 by the processor 502 to the threshold minimum force.

**[0164]** Once the force on the tissue has reached the minimum threshold force, the processor 502 samples 2608 the force applied to the tissue and the power delivered to the end effector 125 by the generator 500 and saves 2612 the samples in a buffer. The processor 502 determines 2610 whether the buffer is full. Samples are saved 2612 in the buffer until the buffer is full, and then the processor 502 utilizes the samples to plot 2614 points on a graph of force versus power. This information is used by the processor 502 to calculate and store 2616 the slope and  $R^2$  values in a database. See FIG. 46, for example, for a graphical representation of the linear sections of power versus force plots including the coefficient of determination  $R^2$ . Once stored in the database, the processor 502 compares the force values to a maximum force threshold and determines 2618 when the force reaches a maximum threshold value. Until the force reaches the maximum threshold value, the processor 502 continues along the NO branch and continues sampling 2608 the next samples of power and force. The processor 502 repeats until the force reaches the maximum threshold.

**[0165]** Once the force has reached the maximum threshold, the samples are no longer taken and stored and the processor 502 continues along the YES branch to select 2620 the slope with the highest  $R^2$  value and to calculate the tissue coefficient of friction  $\mu$ . Next, the processor 502 determines 2622 if the maximum slope value  $R^2$  is greater than a predetermined threshold. If  $R^2$  is less than the predetermined threshold, then the confidence level is low and the slope value cannot be used to calculate the tissue coefficient of friction  $\mu$  and identify the tissue type and the processor 502 continues along the YES branch to display 2624 a message indicating that the tissue type has not been identified. In one aspect, the threshold may be selected in the range of 0.6 to 1.00 and may preferably be set to about 0.90, for example. In other aspects, the threshold may be selected using more sophisticated, statistical tests applied to determine the level confidence  $R^2$ . If  $R^2$  is greater than or equal to the predetermined threshold, the calculated tissue coefficient of friction  $\mu$  can be used to identify the tissue and the processor 502 continues along the NO branch where the processor 502 compares 2626 the calculated tissue coefficient of friction  $\mu$  to a database 2628 of stored tissue coefficients of friction  $\mu$  which the stored tissue coefficients of friction  $\mu$  correspond to tissue types. The tissue type is selected 2630 and displayed and used to specify 2632 the power delivery profile for delivering energy from the generator 500 to the end effector 125 of the surgical instrument 108. Normal operation mode is entered 2634 such that the tissue type and related power delivery profile are used to control the end effector 125 for treating the tissue.

**[0166]** In one aspect, the tissue coefficient of friction  $\mu$  and its rate may be employed to determine tissue type and power delivery profile. For example, it has been shown in preliminary work that  $\mu$  is significantly different between porcine bowel, artery and chamois. Furthermore the  $\mu$  values were initial flat but did change as time progressed presumably because of the temperature rise as heat was added. From these observations,  $\mu$  can be used as a tissue differentiator and can be done at lower current to avoid rapid changes in  $\mu$  when the purpose is to characterize the tissue. Also, another parameter of interest would be the rate of change of the tissue coefficient of friction  $\mu$  for fixed conditions of  $\dot{Q}$ ,

$\nu$ , and  $N$ . It will be appreciated that  $\dot{\mu}$  is not simply differentiating the equation  $\mu = \frac{\dot{Q}}{\nu \cdot N}$  with respect time, because the change is due to changes in the tissue itself. For example the  $\mu$  for chamois rapidly rises when heat is delivered compare with carotid arteries or bowel, because it is dry. The rate of change is likely dependent on the percentage of water content in the tissue. In one aspect, the present disclosure provides a technique to estimate both the tissue coefficient of friction  $\mu$  and the rate of change of the tissue coefficient of friction  $\mu$  and compare them with table of know values stored in a database, for example. The tissue selection can be based on the values closet to the estimated values. The power delivery profile could then be optimize for that tissue and delivered to the surgical instrument 108. Estimating the tissue coefficient of friction  $\mu$  and the rate of change of the tissue coefficient of friction  $\mu$  may be able to further differentiate tissue types. Another aspect of the present disclosure includes tracking the tissue coefficient of friction  $\mu$  as the seal/transection progresses. Key changes in this parameter may signal a need to modify the power delivery profile.

**[0167]** FIG. 48 illustrates a logic flow diagram 2700 of another form of a method for dynamically changing the energy

delivered to a surgical instrument 108 based on a determination of tissue type being treated by the surgical instrument 108. As described herein, the logic flow diagram 2700 may be implemented in the generator 500, the multifunction surgical instrument 108, or a combination thereof. With reference now to the logic flow diagram 2700 shown in FIG. 48 and the surgical system 100 of FIG. 1, the processor 502 initiates a power evaluation mode of the system to determine 2702 the power profile for the generator 500. The end effector 125 of the surgical instrument 108 is then engaged with the tissue to be treated and the end effector 125 of the surgical instrument 108 is activated 2704 with energy delivered from the generator 500. The tissue coefficient of friction  $\mu$  and the processor 502 measures 2706 the rate of change (e.g., rate of rise) of the tissue coefficient of friction  $\mu$  over a short period of time. The processor 502 compares 2708 the measured tissue coefficient of friction  $\mu$  and the rate of change of the tissue coefficient of friction  $\mu$  to values stored in a tissue information database 2714.

**[0168]** If the system is in learning mode, in which the tissue information database 2714 information regarding tissue type and tissue coefficients of friction  $\mu$  are being updated, the tissue type can be visually identified 2710. The processor 502 updates 2712 the tissue information database 2714 of tissue coefficient of friction  $\mu$  and tissue type information to increase the accuracy of the tissue information database 2714.

**[0169]** If the system is not in learning mode, then the processor 502 selects 2716 a power delivery profile based on the determined tissue and surgical instrument type 2718 being used. During the treatment of the tissue, the processor 502 continues to calculate and monitor 2720 the tissue coefficient of friction  $\mu$  and the rate of change of the tissue coefficient of friction  $\mu$  to allow the processor 502 to dynamically update the power delivery profile. If an error is found, the power delivery profile is modified 2724. The processor 502 then determines 2722 if the correct pattern of power and delivery profile is being used. If no error is found, the current power delivery profile continues 2726. Otherwise, the processor 502 modifies 2724 the power and delivery profile. The processor 502 continues the process of verification and correction of the power delivery profile during the entire surgical procedure to optimize the energy being delivered to the surgical instrument 108 based on the type of tissue being treated.

#### **Techniques For Controlling A Generator Based On The Hydration Level Of Tissue**

**[0170]** In one aspect, the present disclosure provides a technique for controlling the power output from a generator, such as any one of the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), or a surgical instrument, such as the surgical instrument 108 (FIGS. 1-3). The power delivered to the end effector of the surgical instrument can be controlled based on the hydration level of the tissue to allow the energy profile from the generator to be dynamically changed based on the hydration level of the tissue being treated by the end effector of the surgical instrument. For conciseness and clarity of disclosure, the techniques for controlling the power output from a generator based on the hydration level of tissue will be described with reference to the multifunction surgical instrument 108 of FIG. 2 coupled to the generator 500 of FIG. 8, although it will be appreciated that other configurations of instruments, generators, and end effectors described herein may be readily substituted without departing from the scope of the present disclosure.

**[0171]** In order to determine the hydration level of the tissue being treated by the end effector 125 of the surgical instrument 108, the tissue coefficient of friction  $\mu$  as described herein can be used. As explained above, a force measurement is taken using any of the methods described herein to calculate the tissue coefficient of friction  $\mu$  to determine the hydration level of the tissue. The calculated tissue coefficient of friction  $\mu$  is compared to a database of tissue coefficients of friction  $\mu$  that correlate each tissue coefficient with a hydration level of the tissue. The calculated tissue coefficient of friction  $\mu$  and its related tissue hydration level are used by the technique to control the energy being delivered from the generator 500 to the surgical instrument 108.

**[0172]** The hydration level of the tissue can be indicative of a variety of conditions within the tissue. During a coagulation cycle in which RF energy is being used to coagulate the tissue, a decrease in the tissue hydration level can indicate that the tissue is nearing the end of the coagulation cycle. In addition, the hydration level of the tissue can vary during the coagulation cycle such that the RF energy being delivered from the generator 500 can dynamically change during the coagulation cycle based on the calculated hydration levels during the cycle. For example, as a coagulation cycle progresses, the hydration level of the tissue will decrease such that the technique used to control the energy delivered from the generator 500 can decrease the power as the cycle progresses. Thus, the hydration level of the tissue can be an indicator of the progress through a coagulation cycle of the tissue by the end effector of the surgical instrument 108.

**[0173]** FIG. 49 illustrates a logic flow diagram 2800 of one form of a method for dynamically changing the energy delivered to a surgical instrument 108 based on a determination of the hydration level of tissue being treated by the surgical instrument 108. As described herein, the logic flow diagram 2800 may be implemented in the generator 500, the multifunction surgical instrument 108, or a combination thereof. With reference now to the logic flow diagram 2800 shown in FIG. 49, the processor 502 determines the hydration level of the tissue by measuring the tissue coefficient of friction  $\mu$  as described herein. Initially, the end effector 125 of the surgical instrument 108 is closed on tissue and the end effector 125 is activated 2802 with energy from the generator 500. The processor 502 measures 2804 the tissue coefficient of friction  $\mu$ , as described herein. The processor 502 compares 2806 the tissue coefficient of friction  $\mu$  and

the rate of change of the tissue coefficient of friction  $\mu$  to values stored in a database to determine the hydration level of the tissue. The processor 502 selects 2808 a power delivery profile based on the hydration level of the tissue. The processor 502 monitors 2810 the tissue coefficient of friction  $\mu$  and the rate of change of the tissue coefficient of friction  $\mu$  during treatment of the tissue to monitor the changes in the tissue hydration level. This allows the processor 502 to dynamically change 2812 the power delivery profile, such that power delivered from the generator 500 can dynamically change during tissue treatment.

### ***Techniques For Switching Between RF And Ultrasonic Energy Based On Tissue Type***

[0174] In another aspect, the present disclosure provides a technique for controlling energy delivered by a generator and/or a surgical instrument by dynamically switching between RF and ultrasonic energy. The technique includes controlling the power output from a generator, such as any one of the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), or a surgical instrument, such as the surgical instrument 108 (FIGS. 1-3). In one aspect, RF and ultrasonic energy can be dynamically switched based on the type of tissue being treated by an end effector of a surgical instrument and various characteristics of the tissue. The power delivered to the end effector of the surgical instrument can include an input that represents the tissue type to allow the energy profile from the generator to be dynamically changed between RF and ultrasonic energy during the procedure based on the type of tissue being effected by the end effector of the surgical instrument. For conciseness and clarity of disclosure, techniques for dynamically switching between different energy modalities, such as, for example RF energy and ultrasonic energy, will be described with reference to the multifunction surgical instrument 108 of FIG. 2 coupled to the generator 500 of FIG. 8, although it will be appreciated that other configurations of instruments, generators, and end effectors described herein may be readily substituted without departing from the scope of the present disclosure.

[0175] In order to determine the type of tissue being treated by the end effector 125 of the surgical instrument 108, a tissue coefficient of friction  $\mu$  and/or the rate of change of the tissue coefficient of friction  $\mu$  can be calculated and compared to a database of tissue coefficient of friction  $\mu$  that correlates each tissue coefficient of friction  $\mu$  with a tissue type, as explained above. The calculated tissue coefficient of friction  $\mu$  and its related tissue type are used by a technique to control the type (RF or ultrasonic) and power level of energy being delivered from the generator 500 to the surgical instrument 108. Tissue types include, without limitation, tissue with a muscular structure, tissue with a vascular structure, tissue with a thin mesentery structure.

[0176] FIG. 50 illustrates a logic flow diagram 2900 of one aspect of a method of dynamically changing the between RF energy delivered from the ENERGY2/RETURN output of the generator 500 and ultrasonic energy delivered from the ENERGY1/RETURN output of the generator 500 to the surgical instrument 108 based on a determination of tissue type being treated by the surgical instrument 108. As described herein, the logic flow diagram 2900 may be implemented in the generator 500, the multifunction surgical instrument 108, or a combination thereof. With reference now to the logic flow diagram 2900 shown in FIG. 50, the processor 502 initiates a characterization mode of the generator 500 to determine 2902 the type of tissue being treated by the surgical instrument 108. The end effector 125 is positioned such that tissue is positioned between the clamp arm 145 and the ultrasonic blade 149 of the end effector 125. The end effector 125 applies a force to the tissue. The force applied to the tissue by the end effector 125 is measured and compared 2904 by the processor 502 to a threshold minimum force. If the processor 502 determines that the force applied to the tissue is below a minimum threshold force, the force applied to the tissue is increased 2906. Once the force on the tissue has reached the minimum threshold force, the processor 502 samples 2908 the force applied to the tissue and the power delivered to the end effector 125 by the generator 500 and stores the samples in a buffer.

[0177] The processor 502 determines 2901 if the buffer is full. If the buffer is not full, the processor 502 enters 2912 the most recent samples into the buffer. If the buffer is full, the processor 502 enters the most recent samples into the buffer and drops 2914 the first point (e.g., FIFO method for organizing and manipulating a data buffer). The processor 502 calculates and stores 2916 the slope and  $R^2$  values and the force is tested to determine 2918 if the force applied to the tissue is greater than a maximum force threshold. If the force applied to the tissue is not greater than the maximum threshold, the processor 502 samples 2908 the force applied to the tissue and the power delivered to the end effector 125 by the generator 500 and saves the samples in a buffer. The processor 502 repeats until the force applied to the tissue is greater than the maximum threshold. The processor 502 selects the slope with the highest  $R^2$  and calculates 2920 the tissue coefficient of friction  $\mu$ . If the processor 502 determines 2922 that  $R^2$  is less than a predetermined threshold, the processor 502 continues along the YES branch and the system displays 2924 that the tissue type was not identified. If  $R^2$  is greater than or equal to the threshold, the processor 502 continues along the NO branch and compares 2926 the calculated tissue coefficient of friction  $\mu$  to values stored in a tissue information database 2928.

[0178] In one aspect, the threshold may be selected in the range of 0.6 to 1.00 and may preferably be set to about 0.90, for example. In other aspects, the threshold may be selected using more sophisticated, statistical tests applied to determine the level confidence  $R^2$ . In one aspect, the threshold may be selected between 0.6 to 1.00 and may preferably be set to about 0.90. In other aspects, the threshold may be selected using more sophisticated, statistical tests applied

to determine the level confidence  $R^2$ . The processor 502 then selects 2930 and displays the tissue type and specifies 2932 the power delivery profile for delivering either ultrasonic energy (ENERGY1/RETURN) or RF energy (ENERGY2/RETURN) from the generator 500 to the end effector 125 of the surgical instrument 108. Normal operation mode is entered 2934 such that the tissue type and related power delivery profile are used to control the end effector for treating the tissue. Using the tissue type determined in accordance with the logic flow diagram 2900, the processor 502 can provide optimized energy delivery for that particular tissue. Tissue types that are determined to be more vascular or muscular would require more RF energy at the beginning of the cycle to provide more sealing energy either before the application of the ultrasonic energy or at a higher ratio of RF vs. ultrasonic energy. And tissue types that are less vascular would require less RF energy at the beginning of the cycle and the system would change the energy delivery toward a cutting profile using ultrasonic energy. Energy may be applied to the tissue until the tissue meets or exceeds a predetermined threshold impedance, which may be referred to as the termination impedance, and which is the tissue impedance that corresponds to the impedance of tissue when the tissue seal is complete.

#### **Techniques For Dynamically Changing The Energy Delivered From A Generator Based On Aperture Defined By The End Effector And Energy Parameters**

**[0179]** In one aspect, the present disclosure provides a technique for dynamically changing the energy delivered from a generator based on an aperture defined by an end effector. According to one aspect, the techniques for dynamically changing the energy delivered from a generator, such as any one of the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), or a surgical instrument, such as the surgical instrument 108 (FIGS. 1-3). For conciseness and clarity of disclosure, techniques for dynamically changing energy delivered from a generator based on an aperture defined by an end effector and energy parameters will be described with reference to the multifunction surgical instrument 108 of FIG. 2 coupled to the generator 500 of FIG. 8, although it will be appreciated that other configurations of instruments, generators, and end effectors described herein may be readily substituted without departing from the scope of the present disclosure.

**[0180]** The energy profile from the generator 500 can be dynamically changed during the procedure between RF and ultrasonic energy based on the aperture defined by the end effector 125 that is clamping on a tissue. This allows the generator 500 to switch from RF energy to ultrasonic energy based on the amount of clamping force exerted by the end effector 125 on the tissue being treated by the surgical instrument 108. The aperture defined by the clamp arm 145 is related to the creation of a proper coagulation seal, for example, when RF energy is being delivered from the generator 500 to the end effector 125 of the surgical instrument 108, such that RF energy should be used when there is a sufficient closure of the end effector 108 on the tissue. Thus, when the aperture defined by the end effector 108 is too large and there is not a sufficient clamping force exerted on the tissue for proper coagulation, only ultrasonic energy is delivered to the ultrasonic blade 149 of the end effector 125 through the ENERGY1/RETURN output of the generator 500.

**[0181]** For example, misleading information regarding various tissue measurements and/or characteristics can be transmitted to the generator 500 for use in the techniques for controlling the energy delivered depending on the aperture defined by the end effector 125. One such measurement that can be affected by the aperture defined by the end effector 125 is tissue impedance. Tissue impedance measured by an instrument as described herein will depend on several factors including the clamping force or pressure applied by the end effector 125 and the distance between electrodes in the case of RF energy or the clamp arm 145 aperture in the case of ultrasonic energy. Many techniques that determine the end of the coagulation cycle depend on the tissue impedance. A clamp arm 145 that is not fully closed on the ultrasonic blade 149 can lead to a tissue impedance determination that may not properly represent the condition of the tissue. An incorrect tissue impedance determination can lead to early termination of a coagulation cycle as the generator 500 may switch from RF to ultrasonic energy before a proper coagulation seal has been achieved. In one aspect, the tissue impedance may be determined by the processor 502 by dividing the output of the second voltage sensing circuit 524 coupled across the terminals labeled ENERGY2/RETURN by the output of the current sensing circuit 514 is disposed in series with the RETURN leg of the secondary side of the power transformer 508 as shown in FIG. 8. The outputs of the voltage sensing circuit 512 are provided to an isolation transformer and ADC 516 and the output of the current sensing circuit 514 is provided to another isolation transformer and ADC 518 to provide the voltage and current measurement is digital form to the processor 502.

**[0182]** The aperture defined by the end effector 125 can be determined using a variety of techniques. In various aspects, the aperture defined by the end effector 125 may be determined by detecting the pivot angle of the end effector 125. This can be accomplished using a potentiometer, a Hall effect sensor (in a manner described in connection with FIG. 19), an optical encoder, an optical IR sensor, an inductance sensor, or combinations thereof. In another aspect, the proximity of first and second components of an end effector is measured to determine the aperture defined by the end effector using, for example, the Hall effect sensor, an optical encoder, an optical IR sensor, an inductance sensor, or combinations thereof. In another aspect, the surgical instrument 108 may be configured to detect the aperture defined by the end effector 125 by measuring a change in a tissue impedance of the tissue interacting with the end effector. In

another aspect, the surgical instrument 108 may be configured to detect the aperture defined by the end effector 125 by measuring a load applied by the end effector 125 on the tissue as ultrasonic energy is pulsed to the end effector 125. In another aspect, the surgical instrument 108 includes a switch or other mechanism for closing the end effector 125 that can detect the aperture defined by the end effector 125. In another aspect, the aperture defined by the end effector 125 may be determined based on linear displacement or stroke of the closure mechanism located either in the shaft 129 or the handle 109 of the surgical instrument 108. In another form, the aperture defined by the end effector 125 may be determined by the angular displacement of the trigger mechanism located in the handle 109 of the surgical instrument 108.

**[0183]** FIG. 51 is a logic flow diagram 3000 of one aspect of a technique for dynamically changing the energy delivered from a generator based on aperture defined by the end effector and energy parameters. As described herein, the logic flow diagram 3000 may be implemented in the generator 500, the surgical instrument 108, or a combination thereof. With reference now to the logic flow diagram 3000 shown in FIG. 51 and the surgical system 100 of FIG. 1 with the combination electrosurgical and ultrasonic multifunction surgical instrument 108 and the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), the aperture defined by the end effector 125 is determined as described herein. Initially, the end effector 125 of the surgical instrument 108 is closed on tissue, as shown in FIGS. 42A-B, 43A-B, and the end effector 125 is activated 3002 with energy from the generator 500, for example. The aperture defined by the end effector 125 is determined 3004 by the processor 502 using any of the techniques described herein. Once the aperture defined by the end effector 125 is determined 3004, the processor 502 signals the waveform generator 504 to deliver energy from the generator 500 and controls switching the output of the generator 500 between RF energy ENERGY2 and ultrasonic energy (ENERGY1/RETURN) based on the determined aperture defined by the end effector 125. Accordingly, the processor 502 selects 3006 a power delivery profile based on the aperture defined by the end effector 125. The processor 502 continues to monitor 3008 the aperture defined by the end effector 125 during the tissue treatment process and dynamically changes 3010 the energy delivered from the generator 500 based on the changing aperture defined by the end effector 125.

**[0184]** In another aspect, a technique for controlling the power output from the generator, such as the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), or the surgical instrument 108, such as the surgical instrument 108 (FIGS. 1-3) that is delivered to the end effector 125 of the surgical instrument 108 can include an input that includes energy parameters based on an aperture defined by the end effector 125 of the surgical instrument 108. The energy profile from the generator 500 can be dynamically changed during the procedure between RF and ultrasonic energy using the energy parameters based on the aperture defined by the end effector 125 that is clamping on tissue as shown in FIGS. 42A-B, 43A-B. This allows the generator 500 to switch from RF energy ENERGY2 to ultrasonic energy (ENERGY1/RETURN) based on the clamping force applied by the end effector 125 to the tissue being treated by the surgical instrument 108. As explained above, the aperture defined by the clamp arm 145 is related to creating a proper coagulation seal, for example, when RF energy is being delivered from the generator 500 to the end effector 125 of the surgical instrument 108, such that RF energy should be used when there is a sufficient closure of the end effector 125 on the tissue. Thus, when the aperture defined by the end effector 125 is too large and the clamping force applied to the tissue is insufficient for proper coagulation, only ultrasonic energy is delivered to the end effector 125 by the generator 500.

**[0185]** The aperture defined by the end effector 125 can be determined using any of the methods described herein. For example, the surgical instrument 108 can include an aperture defined by a sensor located in the end effector 125 that can be fed through a connector to the ASIC in the handle 109 of the surgical instrument 108. The surgical instrument 108 also can include a sensor in the handle 109 of the surgical instrument 108 that is configured to detect the aperture defined by the end effector 125.

**[0186]** Energy parameters are configured to be loaded into the generator 500, and can include a plurality of different parameters, including but not limited to voltage, current, power, and one or more techniques for use in treating tissue. These parameters can be related to the RF energy and ultrasonic energy that can be delivered from the generator 500. The energy parameters can include information such as maximum and/or minimum values to be used to control the energy delivered from the generator 500. The energy parameters can be stored in a variety of locations, including an EEPROM on the surgical instrument or some other non-volatile memory. In addition, there can be multiple sets of energy parameters. For example, there can be a first set of energy parameters that are used to optimize tissue transection, and a second set of energy parameters that are used to optimize tissue spot coagulation. It will be understood that there can be any number of set of energy parameters that correspond to various types of tissue treatments to allow the generator to switch between the various sets of energy parameters based on the necessary tissue treatments.

**[0187]** When the end effector 125 of the surgical instrument 108 is activated, one or more of the various techniques described herein are used to detect the aperture defined by the end effector 108. In one aspect, when the end effector 125 is closed around the tissue, the generator 500 can utilize the energy parameters for optimizing tissue transection. When the end effector 125 has a larger aperture and is not clamped on the tissue, the generator 500 can utilize the energy parameters for optimizing spot coagulation of the tissue.

**[0188]** FIG. 52 is a logic flow diagram 3200 of one aspect of a technique for dynamically changing the energy delivered

from a generator based on aperture defined by the end effector and energy parameters. As described herein, the logic flow diagram 3200 may be implemented in the generator 500, the multifunction surgical instrument 108, or a combination thereof. With reference now to the logic flow diagram 3100 shown in FIG. 52, with the combination electrosurgical and ultrasonic instrument 108 and the generator 500, initially the aperture defined by the end effector 125 is determined by the processor 502 as described herein. The end effector 125 of the surgical instrument 108 is closed on tissue, as shown in FIGS. 42A-B, 43A-B, and the end effector 125 is activated with energy from the generator 500. The aperture defined by the end effector 125 is determined 3104 using any of the techniques described herein. The processor 502 signals the waveform generator 504 to deliver energy from the generator 500 to the end effector 125 and switch the generator 500 output between to switch between RF and ultrasonic energy based on the aperture defined by the end effector 125 by using one of the sets of energy parameters that was previously loaded into the generator 500. Accordingly, energy parameters are communicated 3106 to the generator 500 based on the previously determined 3104 aperture defined by the end effector 125. The processor 502 monitors 3108 the aperture defined by the end effector 125 during the tissue treatment process such that the energy delivered from the generator 500 can be dynamically communicated 3110 by the processor 502 during the tissue treatment process based on the changing aperture defined by the end effector 125. This allows the generator 500 to dynamically switch between the various sets of energy parameters based on the changing aperture defined by the end effector 125.

**[0189]** It will be understood that various combinations of information can be used to determine which set of energy parameters are to be used during tissue treatment. For example, the aperture defined by the end effector and calculated tissue impedance can be used to determine which set of energy parameters are needed to control the energy being delivered from the generator.

### Techniques For Dynamic Sensing Of Tissue

**[0190]** In one aspect, the present disclosure provides dynamic tissue sensing techniques. In one aspect, the dynamic tissue sensing techniques may be implemented with a generator, such as any one of the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), or a surgical instrument, such as the surgical instrument 108 (FIGS. 1-3). For conciseness and clarity of disclosure, the dynamic tissue sensing techniques will be described with reference to the multifunction surgical instrument 108 of FIG. 2 coupled to the generator 500 of FIG. 8, although it will be appreciated that other configurations of instruments, generators, and end effectors described herein may be readily substituted without departing from the scope of the present disclosure.

**[0191]** Upon measuring the tissue thickness and the normal force on the ultrasonic blade as shown in and described in connection with FIGS. 42A-B, 43A-B, and measuring the tissue coefficient of friction  $\mu$ , in one aspect a technique is provided that pulses to measure the tissue coefficient of friction  $\mu$  at set time intervals and based on the measured coefficient  $\mu$  over time, adjusts the power delivered by any one of and the generators 102, 200, 300, 400, 500 (FIGS. 1-3 and 4-8), or the surgical instrument (FIGS. 1-3), in order to transect or "cut" tissue efficiently and with good hemostasis. In one aspect, a sensing technique is configured to monitor tissue friction and determine the tissue coefficient of friction  $\mu$ , as described herein, throughout a cut and modulates energy delivered during the cut based on change in the tissue coefficient of friction  $\mu$  over time.

**[0192]** FIG. 53 is a logic flow diagram 3200 of one aspect of a dynamic tissue sensing technique. As described herein, the logic flow diagram 3200 may be implemented in the generator 500, the multifunction surgical instrument 108, or a combination thereof. With reference now to the logic flow diagram 3200 shown in FIG. 53, case specific details for knowledge of tissue type are input 322 into the generator 500. The end effector 125 of the surgical instrument 108 is closed on the tissue, as shown in FIGS. 42A-B, 43A-B, and the tissue coefficient of friction  $\mu$  is checked 3204 by the processor 502 and power is delivered 3206 to the tissue according to known normal force  $N$ , tissue coefficient of friction  $\mu$ , and tissue thickness as determined by processor 502 as described herein. The processor 502 compares 3208 the rate of change of  $\mu/|\mu|$  to known values of  $\mu$ . In one aspect, a look-up table for predefined  $\mu$ /Power Delivery Ratios may be utilized by the processor 502. The processor 502 adjusts 3210 the power delivered by the generator 500 higher or lower based on  $\Delta\mu$ . When  $\mu$  reaches 3212 a known or predetermined threshold, the processor 502 checks 3214 the clamp arm 145 of the end effector 125 for closure against the ultrasonic blade 149. In one aspect, this can be determined based on known normal force  $N$  and known production normal force  $N_p$  that represents a closed end effector jaw. When the jaw closure is complete, the processor 502 control the delivery 3216 of short bursts of power to complete the tissue transection or cut until the cut is completed 3218.

**[0193]** Accordingly, by measuring the tissue coefficient of friction  $\mu$ , the processor 502 pulses to measure the tissue coefficient of friction  $\mu$  at set time intervals and based on the measured coefficient  $\mu$  over time, and adjusts the power delivered by generator 500 to the tissue in order to transect or "cut" tissue efficiently and with good hemostasis. In one aspect, the sensing technique monitors and periodically determines the tissue coefficient of friction  $\mu$ , as described herein, while making the cut and modulates the energy delivered by the generator during the cut based on change in the tissue coefficient of friction  $\Delta\mu$  over time.



**[0194]** While various details have been set forth in the foregoing description, it will be appreciated that the various aspects of the surgical system with user adaptable techniques based on tissue type may be practiced without these specific details. For example, for conciseness and clarity selected aspects have been shown in block diagram form rather than in detail. Some portions of the detailed descriptions provided herein may be presented in terms of instructions that operate on data that is stored in a computer memory. Such descriptions and representations are used by those skilled in the art to describe and convey the substance of their work to others skilled in the art. In general, a technique refers to a self-consistent sequence of steps leading to a desired result, where a "step" refers to a manipulation of physical quantities which may, though need not necessarily, take the form of electrical or magnetic signals capable of being stored, transferred, combined, compared, and otherwise manipulated. It is common usage to refer to these signals as bits, values, elements, symbols, characters, terms, numbers, or the like. These and similar terms may be associated with the appropriate physical quantities and are merely convenient labels applied to these quantities.

**[0195]** Unless specifically stated otherwise as apparent from the foregoing discussion, it is appreciated that, throughout the foregoing description, discussions using terms such as "processing" or "computing" or "calculating" or "determining" or "displaying" or the like, refer to the action and processes of a computer system, or similar electronic computing device, that manipulates and transforms data represented as physical (electronic) quantities within the computer system's registers and memories into other data similarly represented as physical quantities within the computer system memories or registers or other such information storage, transmission or display devices.

**[0196]** It is worthy to note that any reference to "one aspect," "an aspect," "one form," or "an form" means that a particular feature, structure, or characteristic described in connection with the aspect is included in at least one aspect. Thus, appearances of the phrases "in one aspect," "in an aspect," "in one form," or "in an form" in various places throughout the specification are not necessarily all referring to the same aspect. Furthermore, the particular features, structures or characteristics may be combined in any suitable manner in one or more aspects.

**[0197]** Some aspects may be described using the expression "coupled" and "connected" along with their derivatives. It should be understood that these terms are not intended as synonyms for each other. For example, some aspects may be described using the term "connected" to indicate that two or more elements are in direct physical or electrical contact with each other. In another example, some aspects may be described using the term "coupled" to indicate that two or more elements are in direct physical or electrical contact. The term "coupled," however, also may mean that two or more elements are not in direct contact with each other, but yet still cooperate or interact with each other.

**[0198]** Although various forms have been described herein, many modifications, variations, substitutions, changes, and equivalents to those forms may be implemented and will occur to those skilled in the art. Also, where materials are disclosed for certain components, other materials may be used. It is therefore to be understood that the foregoing description and the appended claims are intended to cover all such modifications and variations as falling within the scope of the disclosed forms. The following claims are intended to cover all such modification and variations.

**[0199]** In a general sense, those skilled in the art will recognize that the various aspects described herein which can be implemented, individually and/or collectively, by a wide range of hardware, software, firmware, or any combination thereof can be viewed as being composed of various types of "electrical circuitry." Consequently, as used herein "electrical circuitry" includes, but is not limited to, electrical circuitry having at least one discrete electrical circuit, electrical circuitry having at least one integrated circuit, electrical circuitry having at least one application specific integrated circuit, electrical circuitry forming a general purpose computing device configured by a computer program (e.g., a general purpose computer configured by a computer program which at least partially carries out processes and/or devices described herein, or a microprocessor configured by a computer program which at least partially carries out processes and/or devices described herein), electrical circuitry forming a memory device (e.g., forms of random access memory), and/or electrical circuitry forming a communications device (e.g., a modem, communications switch, or optical-electrical equipment). Those having skill in the art will recognize that the subject matter described herein may be implemented in an analog or digital fashion or some combination thereof.

**[0200]** The foregoing detailed description has set forth various forms of the devices and/or processes via the use of block diagrams, flowcharts, and/or examples. Insofar as such block diagrams, flowcharts, and/or examples contain one or more functions and/or operations, it will be understood by those within the art that each function and/or operation within such block diagrams, flowcharts, or examples can be implemented, individually and/or collectively, by a wide range of hardware, software, firmware, or virtually any combination thereof. In one form, several portions of the subject matter described herein may be implemented via Application Specific Integrated Circuits (ASICs), Field Programmable Gate Arrays (FPGAs), digital signal processors (DSPs), or other integrated formats. However, those skilled in the art will recognize that some aspects of the forms disclosed herein, in whole or in part, can be equivalently implemented in integrated circuits, as one or more computer programs running on one or more computers (e.g., as one or more programs running on one or more computer systems), as one or more programs running on one or more processors (e.g., as one or more programs running on one or more microprocessors), as firmware, or as virtually any combination thereof, and that designing the circuitry and/or writing the code for the software and or firmware would be well within the skill of one of skill in the art in light of this disclosure. In addition, those skilled in the art will appreciate that the mechanisms of the

subject matter described herein are capable of being distributed as a program product in a variety of forms, and that an illustrative form of the subject matter described herein applies regardless of the particular type of signal bearing medium used to actually carry out the distribution. Examples of a signal bearing medium include, but are not limited to, the following: a recordable type medium such as a floppy disk, a hard disk drive, a Compact Disc (CD), a Digital Video Disk (DVD), a digital tape, a computer memory, etc.; and a transmission type medium such as a digital and/or an analog communication medium (e.g., a fiber optic cable, a waveguide, a wired communications link, a wireless communication link (e.g., transmitter, receiver, transmission logic, reception logic, etc.), etc.).

**[0201]** One skilled in the art will recognize that the herein described components (e.g., operations), devices, objects, and the discussion accompanying them are used as examples for the sake of conceptual clarity and that various configuration modifications are contemplated. Consequently, as used herein, the specific exemplars set forth and the accompanying discussion are intended to be representative of their more general classes. In general, use of any specific exemplar is intended to be representative of its class, and the non-inclusion of specific components (e.g., operations), devices, and objects should not be taken limiting.

**[0202]** With respect to the use of substantially any plural and/or singular terms herein, those having skill in the art can translate from the plural to the singular and/or from the singular to the plural as is appropriate to the context and/or application. The various singular/plural permutations are not expressly set forth herein for sake of clarity.

**[0203]** The herein described subject matter sometimes illustrates different components contained within, or connected with, different other components. It is to be understood that such depicted architectures are merely example, and that in fact many other architectures may be implemented which achieve the same functionality. In a conceptual sense, any arrangement of components to achieve the same functionality is effectively "associated" such that the desired functionality is achieved. Hence, any two components herein combined to achieve a particular functionality can be seen as "associated with" each other such that the desired functionality is achieved, irrespective of architectures or intermedial components. Likewise, any two components so associated also can be viewed as being "operably connected," or "operably coupled," to each other to achieve the desired functionality, and any two components capable of being so associated also can be viewed as being "operably couplable," to each other to achieve the desired functionality. Specific examples of operably couplable include but are not limited to physically mateable and/or physically interacting components, and/or wirelessly interactable, and/or wirelessly interacting components, and/or logically interacting, and/or logically interactable components.

**[0204]** In some instances, one or more components may be referred to herein as "configured to," "configurable to," "operable/operative to," "adapted/adaptable," "able to," "conformable/conformed to," etc. Those skilled in the art will recognize that "configured to" can generally encompass active-state components and/or inactive-state components and/or standby-state components, unless context requires otherwise.

**[0205]** In addition, even if a specific number of an introduced claim recitation is explicitly recited, those skilled in the art will recognize that such recitation should typically be interpreted to mean at least the recited number (e.g., the bare recitation of "two recitations," without other modifiers, typically means at least two recitations, or two or more recitations). Furthermore, in those instances where a convention analogous to "at least one of A, B, and C, etc." is used, in general such a construction is intended in the sense one having skill in the art would understand the convention (e.g., "a system having at least one of A, B, and C" would include but not be limited to systems that have A alone, B alone, C alone, A and B together, A and C together, B and C together, and/or A, B, and C together, etc.). In those instances where a convention analogous to "at least one of A, B, or C, etc." is used, in general such a construction is intended in the sense one having skill in the art would understand the convention (e.g., "a system having at least one of A, B, or C" would include but not be limited to systems that have A alone, B alone, C alone, A and B together, A and C together, B and C together, and/or A, B, and C together, etc.). It will be further understood by those within the art that typically a disjunctive word and/or phrase presenting two or more alternative terms, whether in the description, claims, or drawings, should be understood to contemplate the possibilities of including one of the terms, either of the terms, or both terms unless context dictates otherwise. For example, the phrase "A or B" will be typically understood to include the possibilities of "A" or "B" or "A and B."

**[0206]** Although various forms have been described herein, many modifications, variations, substitutions, changes, and equivalents to those forms may be implemented and will occur to those skilled in the art. Also, where materials are disclosed for certain components, other materials may be used. It is therefore to be understood that the foregoing description and the appended claims are intended to cover all such modifications and variations as falling within the scope of the disclosed forms. The following claims are intended to cover all such modification and variations.

**[0207]** In summary, numerous benefits have been described which result from employing the concepts described herein. The foregoing description of the one or more forms has been presented for purposes of illustration and description. It is not intended to be exhaustive or limiting to the precise form disclosed. Modifications or variations are possible in light of the above teachings. The one or more forms were chosen and described in order to illustrate principles and practical application to thereby enable one of ordinary skill in the art to utilize the various forms and with various modifications as are suited to the particular use contemplated. It is intended that the claims submitted herewith define the

overall scope.

## Claims

1. A surgical instrument (108) for coagulating and dissecting tissue, the surgical instrument comprising:

a processor;

an end effector (125) at a distal end of the surgical instrument, the end effector configured to interact with tissue, the end effector comprising:

a clamp arm (145);

an ultrasonic blade (149);

a force sensor in communication with the processor and configured to measure a force applied to tissue located between the clamp arm and the ultrasonic blade; and

a temperature sensor in communication with the processor;

an ultrasonic transducer (120) acoustically coupled to the ultrasonic blade and configured to receive a drive signal from a generator to cause ultrasonic motion of the ultrasonic blade and deliver energy to the ultrasonic blade;

**characterized in that** the processor is configured to:

determine a type of tissue or the hydration level of tissue interacting with the end effector based on a tissue coefficient of friction, wherein the tissue coefficient of friction is determined based on the force applied to the tissue by the end effector, the ultrasonic motion of the ultrasonic blade, and a rate of heat generated by the end effector; and

dynamically control the drive signal delivered to the ultrasonic transducer based on the type of tissue or the hydration level of tissue interacting with the end effector;

wherein the tissue coefficient of friction is determined based on the following expression:

$$\mu = \frac{\dot{Q}}{v \cdot N}$$

wherein  $\dot{Q}$  is rate of heat generation,  $v$  is velocity of the ultrasonic motion of the ultrasonic blade, and  $N$  is force applied to the tissue by the end effector.

2. The surgical instrument of claim 1, wherein the clamp arm comprises an electrode, wherein the processor is further configured to:

dynamically control delivery of radio frequency (RF) energy to the electrode based on the type of tissue interacting with the end effector; and

switch between delivery of the RF energy and the ultrasonic drive signal based on the type of tissue interacting with the end effector.

3. The surgical instrument of any one of claims 1 or 2, wherein the force sensor comprises a strain gauge coupled to the end effector, optionally wherein the force measurement is communicated from the strain gauge to the processor.

4. The surgical instrument of any one of claims 1 or 2, wherein the processor is configured to:

store two data points of the measured force and power delivered to the end effector in a buffer;

determine a slope value of the power versus force based on the two data points;

determine a statistical measure  $R^2$  of the slope value relative to a fitted regression line; and

determine the tissue coefficient of friction based on the slope value;

optionally wherein the processor is further configured to:

compare  $R^2$  to a threshold value; and

select a slope value associated with an  $R^2$  value that is greater than or equal to the threshold value to determine the tissue coefficient of friction.

5. The surgical instrument of any one of claims 1 or 2, wherein the processor is configured to:

compare the tissue coefficient of friction to tissue coefficients of friction stored in a tissue information database; determine the type of tissue based on the tissue coefficient of friction; and deliver the drive signal to the ultrasonic transducer based on the tissue coefficient of friction.

6. The surgical instrument of any one of claims 1 or 2, further comprising a generator (500) configured to deliver radio frequency (RF) energy to the electrode and/or the drive signal to the ultrasonic transducer.

7. A generator (500) for delivering energy to a surgical instrument (108) for coagulating and dissecting tissue, the surgical instrument comprising an end effector (125) at a distal end thereof, the end effector configured to interact with tissue, the end effector comprising a clamp arm (145), an ultrasonic blade (149), a force sensor in communication with the processor and configured to measure a force applied to tissue located between the clamp arm and the ultrasonic blade, and a temperature sensor in communication with a processor, an ultrasonic transducer (120) acoustically coupled to the ultrasonic blade and configured to receive a drive signal from the generator to cause ultrasonic motion of the ultrasonic blade and deliver energy to the ultrasonic blade, the generator comprising:

a drive circuit configured to deliver a drive signal to an ultrasonic transducer; and **characterized by** a processor (502) configured to control energy delivered by the drive circuit to the ultrasonic transducer based on a type of tissue or the hydration level of tissue interacting with the end effector; wherein the processor is configured to:

receive a measurement of force applied to tissue by the end effector, ultrasonic motion of the ultrasonic blade, and a rate of heat generated by the end effector; determine the type of tissue or a hydration level of tissue interacting with the end effector based on a tissue coefficient of friction, wherein the tissue coefficient of friction is determined based on the force applied to the tissue by the end effector, the ultrasonic motion of the ultrasonic blade, and a rate of heat generated by the end effector; and dynamically control the drive signal delivered to the ultrasonic transducer based on the type of tissue interacting with the end effector;

wherein the tissue coefficient of friction is determined based on the following expression:

$$\mu = \frac{\dot{Q}}{v \cdot N}$$

wherein  $\dot{Q}$  is rate of heat generation,  $v$  is velocity of the ultrasonic motion of the ultrasonic blade, and  $N$  is force applied to the tissue by the end effector.

8. The generator of claim 7, wherein the clamp arm comprises an electrode, wherein the generator further comprises a second drive circuit configured to deliver radio frequency (RF) energy to the electrode, wherein the processor is configured to control energy delivered by the second drive circuit to the electrode based on a type of tissue interacting with the end effector, and wherein the processor is configured to dynamically control delivery of RF energy to the electrode based on the type of tissue interacting with the end effector and switch between delivery of the RF energy and the ultrasonic drive signal based on the type of tissue interacting with the end effector.

9. The generator of claim 7, wherein the processor is configured to:

store two data points of the measured force and power delivered to the ultrasonic transducer in a buffer; determine a slope value of the power versus force based on the two data points; determine a statistical measure  $R^2$  of the slope value relative to a fitted regression line; and determine the tissue coefficient of friction based on the slope value; optionally wherein the processor is further configured to:

compare  $R^2$  to a threshold value; and  
select a slope value associated with an  $R^2$  value that is greater than or equal to the threshold value to  
determine the tissue coefficient of friction.

10. The generator of claim 7, wherein the processor is configured to:

compare the tissue coefficient of friction to tissue coefficients of friction stored in a tissue information database;  
determine the tissue type based on the tissue coefficient of friction; and  
deliver the drive signal to the end effector based on the tissue coefficient of friction.

11. The generator of claim 7, further comprising a second drive circuit configured to deliver radio frequency (RF) energy to the electrode.

12. The surgical instrument of claim 1 or the generator of claim 7, wherein the processor is configured to control power of the drive signal delivered to the ultrasonic transducer based on the tissue coefficient of friction.

13. The surgical instrument of claim 1 or the generator of claim 7, wherein the processor is configured to control power of the drive signal delivered to the ultrasonic transducer based on the hydration level of the tissue, optionally wherein the power of the drive signal delivered to the ultrasonic transducer is decreased as the hydration level of the tissue decreases.

14. The surgical instrument of claim 2 or the generator of claim 8, wherein either the processor is configured to control delivery of the ultrasonic drive signal to the ultrasonic transducer if the tissue type is one having a muscular structure, or the processor is configured to control delivery of the RF energy to the electrode if the tissue type is one having a vascular structure, or the processor is configured to control delivery of a drive signal to the ultrasonic transducer corresponding to a rapid cutting motion if the tissue type is one having a thin mesentery structure.

## Patentansprüche

1. Chirurgisches Instrument (108) zum Koagulieren und Zerteilen von Gewebe, wobei das chirurgische Instrument umfasst:

einen Prozessor;  
einen Endeffektor (125) an einem distalen Ende des chirurgischen Instruments, wobei der Endeffektor ausgestaltet ist, um mit dem Gewebe zu interagieren, wobei der Endeffektor umfasst:

einen Klemmarm (145);  
eine Ultraschallklinge (149);  
einen Kraftsensor in Kommunikation mit dem Prozessor und ausgestaltet, um eine auf das Gewebe, welches sich zwischen dem Klemmarm und der Ultraschallklinge befindet, ausgeübte Kraft zu messen; und  
einen Temperatursensor in Kommunikation mit dem Prozessor;  
einen Ultraschallwandler (120), der akustisch an die Ultraschallklinge gekoppelt ist und ausgestaltet ist, um ein Antriebssignal von einem Generator zu empfangen, um Ultraschallbewegung der Ultraschallklinge zu bewirken und Energie an die Ultraschallklinge abzugeben;

**dadurch gekennzeichnet, dass** der Prozessor ausgestaltet ist zum:

Ermitteln eines Gewebetyps oder des Hydratisierungsniveaus von Gewebe, das mit dem Endeffektor interagiert, basierend auf einem Reibungskoeffizienten des Gewebes, wobei der Reibungskoeffizient des Gewebes basierend auf der Kraft, die durch den Endeffektor auf das Gewebe ausgeübt wird, der Ultraschallbewegung der Ultraschallklinge und einer Wärmerate ermittelt wird, die durch den Endeffektor erzeugt wird; und  
dynamisches Steuern des an den Ultraschallwandler abgegebenen Antriebssignals basierend auf dem Gewebetyp oder dem Hydratisierungsniveau des Gewebes, welches mit dem Endeffektor interagiert; wobei der Reibungskoeffizient des Gewebes basierend auf dem folgenden Ausdruck ermittelt wird:

$$\mu = \frac{\dot{Q}}{v \cdot N}$$

- 5 wobei  $\dot{Q}$  die Rate der Wärmeerzeugung ist,  $v$  die Geschwindigkeit der Ultraschallbewegung der Ultraschall-  
klinge ist, und  $N$  die Kraft ist, die durch den Endeffektor auf das Gewebe ausgeübt wird.
2. Chirurgisches Instrument nach Anspruch 1, wobei der Klemmarm eine Elektrode umfasst, wobei der Prozessor  
10 ferner ausgestaltet ist zum:
- dynamischen Steuern der Abgabe von Hochfrequenz (HF)-Energie an die Elektrode basierend auf dem Gewe-  
betyp, der mit dem Endeffektor interagiert; und  
Schalten zwischen Abgabe der HF-Energie und dem Ultraschallantriebssignal basierend auf dem Gewebetyp,  
15 der mit dem Endeffektor interagiert.
3. Chirurgisches Instrument nach einem der Ansprüche 1 oder 2, wobei der Kraftsensor einen Dehnungsmessstreifen  
umfasst, der an den Endeffektor gekoppelt ist, wobei gegebenenfalls die Kraftmessung von dem Dehnungsmess-  
streifen an den Prozessor kommuniziert wird.
- 20 4. Chirurgisches Instrument nach einem der Ansprüche 1 oder 2, wobei der Prozessor ferner ausgestaltet ist zum:
- Speichern von zwei Datenpunkten der gemessenen Kraft und Leistung, die an den Endeffektor abgegeben  
werden, in einem Puffer;  
Ermitteln eines Neigungswerts von Leistung gegen Kraft basierend auf den zwei Datenpunkten;  
25 Ermitteln eines statistischen Maßes  $R^2$  des Neigungswerts relativ zu einer angepassten Regressionslinie; und  
Ermitteln des Reibungskoeffizienten des Gewebes basierend auf dem Neigungswert;  
wobei der Prozessor ferner ausgestaltet ist zum:
- Vergleichen von  $R^2$  mit einem Schwellenwert; und  
30 Auswählen eines Neigungswerts, der einem  $R^2$ -Wert zugeordnet ist, der größer als oder gleich dem Schwel-  
lenwert ist, um den Reibungskoeffizienten des Gewebes zu ermitteln.
5. Chirurgisches Instrument nach einem der Ansprüche 1 oder 2, wobei der Prozessor ausgestaltet ist zum:
- 35 Vergleichen des Reibungskoeffizienten des Gewebes mit Reibungskoeffizienten von Gewebe, die in einer  
Gewebeinformationsdatenbank gespeichert sind;  
Ermitteln des Gewebetyps basierend auf dem Reibungskoeffizienten des Gewebes; und  
Abgeben des Antriebssignals an den Ultraschallwandler basierend auf dem Reibungskoeffizienten des Gewe-  
bes.
- 40 6. Chirurgisches Instrument nach einem der Ansprüche 1 oder 2, ferner umfassend einen Generator (500), der aus-  
gestaltet ist, um Hochfrequenz (HF)-Energie an die Elektrode und/oder das Antriebssignal an den Ultraschallwandler  
abzugeben.
- 45 7. Generator (500) zur Abgabe von Energie an ein chirurgisches Instrument (108) zum Koagulieren und Zerteilen von  
Gewebe, wobei das chirurgische Instrument einen Endeffektor (125) an einem distalen Ende davon umfasst, wobei  
der Endeffektor zum Interagieren mit Gewebe ausgestaltet ist, der Endeffektor einen Klemmarm (145), eine Ultra-  
schallklinge (149), einen Kraftsensor in Kommunikation mit dem Prozessor und ausgestaltet, um eine auf Gewebe,  
welches sich zwischen dem Klemmarm und der Ultraschallklinge befindet, ausgeübte Kraft zu messen, und einen  
50 Temperatursensor in Kommunikation mit einem Prozessor umfasst, wobei ein Ultraschallwandler (120) akustisch  
an die Ultraschallklinge gekoppelt ist und ausgestaltet ist, um ein Antriebssignal von dem Generator zu empfangen,  
um Ultraschallbewegung der Ultraschallklinge zu bewirken und Energie an die Ultraschallklinge abzugeben, wobei  
der Generator umfasst:
- 55 eine Treiberschaltung, die ausgestaltet ist, um ein Antriebssignal an einen Ultraschallwandler abzugeben; und  
**gekennzeichnet durch** einen Prozessor (502), der ausgestaltet ist, um durch die Treiberschaltung an den  
Ultraschallwandler abgegebene Energie basierend auf einem Gewebetyp oder dem Hydratisierungsniveau des  
Gewebes zu steuern, das mit dem Endeffektor interagiert;

wobei der Prozessor ausgestaltet ist zum:

Empfangen einer Messung der Kraft, die durch den Endeffektor, die Ultraschallbewegung der Ultraschall-  
 klinge und eine Wärmerate, die durch den Endeffektor erzeugt wird, auf Gewebe ausgeübt wird;  
 5 Ermitteln des Gewebetyps oder eines Hydratisierungsniveaus von Gewebe, das mit dem Endeffektor in-  
 teragiert, basierend auf einem Reibungskoeffizienten des Gewebes, wobei der Reibungskoeffizient des  
 Gewebes basierend auf der Kraft, die durch den Endeffektor auf das Gewebe ausgeübt wird, der Ultra-  
 schallbewegung der Ultraschallklinge und einer Wärmerate ermittelt wird, die durch den Endeffektor erzeugt  
 10 wird; und  
 dynamisches Steuern des an den Ultraschallwandler abgegebenen Antriebssignals basierend auf dem  
 Gewebetyp, welcher mit dem Endeffektor interagiert;

wobei der Reibungskoeffizient des Gewebes basierend auf dem folgenden Ausdruck ermittelt wird:

$$\mu = \frac{\dot{Q}}{v \cdot N}$$

wobei  $\dot{Q}$  die Rate der Wärmeerzeugung ist,  $v$  die Geschwindigkeit der Ultraschallbewegung der Ultraschallklinge  
 20 ist, und  $N$  die Kraft ist, die durch den Endeffektor auf das Gewebe ausgeübt wird.

8. Generator nach Anspruch 7, wobei der Klemmarm eine Elektrode umfasst, wobei der Generator ferner eine zweite  
 Treiberschaltung umfasst, die ausgestaltet ist, um Hochfrequenz (HF)-Energie an die Elektrode abzugeben, wobei  
 der Prozessor ausgestaltet ist, um Energie, die durch die zweite Treiberschaltung an die Elektrode abgegeben wird,  
 25 basierend auf einem Gewebetyp abzugeben, der mit dem Endeffektor interagiert, und wobei der Prozessor aus-  
 gestaltet ist, um die Abgabe von HF-Energie an die Elektrode basierend auf einem Gewebetyp, der mit dem Endeffektor  
 interagiert, dynamisch zu steuern und zwischen der Abgabe von HF-Energie und dem Ultraschallantriebssignal  
 basierend auf dem Gewebetyp, der mit dem Endeffektor interagiert, zu schalten.

9. Generator nach Anspruch 7, wobei der Prozessor ausgestaltet ist zum:

Speichern von zwei Datenpunkten der gemessenen Kraft und Leistung, die an den Ultraschallwandler abge-  
 geben wird, in einem Puffer;  
 Ermitteln eines Neigungswerts von Leistung gegen Kraft basierend auf den beiden Datenpunkten;  
 35 Ermitteln eines statistischen Maßes  $R^2$  des Neigungswerts relativ zu einer angepassten Regressionslinie; und  
 Ermitteln des Reibungskoeffizienten des Gewebes basierend auf dem Neigungswert;  
 wobei der Prozessor gegebenenfalls ferner ausgestaltet ist zum:

Vergleichen von  $R^2$  mit einem Schwellenwert; und  
 40 Auswählen eines Neigungswerts, der einem  $R^2$ -Wert zugeordnet ist, der größer als oder gleich dem Schwel-  
 lenwert ist, um den Reibungskoeffizienten des Gewebes zu ermitteln.

10. Generator nach Anspruch 7, wobei der Prozessor ausgestaltet ist zum:

Vergleichen des Reibungskoeffizienten des Gewebes mit den Reibungskoeffizienten von Gewebe, die in einer  
 Gewebeinformationsdatenbank gespeichert sind;  
 Ermitteln des Gewebetyps basierend auf dem Reibungskoeffizienten des Gewebes; und  
 45 Abgeben des Antriebssignals an den Endeffektor basierend auf dem Reibungskoeffizienten des Gewebes.

11. Generator nach Anspruch 7, ferner umfassend eine zweite Treiberschaltung, die ausgestaltet ist, um Hochfrequenz  
 (HF)-Energie an die Elektrode abzugeben.

12. Chirurgisches Instrument nach Anspruch 1 oder Generator nach Anspruch 7, wobei der Prozessor ausgestaltet ist,  
 um die Leistung des Antriebssignals, das an den Ultraschallwandler abgegeben wird, basierend auf dem Reibungs-  
 55 koeffizient des Gewebes abzugeben.

13. Chirurgisches Instrument nach Anspruch 1 oder Generator nach Anspruch 7, wobei der Prozessor ausgestaltet ist,  
 um Leistung des Antriebssignals, das an den Ultraschallwandler abgegeben wird, basierend auf dem Hydratisie-

rungsniveau des Gewebes abzugeben, wobei die Leistung des Antriebssignals, das an den Ultraschallwandler abgegeben wird, gegebenenfalls herabgesetzt wird, wenn das Hydratisierungsniveau des Gewebes zunimmt.

14. Chirurgisches Instrument nach Anspruch 2 oder Generator nach Anspruch 8, wobei entweder der Prozessor ausgestaltet ist, um die Abgabe des Ultraschallantriebssignals an den Ultraschallwandler zu steuern, falls der Gewebetyp einer mit einer Muskelstruktur ist, oder der Prozessor ausgestaltet ist, um die Abgabe der HF-Energie zu der Elektrode zu steuern, falls der Gewebetyp einer mit einer vaskulären Struktur ist, oder der Prozessor ausgestaltet ist, um die Abgabe eines Antriebssignals an den Ultraschallwandler entsprechend einer raschen Schnittbewegung zu steuern, falls der Gewebetyp einer mit einer dünnen Mesenterialstruktur ist.

## Revendications

1. Instrument chirurgical (108) permettant de coaguler et disséquer un tissu, l'instrument chirurgical comprenant :

un processeur ;  
un effecteur terminal (125) au niveau d'une extrémité distale de l'instrument chirurgical, l'effecteur terminal étant conçu pour interagir avec le tissu, l'effecteur terminal comprenant :

un bras (145) de pince ;  
une lame à ultrasons (149) ;  
un capteur de force en communication avec le processeur et conçu pour mesurer une force appliquée à un tissu situé entre le bras de pince et la lame à ultrasons ; et  
un capteur de température en communication avec le processeur ;

un transducteur à ultrasons (120) couplé acoustiquement à la lame à ultrasons et conçu pour recevoir un signal d'attaque en provenance d'un générateur pour produire un mouvement à ultrasons de la lame à ultrasons et émettre de l'énergie vers la lame à ultrasons ;

**caractérisé en ce que** le processeur est conçu pour :

déterminer un type de tissu ou le niveau d'hydratation du tissu interagissant avec l'effecteur terminal sur la base d'un coefficient de frottement de tissu, le coefficient de frottement de tissu étant déterminé sur la base de la force appliquée au tissu par l'effecteur terminal, du mouvement à ultrasons de la lame à ultrasons et d'une puissance thermique produite par l'effecteur terminal ; et  
commander dynamiquement le signal d'attaque émis vers le transducteur à ultrasons sur la base du type de tissu ou du niveau d'hydratation du tissu interagissant avec l'effecteur terminal ;

dans lequel le coefficient de frottement du tissu est déterminé sur la base de l'expression suivante :

$$\mu = \frac{\dot{Q}}{v \cdot N}$$

dans laquelle  $\dot{Q}$  est la production de puissance thermique,  $v$  est la vitesse du mouvement à ultrasons de la lame à ultrasons, et  $N$  est la force appliquée au tissu par l'effecteur terminal.

2. Instrument chirurgical selon la revendication 1, dans lequel le bras de pince comprend une électrode, dans lequel le processeur est en outre conçu pour :

commander dynamiquement l'émission d'une énergie radiofréquence (RF) vers l'électrode sur la base du type de tissu interagissant avec l'effecteur terminal ; et  
commuter entre l'émission de l'énergie RF et du signal d'attaque à ultrasons sur la base du type de tissu interagissant avec l'effecteur terminal.

3. Instrument chirurgical selon l'une quelconque des revendications 1 et 2, dans lequel le capteur de force comprend une jauge de contrainte accouplée à l'effecteur terminal, éventuellement dans lequel la mesure de la force est communiquée de la jauge de contrainte au processeur.



4. Instrument chirurgical selon l'une quelconque des revendications 1 ou 2, dans lequel le processeur est conçu pour :

mémoriser deux points de données de la force mesurée et de l'énergie distribuée vers l'effecteur terminal dans une mémoire tampon ;

déterminer une valeur de pente de l'énergie par rapport à la force sur la base de deux points de données ;

déterminer une mesure statistique  $R^2$  de la valeur de pente par rapport à une droite de régression ajustée ; et

déterminer le coefficient de frottement du tissu sur la base de la valeur de pente ;

éventuellement dans lequel le processeur est en outre conçu pour :

comparer  $R^2$  à une valeur seuil ; et

sélectionner une valeur de pente associée à une valeur  $R^2$  qui est supérieure ou égale à la valeur seuil pour déterminer le coefficient de frottement du tissu.

5. Instrument chirurgical selon l'une quelconque des revendications 1 ou 2, dans lequel le processeur est conçu pour :

comparer le coefficient de frottement de tissu à des coefficients de frottement de tissu mémorisés dans une base de données d'informations de tissu ;

déterminer le type de tissu sur la base du coefficient de frottement de tissu ; et

émettre le signal d'attaque vers le transducteur à ultrasons sur la base du coefficient de frottement de tissu.

6. Instrument chirurgical selon l'une quelconque des revendications 1 ou 2, comprenant en outre un générateur (500) conçu pour émettre une énergie radiofréquence (RF) vers l'électrode et/ou le signal d'attaque vers le transducteur à ultrasons.

7. Générateur (500) permettant d'émettre de l'énergie vers un instrument chirurgical (108) permettant de coaguler et disséquer un tissu, l'instrument chirurgical comprenant un effecteur terminal (125) au niveau d'une extrémité distale de celui-ci, l'effecteur terminal étant conçu pour interagir avec le tissu, l'effecteur terminal comprenant un bras (145) de pince, une lame à ultrasons (149), un capteur de force en communication avec le processeur et conçu pour mesurer une force appliquée à un tissu situé entre le bras de pince et la lame à ultrasons, et un capteur de température en communication avec un processeur, un transducteur à ultrasons (120) couplé acoustiquement à la lame à ultrasons et conçu pour recevoir un signal d'attaque en provenance d'un générateur pour produire un mouvement à ultrasons de la lame à ultrasons et émettre de l'énergie vers la lame à ultrasons, le générateur comprenant :

un circuit d'attaque conçu pour émettre un signal d'attaque vers un transducteur à ultrasons ; et

**caractérisé par** un processeur (502) configuré pour commander une énergie émise par le circuit d'attaque vers le transducteur à ultrasons sur la base d'un type de tissu ou du niveau d'hydratation du tissu interagissant avec l'effecteur terminal ;

dans lequel le processeur est conçu pour :

recevoir une mesure de force appliquée au tissu par l'effecteur terminal, le mouvement à ultrasons de la lame à ultrasons et une puissance thermique produite par l'effecteur terminal ;

déterminer le type de tissu ou un niveau d'hydratation du tissu interagissant avec l'effecteur terminal sur la base d'un coefficient de frottement du tissu, le coefficient de frottement du tissu étant déterminé sur la base de la force appliquée au tissu par l'effecteur terminal, le mouvement à ultrasons de la lame à ultrasons et une puissance thermique produite par l'effecteur terminal ; et

commander dynamiquement le signal d'attaque émis vers le transducteur à ultrasons sur la base du type de tissu interagissant avec l'effecteur terminal ;

dans lequel le coefficient de frottement du tissu est déterminé sur la base de l'expression suivante :

$$\mu = \frac{\dot{Q}}{v \cdot N}$$

dans laquelle  $\dot{Q}$  est la production de puissance thermique,  $v$  est la vitesse du mouvement à ultrasons de la lame à ultrasons et  $N$  est la force appliquée au tissu par l'effecteur terminal.

8. Générateur selon la revendication 7, dans lequel le bras de pince comprend une électrode, le générateur comprenant

en outre un second circuit d'attaque conçu pour émettre une énergie radiofréquence (RF) vers l'électrode, dans lequel le processeur est conçu pour commander une énergie émise par le second circuit d'attaque vers l'électrode sur la base d'un type de tissu interagissant avec l'effecteur terminal, et dans lequel le processeur est conçu pour commander dynamiquement l'émission d'énergie RF vers l'électrode sur la base du type de tissu interagissant avec l'effecteur terminal et commuter entre l'émission de l'énergie RF et du signal d'attaque à ultrasons sur la base du type de tissu interagissant avec l'effecteur terminal.

9. Générateur selon la revendication 7, dans lequel le processeur est conçu pour :

mémoriser deux points de données de la force mesurée et de l'énergie émise vers le transducteur à ultrasons dans une mémoire tampon ;  
déterminer une valeur de pente de la puissance par rapport à la force sur la base des deux points de données ;  
déterminer une mesure statistique  $R^2$  de la valeur de pente par rapport à une droite de régression ajustée ; et  
déterminer le coefficient de frottement de tissu sur la base de la valeur de pente ;  
éventuellement dans lequel le processeur est en outre conçu pour :

comparer  $R^2$  à une valeur seuil ; et  
sélectionner une valeur de pente associée à une valeur  $R^2$  qui est supérieure ou égale à la valeur seuil pour déterminer le coefficient de frottement de tissu.

10. Générateur selon la revendication 7, dans lequel le processeur est conçu pour :

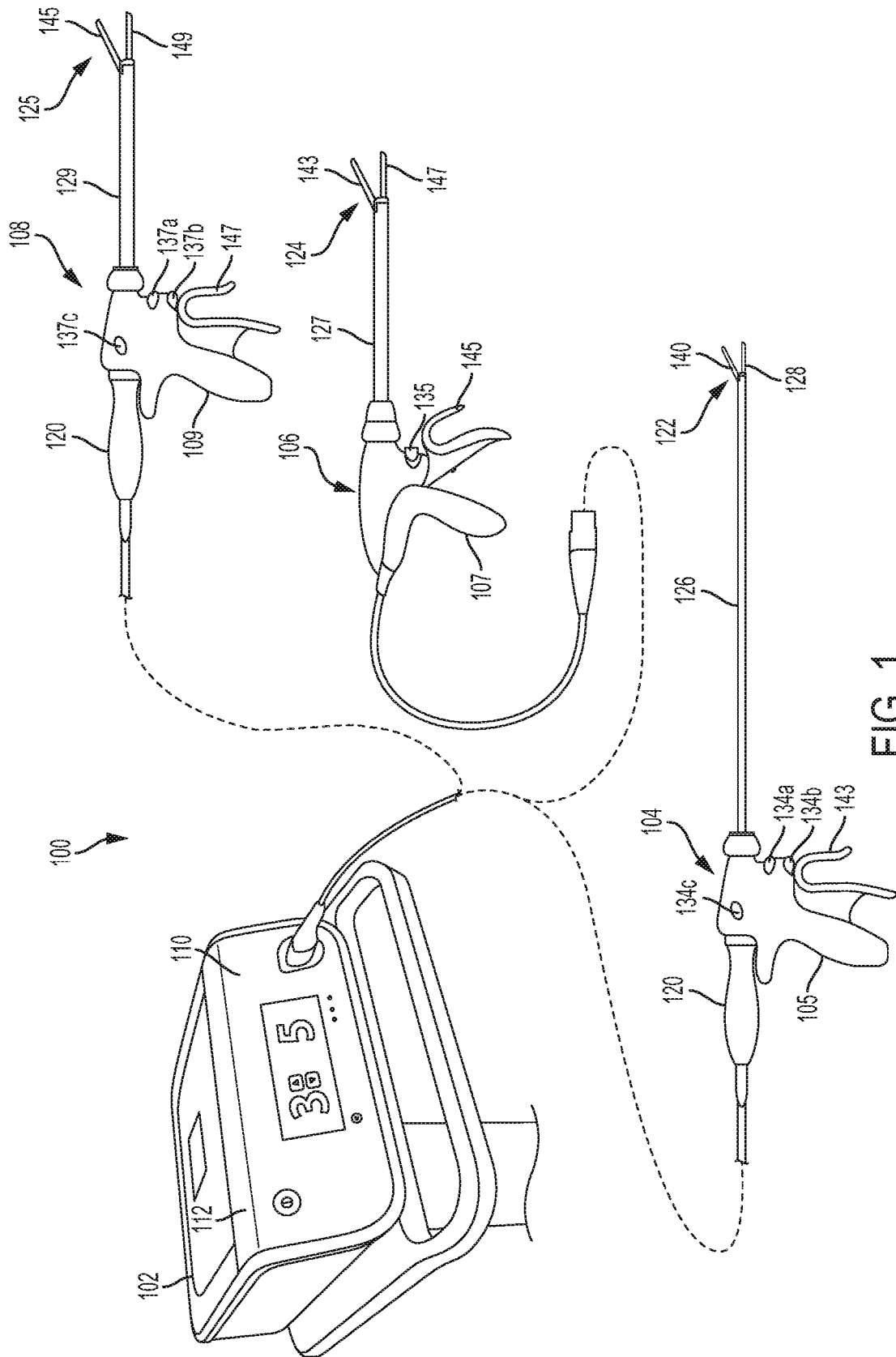
comparer le coefficient de frottement de tissu à des coefficients de frottement de tissu mémorisés dans une base de données d'informations de tissu ;  
déterminer le type de tissu sur la base du coefficient de frottement de tissu ; et  
émettre le signal d'attaque vers l'effecteur terminal sur la base du coefficient de frottement de tissu.

11. Générateur selon la revendication 7, comprenant en outre un second circuit d'attaque conçu pour émettre une énergie radiofréquence (RF) vers l'électrode.

12. Instrument chirurgical selon la revendication 1 ou générateur selon la revendication 7, le processeur étant conçu pour commander l'intensité du signal d'attaque émis vers le transducteur à ultrasons sur la base du coefficient de frottement de tissu.

13. Instrument chirurgical selon la revendication 1 ou générateur selon la revendication 7, dans lequel le processeur est conçu pour commander l'intensité du signal d'attaque émis vers le transducteur à ultrasons sur la base du niveau d'hydratation du tissu, éventuellement dans lequel l'intensité du signal d'attaque émis vers le transducteur à ultrasons est réduite à mesure que le niveau d'hydratation du tissu diminue.

14. Instrument chirurgical selon la revendication 2 ou générateur selon la revendication 8, dans lequel soit le processeur est conçu pour commander l'émission du signal d'attaque à ultrasons vers le transducteur à ultrasons si le type de tissu est un type possédant une structure musculaire, soit le processeur est conçu pour commander l'émission de l'énergie RF vers l'électrode si le type de tissu est un type possédant une structure vasculaire, soit le processeur est conçu pour commander l'émission d'un signal d'attaque vers le transducteur à ultrasons correspondant à un mouvement rapide de coupe si le type de tissu est un type possédant une structure mésentérique mince.



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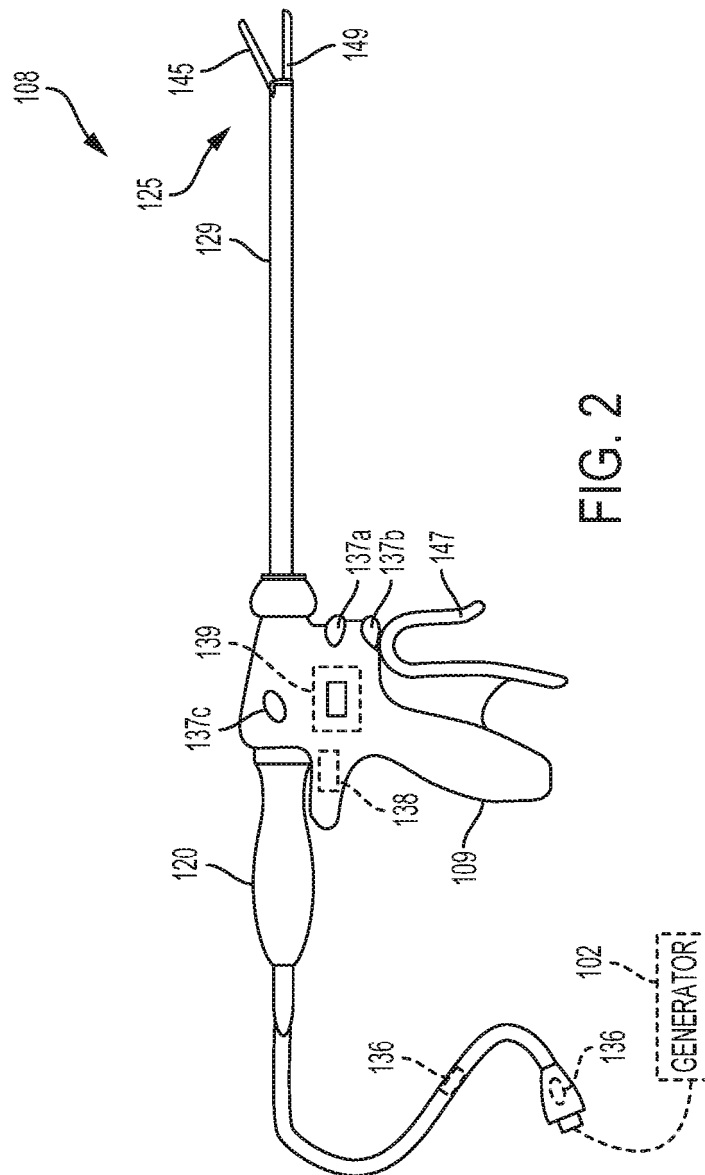


FIG. 2

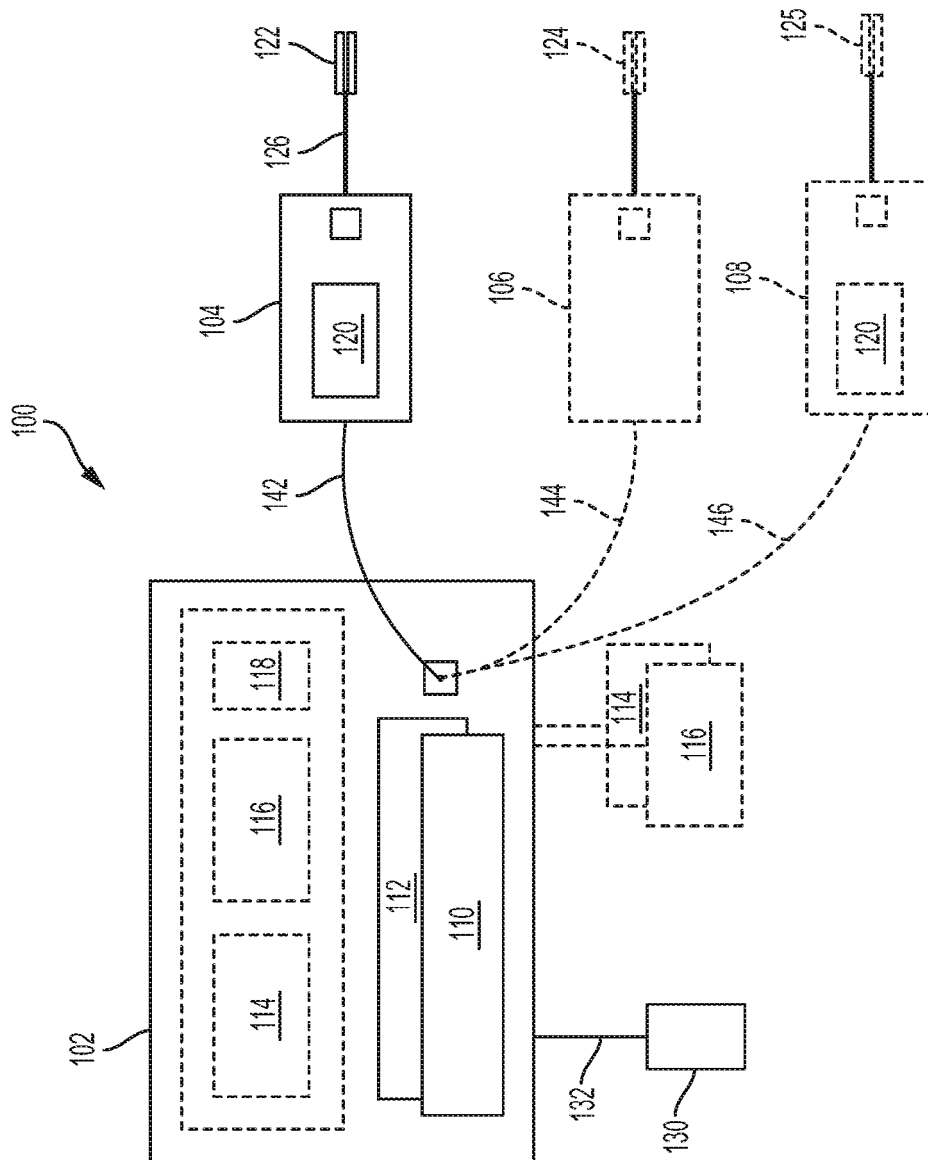


FIG. 3

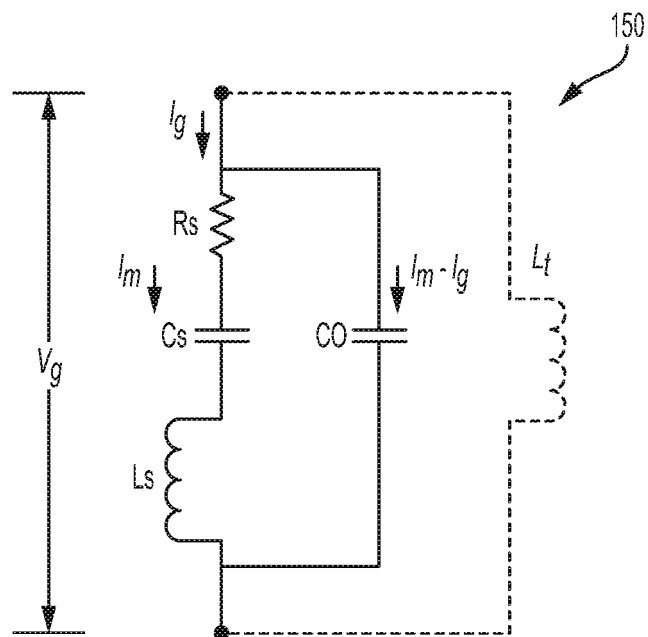
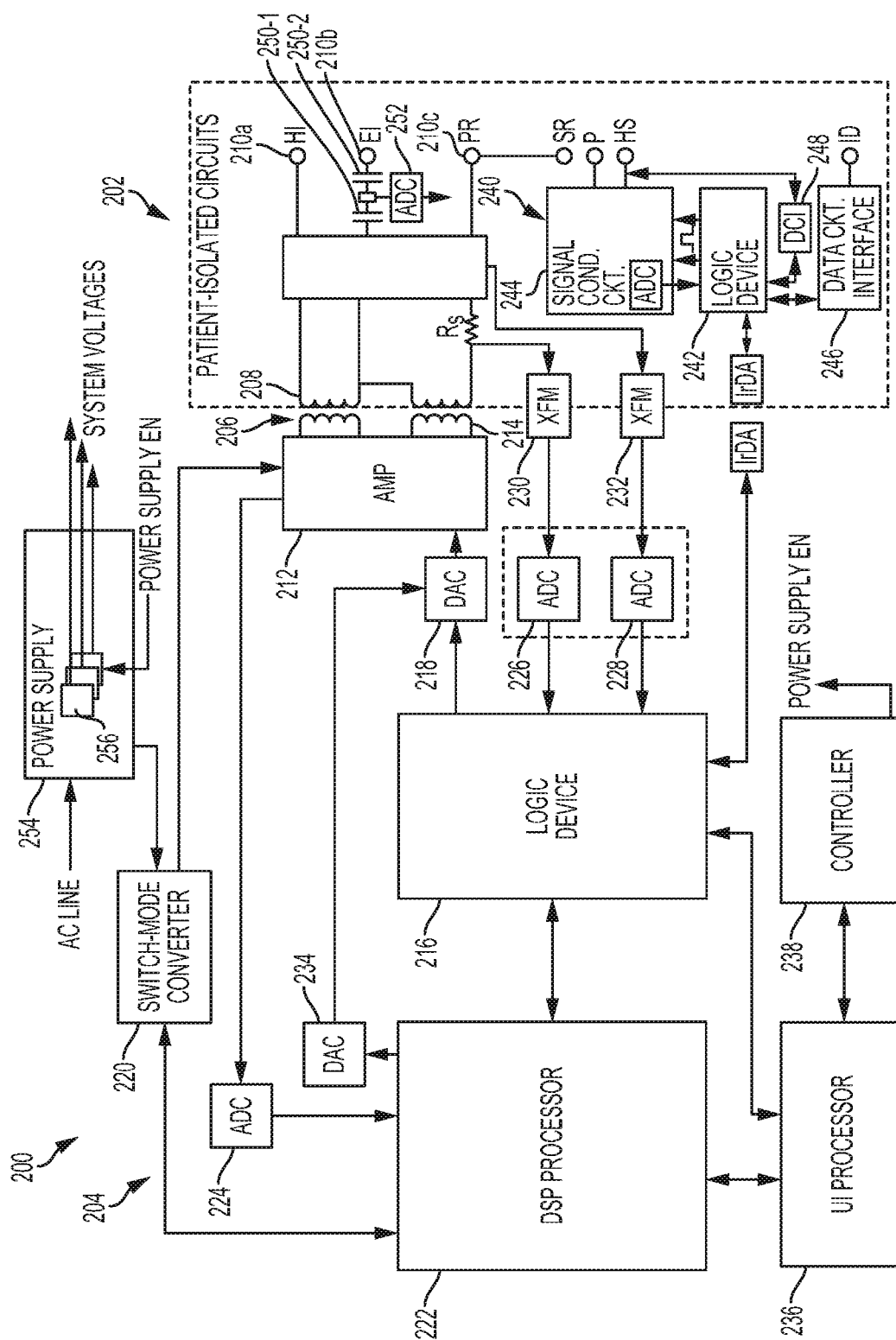


FIG. 4



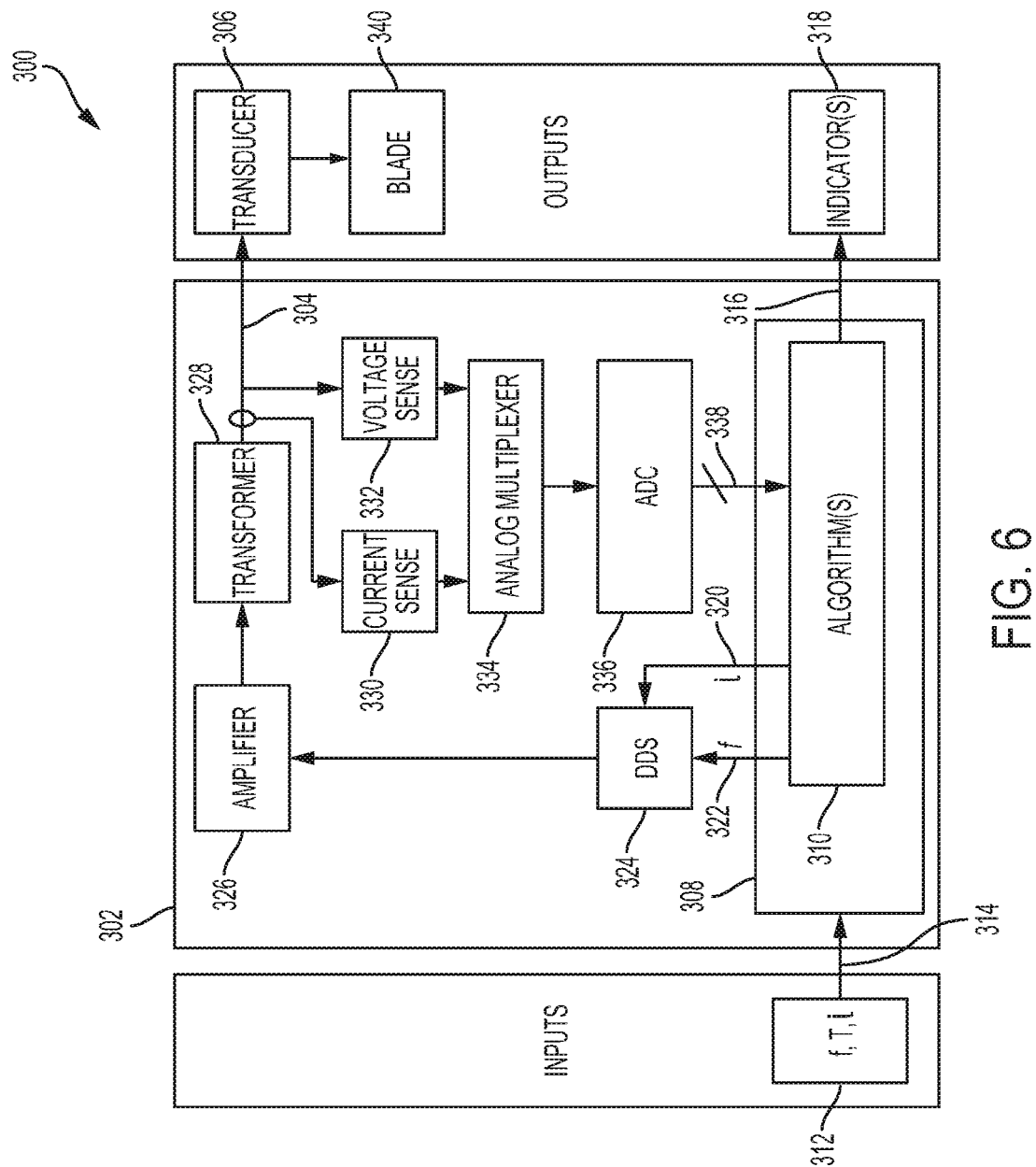


FIG. 6



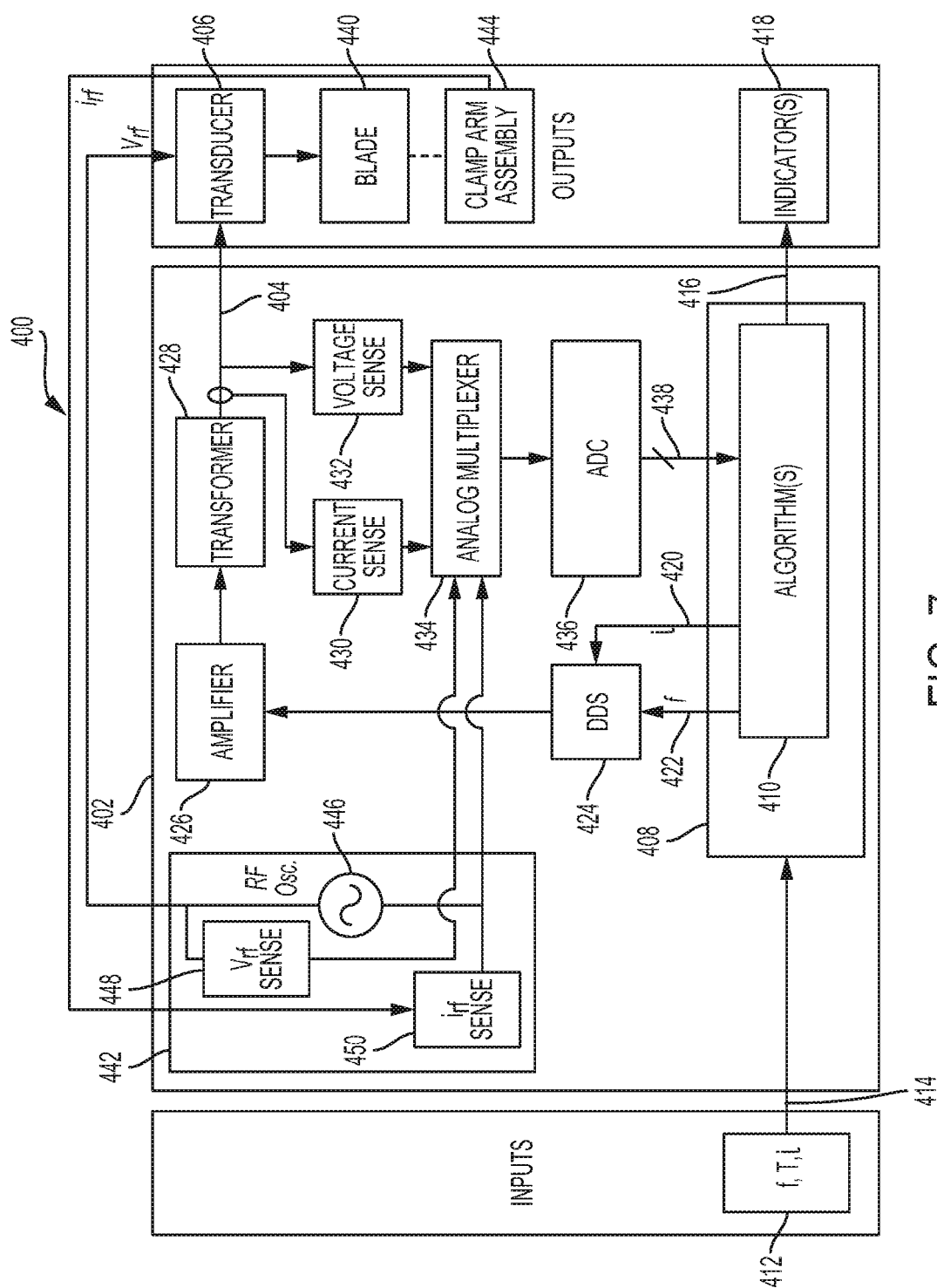


FIG. 7

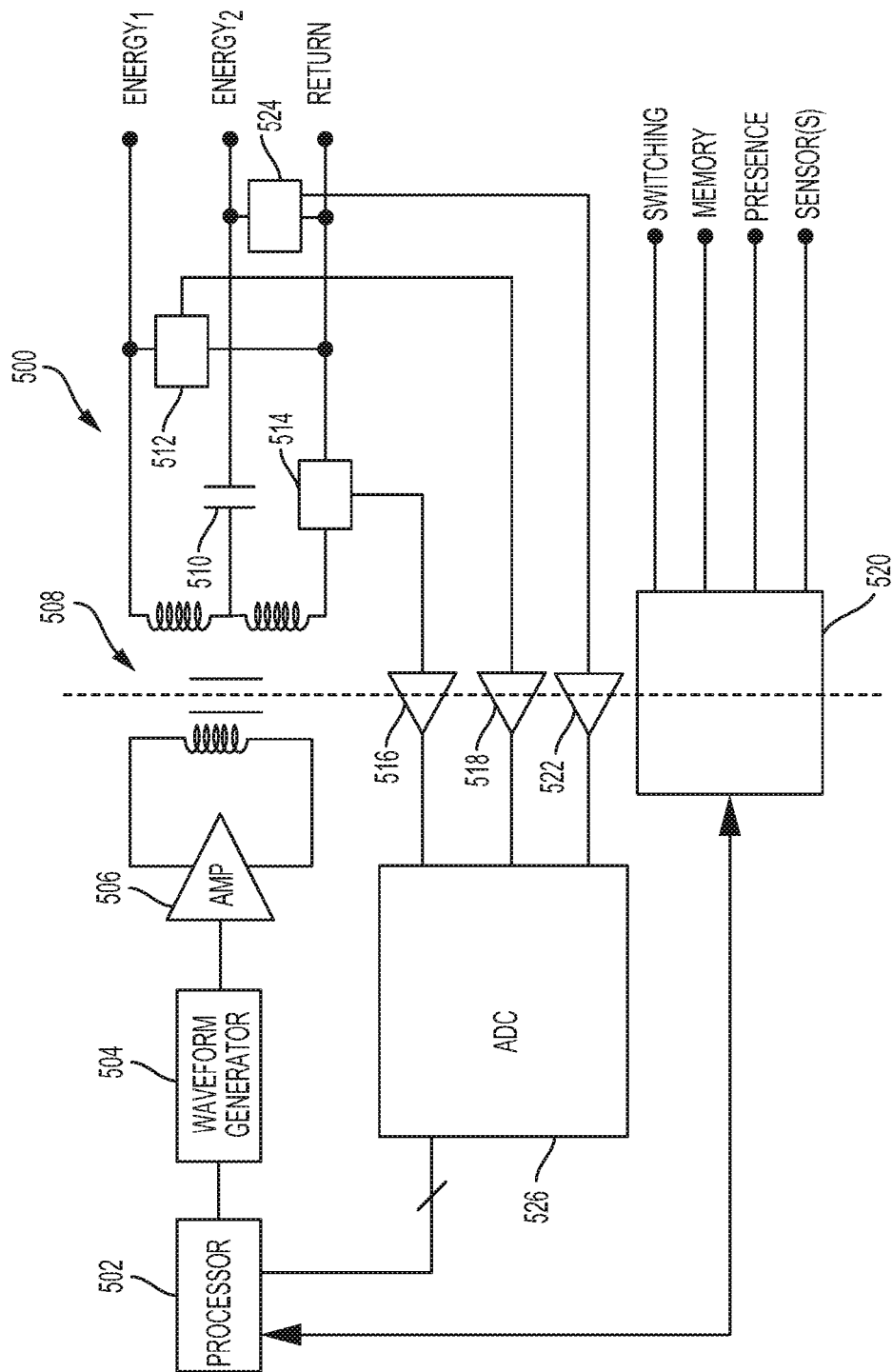


FIG. 8

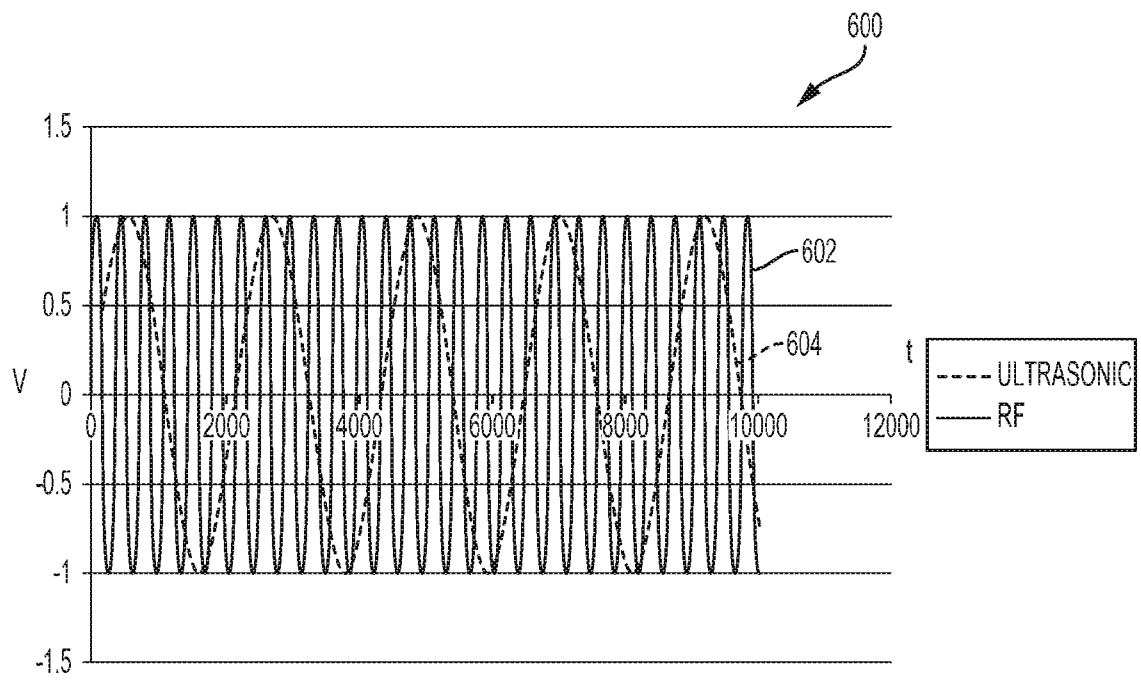


FIG. 9

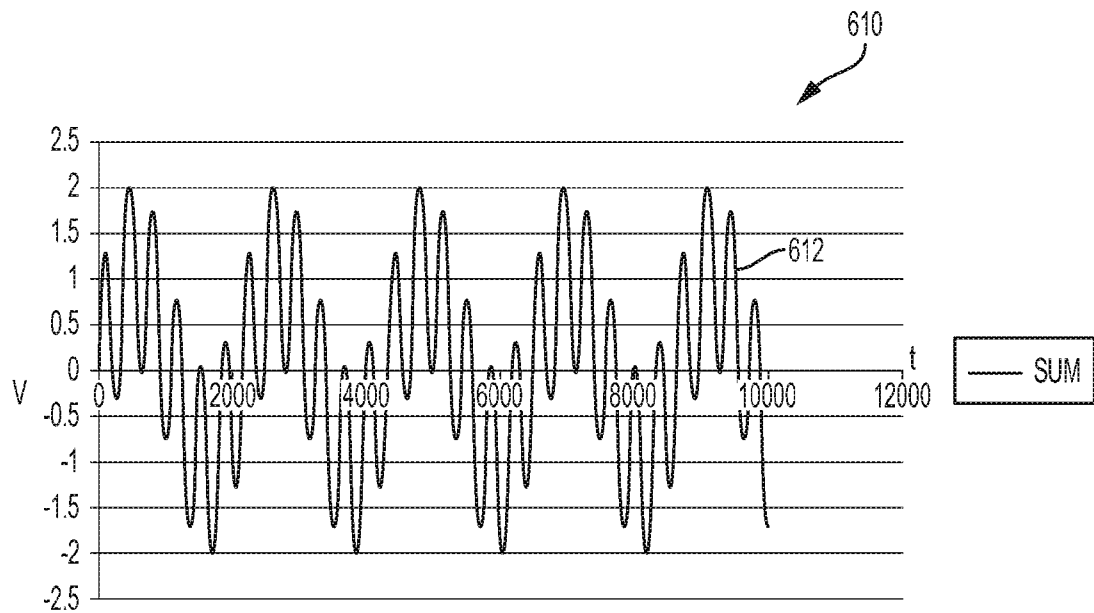


FIG. 10

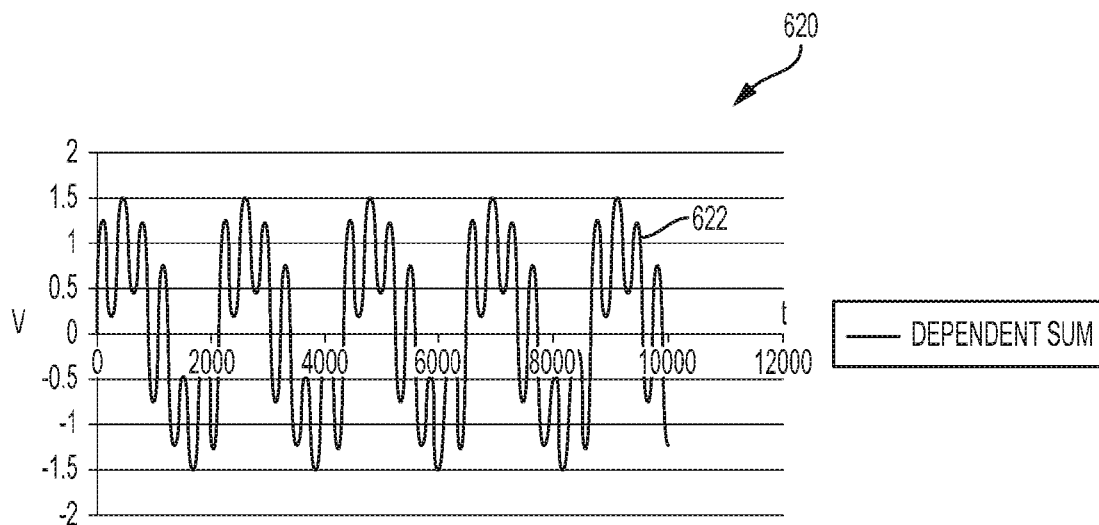


FIG. 11

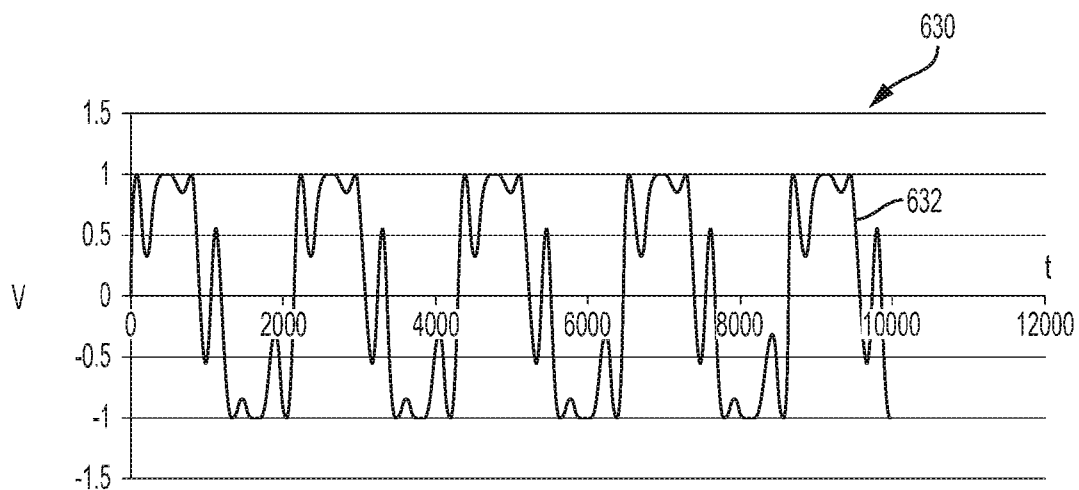


FIG. 12

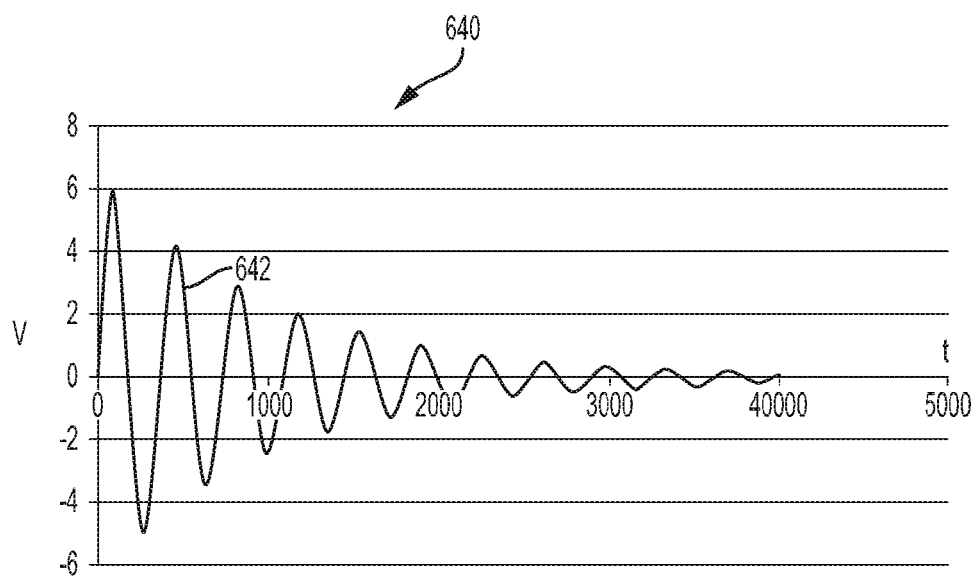


FIG. 13

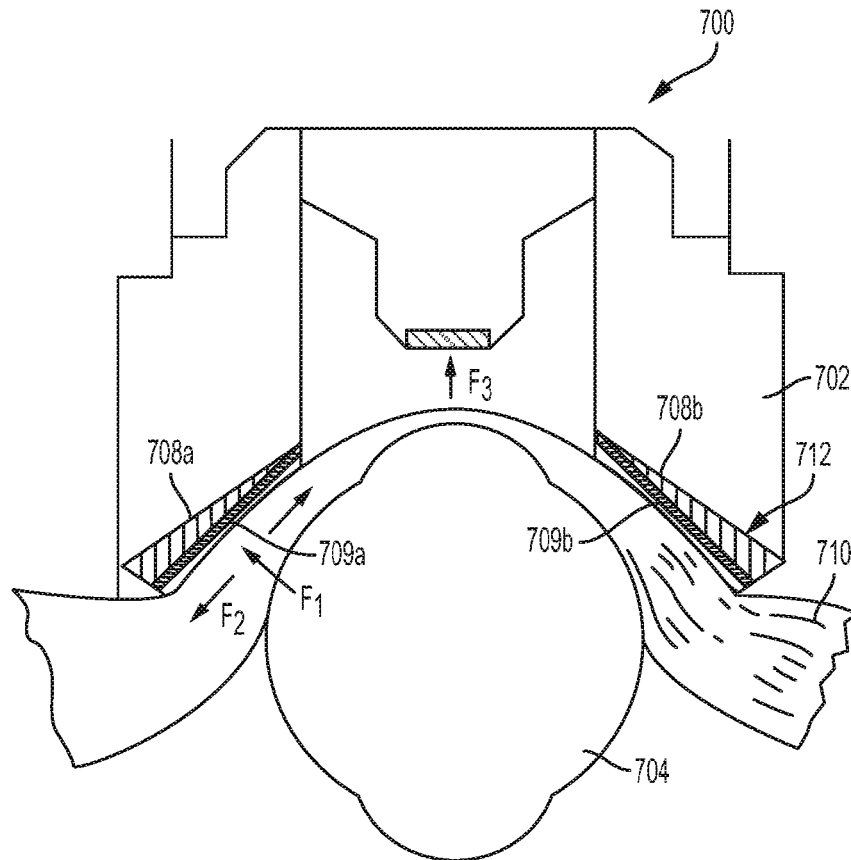


FIG. 14

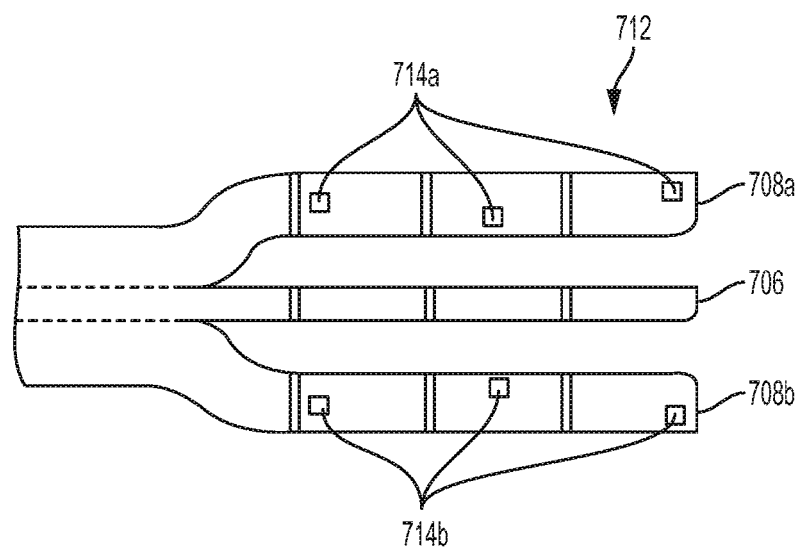


FIG. 15

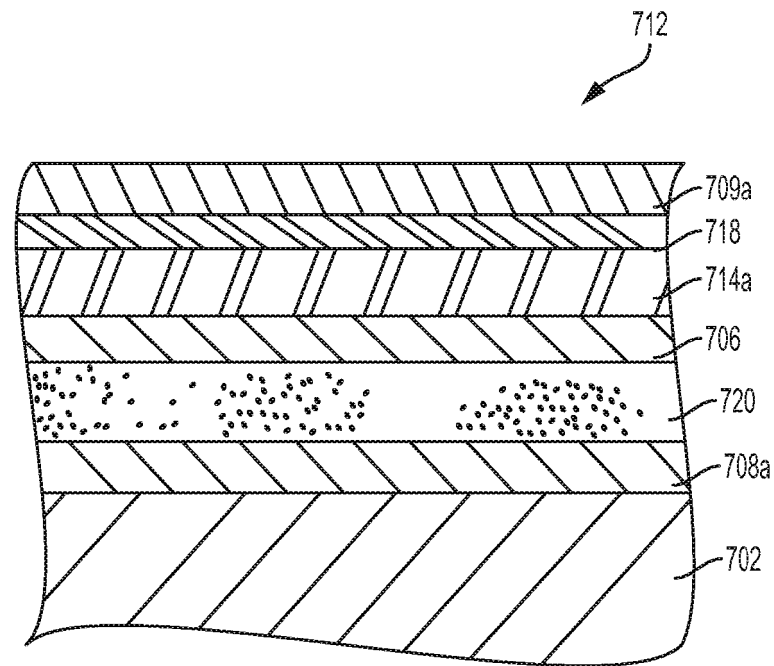


FIG. 16

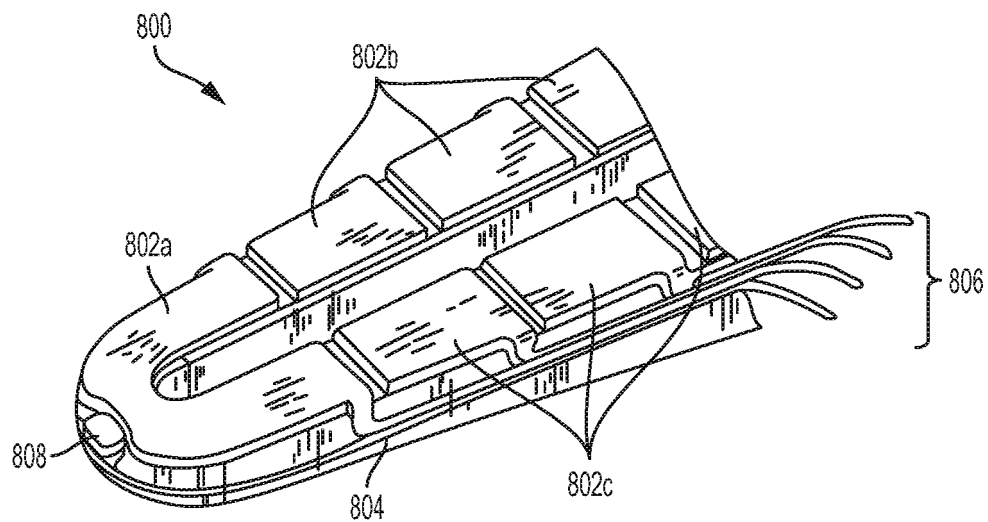
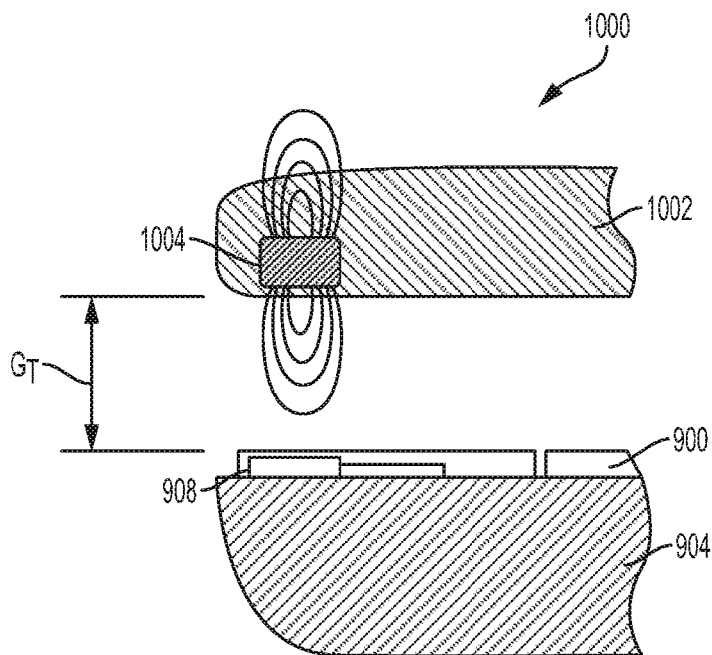
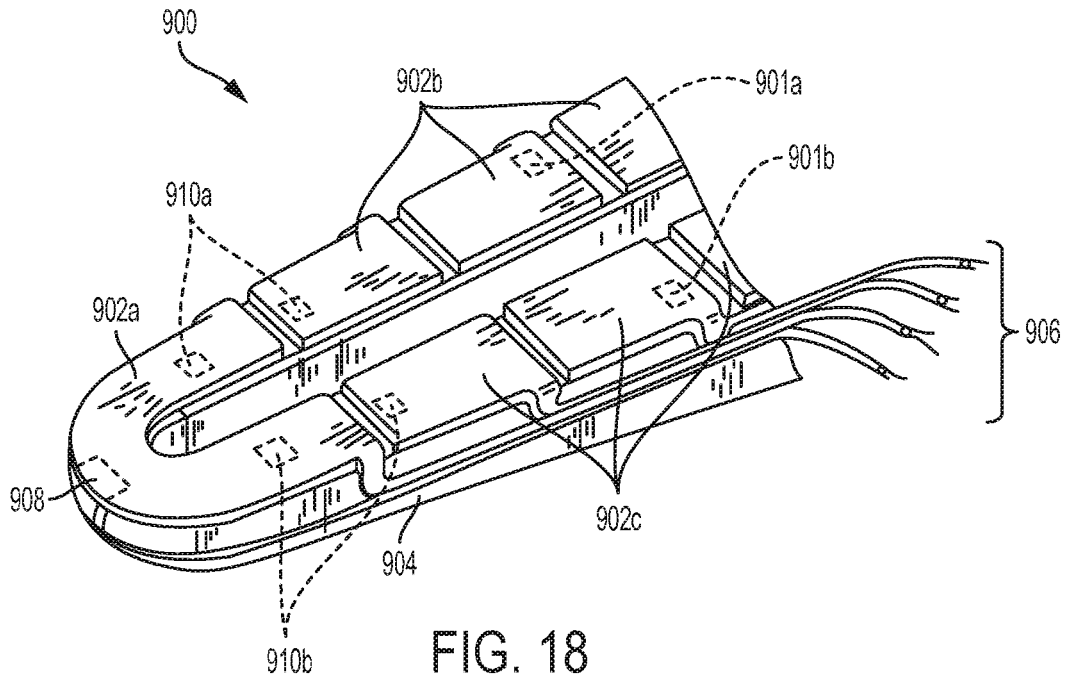


FIG. 17





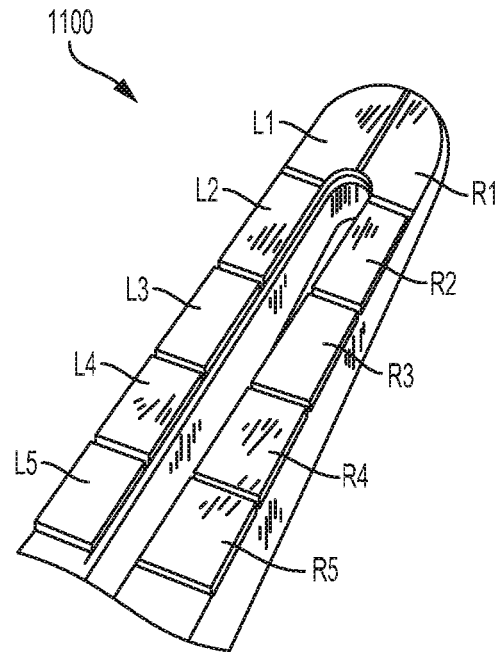


FIG. 20

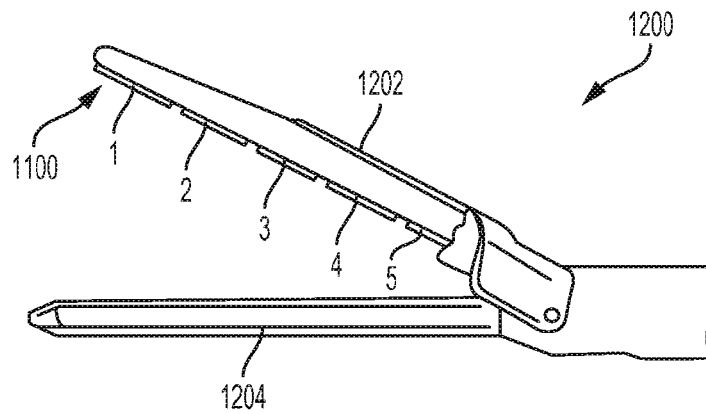


FIG. 21

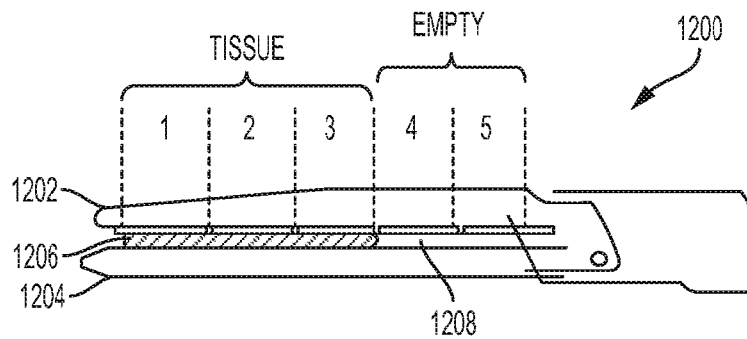


FIG. 22

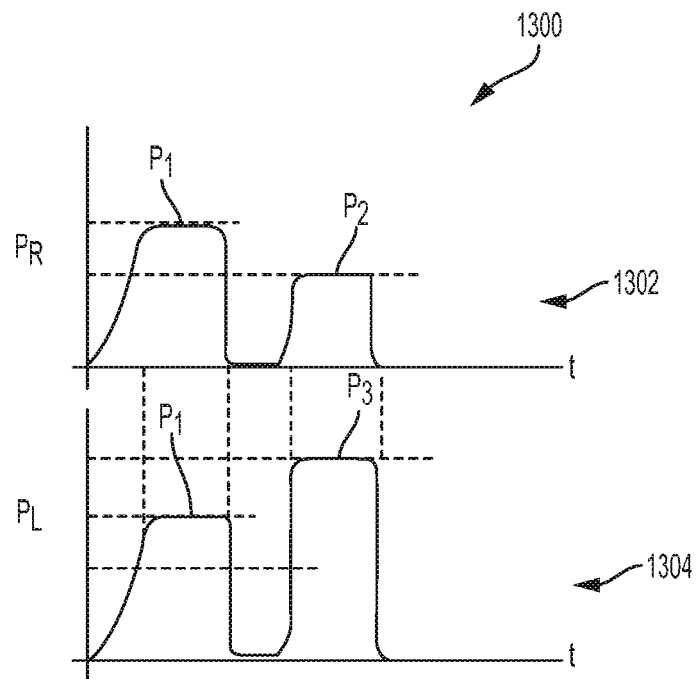


FIG. 23

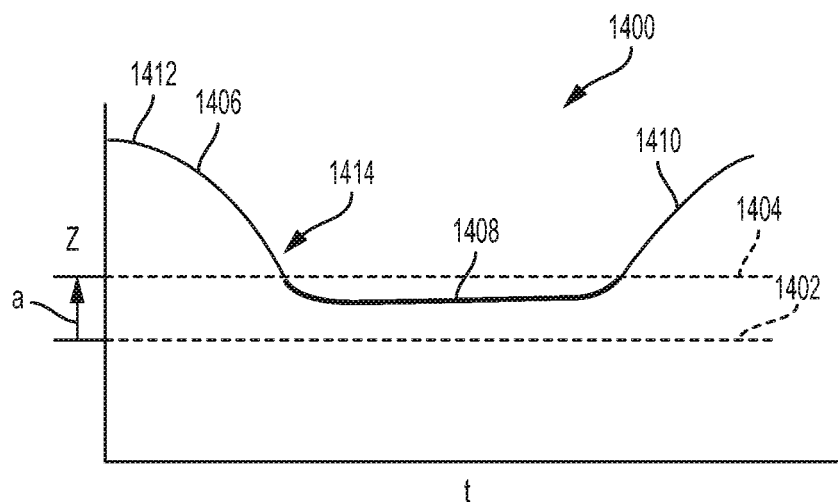


FIG. 24

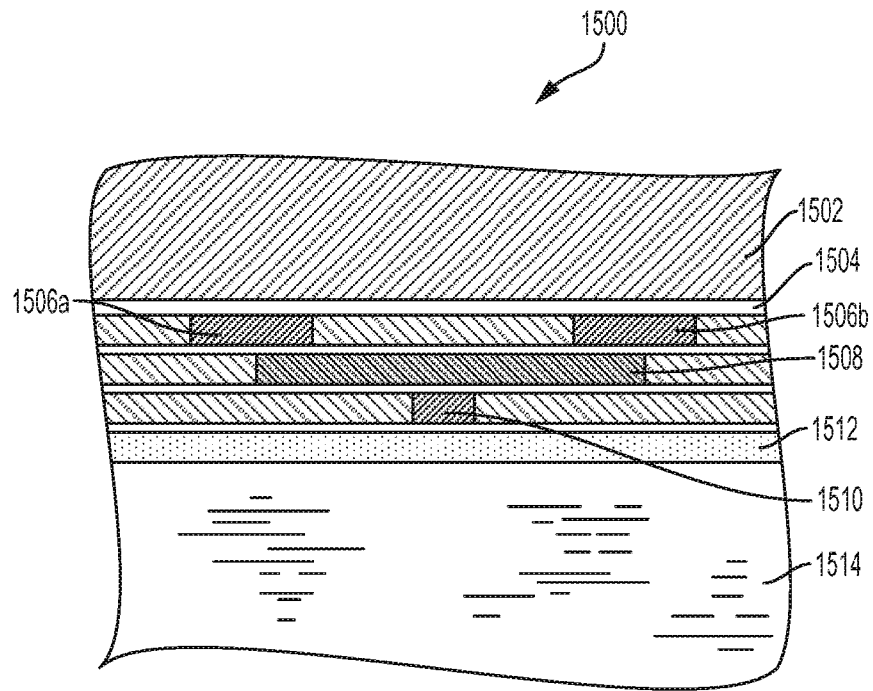


FIG. 25

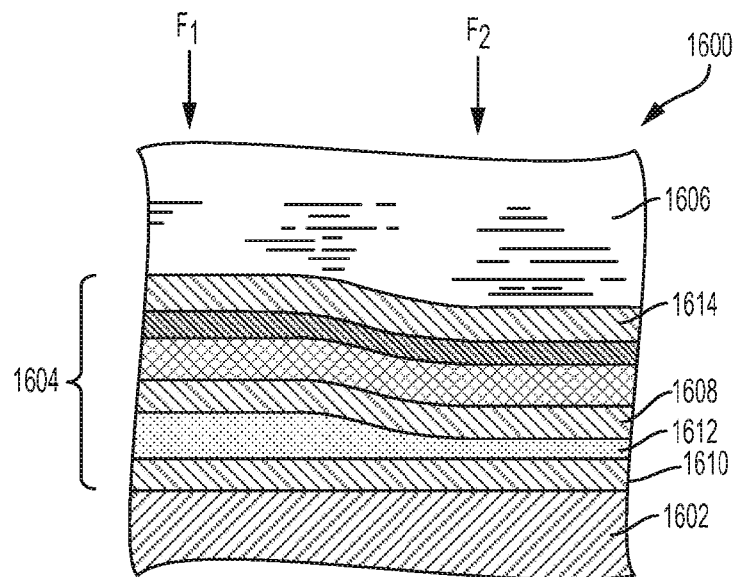


FIG. 26

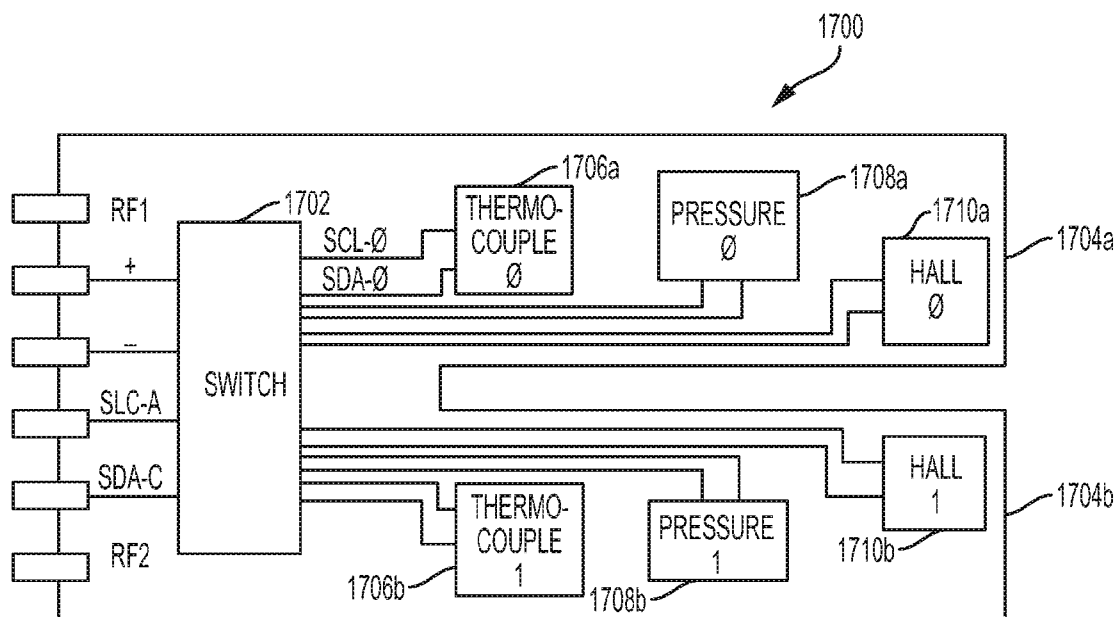


FIG. 27

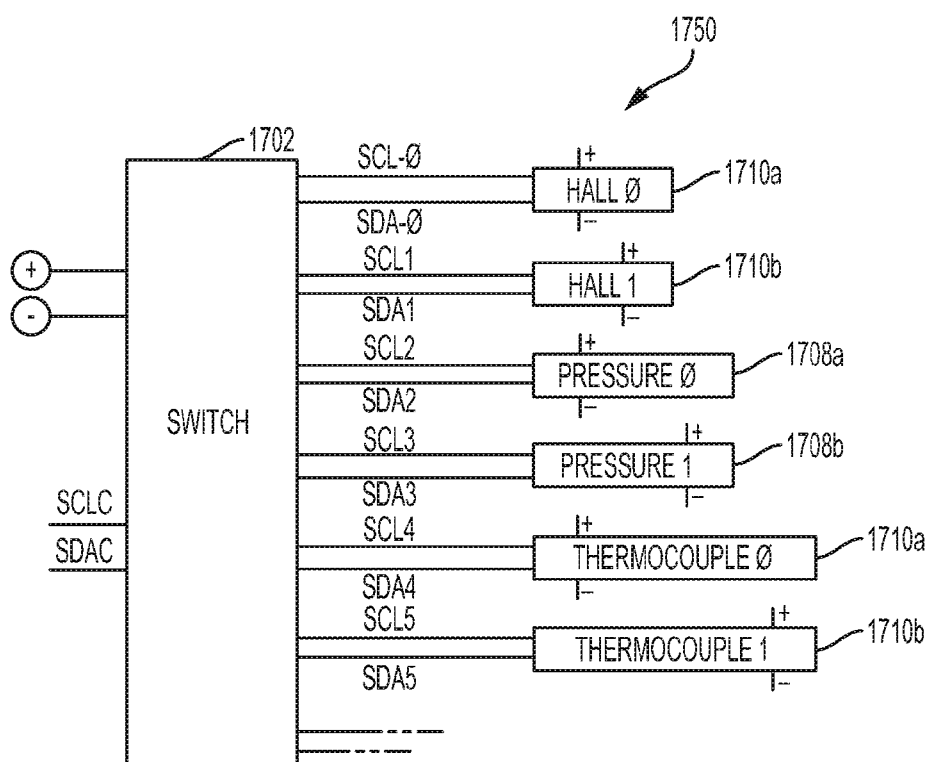


FIG. 28

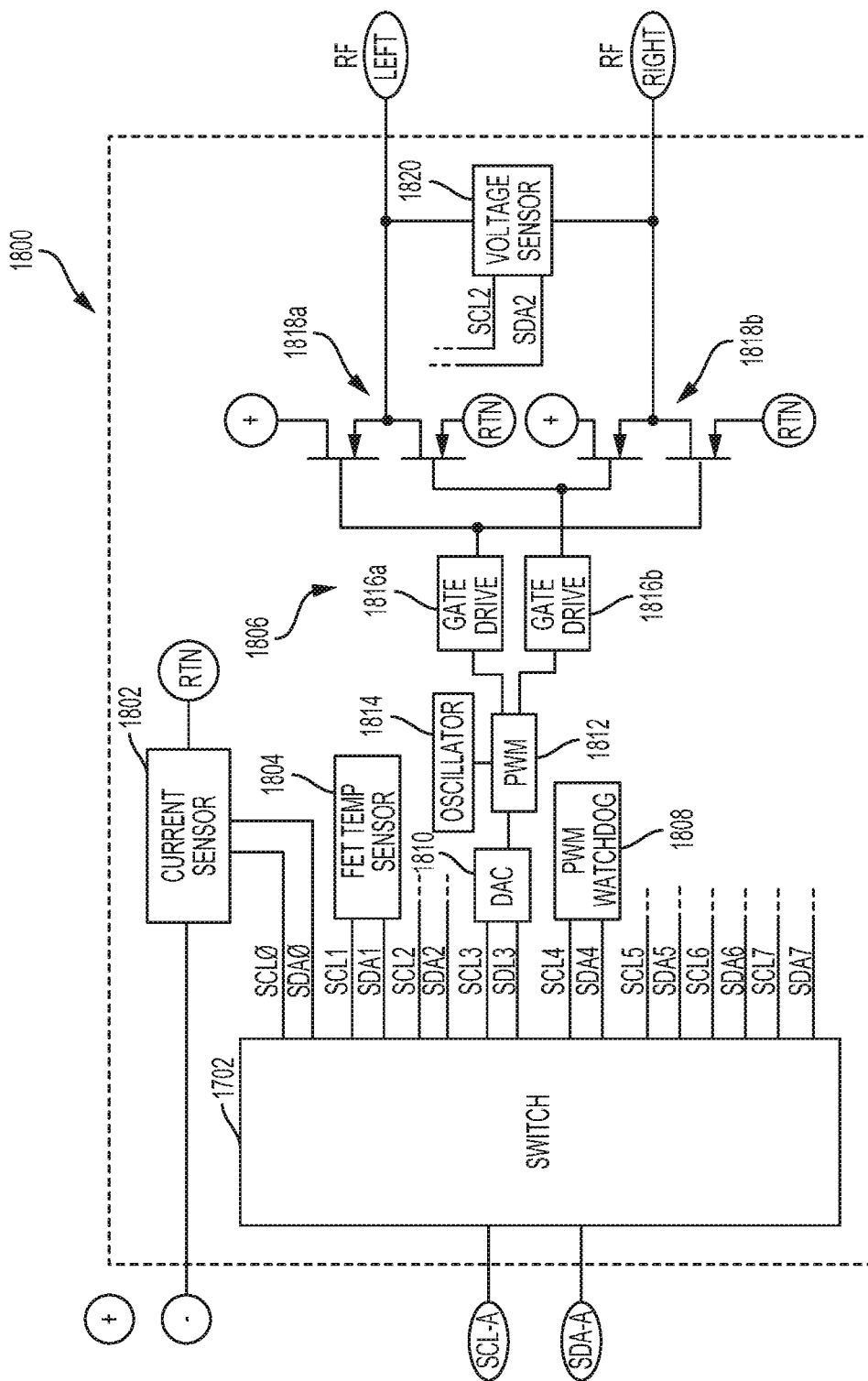


FIG. 29

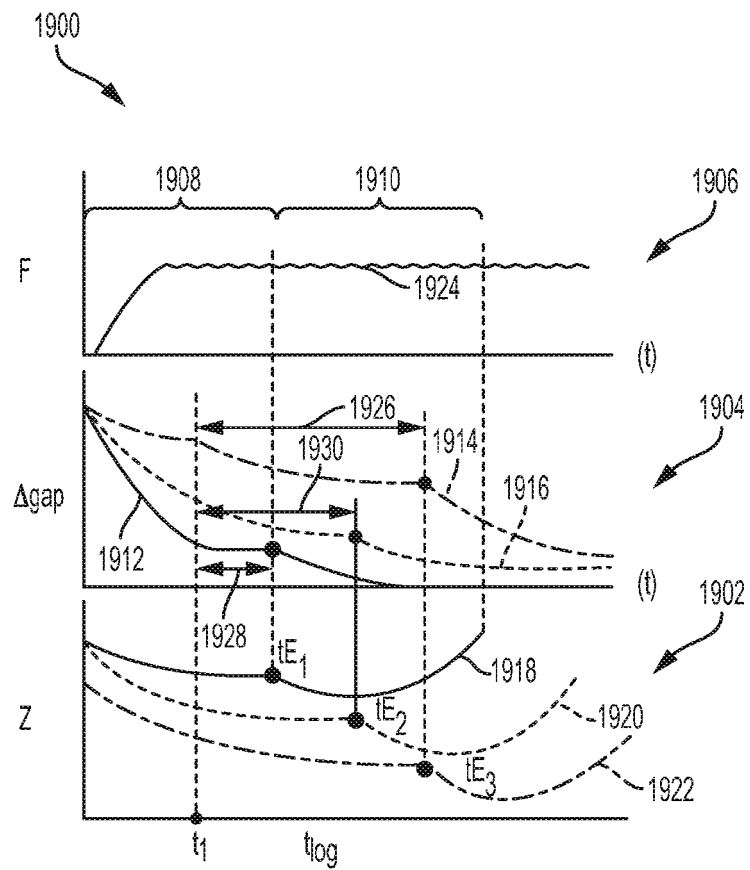


FIG. 30

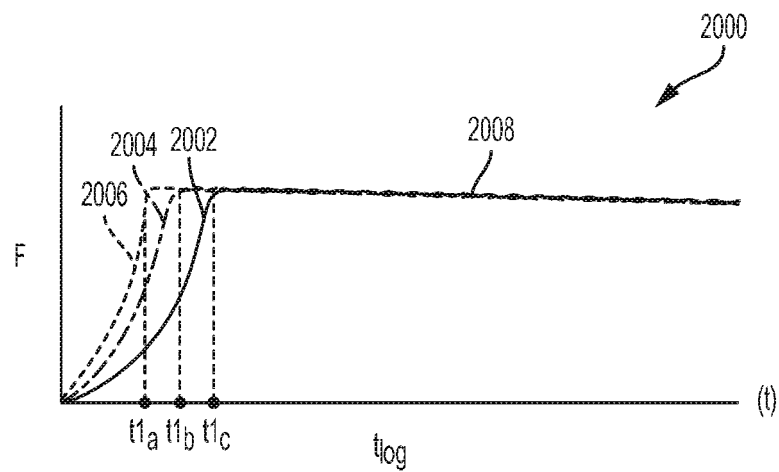


FIG. 31

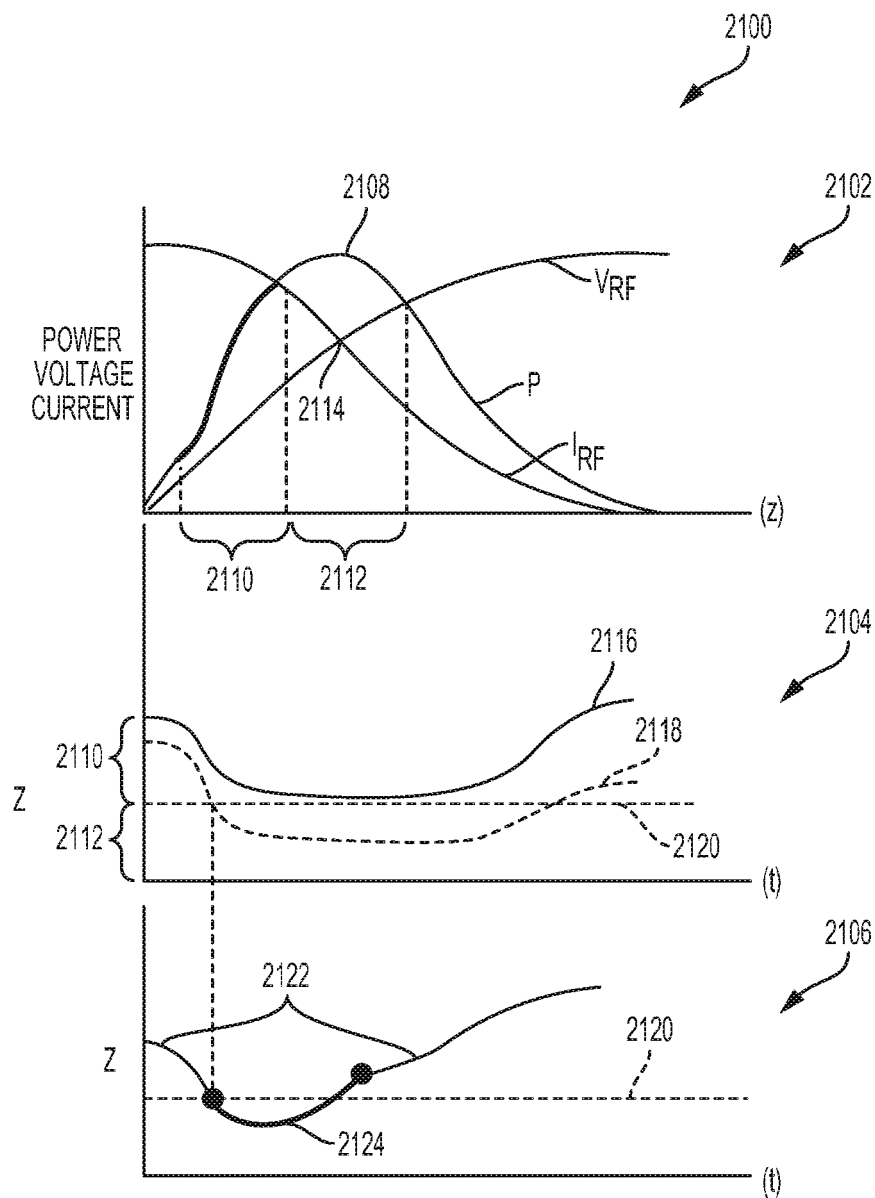


FIG. 32

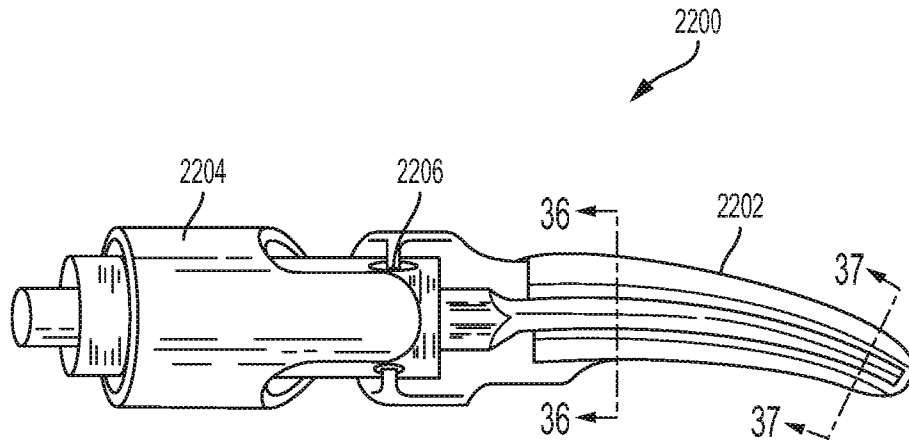


FIG. 33

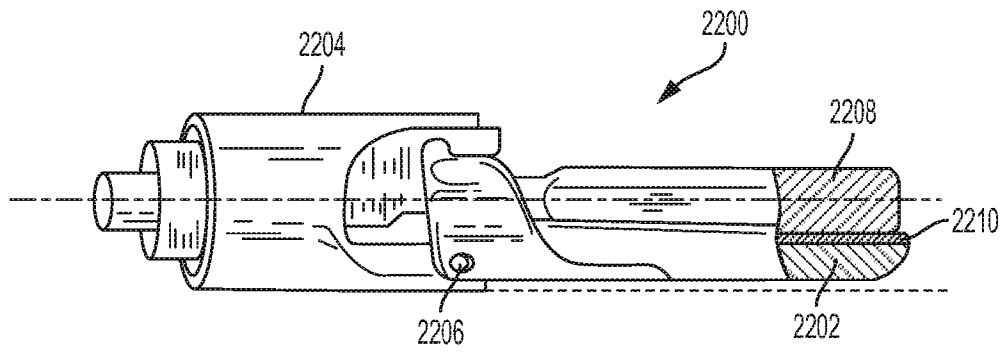


FIG. 34

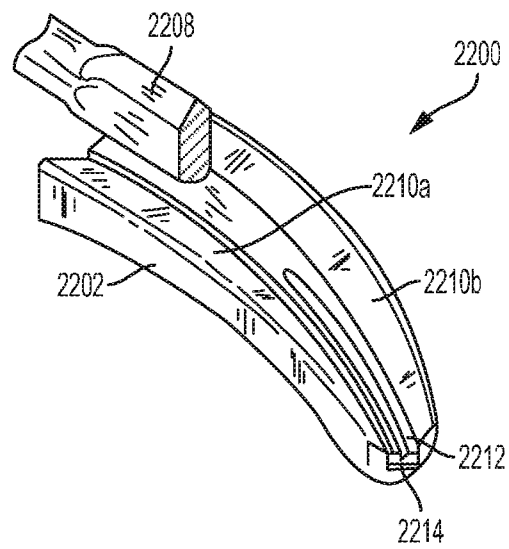


FIG. 35



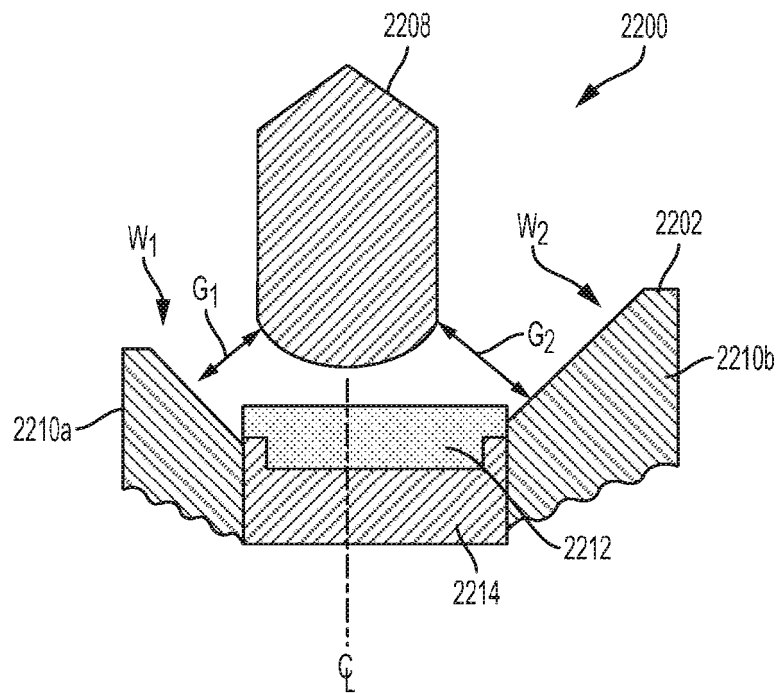


FIG. 36

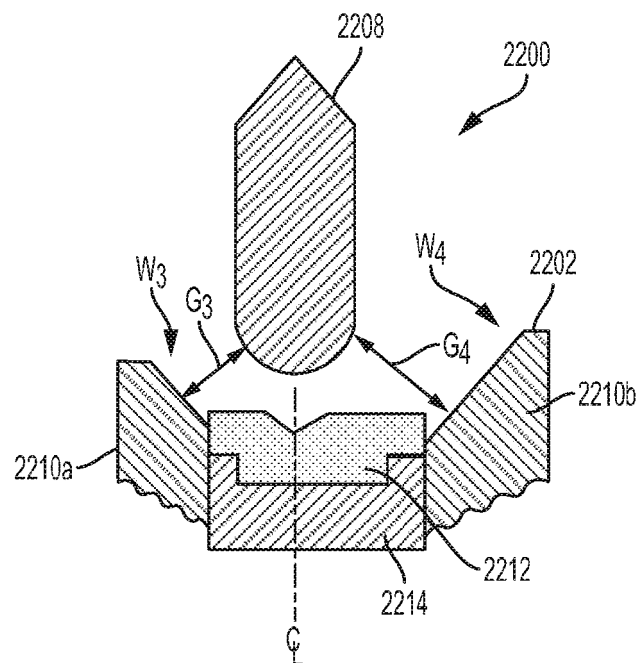


FIG. 37

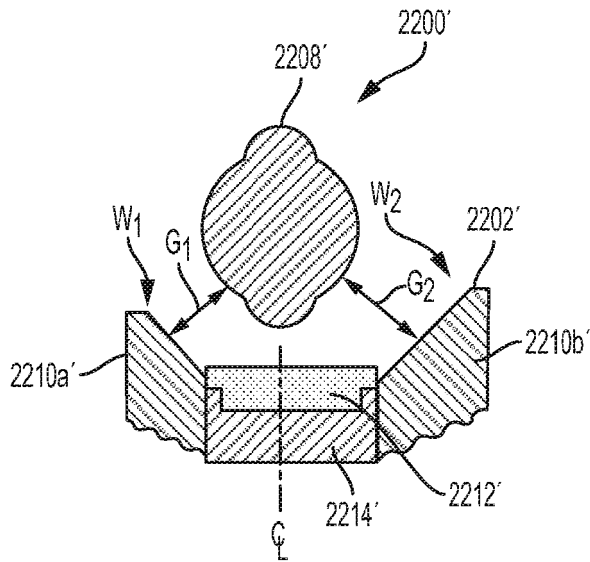


FIG. 38

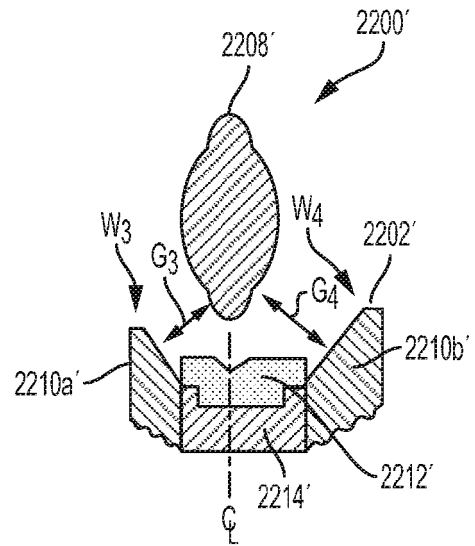


FIG. 39

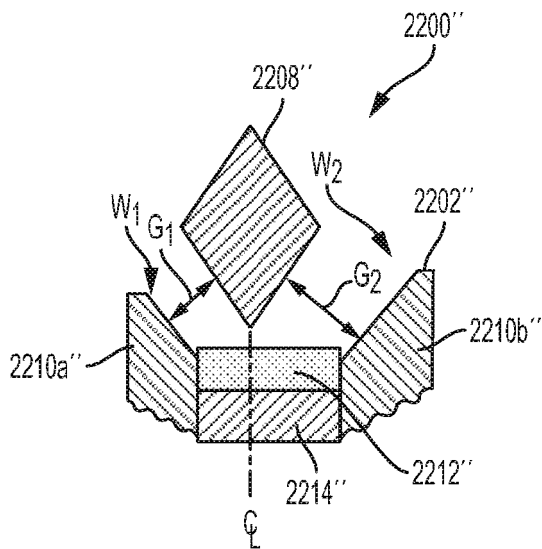


FIG. 40

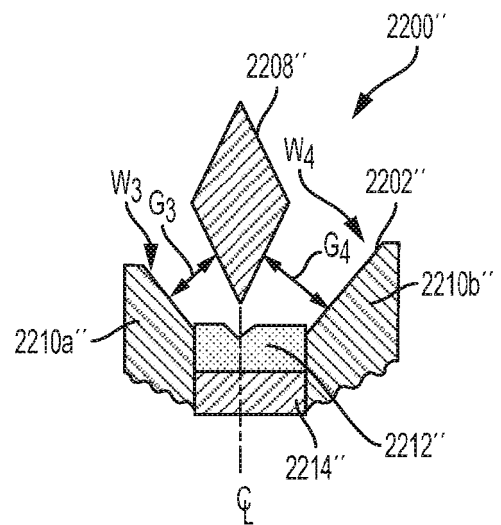


FIG. 41

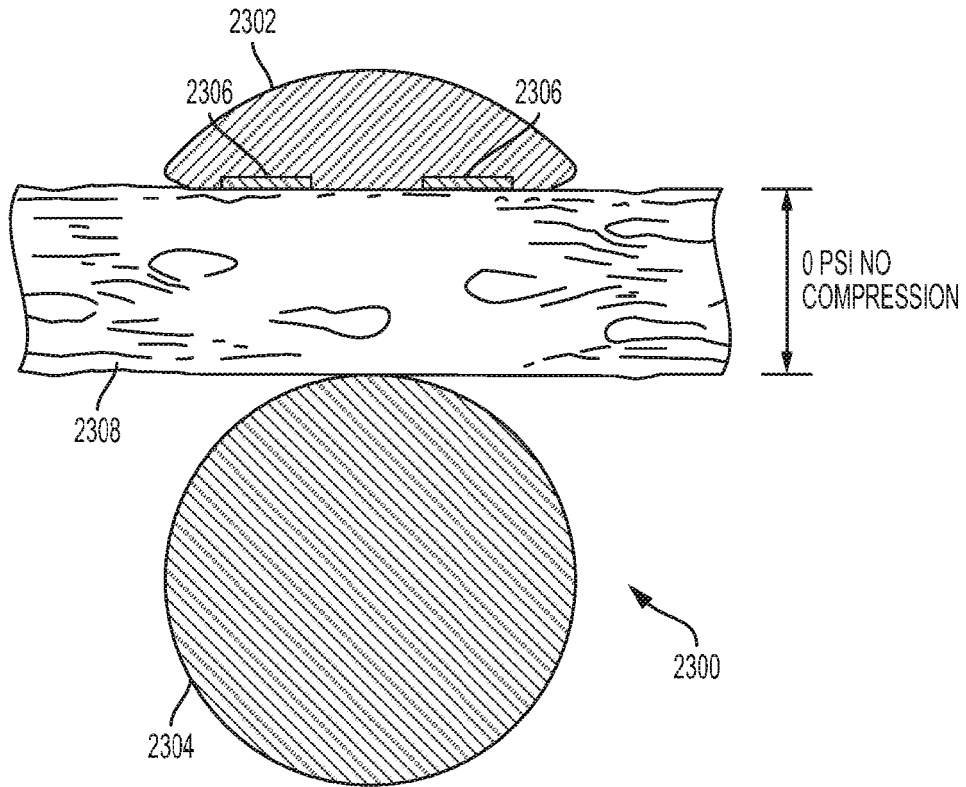


FIG. 42A

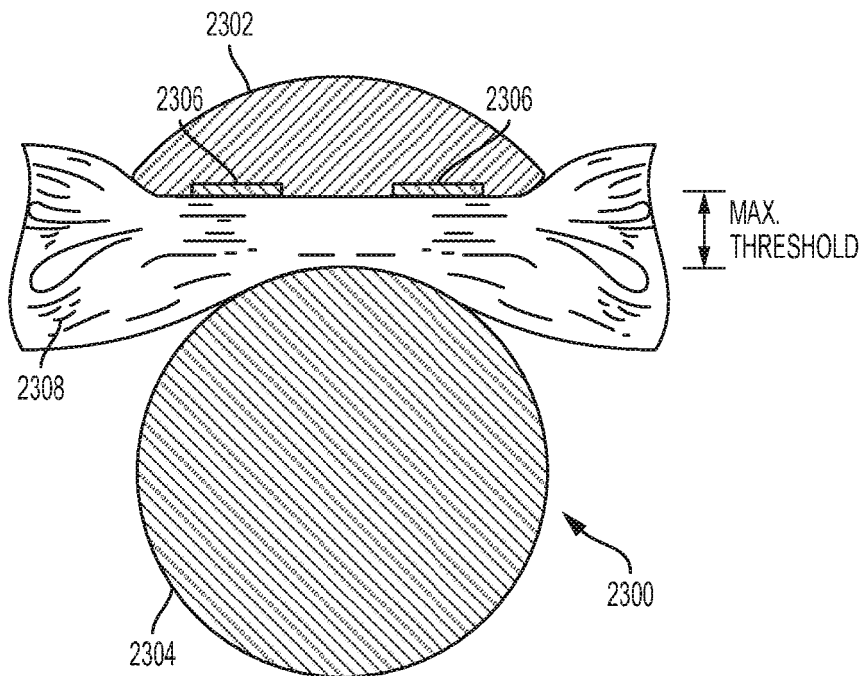
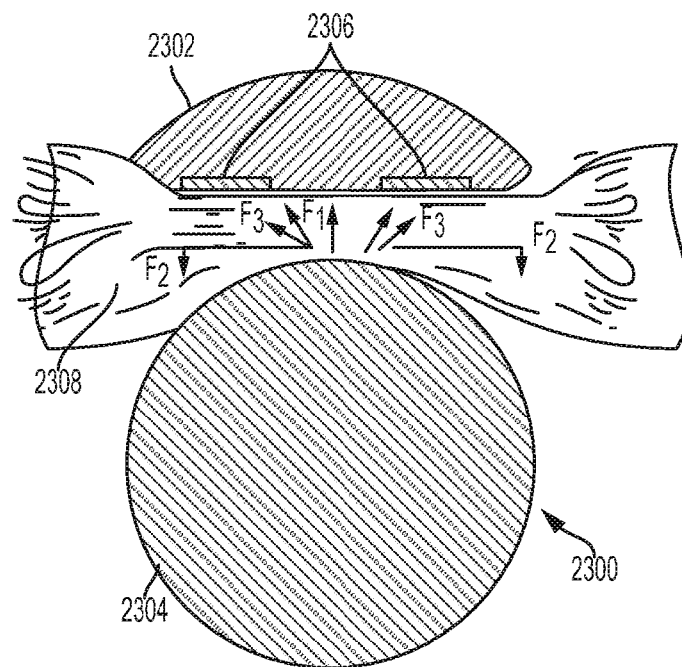
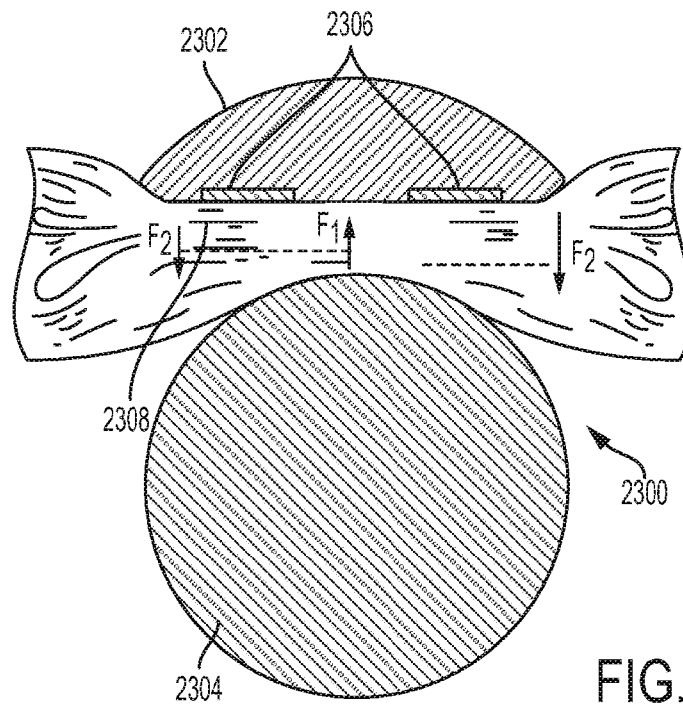


FIG. 42B



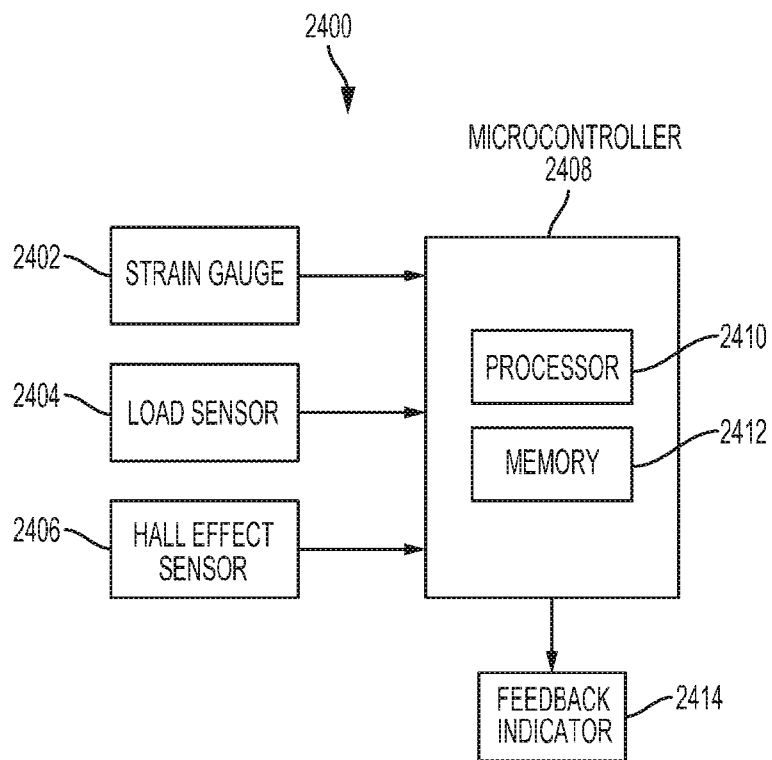


FIG. 44

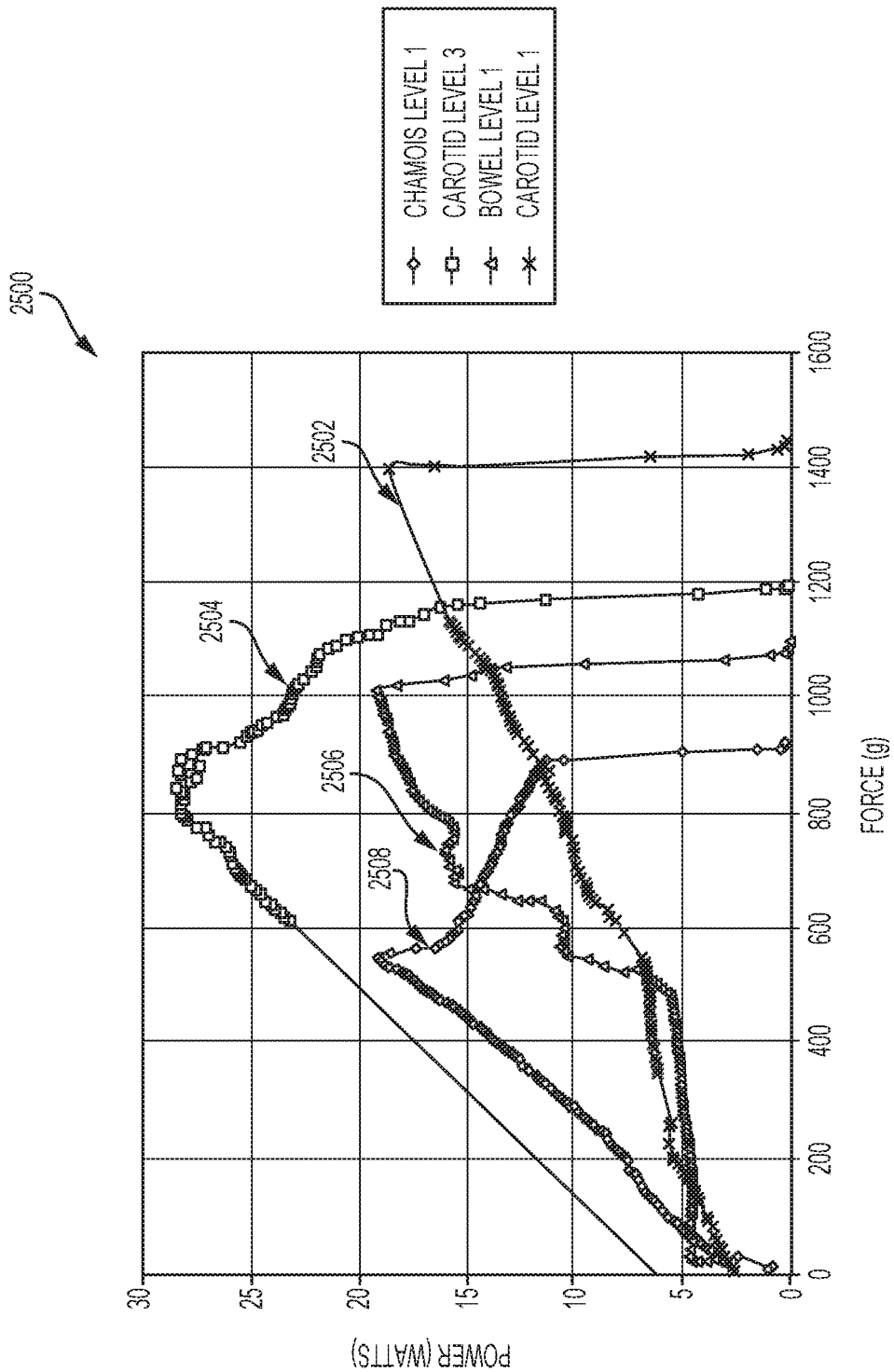


FIG. 45

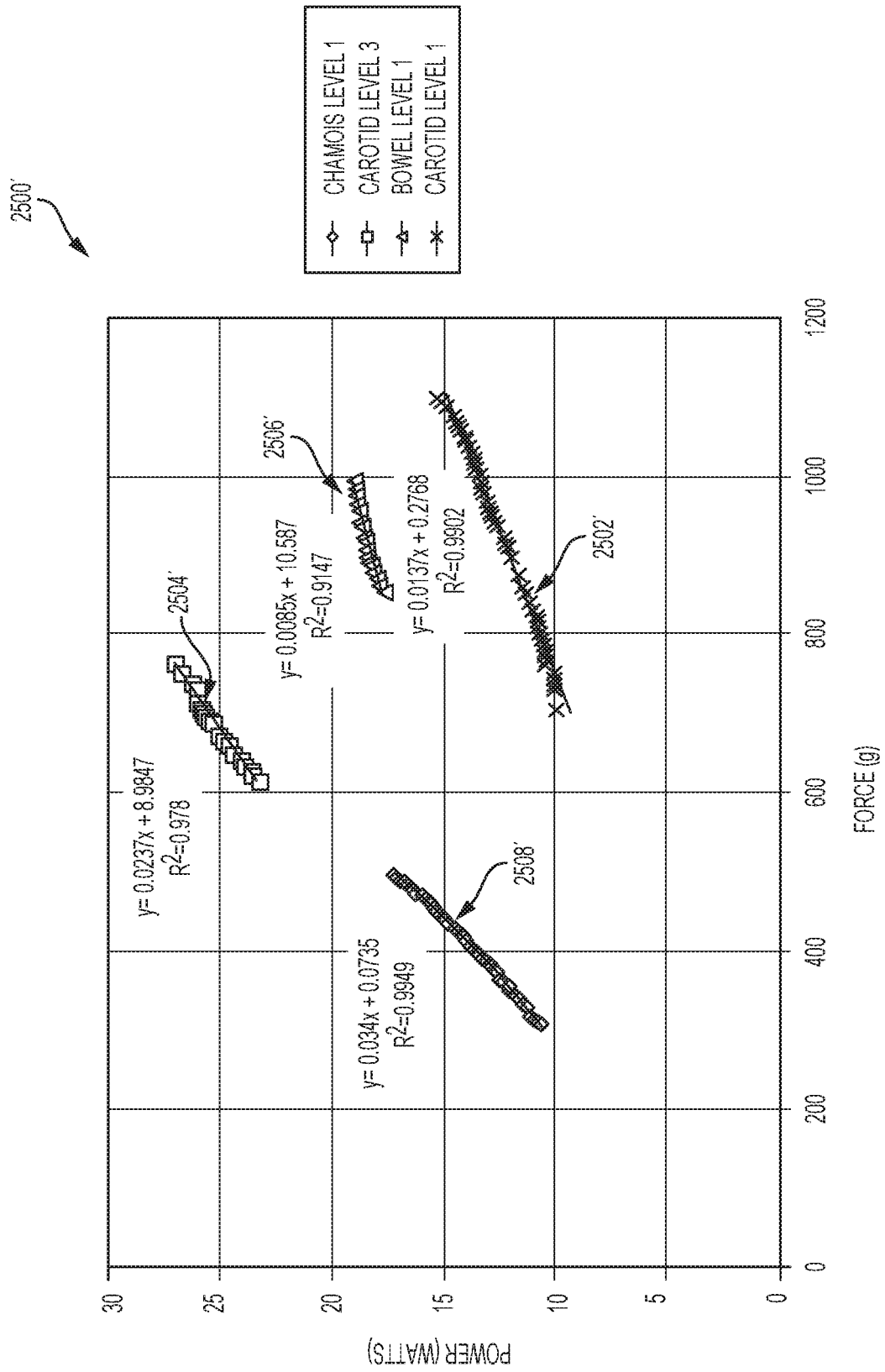


FIG. 46

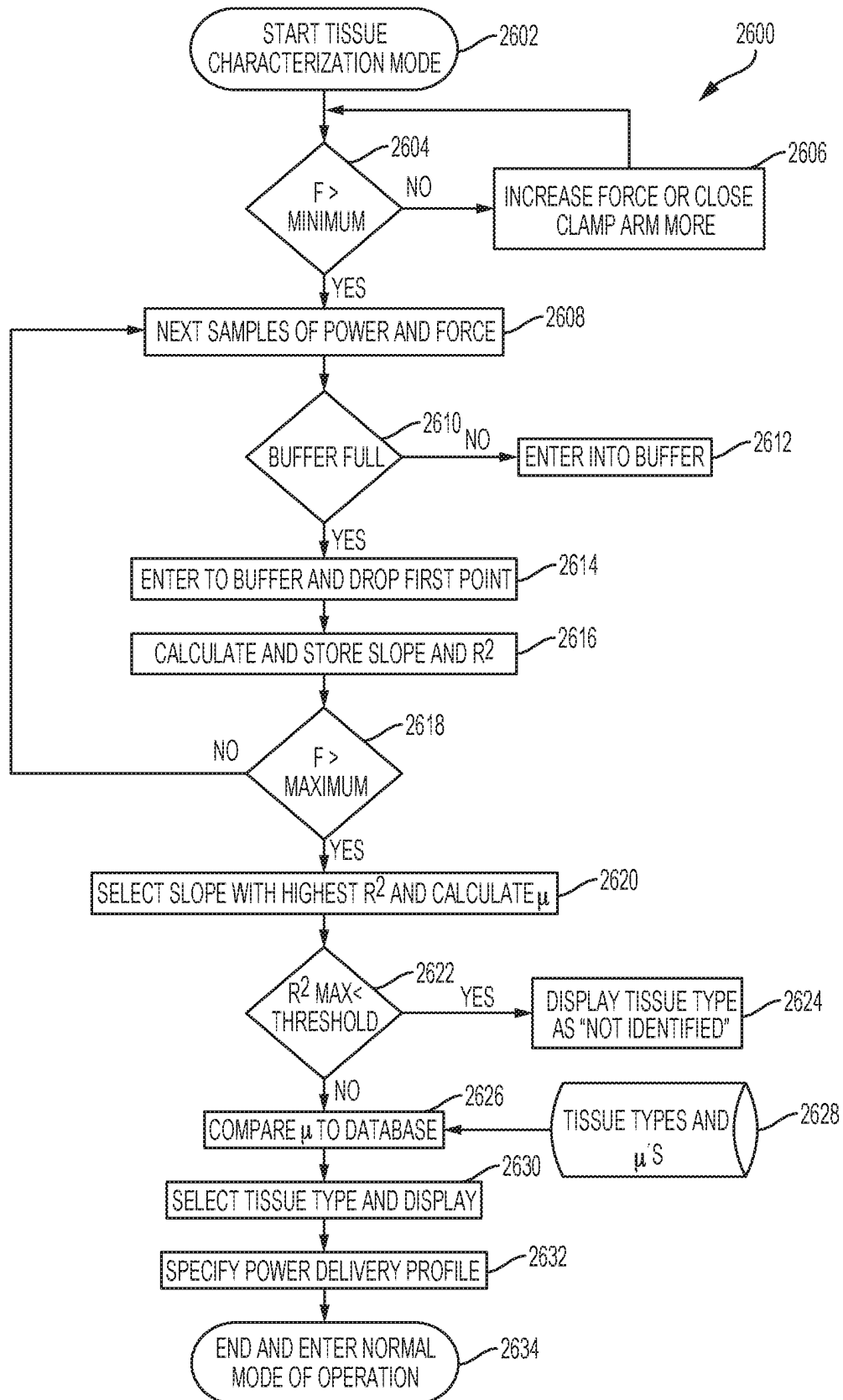


FIG. 47



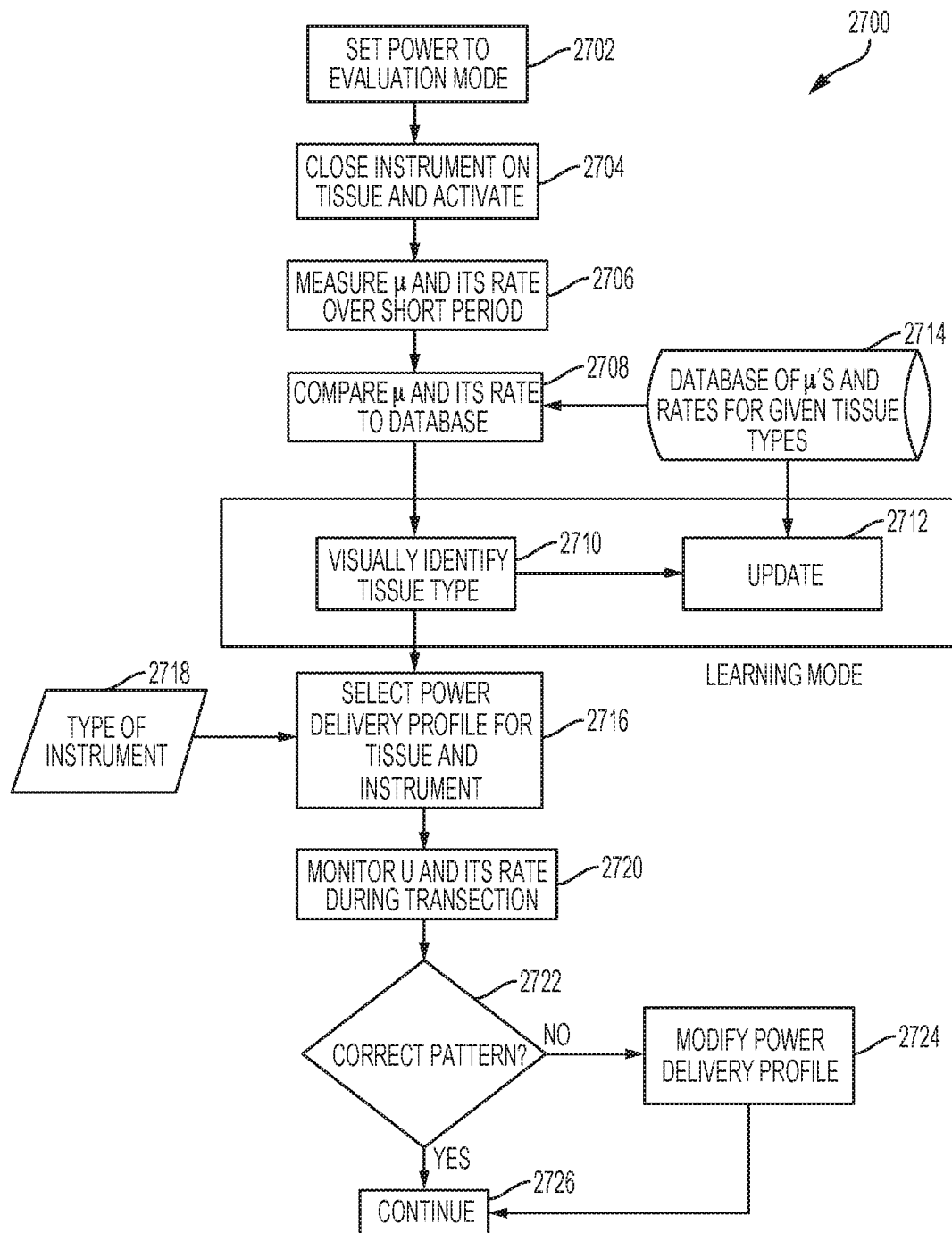


FIG. 48

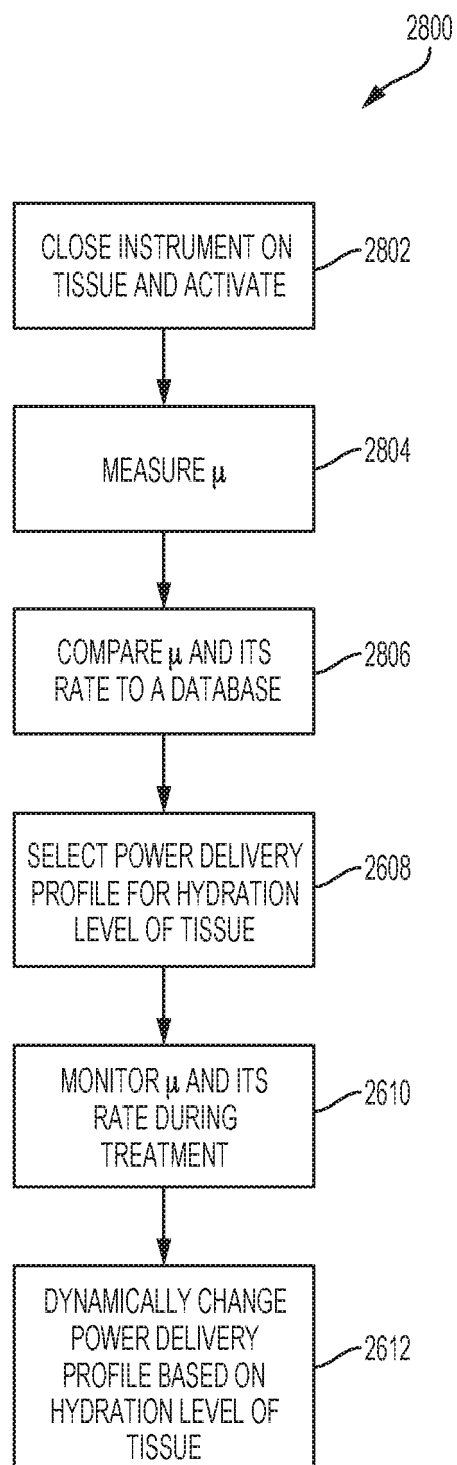


FIG. 49

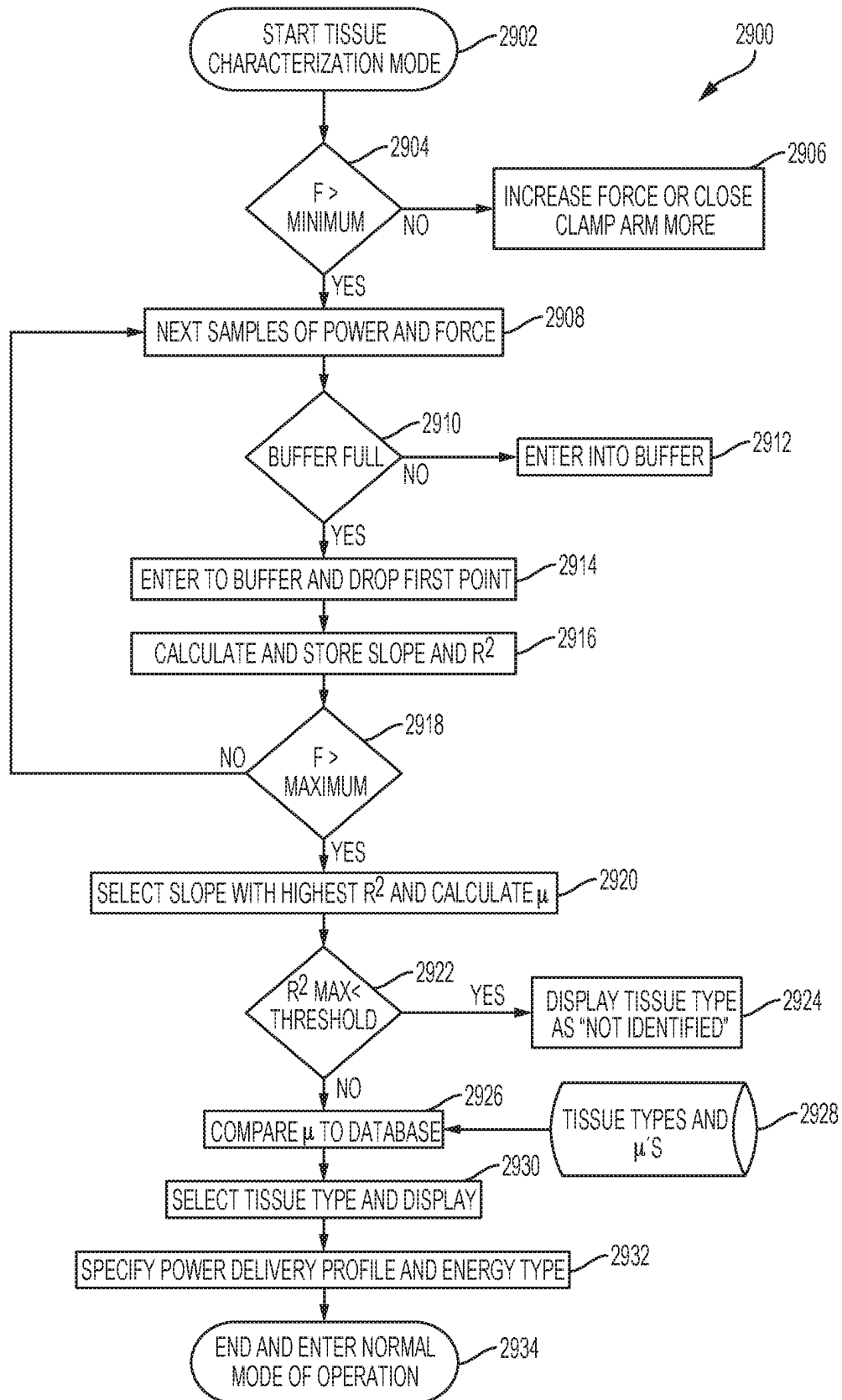


FIG. 50

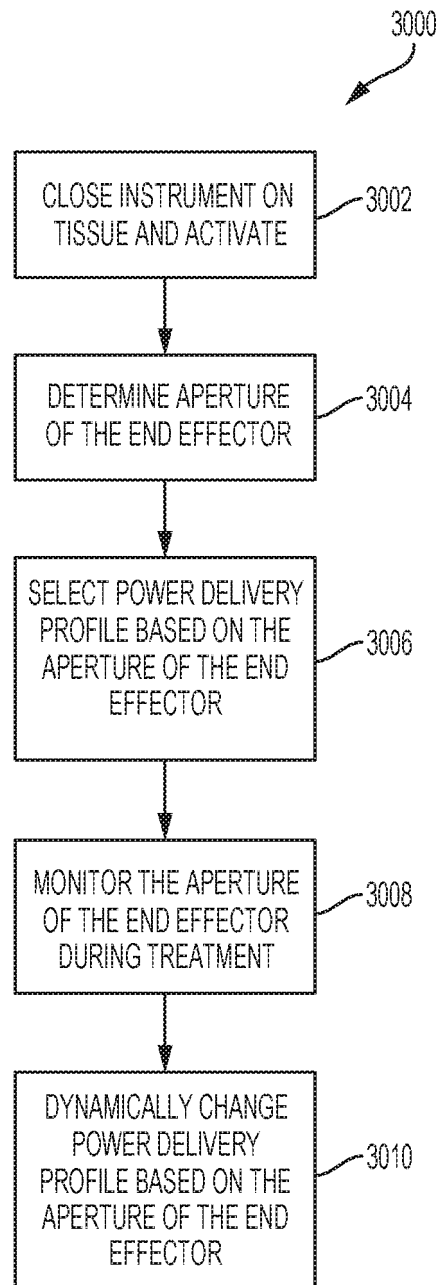


FIG. 51

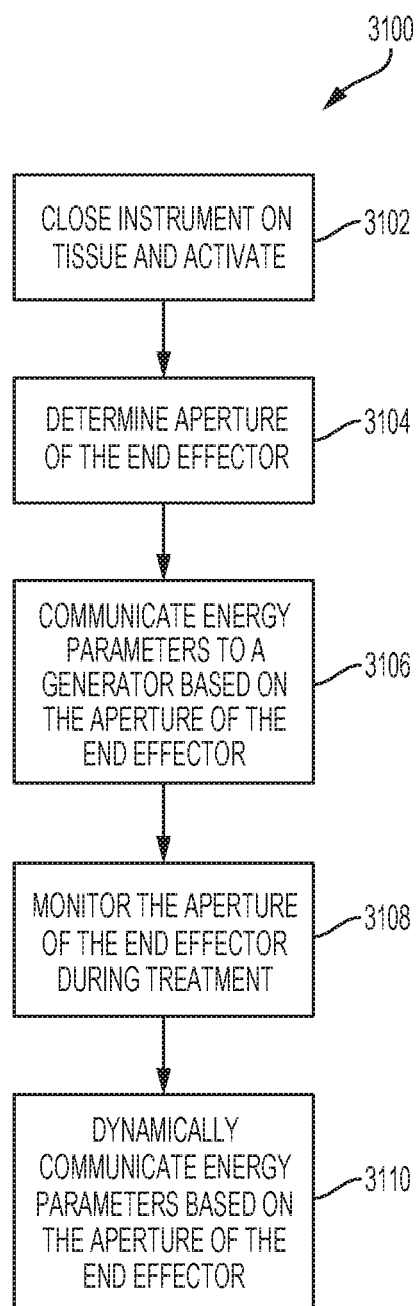


FIG. 52

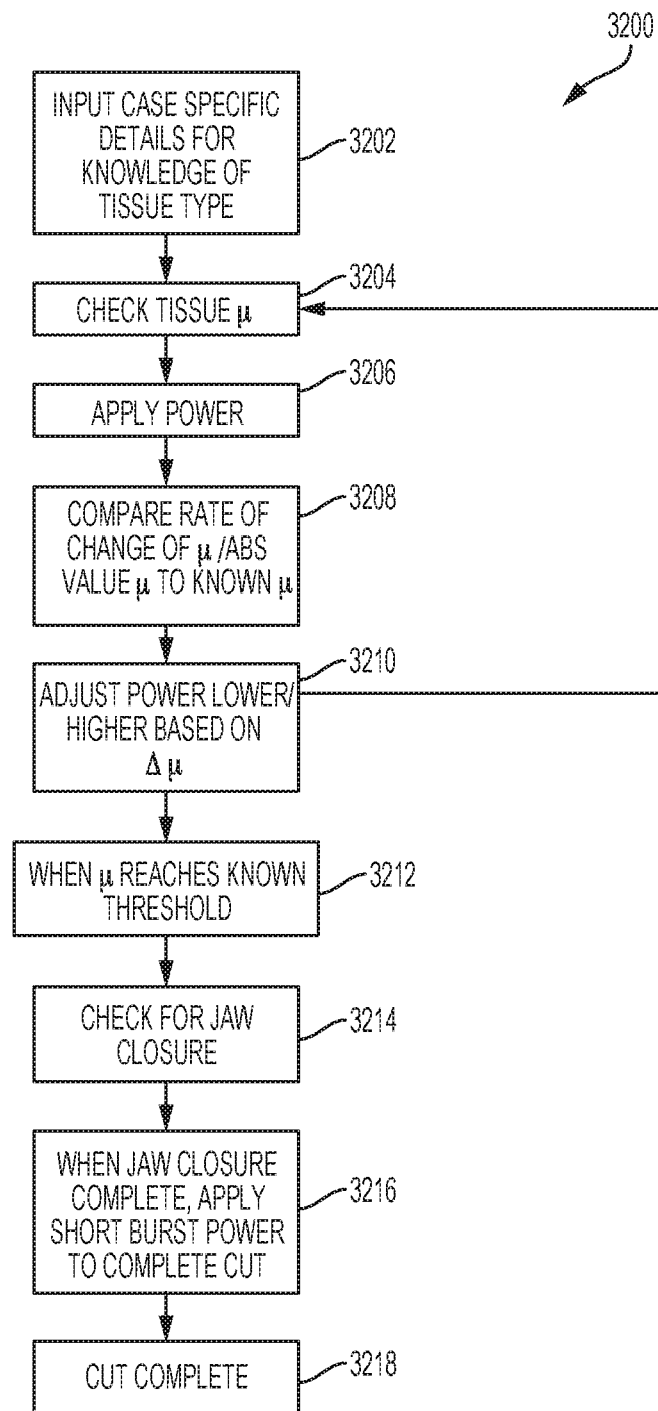


FIG. 53

**REFERENCES CITED IN THE DESCRIPTION**

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专利名称(译)	具有基于组织类型的用户自适应技术的手术系统		
公开(公告)号	<a href="#">EP3316812B1</a>	公开(公告)日	2019-08-07
申请号	EP2016739613	申请日	2016-06-24
[标]申请(专利权)人(译)	ETHICON , LLC		
申请(专利权)人(译)	ETHICON LLC		
当前申请(专利权)人(译)	ETHICON LLC		
[标]发明人	STULEN FOSTER B MADAN ASHVANI K HOUSER KEVIN L		
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IPC分类号	A61B18/14 A61B17/32 A61B18/00 A61B17/00 A61B90/00		
CPC分类号	A61B17/320092 A61B18/1445 A61B2017/00017 A61B2017/00039 A61B2017/00075 A61B2017/00084 A61B2017/320094 A61B2017/320095 A61B2017/320097 A61B2018/00642 A61B2018/00648 A61B2018/00702 A61B2018/00791 A61B2018/00875 A61B2018/0088 A61B2018/00994 A61B2090 /064 A61B2018/00589 A61B2018/00601 A61B2018/0063 A61B2560/0475 A61B2562/0261		
优先权	62/186984 2015-06-30 US 62/235368 2015-09-30 US 62/235466 2015-09-30 US 62/235260 2015-09-30 US 62/279635 2016-01-15 US 62/330669 2016-05-02 US 15/177439 2016-06-09 US		
其他公开文献	EP3316812A1		
外部链接	<a href="#">Espacenet</a>		

#### 摘要(译)

各种形式涉及用于凝固和解剖组织的系统和方法。一种外科器械，包括：末端执行器，被配置为在其远端处密封和切开组织；以及发生器回路，其被配置为将能量输送到末端执行器。力传感器与末端执行器连通并且被配置为测量由末端执行器施加到组织的力。基于与末端执行器相互作用的组织类型的确定，传递到末端执行器的能量是动态的。基于组织系数确定组织类型，该组织系数是基于由末端执行器施加到组织的测量力，末端执行器的超声运动和末端执行器产生的热量计算的。

