



US 20130218013A1

(19) **United States**(12) **Patent Application Publication**
Barthe et al.(10) **Pub. No.: US 2013/0218013 A1**(43) **Pub. Date: Aug. 22, 2013**(54) **ULTRASOUND MEDICAL SYSTEM AND METHOD****Publication Classification**

(71) Applicants: **Peter Barthe**, Phoenix, AZ (US); **Michael Slayton**, Mesa, AZ (US); **Paul Jaeger**, Mesa, AZ (US); **Douglas Mast**, Cincinnati, OH (US); **Inder Makin**, Mesa, AZ (US); **Brian OConner**, Green Cove Springs, FL (US); **Jeffery Messerly**, Cincinnati, OH (US); **Waseem Faidi**, Clifton Park, NJ (US); **Megan Runk**, Cincinnati, OH (US); **Christopher Park**, Oregania, OH (US)

(72) Inventors: **Peter Barthe**, Phoenix, AZ (US); **Michael Slayton**, Mesa, AZ (US); **Paul Jaeger**, Mesa, AZ (US); **Douglas Mast**, Cincinnati, OH (US); **Inder Makin**, Mesa, AZ (US); **Brian OConner**, Green Cove Springs, FL (US); **Jeffery Messerly**, Cincinnati, OH (US); **Waseem Faidi**, Clifton Park, NJ (US); **Megan Runk**, Cincinnati, OH (US); **Christopher Park**, Oregania, OH (US)

(21) Appl. No.: **13/752,151**(22) Filed: **Jan. 28, 2013****Related U.S. Application Data**

(60) Continuation of application No. 12/145,635, filed on Jun. 25, 2008, now abandoned, which is a division of application No. 10/850,984, filed on May 21, 2004, now Pat. No. 7,473,250.

(51) **Int. Cl.**

A61B 5/00 (2006.01)
A61B 18/12 (2006.01)
A61B 18/06 (2006.01)
A61B 18/18 (2006.01)
A61B 5/053 (2006.01)
A61B 8/13 (2006.01)
A61N 7/00 (2006.01)
A61B 18/08 (2006.01)

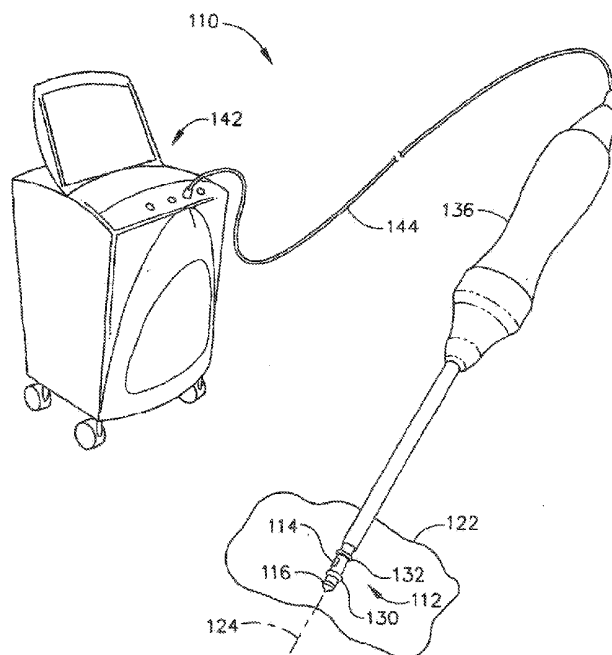
(52) **U.S. Cl.**

CPC *A61B 5/4836* (2013.01); *A61N 7/00* (2013.01); *A61B 18/12* (2013.01); *A61B 18/08* (2013.01); *A61B 18/1815* (2013.01); *A61B 5/053* (2013.01); *A61B 8/13* (2013.01); *A61B 18/06* (2013.01)
 USPC **600/439**; 601/2; 600/547

(57)

ABSTRACT

An ultrasound medical system includes an interstitial end effector. The interstitial end effector is interstitially insertable into patient tissue, includes at least one medical-treatment ultrasound transducer, and includes at least one end-effector-tissue-track ablation device. One method for ultrasonically treating a lesion in a patient includes the steps of obtaining the interstitial end effector and inserting it into the patient creating a tissue track which is surrounded by patient tissue and which ends at the distal end of the inserted interstitial end effector. Other steps include ultrasonically ablating the lesion using the at-least-one medical-treatment ultrasound transducer, using the at-least-one end-effector-tissue-track ablation device to ablate the patient tissue surrounding the tissue track along substantially the entire tissue track, and withdrawing the end effector from the patient.



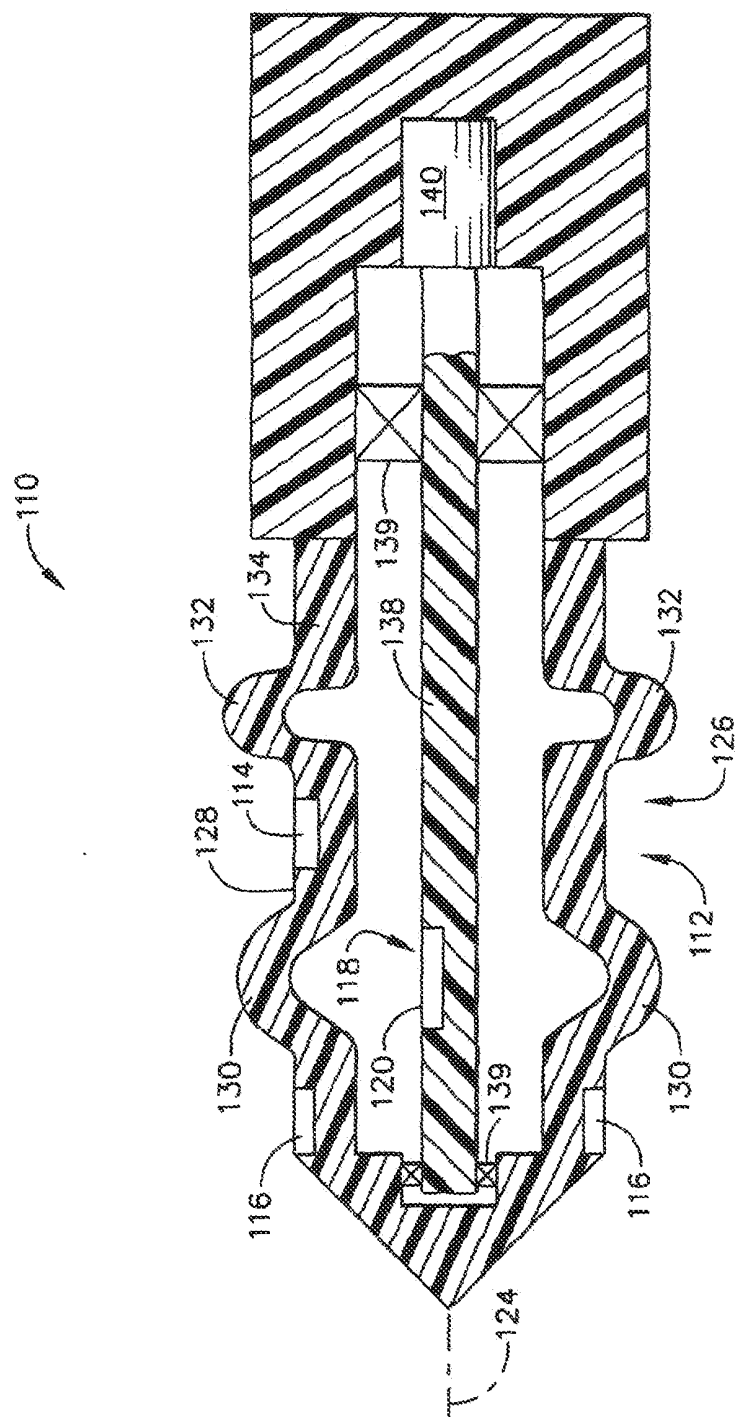


Fig. 2

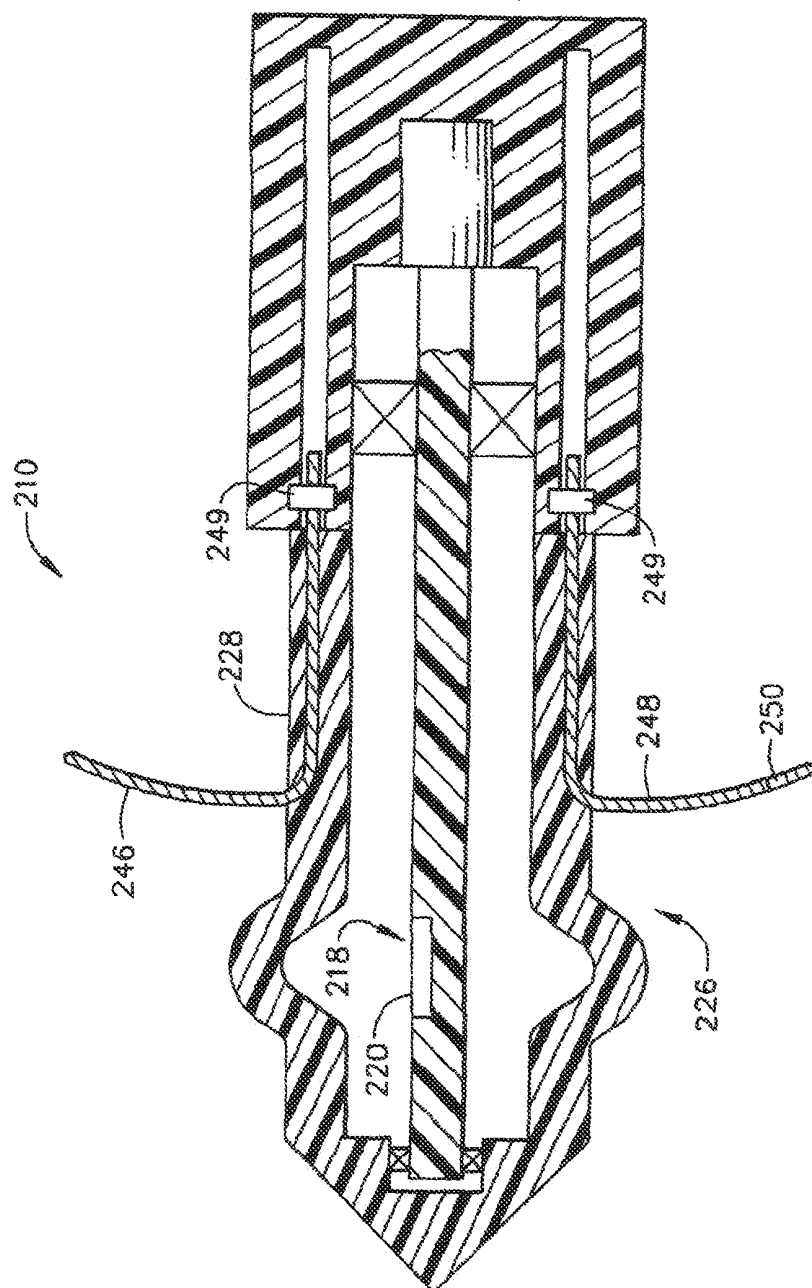


FIG. 3

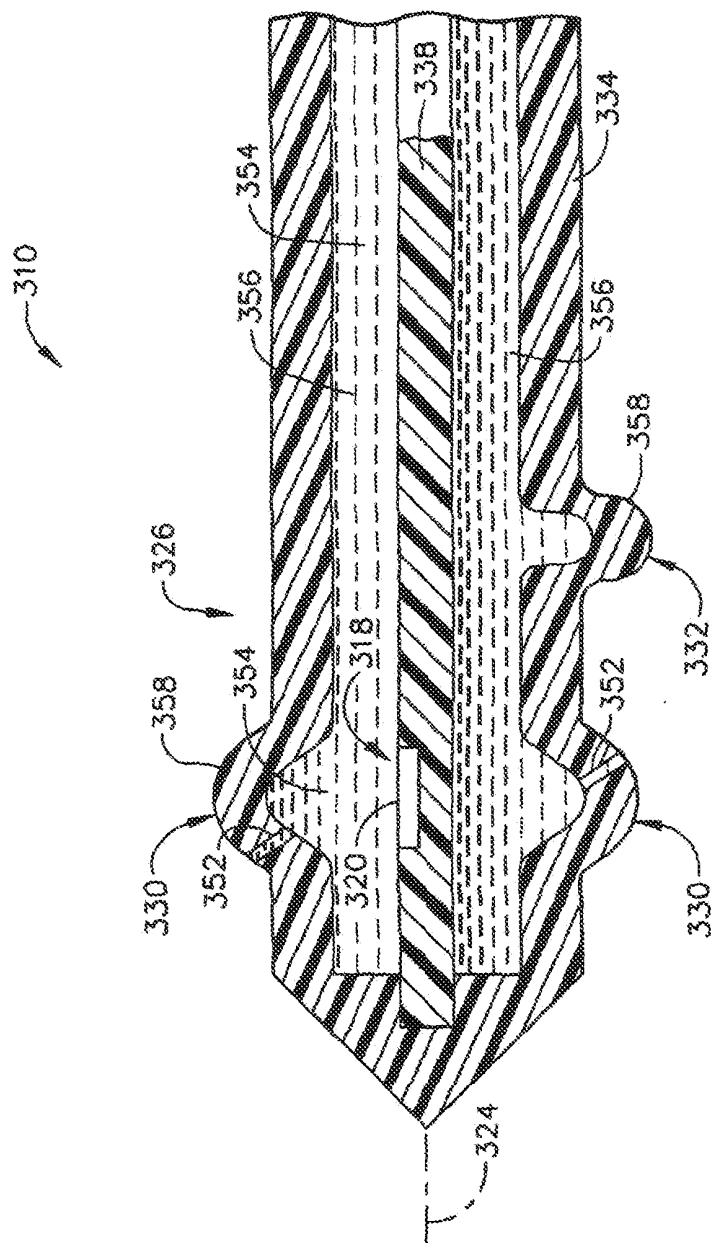


FIG. 4

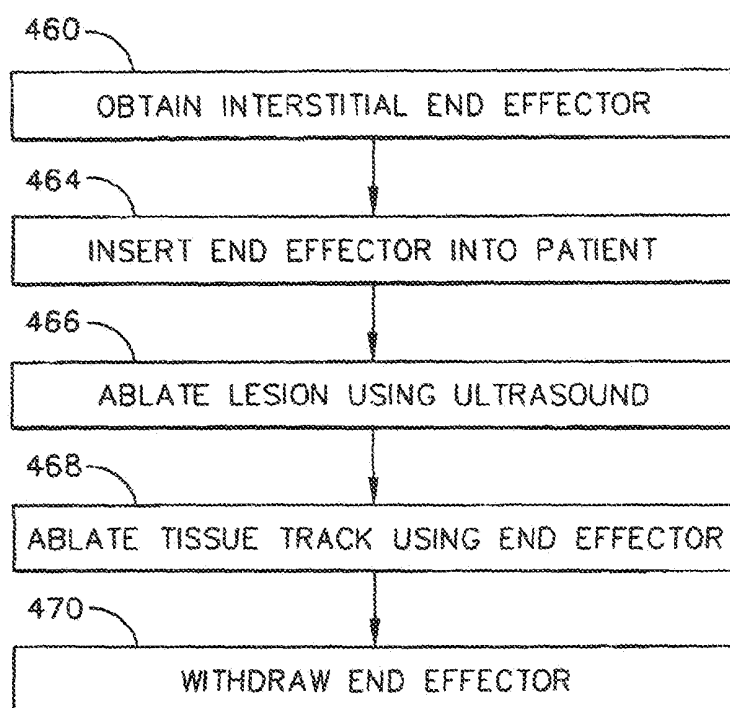
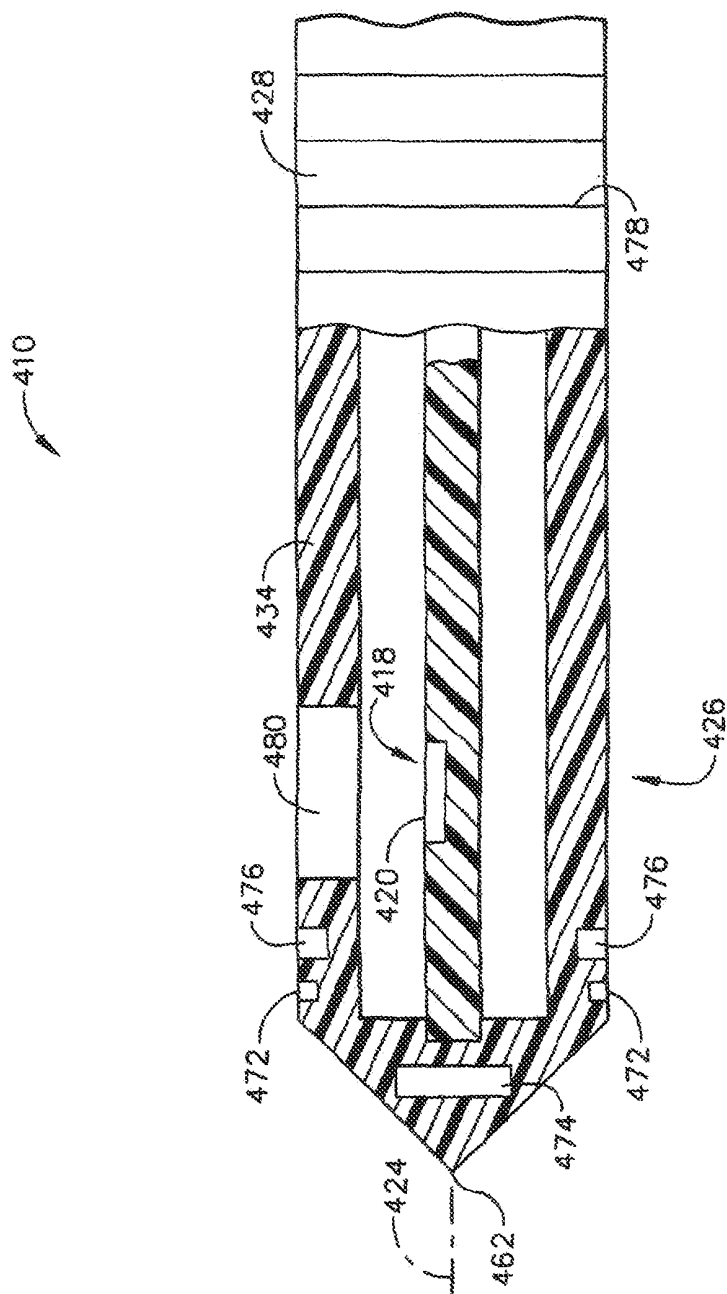


FIG. 5



ULTRASOUND MEDICAL SYSTEM AND METHOD

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application is a continuation application of U.S. patent application Ser. No. 12/145,635, filed on Jun. 25, 2008. The present application claims all the benefit of and priority to U.S. patent application Ser. No. 12/145,635, filed on Jun. 25, 2008, which is a divisional application of U.S. patent application Ser. No. 10/850,984, filed on May 21, 2004, which issued as U.S. Pat. No. 7,473,250, all of which are incorporate by reference herein.

FIELD OF THE INVENTION

[0002] The present invention relates generally to ultrasound, and more particularly to ultrasound medical systems and methods.

BACKGROUND OF THE INVENTION

[0003] Known medical methods include using ultrasound imaging (at low power) of patients to identify patient tissue for medical treatment and include using ultrasound (at high power) to ablate identified patient tissue by heating the tissue.

[0004] Known ultrasound medical systems and methods include deploying an ultrasound end effector having an ultrasound transducer outside the body to break up kidney stones inside the body, endoscopically inserting an ultrasound end effector having an ultrasound transducer in the rectum to medically destroy prostate cancer, laparoscopically inserting an ultrasound end effector having an ultrasound transducer in the abdominal cavity to medically destroy a cancerous liver tumor, intravenously inserting a catheter ultrasound end effector having an ultrasound transducer into a vein in the arm and moving the catheter to the heart to medically destroy diseased heart tissue, and interstitially inserting a needle ultrasound end effector having an ultrasound transducer needle into the tongue to medically destroy tissue to reduce tongue volume to reduce snoring.

[0005] Rotatable ultrasound end effectors are known wherein an ultrasound transducer is non-rotatably attached to a shaft whose distal end is circumferentially and longitudinally surrounded by a sheath having a longitudinal axis and having an acoustic window. Water between the shaft and the sheath provides acoustic coupling between the ultrasound transducer and the acoustic window. The shaft is rotatable about the longitudinal axis with respect to the sheath. The sheath is non-rotatably attached to a handpiece.

[0006] Known medical systems and methods include deploying a radio-frequency (RF) end effector having an RF electrode to thermally ablate patient tissue and to take tissue electric impedance and tissue temperature measurements using electrodes integrated into the shaft or into a tine which also helps stabilize the RF end effector in patient tissue.

[0007] Still, scientists and engineers continue to seek improved ultrasound medical systems and methods.

SUMMARY OF THE INVENTION

[0008] A method of the invention is for ultrasonically treating a lesion in a patient and includes steps a) through e). Step a) includes obtaining an interstitial end effector having a distal end, including a medical ultrasound transducer assembly having at least one medical-treatment ultrasound transducer,

and including at least one end-effector-tissue-track ablation device. Step b) includes inserting the interstitial end effector into the patient creating a tissue track which is surrounded by patient tissue and which ends at the distal end of the inserted interstitial end effector. Step c) includes ultrasonically ablating the lesion using the at-least-one medical-treatment ultrasound transducer. Step d) includes using the at-least-one end-effector-tissue-track ablation device to ablate the patient tissue surrounding the tissue track along substantially the entire tissue track. Step e) includes withdrawing the end effector from the patient.

[0009] An embodiment of the invention is an ultrasound medical system including an interstitial end effector. The interstitial end effector is interstitially insertable into patient tissue, includes at least one medical-treatment ultrasound transducer, and includes at least one end-effector-tissue-track ablation device.

[0010] Several benefits and advantages are obtained from one or more of the embodiments and methods of the invention. In one example, having an interstitial end effector with a medical-treatment ultrasound transducer and an end-effector-tissue-track ablation device allows ultrasonic ablation of a lesion using the medical-treatment ultrasound transducer and allows ablation of patient tissue surrounding the tissue track as the interstitial end effector is withdrawn from the patient to help reduce the possibility of excessive bleeding and/or tissue contamination.

[0011] The present invention has, without limitation, application in conventional interstitial, endoscopic, laparoscopic, and open surgical instrumentation as well as application in robotic-assisted surgery.

BRIEF DESCRIPTION OF THE FIGURES

[0012] FIG. 1 is a perspective view of a first embodiment of the present invention showing an ultrasound medical system which includes an end effector, a handpiece, and a controller;

[0013] FIG. 2 is a schematic cross-sectional view of a first embodiment of the end effector and the handpiece of the ultrasound medical system of FIG. 1 showing a medical ultrasound transducer assembly and two non-ultrasound tissue-property-measuring sensors;

[0014] FIG. 3 is a view, as in FIG. 2, but of a second embodiment of a handpiece and of an end effector having a medical ultrasound transducer assembly and two tines;

[0015] FIG. 4 is a view, as in FIG. 2, but of a third embodiment of an end effector having a medical ultrasound transducer assembly supported by a shaft and having a surrounding sheath, wherein the sheath includes two balloon portions;

[0016] FIG. 5 is a block diagram view of a method of the invention for ultrasonically treating a lesion in a patient; and

[0017] FIG. 6 is a schematic view, partially in cross-section, of a fourth embodiment of an end effector which has a medical-treatment ultrasound transducer and three end-effector-tissue-track ablation devices and which can be used in one employment of the method of FIG. 5.

DETAILED DESCRIPTION OF THE INVENTION

[0018] Before explaining the present invention in detail, it should be noted that the invention is not limited in its application or use to the details of construction and arrangement of parts and/or steps illustrated in the accompanying drawings and description. The illustrative embodiments and methods of the invention may be implemented or incorporated in other

embodiments, methods, variations and modifications, and may be practiced or carried out in various ways. Furthermore, unless otherwise indicated, the terms and expressions employed herein have been chosen for the purpose of describing the illustrative embodiments and methods of the present invention for the convenience of the reader and are not for the purpose of limiting the invention.

[0019] It is understood that any one or more of the following-described embodiments, methods, examples, etc. can be combined with any one or more of the other following-described embodiments, methods, examples, etc.

[0020] Referring now to FIGS. 1-2 of the drawings, a first embodiment of the present invention is an ultrasound medical system 110 comprising an ultrasound end effector 112 and at least one non-ultrasound tissue-property-measuring sensor 114 and 116. The ultrasound end effector 112 includes a medical ultrasound transducer assembly 118 having at least one medical-treatment ultrasound transducer 120. The at-least-one non-ultrasound tissue-property-measuring sensor 114 and 116 is supported by the ultrasound end effector 112 and is disposable in contact with patient tissue 122.

[0021] It is noted that a medical-treatment ultrasound transducer includes a medical-treatment-only ultrasound transducer and a medical-imaging-and-treatment ultrasound transducer. In one arrangement, an ultrasound transducer has a single transducer element, and in another arrangement, an ultrasound transducer has a plurality (also called an array) of transducer elements. It is also noted that a medical ultrasound transducer assembly having at least one medical-treatment ultrasound transducer can also have at least one medical-imaging ultrasound transducer.

[0022] In one example of the embodiment of FIGS. 1-2, the ultrasound end effector 112 includes a longitudinal axis 124. In this example, the at-least-one non-ultrasound tissue-property-measuring sensor 114 and 116 includes a first non-ultrasound tissue-property-measuring sensor 114 and a second non-ultrasound tissue-property-measuring sensor 116. The at-least-one medical-treatment ultrasound transducer 120 is disposed longitudinally between the first and second non-ultrasound tissue-property-measuring sensors 114 and 116.

[0023] In one variation of the embodiment of FIGS. 1-2, the at-least-one non-ultrasound tissue-property-measuring sensor (e.g., 114) measures tissue temperature. In one modification, the at-least-one non-ultrasound tissue-property-measuring sensor (e.g., 114) is chosen from the group consisting of a thermistor, a thermocouple, and combinations thereof. In another variation, the at-least-one non-ultrasound tissue-property-measuring sensor (e.g., 116) measures tissue electric impedance. In one modification, the at-least-one non-ultrasound tissue-property-measuring sensor (e.g., 116) is chosen from the group consisting of a monopolar electrode, a bipolar electrode, and combinations thereof. It is noted that tissue temperature and/or tissue electric impedance is a measure of the degree of ultrasonic ablation of patient tissue, as can be appreciated by those skilled in the art.

[0024] In one construction of the embodiment of FIGS. 1-2, the ultrasound end effector 112 is an ultrasound interstitial end effector 126 which is interstitially insertable into patient tissue 122 and which has an exterior surface 128. The at-least-one non-ultrasound tissue-property-measuring sensor 114 and 116 is attached to the ultrasound interstitial end effector 126 and is fixedly disposed substantially flush with the exterior surface 128. In one arrangement, the exterior surface 128 includes at least one balloon portion 130 and 132 which is

expandable and contractible and which is expandable against patient tissue 122 to provide at least some stabilization of the ultrasound interstitial end effector 126 with respect to patient tissue 122. In one variation, the exterior surface 128 is the exterior surface of a sheath 134 and has first and second balloon portions 130 and 132, wherein the first balloon portion 130 surrounds the medical ultrasound transducer assembly 118 and acts as an acoustic window, and wherein the second balloon portion 132 is longitudinally spaced apart from the medical ultrasound transducer assembly 118. An acoustic coupling medium, such as water, is disposable between the medical ultrasound transducer assembly 118 and the first balloon portion 130 and has been omitted from FIG. 2 for clarity. In one modification, the first balloon portion 130 is omitted and the sheath 134 terminates before the medical ultrasound transducer assembly 118 which is exposed to patient tissue. In another modification, the second balloon portion 132 is omitted. In one employment, the at-least-one balloon portion 130 and 132 is contracted during tissue insertion and withdrawal of the ultrasound interstitial end effector 126. Other constructions, arrangements, variations, and modifications are left to the artisan.

[0025] In one enablement of the embodiment of FIGS. 1-2, the ultrasound end effector 112 is an ultrasound interstitial end effector 126 which is interstitially insertable into patient tissue 122 and which has an exterior surface 128. In this enablement, the at-least-one non-ultrasound tissue-property-measuring sensor 114 and 116 is deployable to extend away from the exterior surface into patient tissue 128 to provide at least some stabilization of the ultrasound interstitial end effector 126 with respect to patient tissue 122 and is retrievable to retract back toward the exterior surface 128. In one arrangement, the at-least-one non-ultrasound tissue-property-measuring sensor 114 and 116 is storable inside the exterior surface.

[0026] In one implementation of the embodiment of FIGS. 1-2, the ultrasound medical system 110 also includes a handpiece 136 operatively connected to the ultrasound end effector 112. The ultrasound end effector 112 has a longitudinal axis 124 and a shaft 138, and the medical ultrasound transducer assembly 118 is supported by the shaft 138. The shaft 138 is rotatable with respect to the handpiece 136 about the longitudinal axis 124 and is supported by bearings 139. In one variation, a motor 140 rotates the shaft 138. In one arrangement, the ultrasound medical system 110 includes a controller 142 operatively connected to the handpiece 136 via a cable 144.

[0027] A second embodiment of the present invention, shown in FIG. 3, is an ultrasound medical system 210 comprising an ultrasound end effector 226. The ultrasound end effector 226 has an exterior surface 228. The ultrasound end effector 226 includes a medical ultrasound transducer assembly 218 having at least one medical-treatment ultrasound transducer 220, and includes at least one tine 246 and 248. The at-least-one tine 246 and 248 is deployable to extend away from the exterior surface into patient tissue to provide at least some stabilization of the ultrasound end effector 226 with respect to patient tissue and is retrievable to retract back toward the exterior surface 228.

[0028] In one example of the embodiment of FIG. 3, the ultrasound end effector 226 is insertable into a patient. In one variation, the ultrasound end effector 226 is an ultrasound interstitial end effector which is interstitially insertable into patient tissue. In other variations, the ultrasound end effector

is insertable into a patient in an endoscopic, laparoscopic, and/or open surgical manner. In another example, the ultrasound end effector is disposable on the outside of a patient. Other examples and variations are left to the artisan.

[0029] In one enablement of the embodiment of FIG. 3, the at-least-one tine **246** and **248** includes a plurality of tines. In one example of the embodiment of FIG. 3, the at-least-one tine **246** and **248** is storable inside the exterior surface. It is noted that construction of deployable tines **246** and **248** in an ultrasound end effector **226** is within the level of skill of the artisan. In one arrangement, such deployment is accomplished using one or more of cables, levers, motors **249**, gearing, push rods and the like to move a tine partially out of, and back into, a lumen in the end effector. In one choice of materials, the tine comprises or consists essentially of Nitinol wire or nichrome wire.

[0030] In one employment of the embodiment of FIG. 3, the at-least-one tine (e.g., **246**) acts as an element chosen from the group consisting of an electrode, a thermistor, a thermocouple, an acoustic reflector, an acoustic absorber, an acoustic emitter, an acoustic receiver, a radio-frequency (RF) heater, a resistive heater, and combinations thereof. In another employment, the at-least-one tine (e.g., **248**) includes a component **250** chosen from the group consisting of an electrode, a thermistor, a thermocouple, an acoustic reflector, an acoustic absorber, an acoustic emitter, an acoustic receiver, a radio-frequency (RE) heater, a resistive heater, and combinations thereof.

[0031] The embodiment, examples, constructions, implementations, etc. of the embodiment of FIGS. 1-2 are equally applicable to the embodiment, constructions, implementations, etc. of FIG. 3. In one construction of the embodiment of FIG. 3, the exterior surface **228** is like the exterior surface **128** of a previously-illustrated and described construction of the embodiment of FIGS. 1-2 including at least one balloon portion which is expandable and contractible, and which is expandable against patient tissue to provide at least some stabilization of the ultrasound end effector with respect to patient tissue. In one implementation of the embodiment of FIG. 3, the ultrasound medical system **210** also includes, like a previously-illustrated and described implementation of the embodiment of FIGS. 1-2, a handpiece operatively connected to the ultrasound end effector, wherein the ultrasound end effector has a longitudinal axis and a shaft, wherein the medical ultrasound transducer assembly is supported by the shaft, and wherein the shaft is rotatable with respect to the handpiece about the longitudinal axis.

[0032] One method, using the embodiment of FIG. 3 and enablements, examples, employments, and constructions thereof, is for ultrasonically treating a lesion in a patient and includes steps a) through f), Step a) includes obtaining the ultrasound medical system **210**. Step b) includes inserting the ultrasound end effector **226** into patient tissue. Step c) includes deploying the plurality of tines **246** and **248** to extend sway from the exterior surface **228** into the patient tissue. Step d) includes ultrasonically ablating the lesion using the at-least-one medical-treatment ultrasound transducer **220**. Step e) includes retrieving the plurality of tines **246** and **248** to retract back toward the exterior surface and storing the plurality of tines **246** and **248** inside the exterior surface **228**. Step f) includes withdrawing the ultrasound end effector **226** from the patient tissue. Another method also includes the step of employing the plurality of tines **246** and **248** to each act as the element or using each component **250**.

An additional method also includes the step of expanding the at-least-one balloon portion against patient tissue and contracting the at-least-one balloon portion.

[0033] A third embodiment of the present invention, shown in FIG. 4, is an ultrasound medical system **310** comprising an ultrasound end effector **326** including a shaft **338**, a sheath **334**, and a medical ultrasound transducer assembly **318**. The medical ultrasound transducer assembly **318** is supported by the shaft **338** and has at least one medical-treatment ultrasound transducer **320**. The sheath **334** surrounds the shaft **338**. The sheath **334** includes at least one balloon portion **330** and **332** which is expandable against patient tissue to provide at least some stabilization of the ultrasound end effector **326** with respect to patient tissue.

[0034] In one example of the embodiment of FIG. 4, the ultrasound end effector **326** is insertable into a patient. In one variation, the ultrasound end effector **326** is an ultrasound interstitial end effector which is interstitially insertable into patient tissue. In other variations, the ultrasound end effector is insertable into a patient in an endoscopic, laparoscopic, and/or open surgical manner. In another example, the ultrasound end effector is disposable on the outside of a patient. Other examples and variations are left to the artisan.

[0035] In one construction of the embodiment of FIG. 3, the ultrasound end effector **326** has a longitudinal axis **324**, and the at-least-one balloon portion (e.g., **330**) acts as an acoustic window and is disposed to longitudinally overlap the at-least-one medical-treatment ultrasound transducer **320**. In one variation of this construction, the at-least-one balloon portion (e.g., **330**) includes at least one through hole **352**. In one modification, the at-least-one balloon portion (e.g., **330**) includes a plurality of through holes **352** creating a "weeping" balloon portion, wherein some of the acoustic coupling medium inside the sheath **334** extends and/or flows outside the sheath acoustic window providing improved acoustic coupling between the at-least-one medical-treatment ultrasound transducer **320** and patient tissue.

[0036] In one arrangement of the embodiment of FIG. 3, the at-least-one balloon portion (e.g., **330**) includes at least one through hole **352** and the ultrasound end effector **326** is adapted to dispense a drug **354** through the at-least-one through hole **352** to patient tissue. In one variation, the drug **354** is carried in a liquid acoustic coupling medium **356**, such as water, disposed between the medical ultrasound transducer assembly **318** and the at-least-one balloon portion **330** whose pressure is controlled (such as by a pump in a handpiece operatively connected to the ultrasound end effector) to expand and contract the at-least-one balloon portion **330**. In one variation, the drug **354** is at least potentiated (i.e., has its medical effect increased and/or activated) by ultrasound emitted from the at-least-one medical-treatment ultrasound transducer **320**.

[0037] In the same or another arrangement of the embodiment of FIG. 3, the ultrasound end effector **326** has a longitudinal axis **324**, and the at-least-one balloon portion (e.g., **332**) is disposed longitudinally apart from the at-least-one medical-treatment ultrasound transducer **320**. In one variation of the embodiment of FIG. 3, the at-least-one balloon portion (e.g., **330**) is a fully-circumferential balloon portion. In a different variation, the at-least-one balloon portion (e.g., **332**) is a blister balloon portion. In one example of the embodiment of FIG. 3, the at-least-one balloon portion **330** and **332** includes an outside surface **358** having a roughness average at least equal to 0.005-inch. In one variation, the

outside surface includes ribs. Such surface roughness improves stabilization of the ultrasound end effector 326 with respect to patient tissue when the at-least-one balloon portion 330 and 332 is expanded against the patient tissue.

[0038] The embodiments, constructions, implementations, etc. of the embodiments of FIGS. 1-2 and 3 are equally applicable to the embodiment, constructions, implementations, etc. of the embodiment of FIG. 4. In one implementation of the embodiment of FIG. 3, the ultrasound medical system 310 also includes a controller, like the controller of the previously-illustrated and described arrangement of the embodiment of FIGS. 1-2, wherein the controller is operatively connected to the medical ultrasound transducer assembly, wherein the medical ultrasound transducer assembly is a medical-imaging-and-treatment ultrasound transducer assembly, and wherein the controller determines if the at-least-one balloon portion is acoustically coupled to, or acoustically decoupled from, patient tissue from ultrasonically imaging a balloon-tissue area using the medical-imaging-and-treatment ultrasound transducer assembly.

[0039] One method of the invention for ultrasonically treating a lesion in a patient is shown in block diagram form in FIG. 5, and an embodiment of an ultrasound medical system which can be used in performing the method is shown in FIG. 6. The method includes steps a) through e). Step a) is labeled as "Obtain Interstitial End Effector" in block 460 of FIG. 5. Step a) includes obtaining an interstitial end effector 426 including a distal end 462 and including a medical ultrasound transducer assembly 418 having at least one medical-treatment ultrasound transducer 420 and at least one end-effector-tissue-track ablation device 472, 474 and 476. It is noted that the distal end of an interstitial end effector is an end having a tissue-piercing tip. Step b) is labeled as "Insert End Effector Into Patient" in block 464 of FIG. 5. Step b) includes inserting the interstitial end effector 426 into the patient creating a tissue track which is surrounded by patient tissue and which ends at the distal end 462 of the inserted interstitial end effector 426. Step c) is labeled as "Ablate Lesion Using Ultrasound" in block 466 of FIG. 5. Step c) includes ultrasonically ablating the lesion using the at-least-one medical-treatment ultrasound transducer 420. Step d) is labeled as "Ablate Tissue Track Using End Effector" in block 468 of FIG. 5. Step d) includes using the at-least-one end-effector-tissue-track ablation device 472, 474 and 476 to ablate the patient tissue surrounding the tissue track along substantially the entire tissue track. Step e) is labeled as "Withdraw End Effector" in block 470 of FIG. 5. Step e) includes withdrawing the interstitial end effector 426 from the patient.

[0040] It is noted that creating a tissue track requires that the interstitial end effector 426 be interstitially inserted into patient tissue. It is also noted that the interstitial end effector 426 can be equipped with a retractable tip shield (not shown) for initial endoscopic or laparoscopic patient entry followed by interstitial insertion into patient tissue.

[0041] In one extension of the method of FIG. 5, there is included the step of using the at-least-one end-effector-tissue-track ablation device (e.g., 474) to ablate the patient tissue at the distal end 462 of the inserted interstitial end effector 426.

[0042] In one implementation of the method of FIG. 5, the at-least-one end-effector-tissue-track ablation device (e.g., 474) includes a non-ultrasound energy source, and step d) uses the non-ultrasound energy source to ablate the patient tissue surrounding the tissue track. In one variation, the non-ultrasound energy source is chosen from the group consisting

of a resistive heat energy source, a hot liquid energy source, a monopolar radio-frequency (RF) energy source, a bipolar radio-frequency (RE) energy source, a capacitive heat energy source, a microwave energy source, and combinations thereof.

[0043] In another implementation of the method, the at-least-one end-effector-tissue-track ablation device (e.g., 476) includes a tissue-ablating chemical agent, and step d) uses the tissue-ablating chemical agent to ablate the patient tissue surrounding the tissue track. In one variation, the tissue-ablating chemical agent is chosen from the group consisting of fibrin, alcohol, an acidic fluid, a chemotherapeutic agent, and combinations thereof.

[0044] In a further implementation of the method, step d) uses the medical ultrasound transducer assembly 418 to ultrasonically ablate the patient tissue surrounding the tissue track. In one variation, step d) ultrasonically ablates at a higher ultrasound frequency than does step c).

[0045] In the same or another extension of the method of FIG. 5, there is included the step of monitoring (such as for acoustic coupling and/or for tissue ablation) the patient tissue surrounding the tissue track during step d). In one variation, the monitoring is chosen from the group consisting of B-mode ultrasonic image monitoring, tissue temperature monitoring, tissue electric impedance, and combinations thereof.

[0046] In the same or another extension of the method of FIG. 5, there are included, after step b) and before step c), the step of stabilizing (such as by using a balloon, a tine and/or suction) the interstitial end effector 426 with respect to the patient tissue surrounding the tissue track and, after step c) and before step d), the step of releasing the stabilizing of the interstitial end effector 426 with respect to the patient tissue surrounding the tissue track.

[0047] In one application of the method of FIG. 5, step e) includes stepwise withdrawing the interstitial end effector 426 from the patient using a plurality of positional steps, and step d) includes ablating the patient tissue surrounding the tissue track for a predetermined time with the interstitial end effector at each positional step.

[0048] A fourth embodiment of the present invention, shown in FIG. 6, is an ultrasound medical system 410 comprising an interstitial end effector 426 which is interstitially insertable into patient tissue, which includes at least one medical-treatment ultrasound transducer 420, and which includes at least one end-effector-tissue-track ablation device 472, 474 and 476.

[0049] In one enablement of the embodiment of FIG. 6, the ultrasound medical system 410 includes a controller (such as the controller 142 illustrated in FIG. 1) which is operatively connected to the at-least-one medical-treatment ultrasound transducer 420 to ultrasonically ablate a lesion in patient tissue of the patient and which is operatively connected to the at-least-one end-effector-tissue-track ablation device 472, 474 and 476 to ablate patient tissue surrounding the interstitial end effector 426 during withdrawal of the interstitial end effector 426 from the patient.

[0050] In one application of the embodiment of FIG. 6, the at-least-one end-effector-tissue-track ablation device 472, 474, 476 includes a cylindrical ultrasound transducer 472. In the same or a different application, the at-least-one end-effector-tissue-track ablation device and the at-least-one medical-treatment ultrasound transducer are a single rotatable ultrasound transducer (such as ultrasound transducer 420

made rotatable such as in a previously illustrated and described implementation of the embodiment of FIGS. 1-2). Other applications of an end-effector-tissue-track ablation device involving ultrasound are left to the artisan.

[0051] In another application of the embodiment of FIG. 6, the at-least-one end-effector-tissue-track ablation device 472, 474 and 476 includes a device 474 which uses a non-ultrasound energy source. In one variation, the non-ultrasound energy source is chosen from the group consisting of a resistive heat energy source, a hot liquid energy source, a monopolar radio-frequency (RF) energy source, a bipolar radio-frequency (RF) energy source, a capacitive heat energy source, a microwave energy source, and combinations thereof.

[0052] In a further application of the embodiment of FIG. 6, the at-least-one end-effector-tissue-track ablation device 472, 474 and 476 includes a device 476 which releases a tissue-ablating chemical agent. In one variation, the tissue-ablating chemical agent is chosen from the group consisting of fibrin, alcohol, an acidic fluid, a chemotherapeutic agent, and combinations thereof.

[0053] In one construction of the embodiment of FIG. 6, the interstitial end effector 426 has a length and an exterior surface 428 and includes position markings 478 on the exterior surface 428 along at least a portion of its length. Such position markings allow a user to withdraw the interstitial end effector 426 from patient tissue in positional steps while ablating patient tissue surrounding the end-effector tissue track for a predetermined dwell time at each positional step. In the same or a different construction, the interstitial end effector 426 has a longitudinal axis 424 and a distal end 462, and wherein the at-least-one end-effector-tissue-track ablation device 472, 474 and 476 includes an end-effector-tissue-track ablation device (such as 474) which is disposed proximate the distal end 462. It is noted that the distal end of an interstitial end effector is an end having a tissue-piercing tip. In the same or a different construction, the interstitial end effector 426 includes a tissue-ablating component (such as 474) adapted (such as by having a resistive heat energy source) to ablate (such as to thermally ablate) patient tissue longitudinally forward of the distal end 462.

[0054] In one variation, the ultrasound interstitial end effector includes a sheath 434 surrounding the medical-treatment ultrasound transducer 120 and having an acoustic window 480. In one modification, the entire sheath acts as an acoustic window. In another modification, the acoustic window is a thinner portion of the sheath. In a further modification, the acoustic window is a separate material(s) from the material(s) of the non-acoustic-window portion(s) of the sheath. Acoustic window component materials are known to those skilled in the art. Other modifications are left to the artisan.

[0055] It is noted that the embodiments, constructions, implementations, etc. of the embodiments of FIGS. 1-2, 3 and 4 are equally applicable to the embodiment, constructions, implementations, etc. of the embodiment of FIG. 6.

[0056] Several benefits and advantages are obtained from one or more of the embodiments and method of the invention. In one example, having an interstitial end effector with a medical-treatment ultrasound transducer and an end-effector-tissue-track ablation device allows ultrasonic ablation of a lesion using the medical-treatment ultrasound transducer and allows ablation of patient tissue surrounding the tissue track

as the interstitial end effector is withdrawn from the patient to help reduce the possibility of excessive bleeding and/or tissue contamination.

[0057] While the present invention has been illustrated by a description of several embodiments and methods, it is not the intention of the applicants to restrict or limit the spirit and scope of the appended claims to such detail. Numerous other variations, changes, and substitutions will occur to those skilled in the art without departing from the scope of the invention. For instance, the ultrasound medical system of the invention has application in robotic assisted surgery taking into account the obvious modifications of such systems, components and methods to be compatible with such a robotic system. It will be understood that the foregoing description is provided by way of example, and that other modifications may occur to those skilled in the art without departing from the scope and spirit of the appended Claims.

1. An ultrasound treatment system comprising:

a probe comprising a treatment ultrasound transducer configured to treat patient tissue;

a non-ultrasound tissue-property measuring sensor coupled to the probe; and

a controller configured to communicate with the probe and the non-ultrasound tissue-property measuring sensor; wherein the non-ultrasound tissue-property measuring sensor is positioned to be in contact with the patient tissue

wherein the controller is configured to control the probe and to receive feedback from the non-ultrasound tissue-property measuring sensor.

2. The ultrasound treatment system according to claim 1, wherein the non-ultrasound tissue-property measuring sensor is configured to measure tissue temperature.

3. The ultrasound treatment system according to claim 2, wherein the non-ultrasound tissue-property measuring sensor is at least one of a thermistor, and a thermocouple.

4. The ultrasound treatment system according to claim 1, wherein the non-ultrasound tissue-property measuring sensor is configured to measure tissue electrical impedance.

5. The ultrasound treatment system according to claim 4, wherein the non-ultrasound tissue-property measuring sensor is at least one of a monopolar electrode, and a bipolar electrode.

6. The ultrasound treatment system according to claim 1, wherein the non-ultrasound tissue-property measuring sensor is configured to a measure of the degree of ultrasonic ablation of the patient tissue.

7. The ultrasound treatment system according to claim 1, wherein the non-ultrasound tissue-property measuring sensor is configured to a measure of the degree of acoustic coupling to patient tissue.

8. The ultrasound treatment system according to claim 1, further comprising a tissue-ablating chemical agent configured to ablate the patient tissue.

9. The ultrasound treatment system according to claim 1, wherein the tissue-ablating chemical agent is at least one of fibrin, alcohol, an acidic fluid, and a chemotherapeutic agent.

10. The ultrasound treatment system according to claim 1, further comprising a non-ultrasound energy source configured to ablate the patient tissue.

11. The ultrasound treatment system according to claim 10, wherein the non-ultrasound energy source is at least one of a resistive heat energy source, a hot liquid energy source, a

monopolar radio-frequency energy source, a bipolar radio-frequency energy source, a capacitive heat energy source, and a microwave energy source.

12. The ultrasound treatment system according to claim **1**, wherein the probe is configured to dispense a drug to the patient tissue.

13. The ultrasound treatment system according to claim **12**, wherein the drug is at least potentiated by ultrasound energy emitted from the treatment ultrasound transducer.

14. An ultrasound treatment system comprising:

a hand piece comprising a treatment ultrasound transducer configured to treat patient tissue; and

a non-ultrasound tissue-property measuring sensor configured to measure tissue electrical impedance;

wherein the non-ultrasound tissue-property measuring sensor is positioned to be in contact with the patient tissue

wherein the tissue electrical impedance reported by the sensor is a measure of the degree of ultrasonic ablation of the patient tissue.

15. The ultrasound treatment system according to claim **14**, further comprising a controller configured to control the treatment ultrasound transducer and to receive the tissue electrical impedance reported by the non-ultrasound tissue-property measuring sensor.

16. The ultrasound treatment system according to claim **14**, wherein the non-ultrasound tissue-property measuring sensor is at least one of a monopolar electrode, and a bipolar electrode.

17. The ultrasound treatment system according to claim **14**, further comprising at least one imaging ultrasound transducer positioned in the hand piece.

18. The ultrasound treatment system according to claim **14**, wherein the non-ultrasound tissue-property measuring sensor is configured to monitor the patient tissue during treatment.

19. An ultrasound treatment system comprising:

a hand-held probe comprising at least one ultrasound transducer configured to treat patient tissue; and

a non-ultrasound tissue-property measuring sensor supported by the hand-held probe and positioned to be in contact with the patient tissue, the non-ultrasound tissue-property measuring sensor is configured to measure tissue electrical impedance and to report a measure of measure of the degree of ultrasonic ablation of the patient tissue as a function of the measure tissue electrical impedance.

20. The ultrasound treatment system according to claim **14**, further comprising a controller configured to control the treatment ultrasound transducer and to receive the tissue electrical impedance reported by the non-ultrasound tissue-property measuring sensor.

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专利名称(译)	超声医疗系统和方法		
公开(公告)号	US20130218013A1	公开(公告)日	2013-08-22
申请号	US13/752151	申请日	2013-01-28
[标]申请(专利权)人(译)	BARTHE PETER 斯雷顿MICHAEL JAEGER PAUL MAST DOUGLAS MAKIN因德尔 OCONNER BRIAN 梅瑟利JEFFERY FAIDI克瓦 RUNK MEGAN PARK CHRISTOPHER		
申请(专利权)人(译)	BARTHE , PETER 斯雷顿 , MICHAEL JAEGER , PAUL MAST , DOUGLAS MAKIN , 因德尔 OCONNER , BRIAN 梅瑟利 , 杰弗瑞 FAIDI , 克瓦 RUNK , 梅根 PARK , CHRISTOPHER		
当前申请(专利权)人(译)	BARTHE , PETER 斯雷顿 , MICHAEL JAEGER , PAUL MAST , DOUGLAS MAKIN , 因德尔 OCONNER , BRIAN 梅瑟利 , 杰弗瑞 FAIDI , 克瓦 RUNK , 梅根 PARK , CHRISTOPHER		
[标]发明人	BARTHE PETER SLAYTON MICHAEL JAEGER PAUL MAST DOUGLAS MAKIN INDER OCONNER BRIAN MESSERLY JEFFERY FAIDI WASEEM RUNK MEGAN PARK CHRISTOPHER		
发明人	BARTHE, PETER SLAYTON, MICHAEL JAEGER, PAUL MAST, DOUGLAS MAKIN, INDER		

OCONNER, BRIAN
MESSERLY, JEFFERY
FAIDI, WASEEM
RUNK, MEGAN
PARK, CHRISTOPHER

IPC分类号	A61B5/00 A61B18/12 A61B18/06 A61B18/18 A61B5/053 A61B8/13 A61N7/00 A61B18/08 A61B17/22 A61B19/00 A61H1/00
CPC分类号	A61B17/2202 A61B2019/5276 A61B5/053 A61B18/1815 A61N7/00 A61B18/08 A61B18/06 A61B8/13 A61B5/4836 A61B18/12 A61B8/4281 A61B8/4483 A61B2090/378
外部链接	Espacenet USPTO

摘要(译)

超声医疗系统包括间质末端执行器。间质末端执行器可间隙地插入患者组织中，包括至少一个医疗超声换能器，并且包括至少一个末端执行器-组织-轨道消融装置。用于超声治疗患者病变的一种方法包括以下步骤：获得间质末端执行器并将其插入患者体内，形成组织轨道，该组织轨道被患者组织包围并终止于插入的间质末端执行器的远端。其他步骤包括使用至少一个医疗处理超声换能器超声消融病变，使用至少一个末端执行器-组织-轨道消融装置沿基本上整个组织消融围绕组织轨道的患者组织跟踪并从患者身上撤回末端执行器。

