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Mickelsen

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(54) **CATHETERS, CATHETER SYSTEMS, AND METHODS FOR PUNCTURING THROUGH A TISSUE STRUCTURE AND ABLATING A TISSUE REGION**

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CPC *A61B 34/73* (2016.02); *A61B 18/1492* (2013.01); *A61M 25/0127* (2013.01);
(Continued)

(58) **Field of Classification Search**
CPC A61B 18/1492; A61B 18/1815; A61M 25/0606; A61M 2025/0089; A61M 25/0169

See application file for complete search history.

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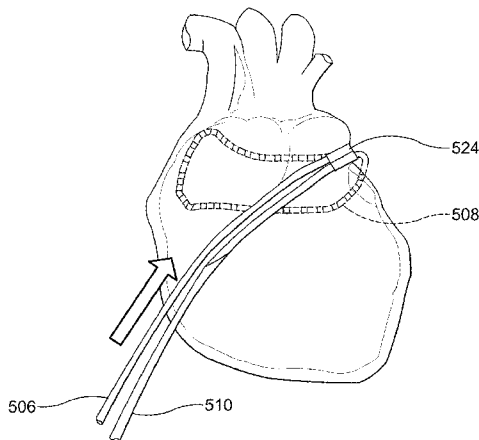
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(57) **ABSTRACT**

A percutaneous catheter system for use within the human body and an ablation catheter for ablating a selected tissue region within the body of a subject. The percutaneous catheter system can include two catheters that are operatively coupled to one another by magnetic coupling through a tissue structure. The ablation catheter can include electrodes positioned within a central portion. The ablation catheter is positioned such that the central portion of a flexible shaft at least partially surrounds the selected tissue region. Each electrode of the ablation catheter can be activated independently to apply ablative energy to the selected tissue region. The ablation catheter can employ high impedance structures to change the current density at specific points. Methods of puncturing through a tissue structure using the percutaneous catheter system are disclosed. Also disclosed are methods for ablating a selected tissue region using the ablation catheter.

19 Claims, 43 Drawing Sheets



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	<i>A61B 18/14</i>	(2006.01)			
	<i>A61M 25/01</i>	(2006.01)			
	<i>A61M 25/06</i>	(2006.01)			
	<i>A61M 25/09</i>	(2006.01)			
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	<i>A61M 25/00</i>	(2006.01)			
(52)	U.S. Cl.				
	CPC <i>A61M 25/0169</i> (2013.01); <i>A61M 25/0606</i>				
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	(2013.01); <i>A61B 2018/00577</i> (2013.01); <i>A61B</i>				
	<i>2018/00613</i> (2013.01); <i>A61B 2018/00839</i>				
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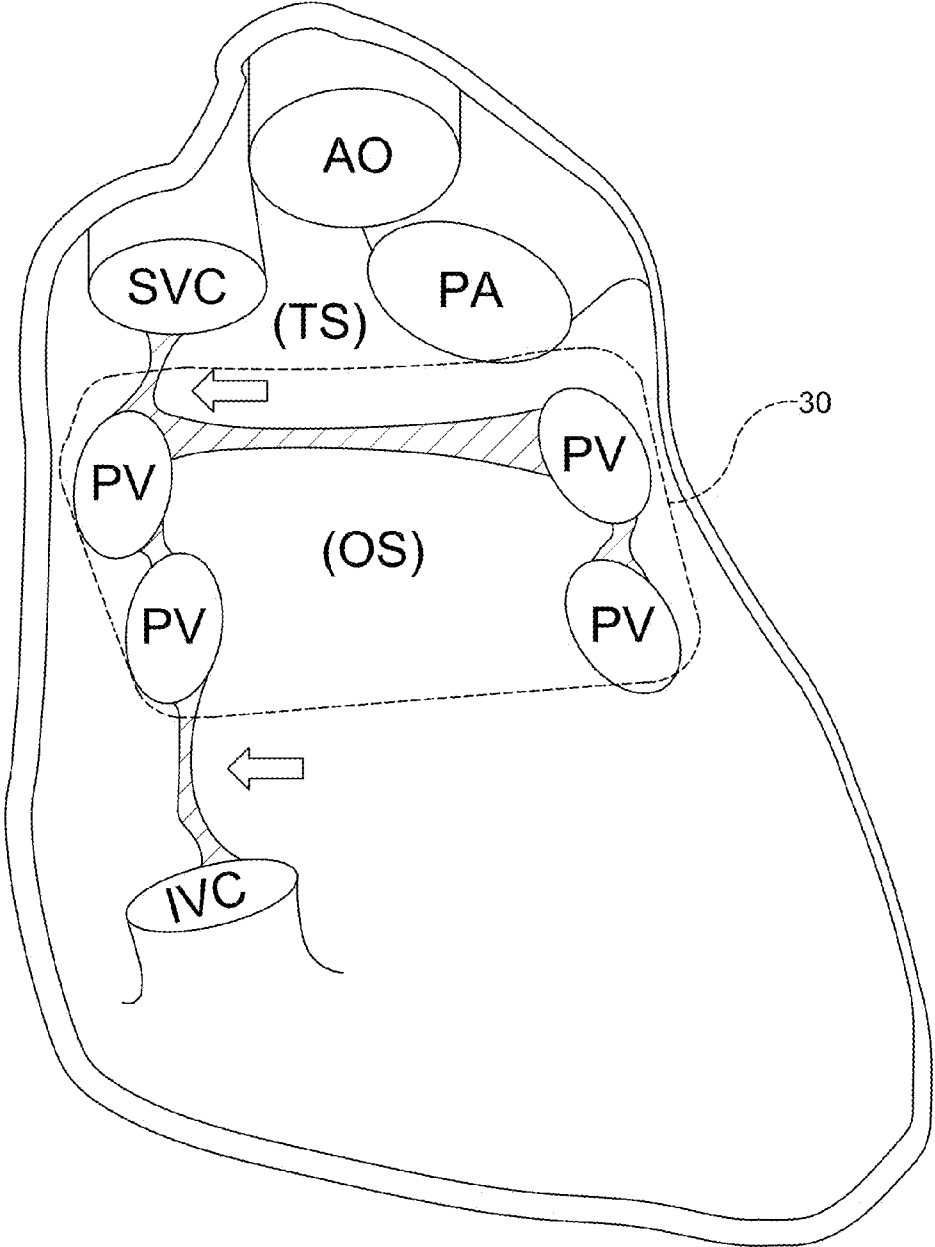


FIG. 1

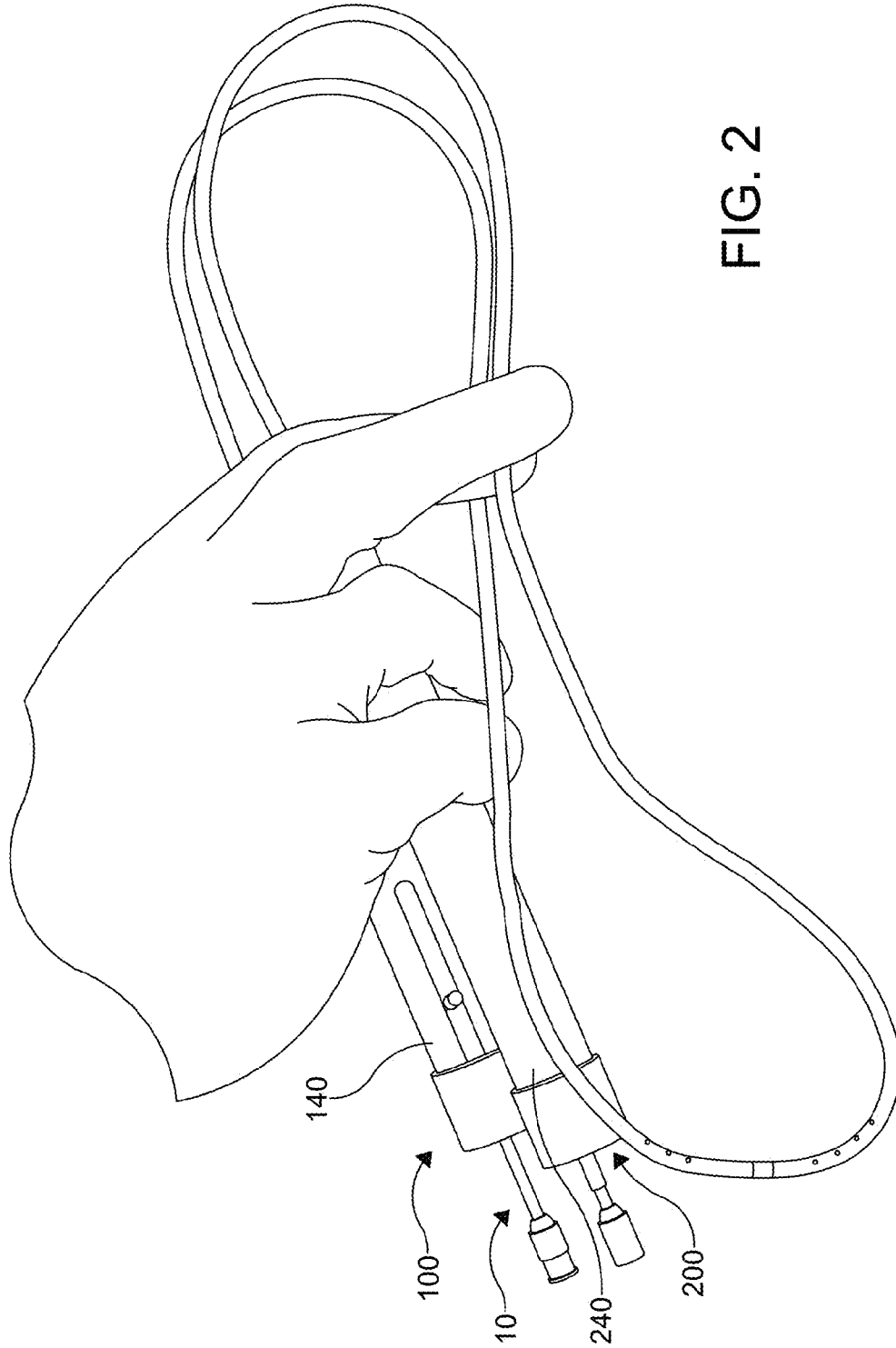


FIG. 2

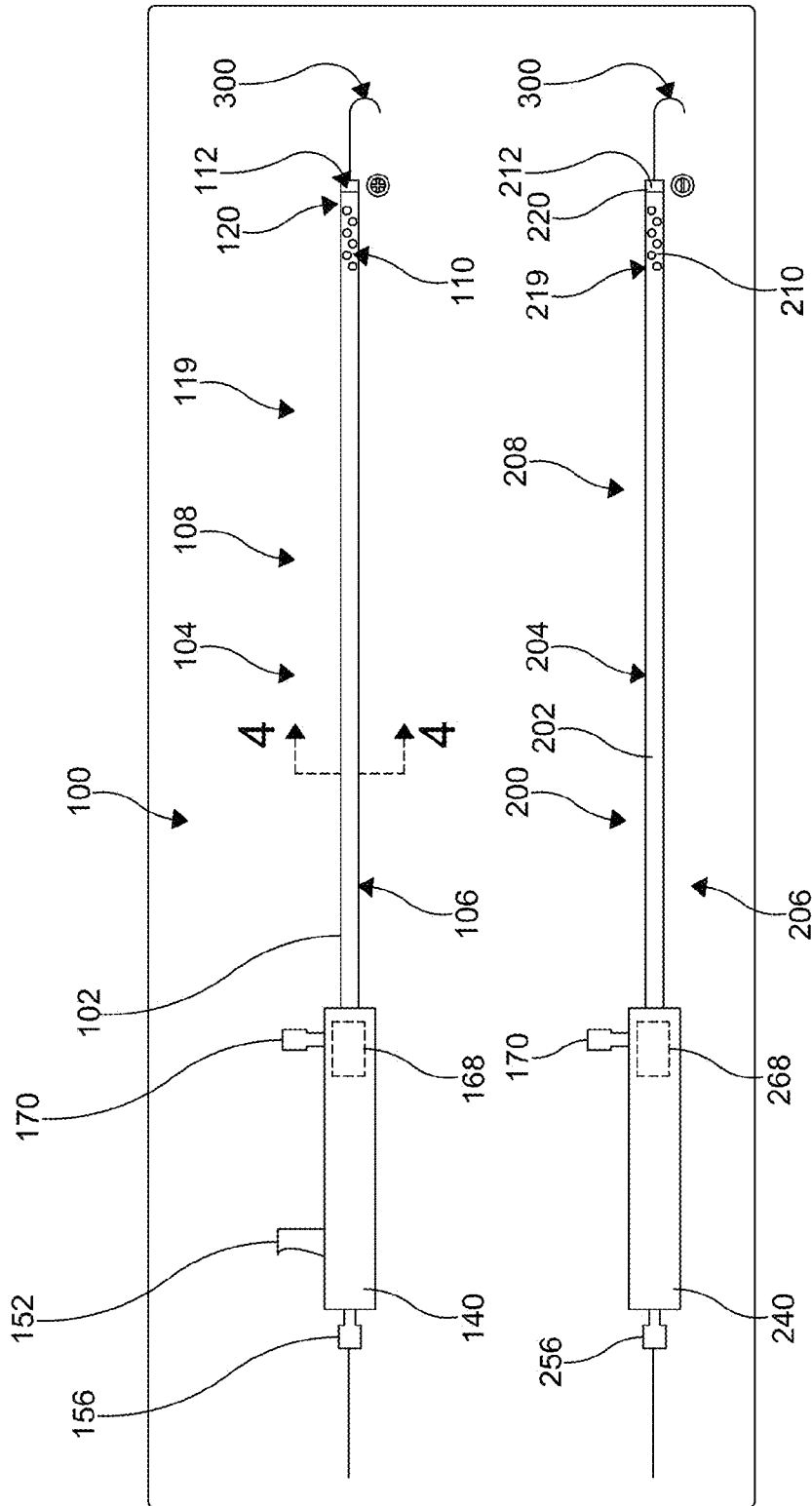


FIG. 3

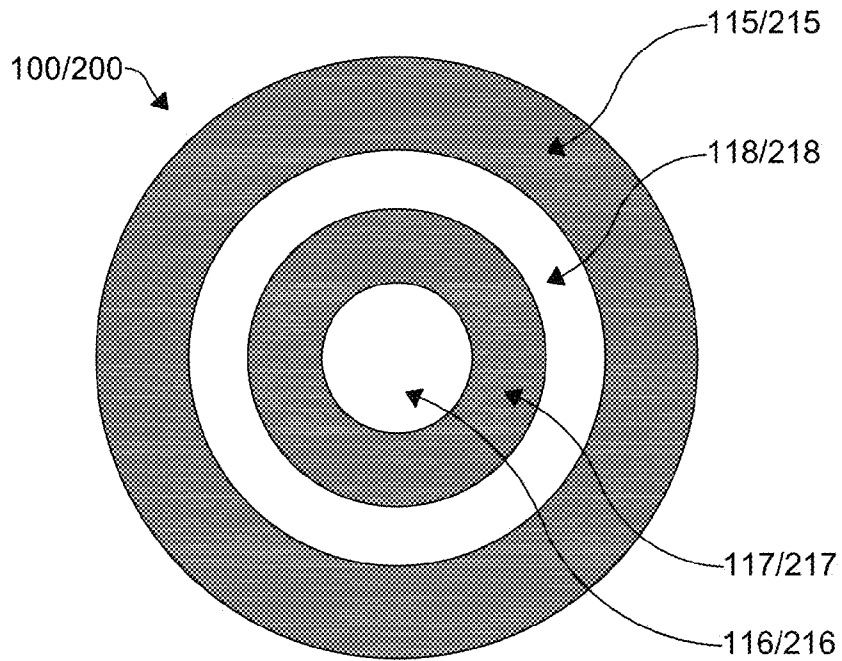


FIG. 4

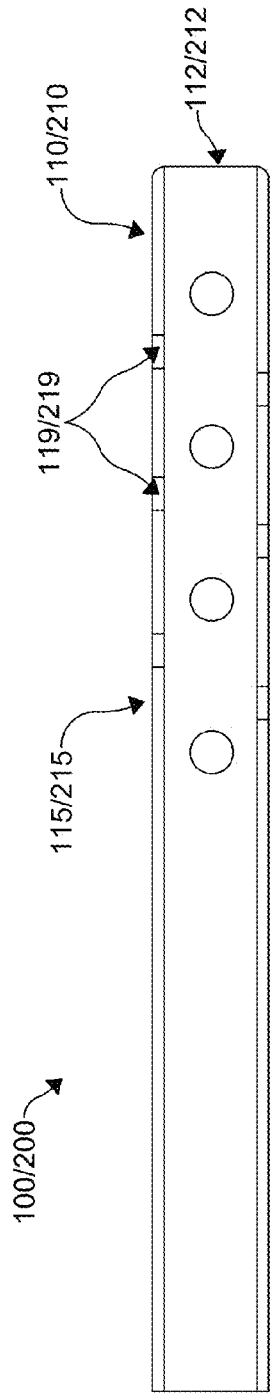


FIG. 5a

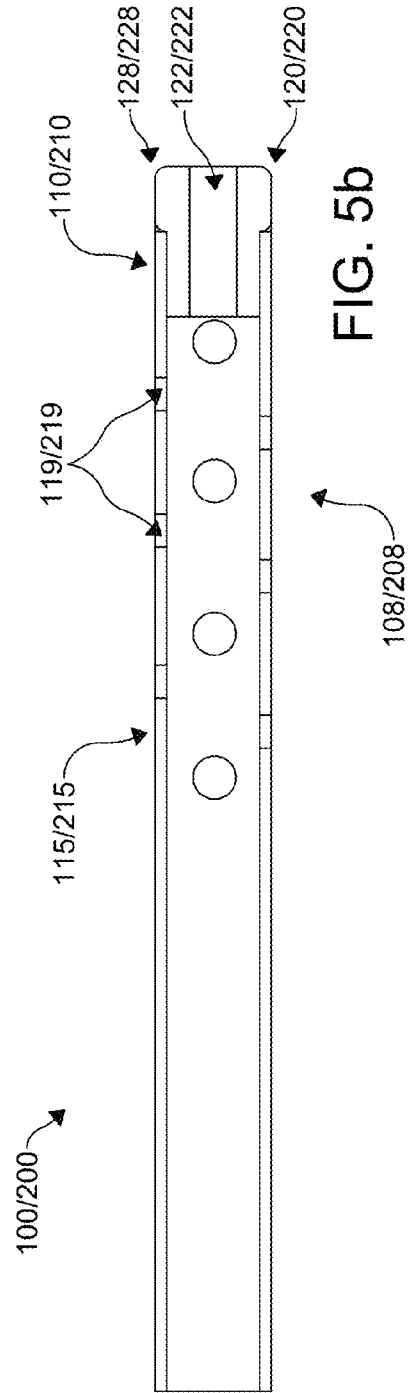


FIG. 5b

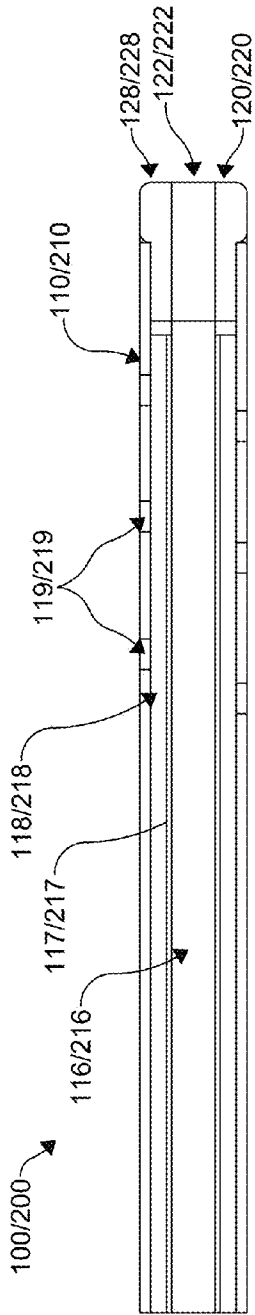


FIG. 5c

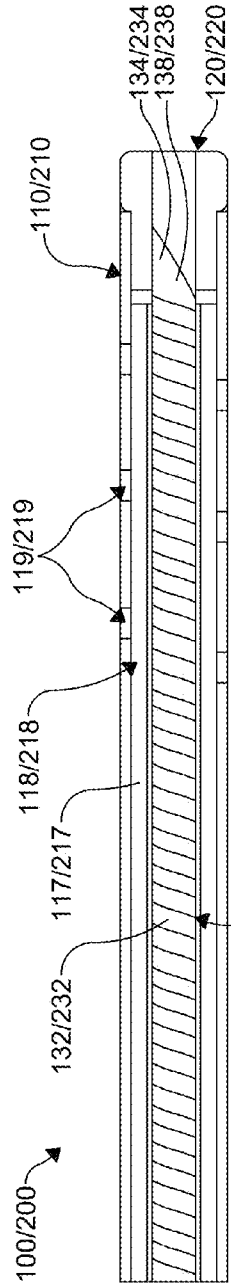


FIG. 5d

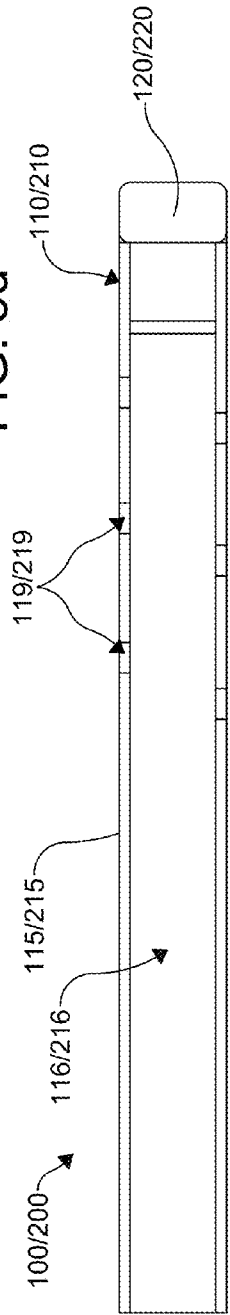


FIG. 5e

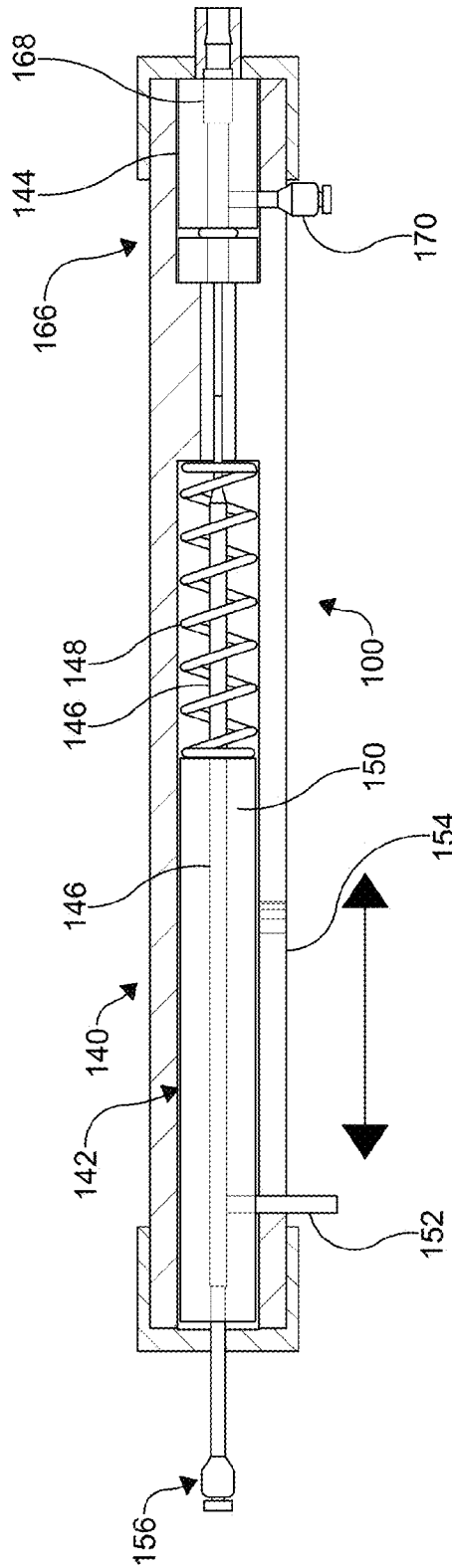


FIG. 6

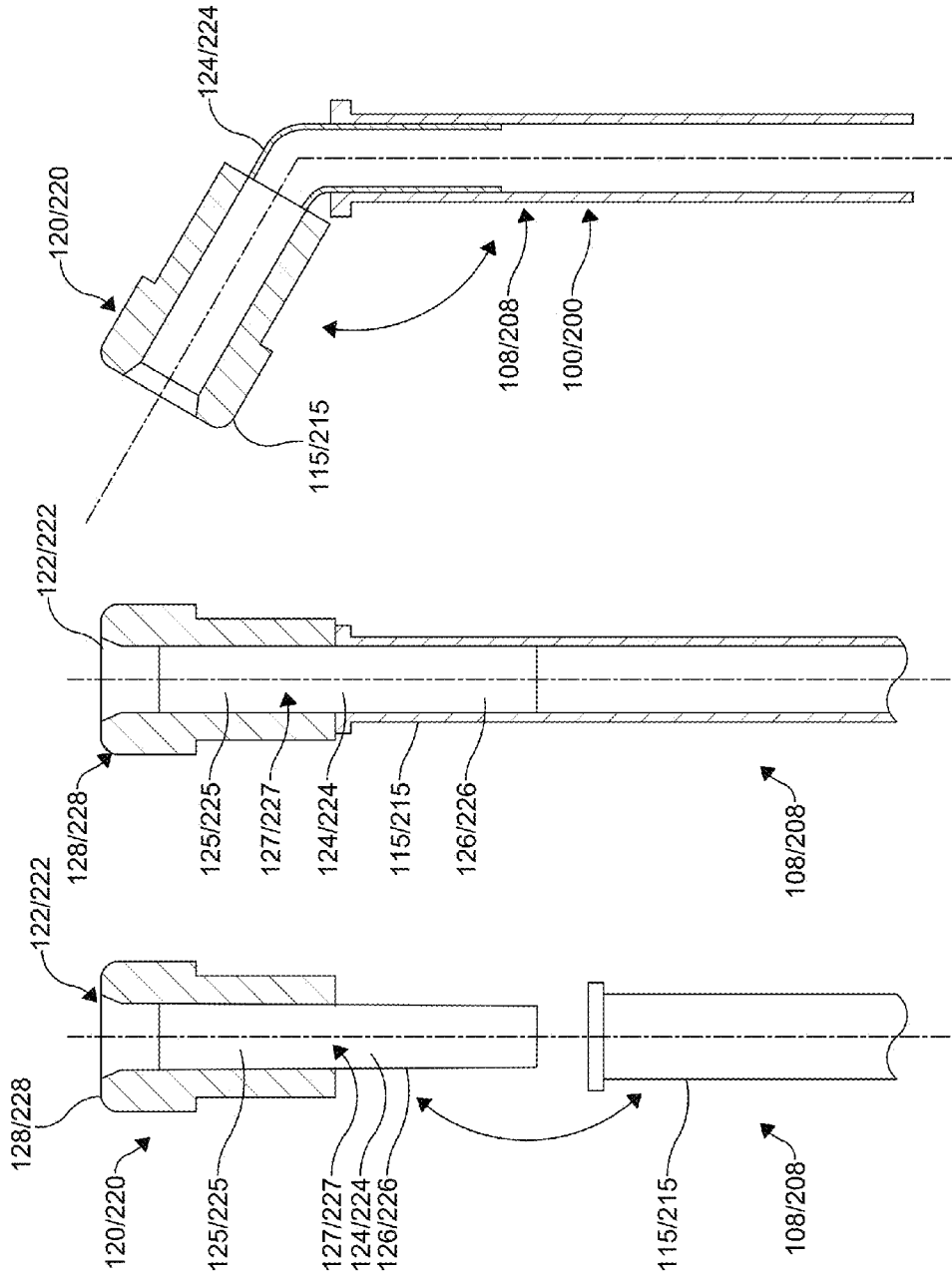


FIG. 8

FIG. 7b

FIG. 7a

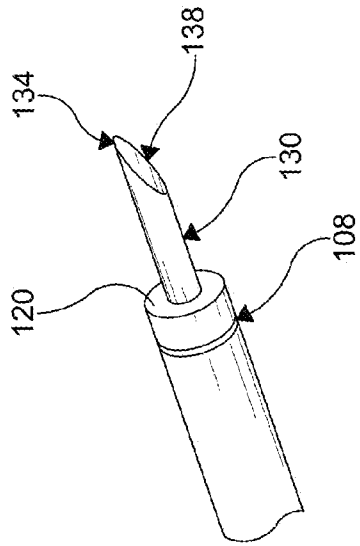


FIG. 9

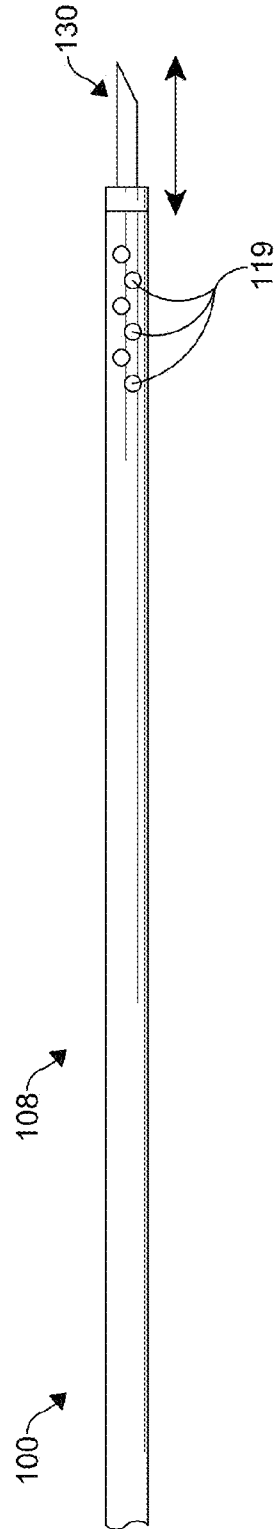
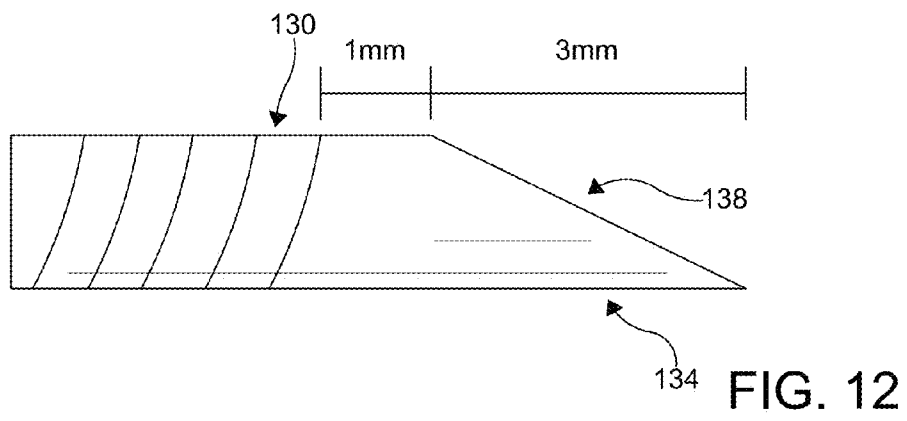
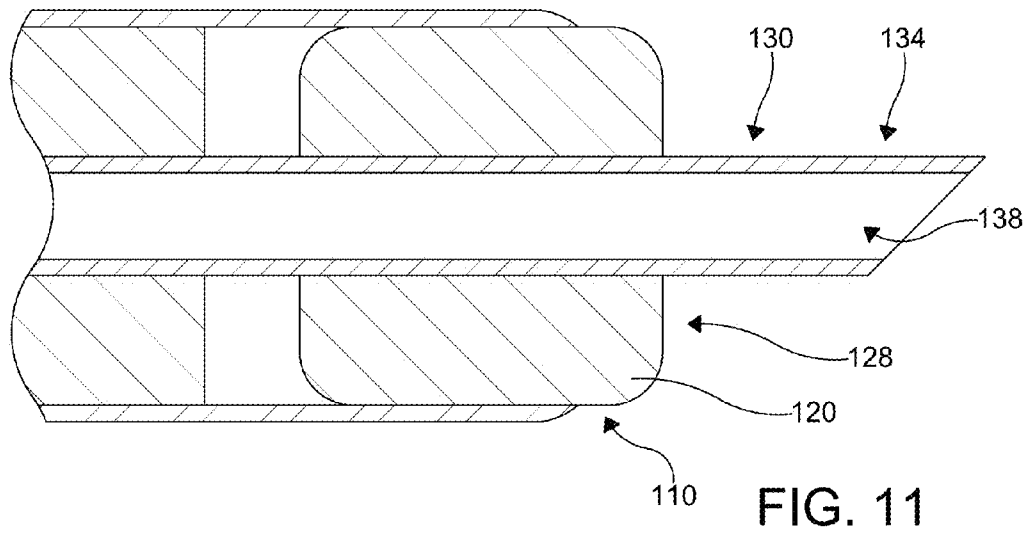


FIG. 10



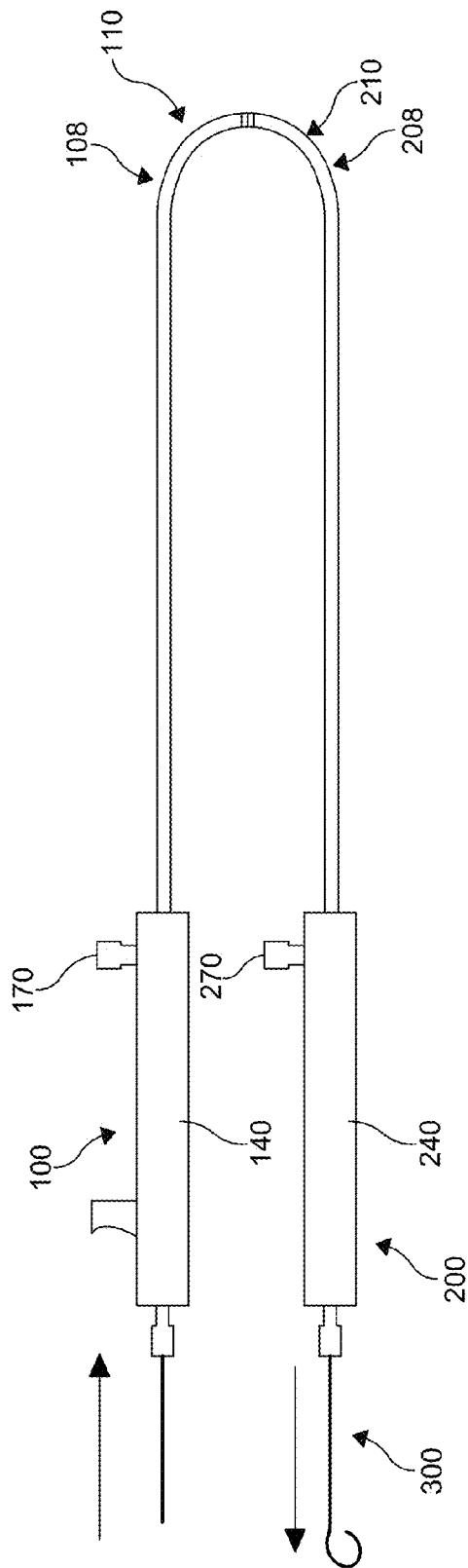
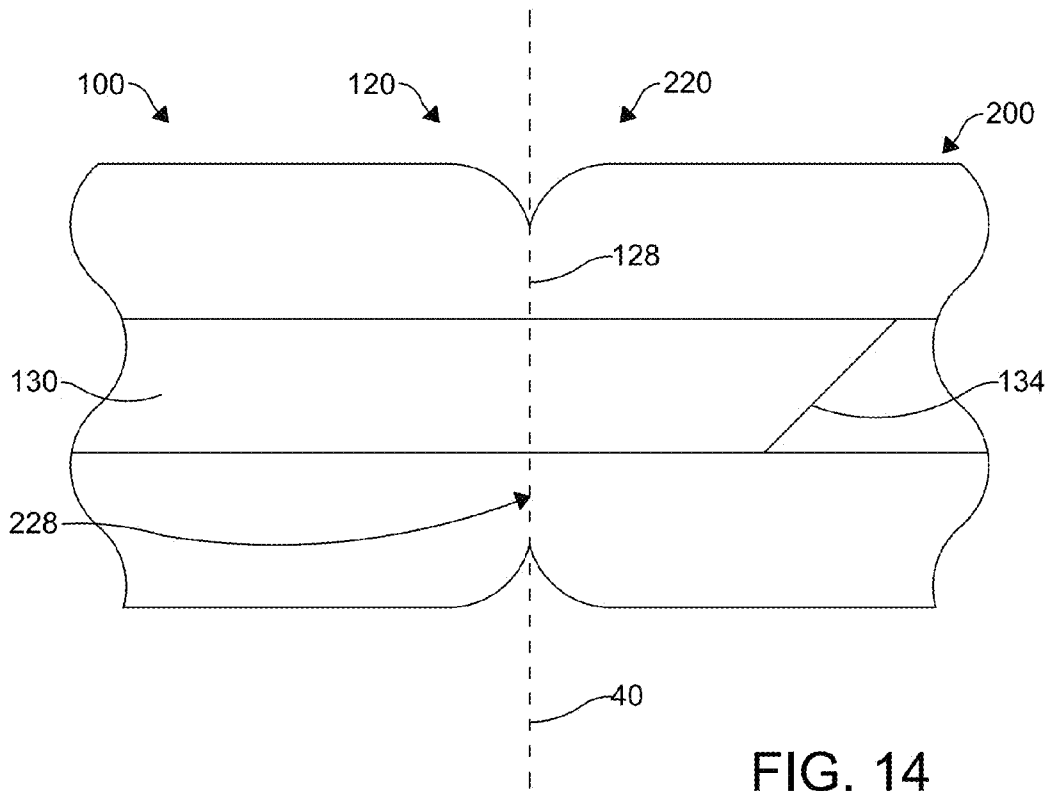


FIG. 13



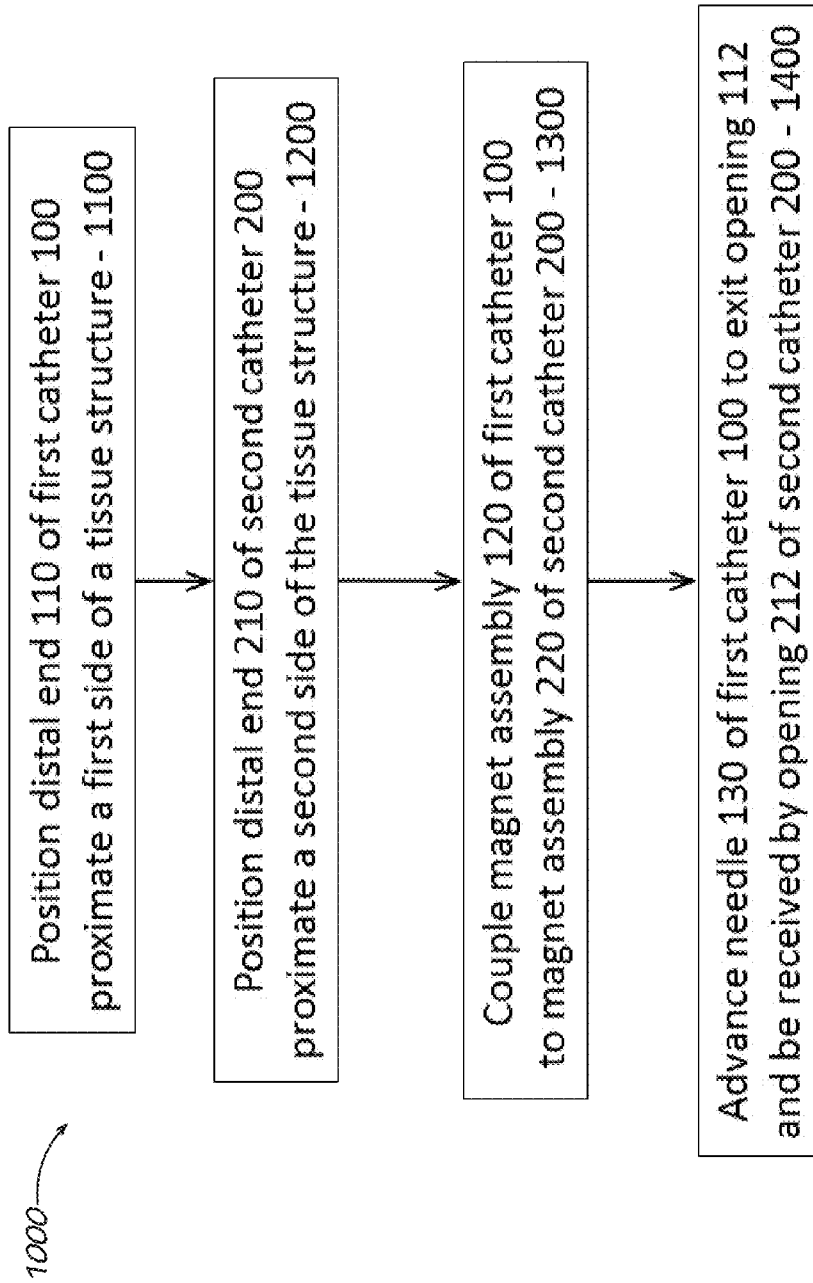


FIG. 15

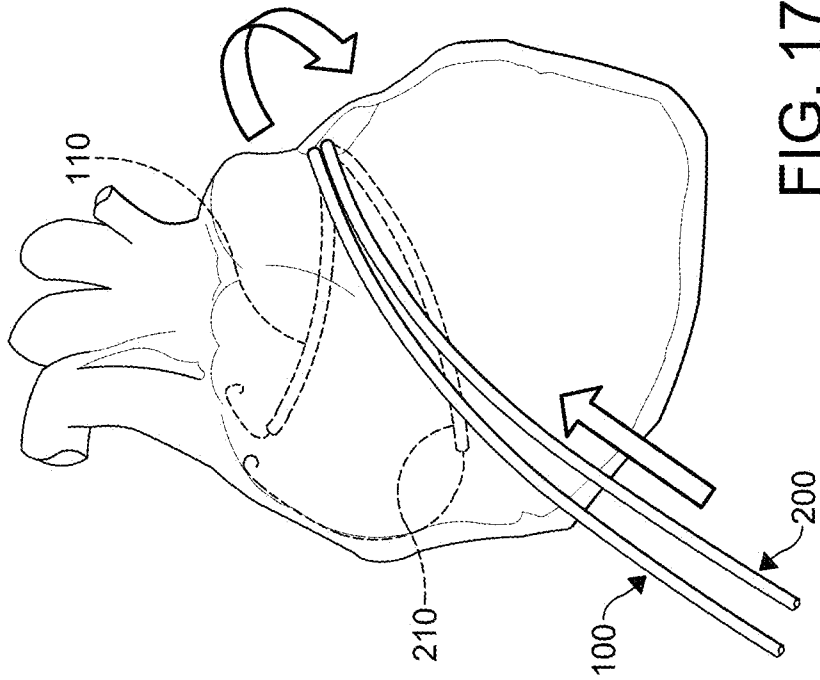


FIG. 17

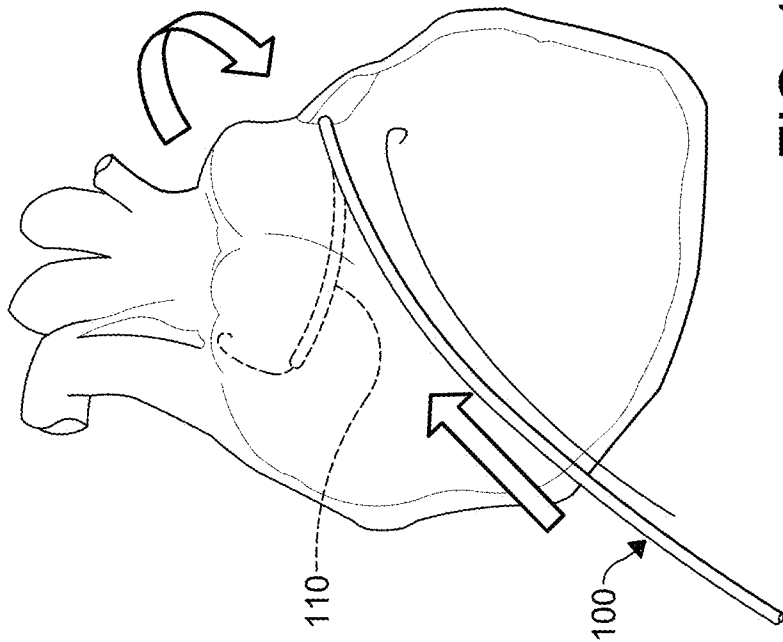


FIG. 16

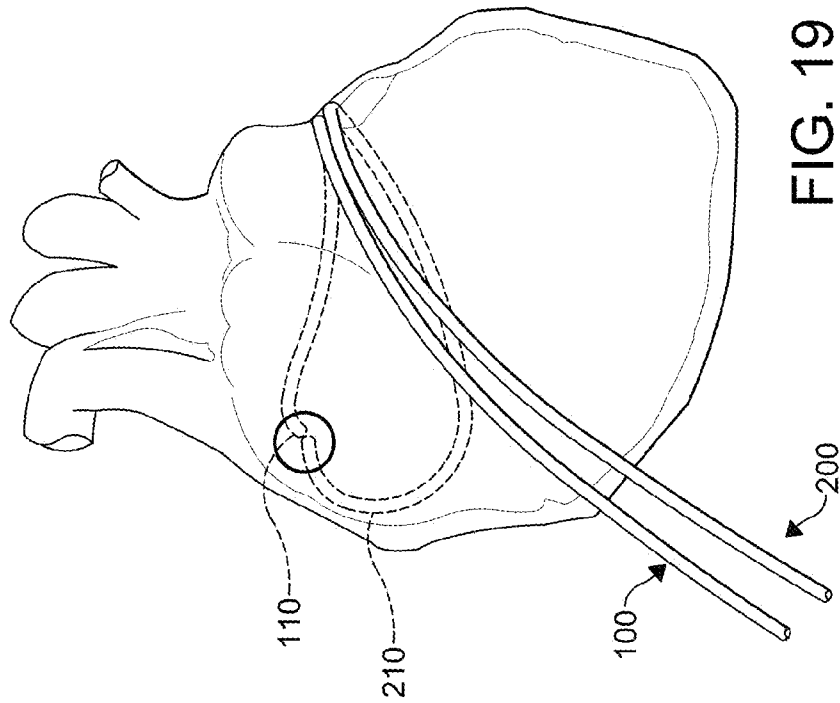


FIG. 19

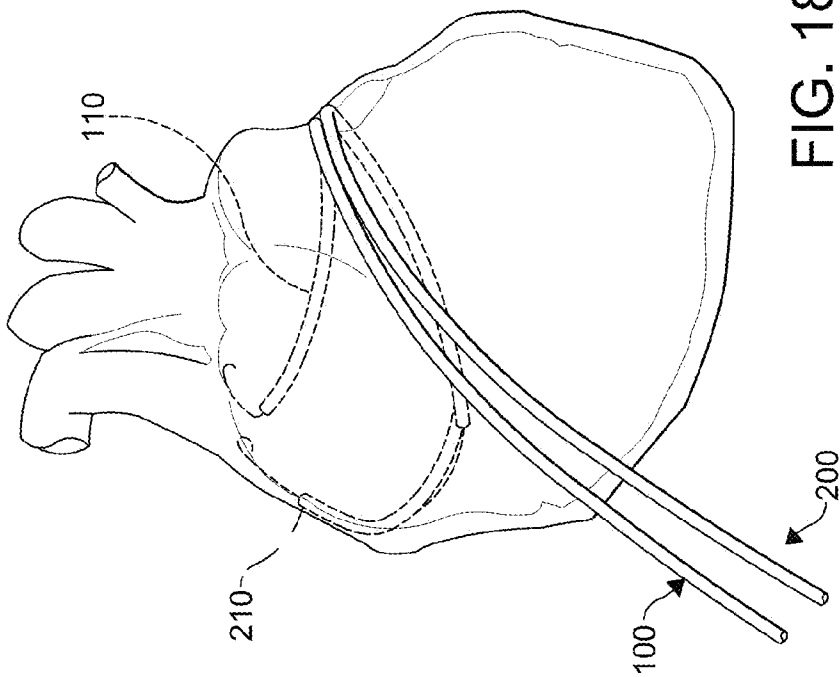
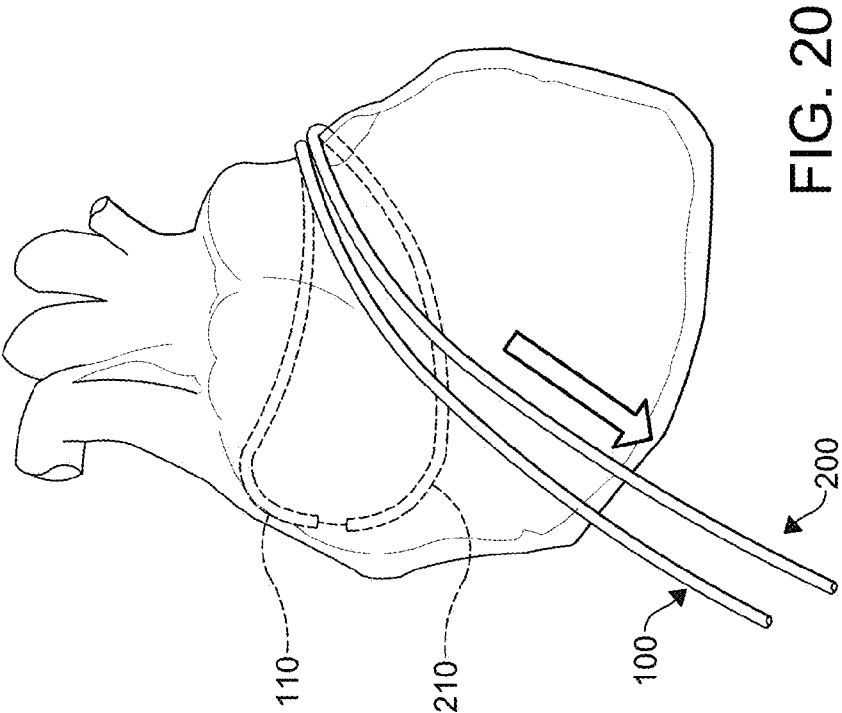
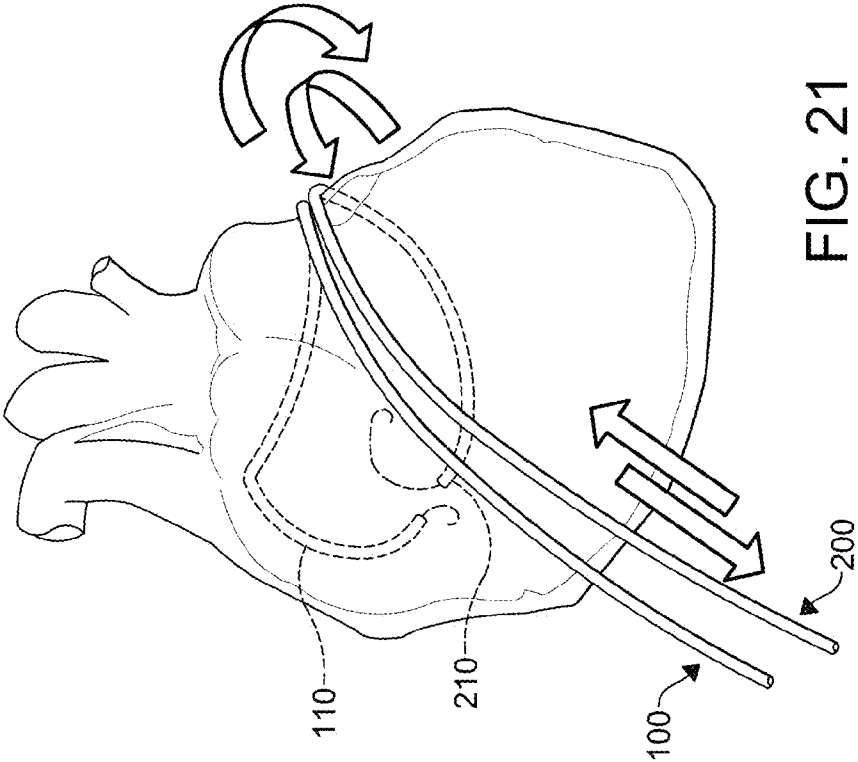
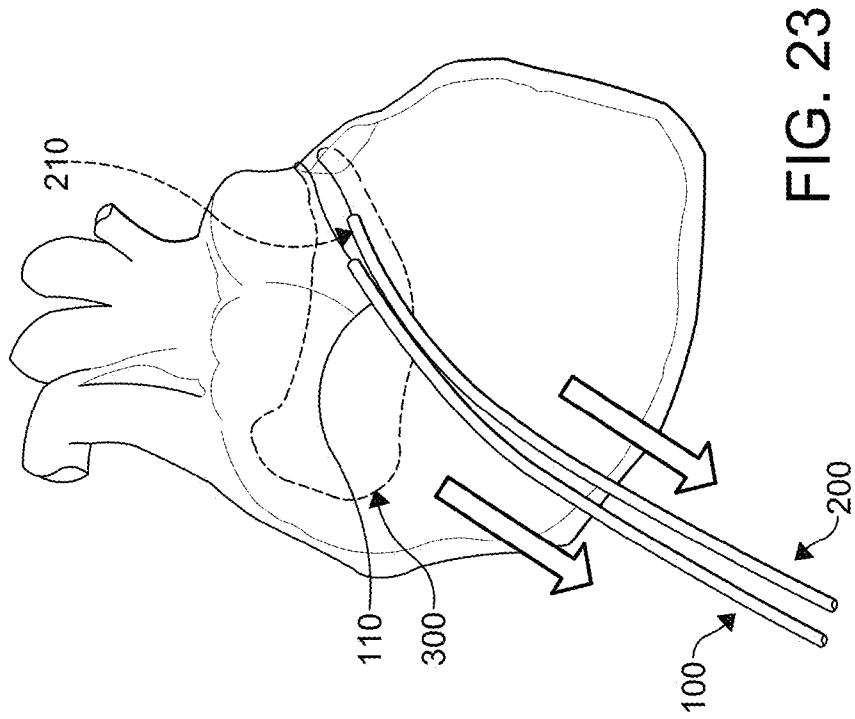
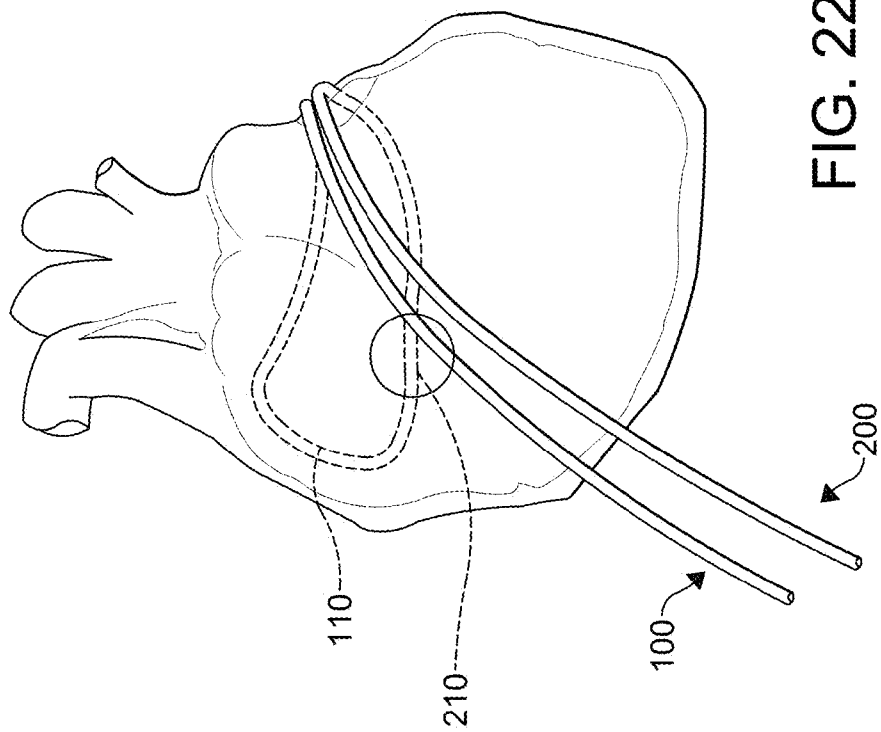


FIG. 18





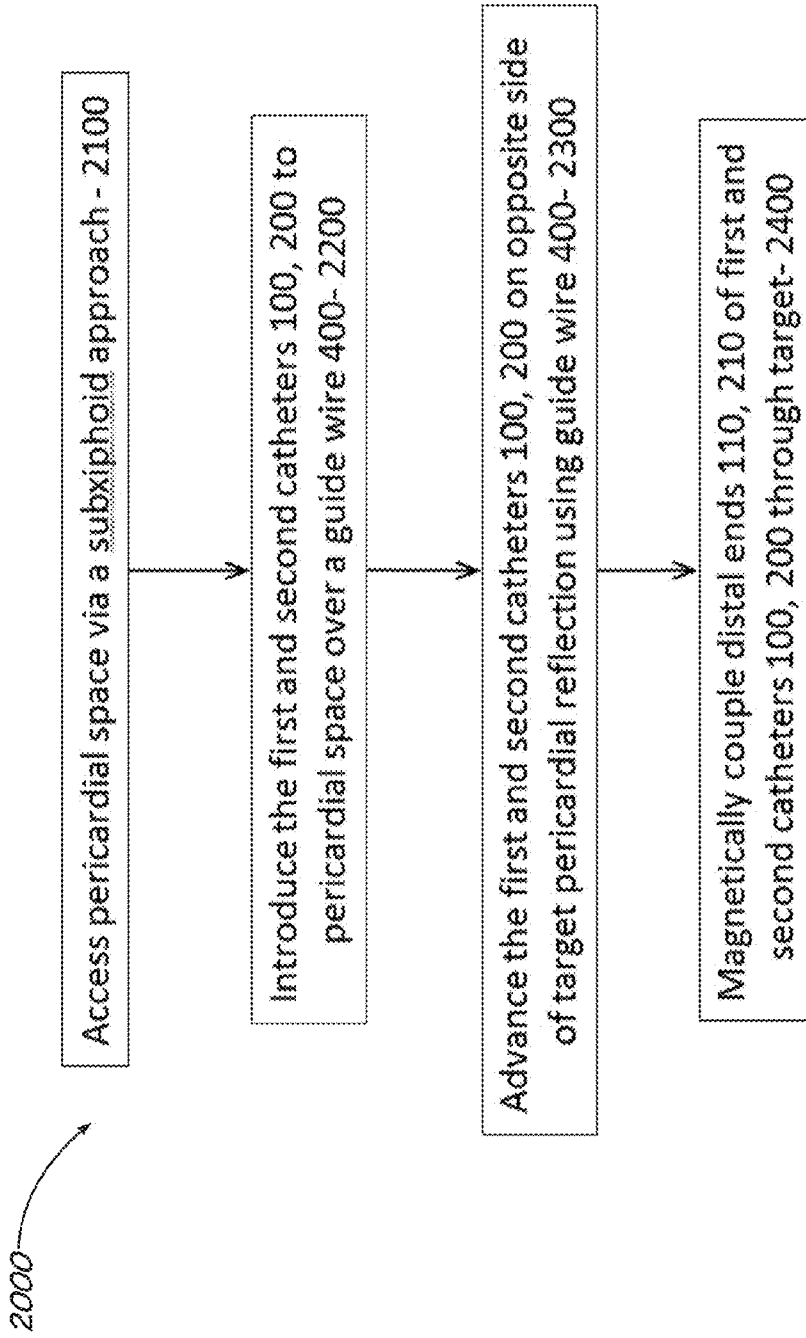


FIG. 24

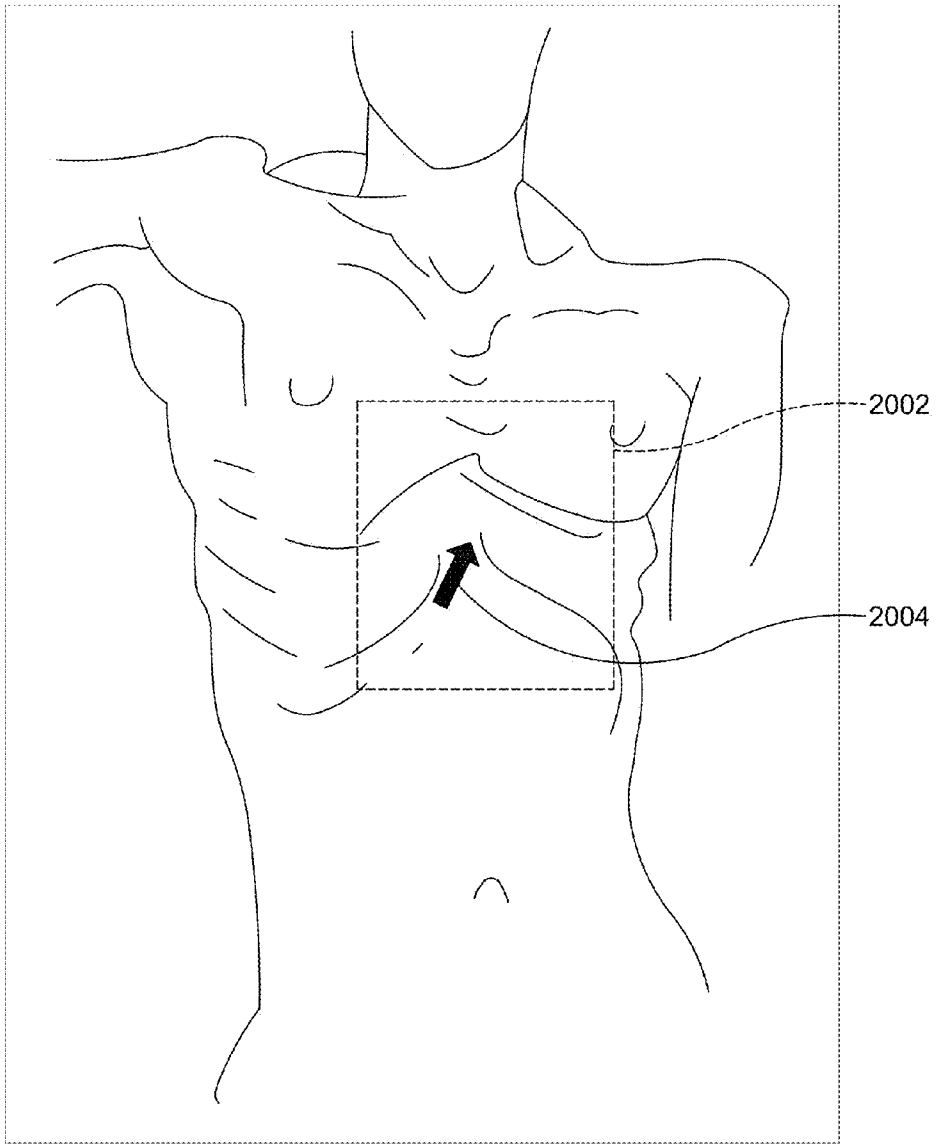


FIG. 25

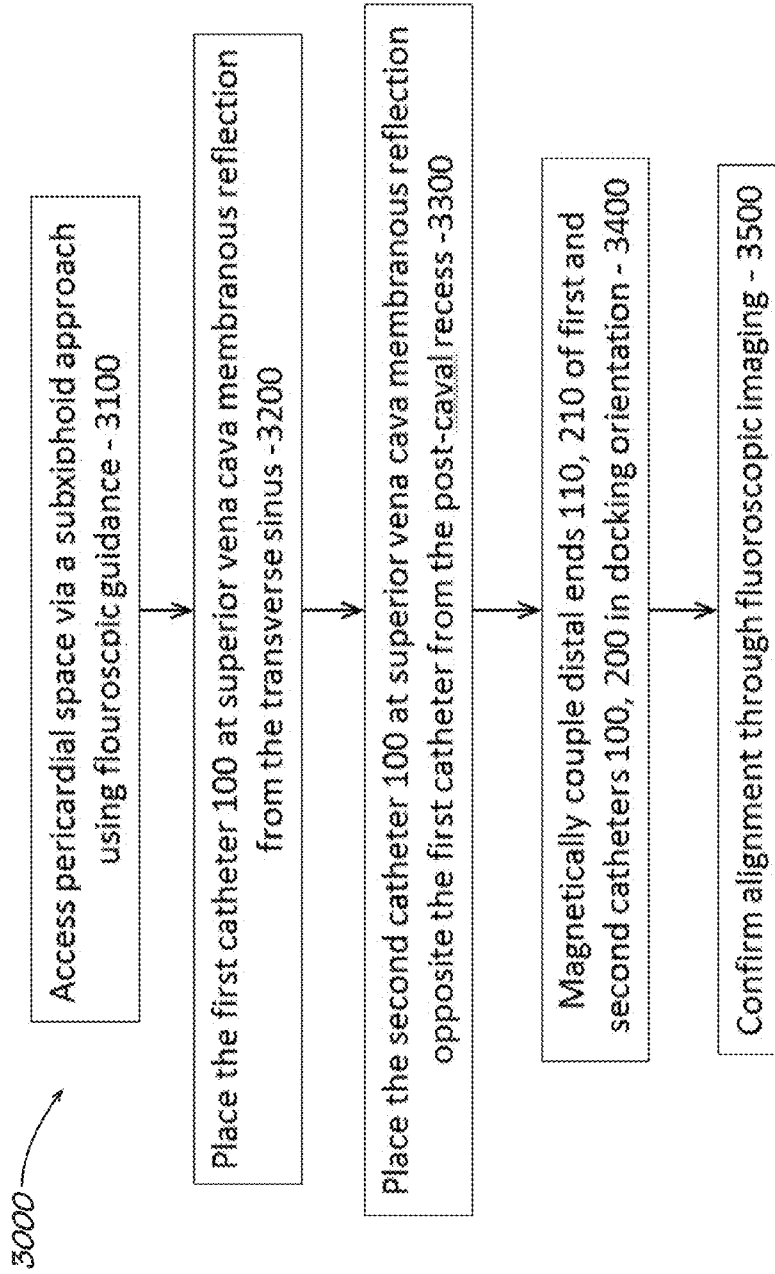


FIG. 26

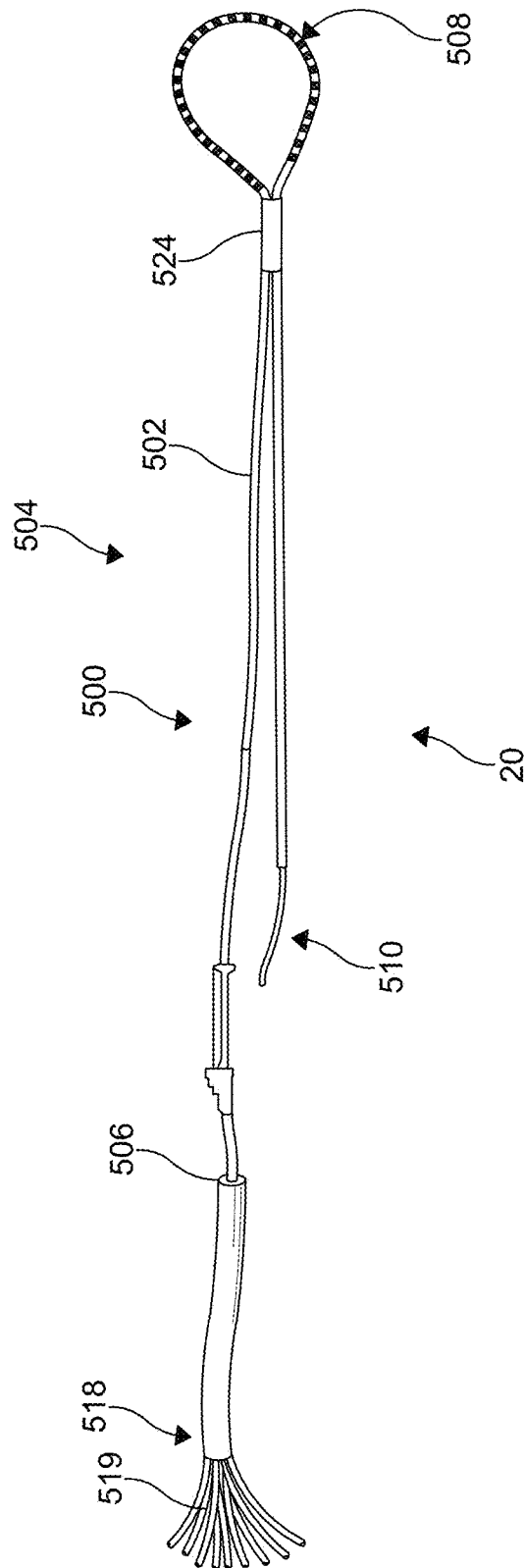


FIG. 27

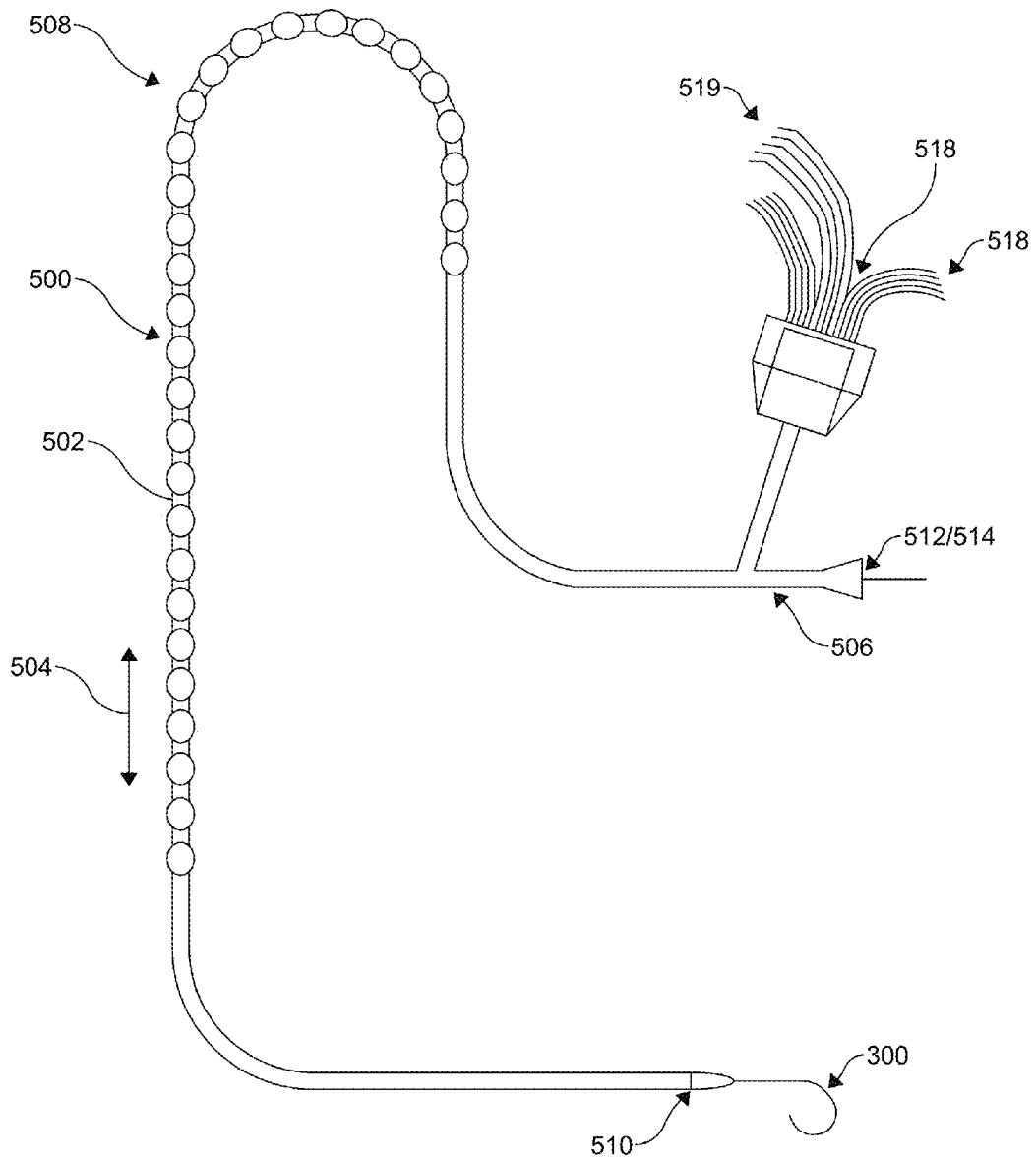


FIG. 28

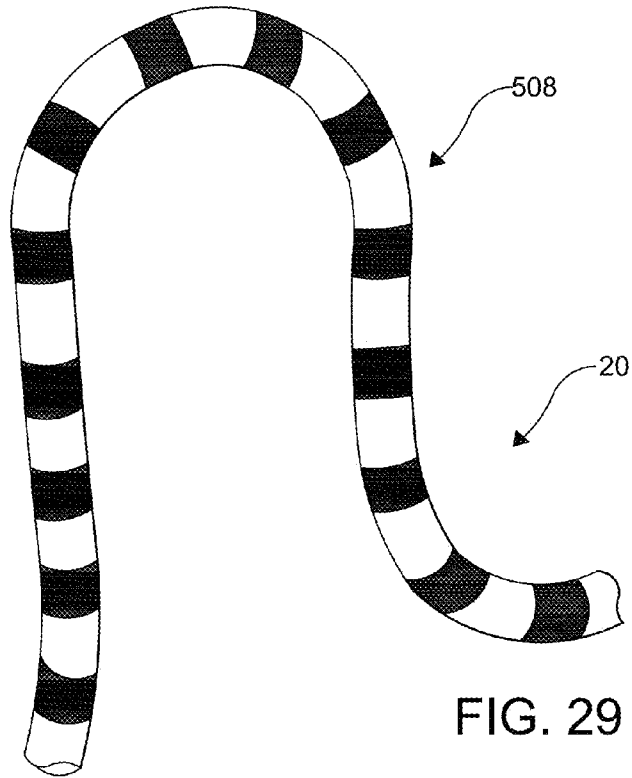


FIG. 29

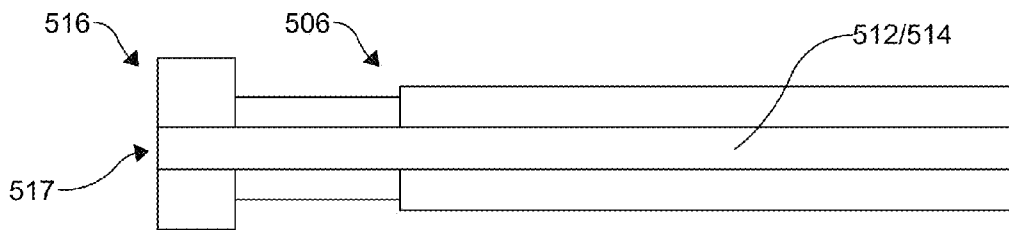


FIG. 30

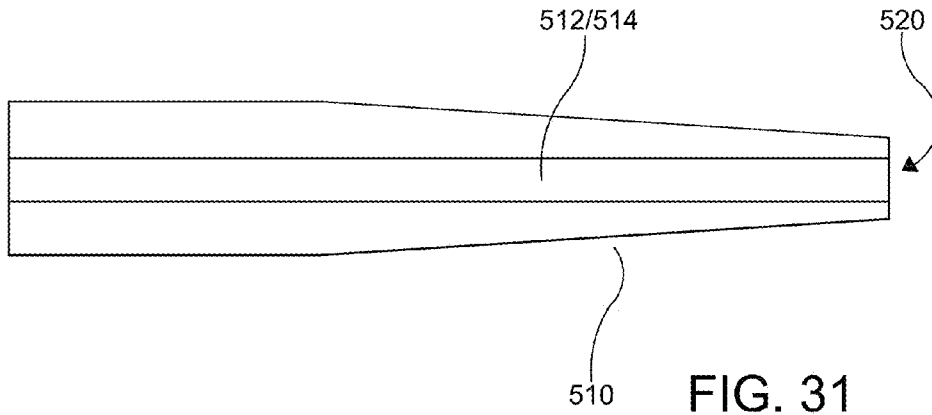


FIG. 31

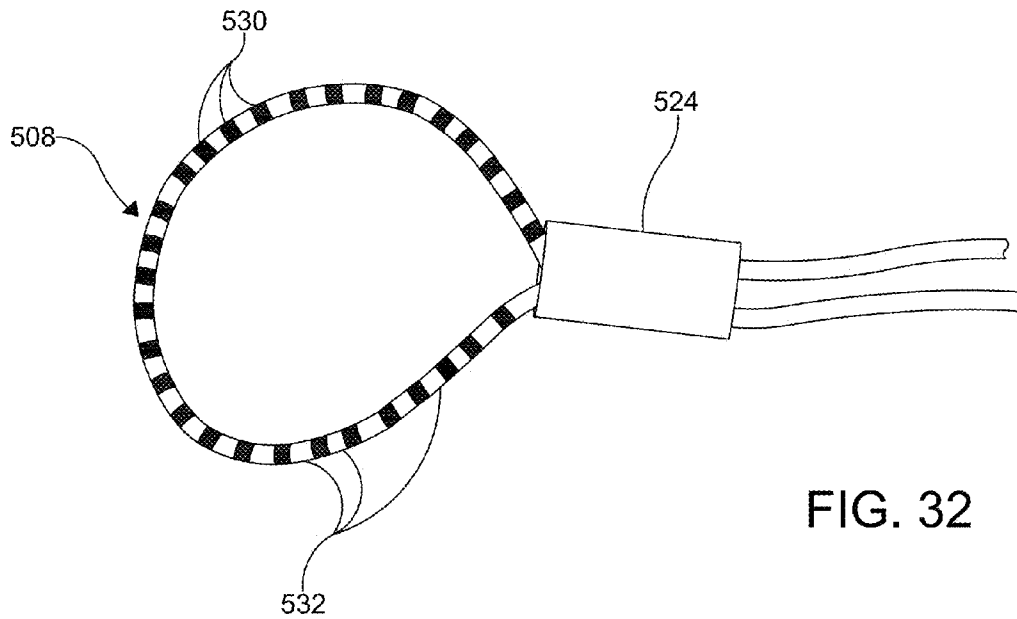


FIG. 32



FIG. 33

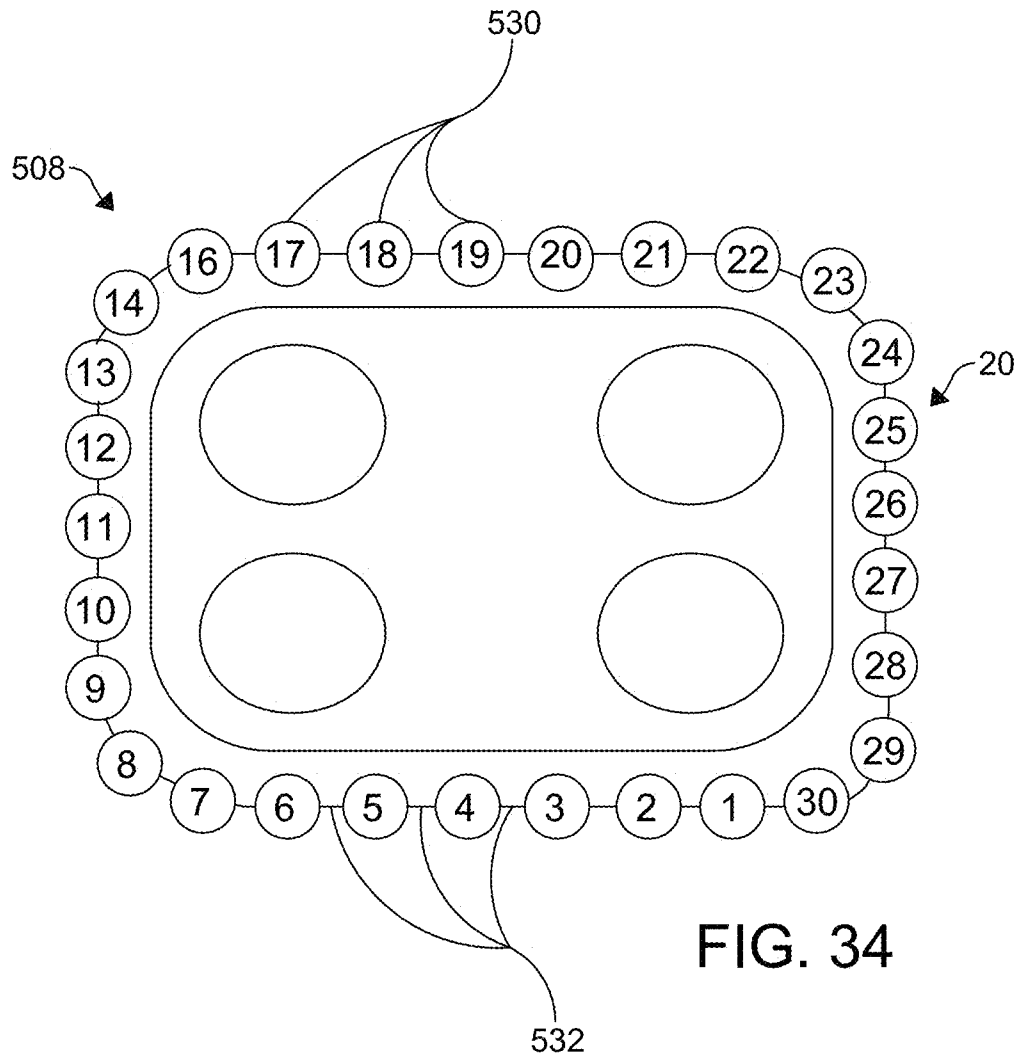


FIG. 34

4000 →

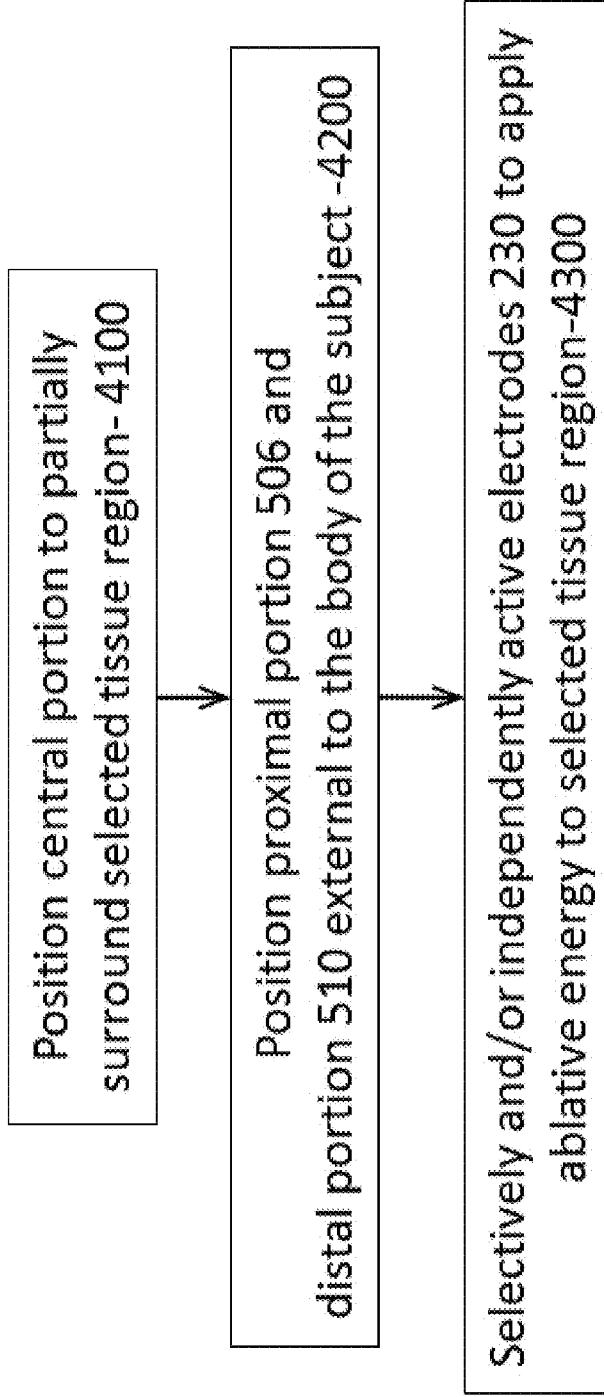


FIG. 35

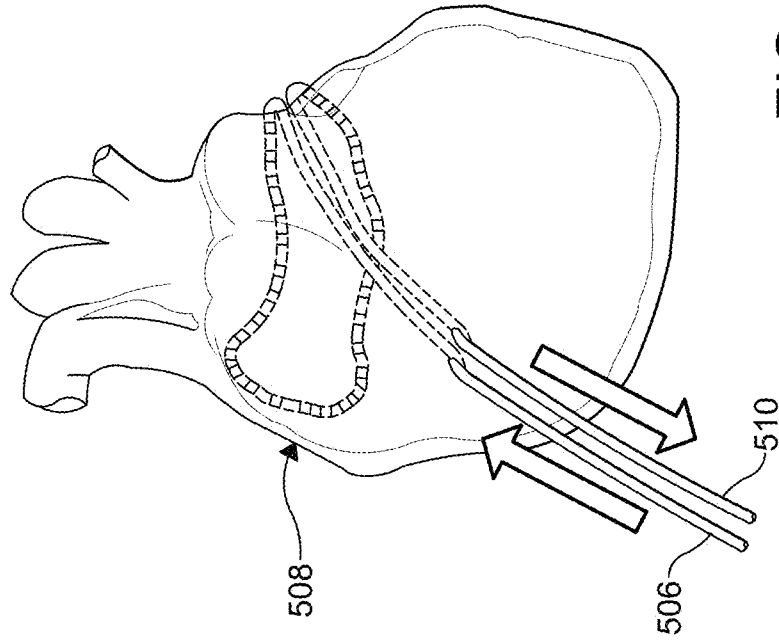


FIG. 37

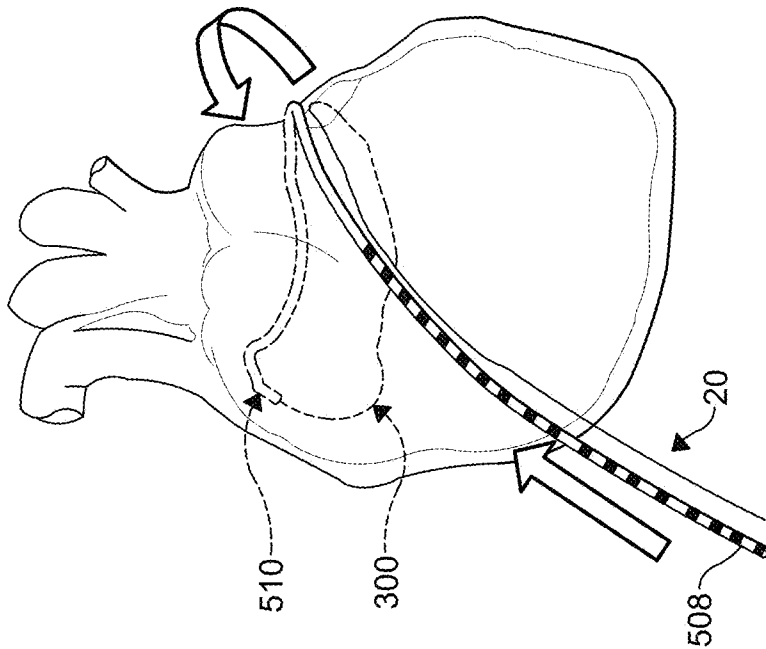


FIG. 36

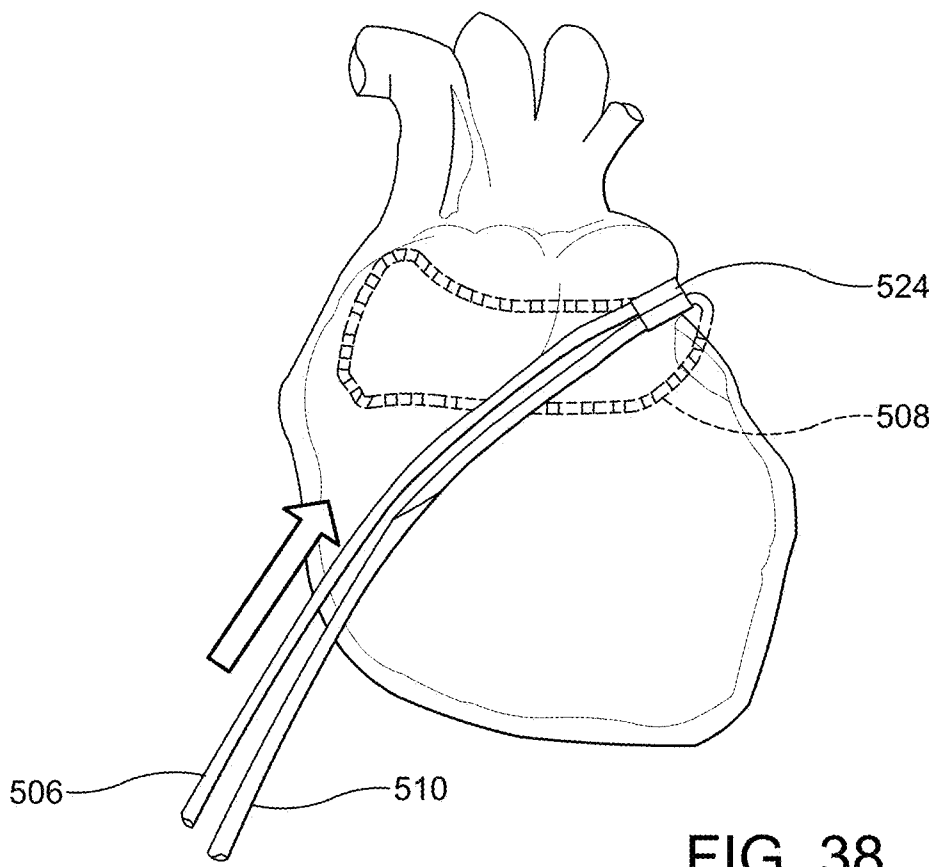


FIG. 38

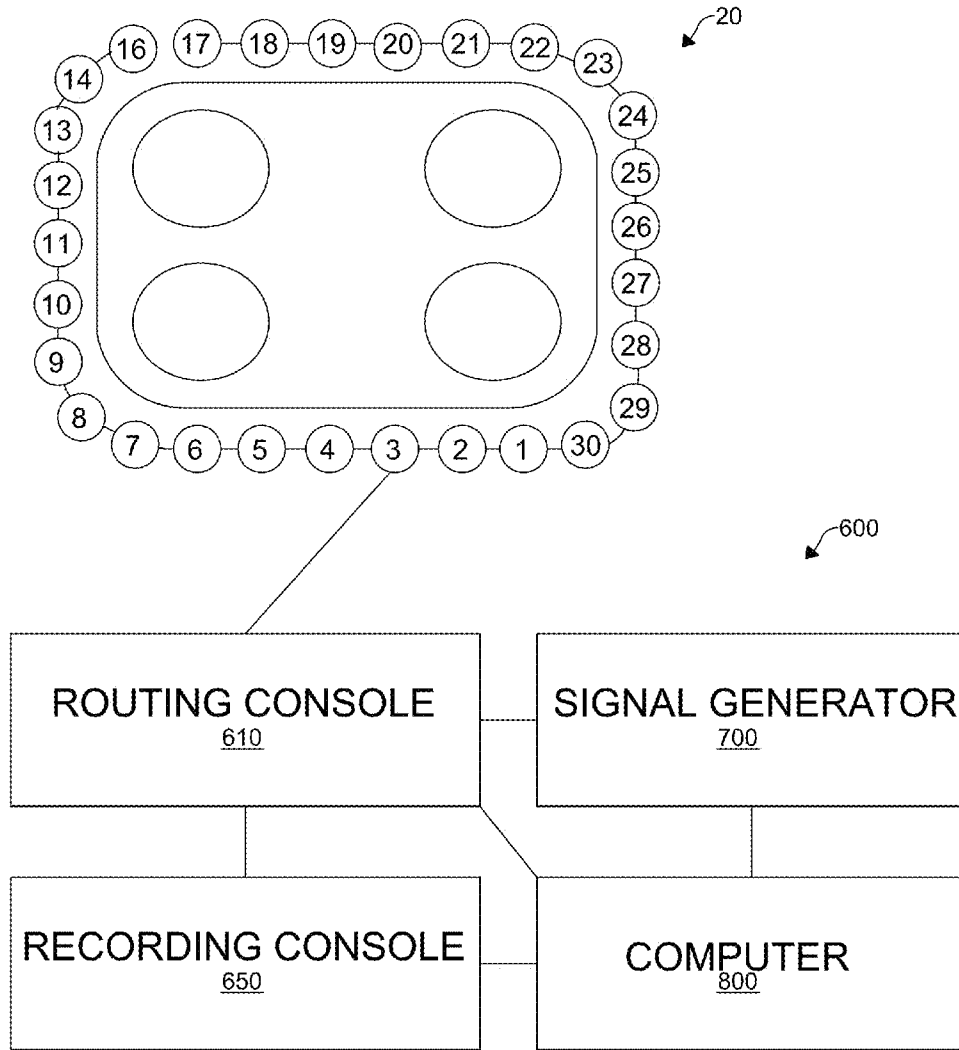


FIG. 39

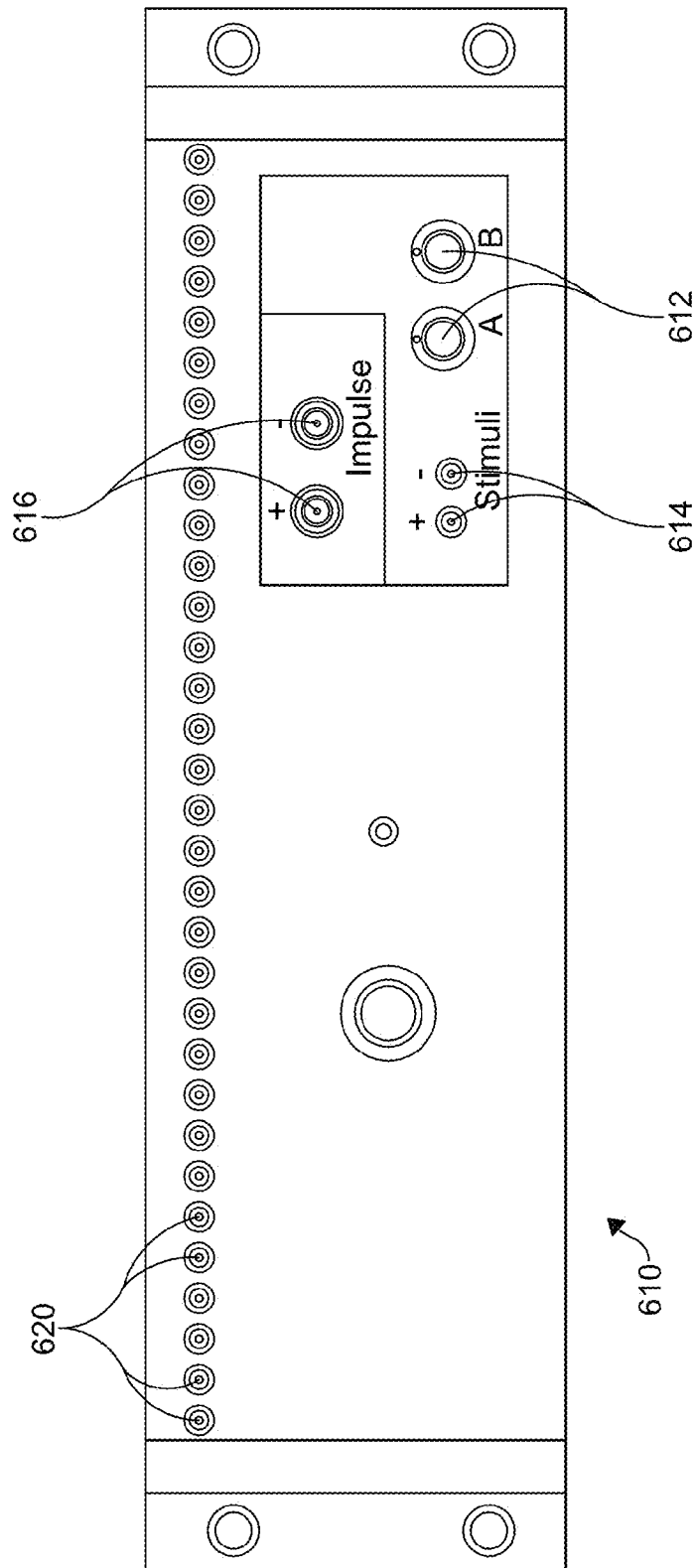


FIG. 40

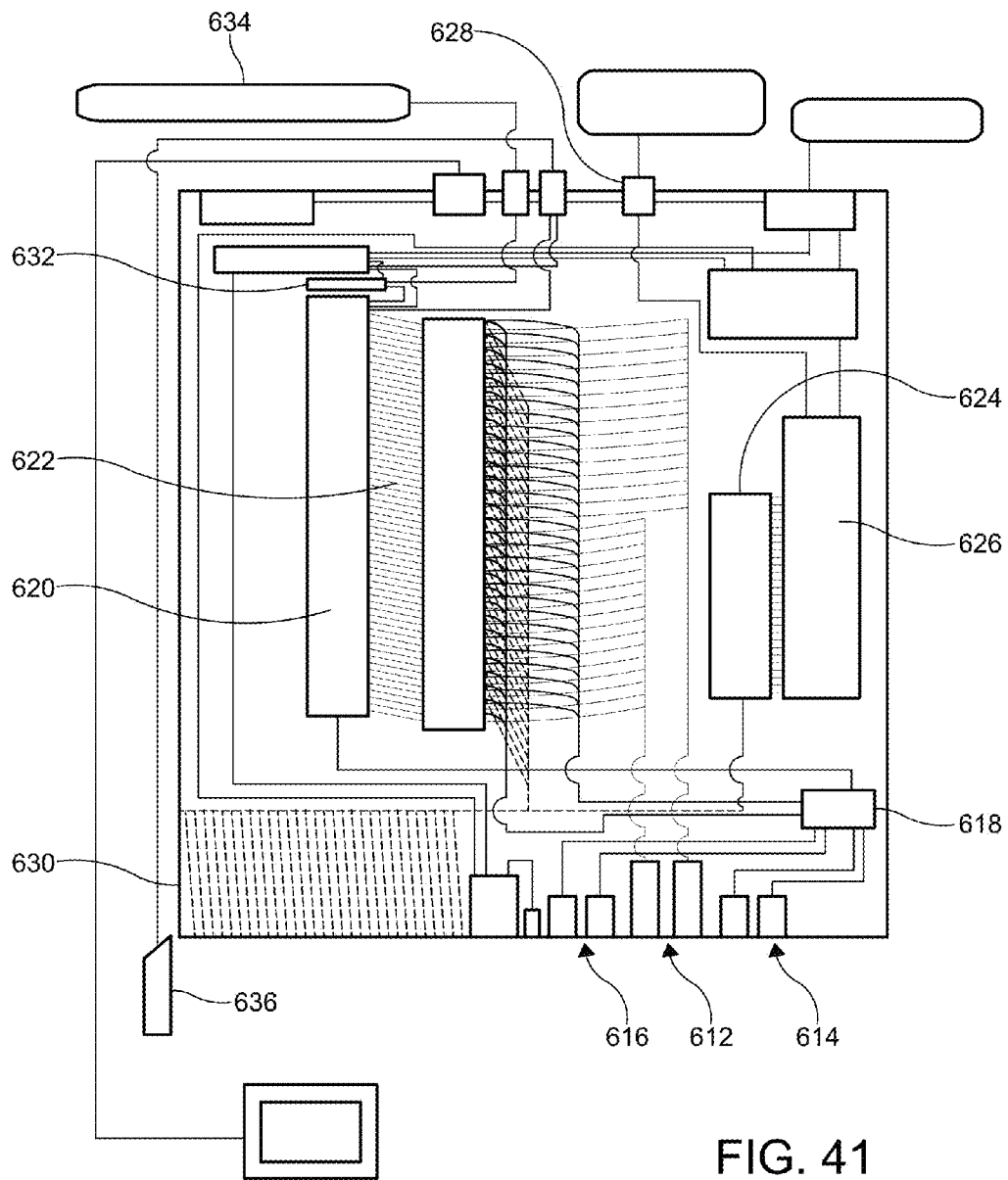


FIG. 41

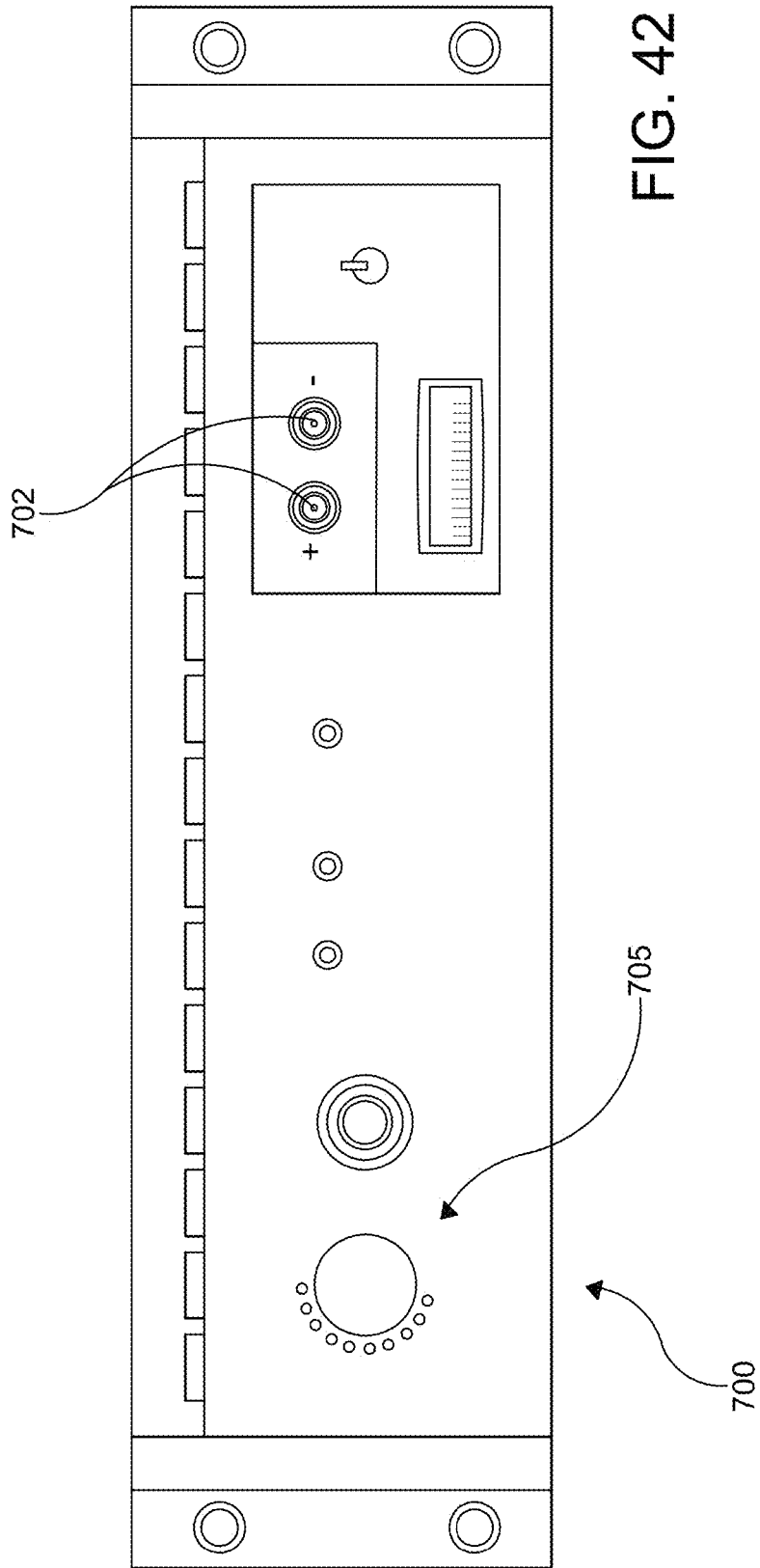


FIG. 42

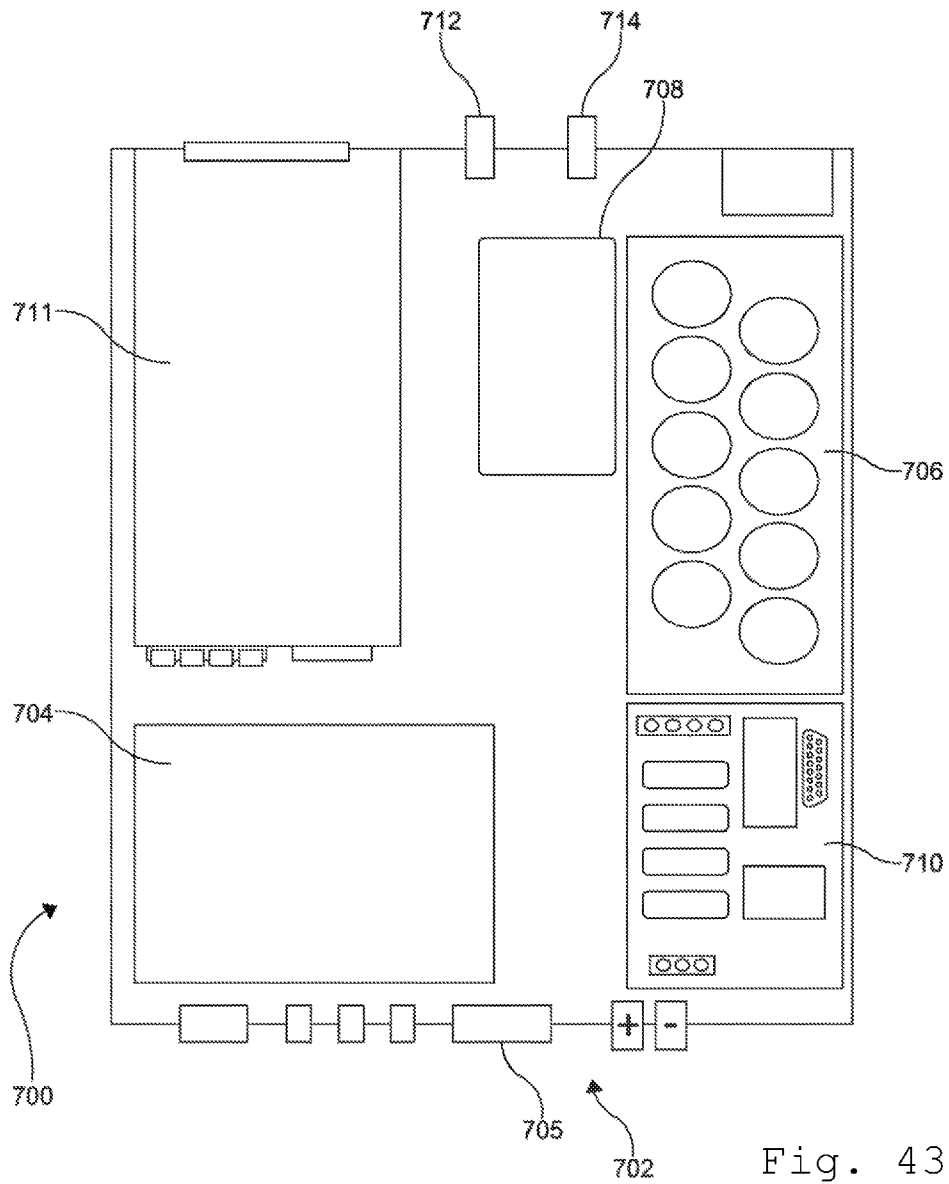


Fig. 43

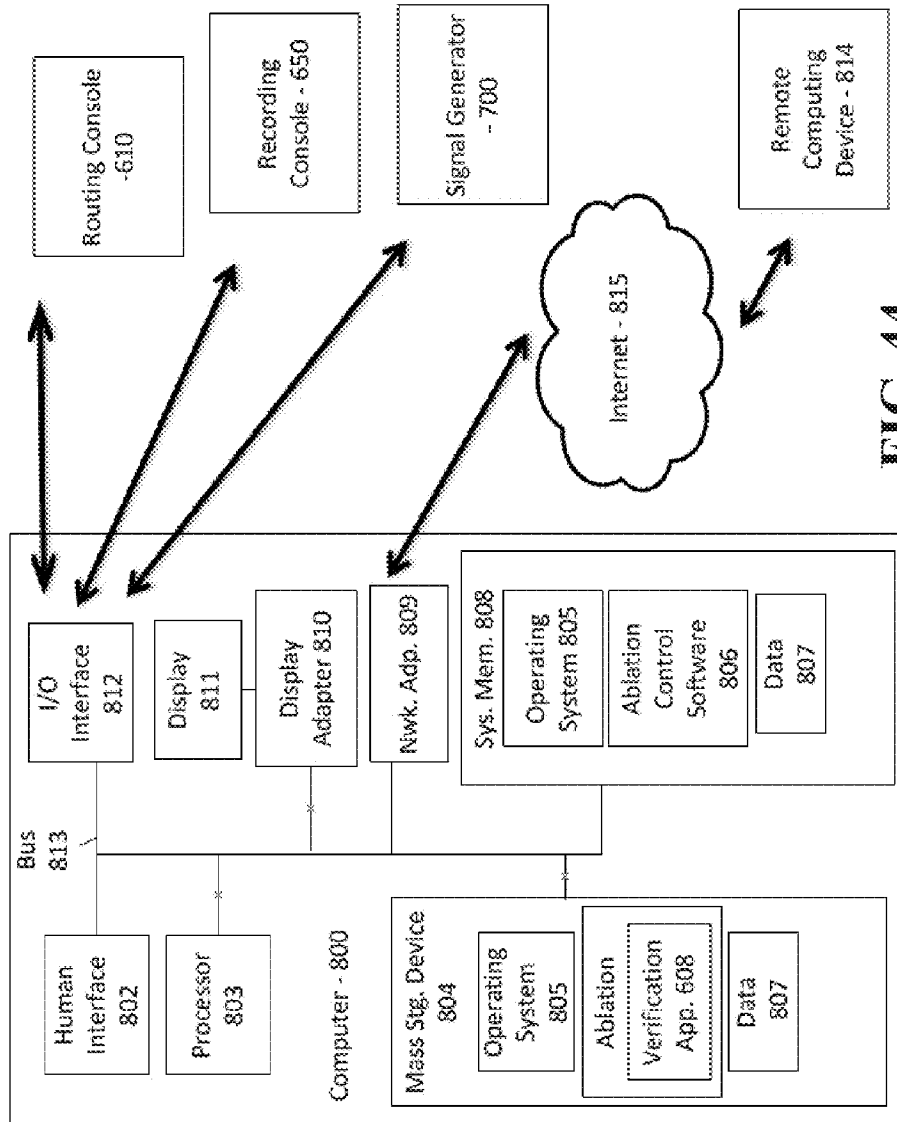


FIG. 44

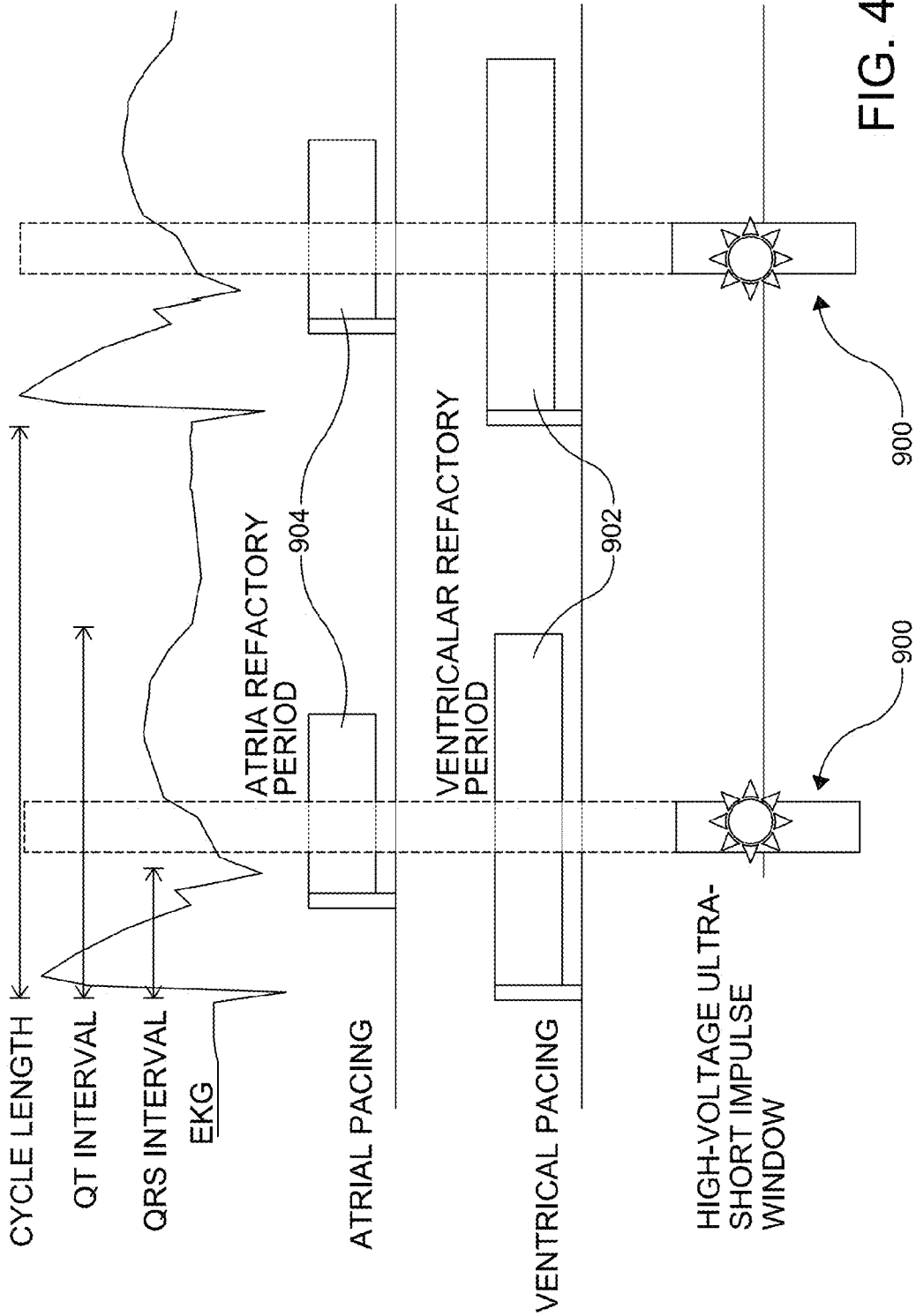


FIG. 45

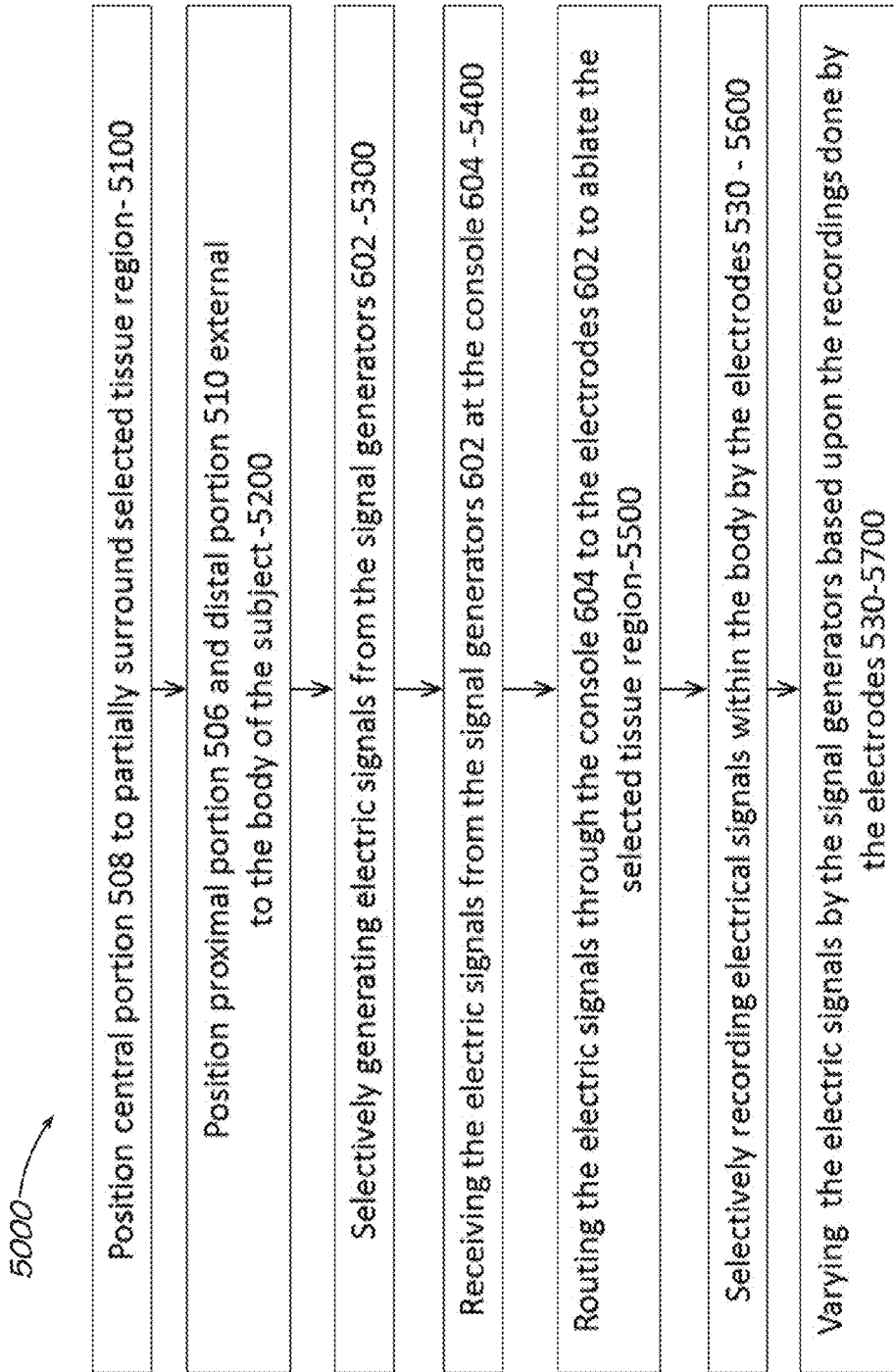


FIG. 46

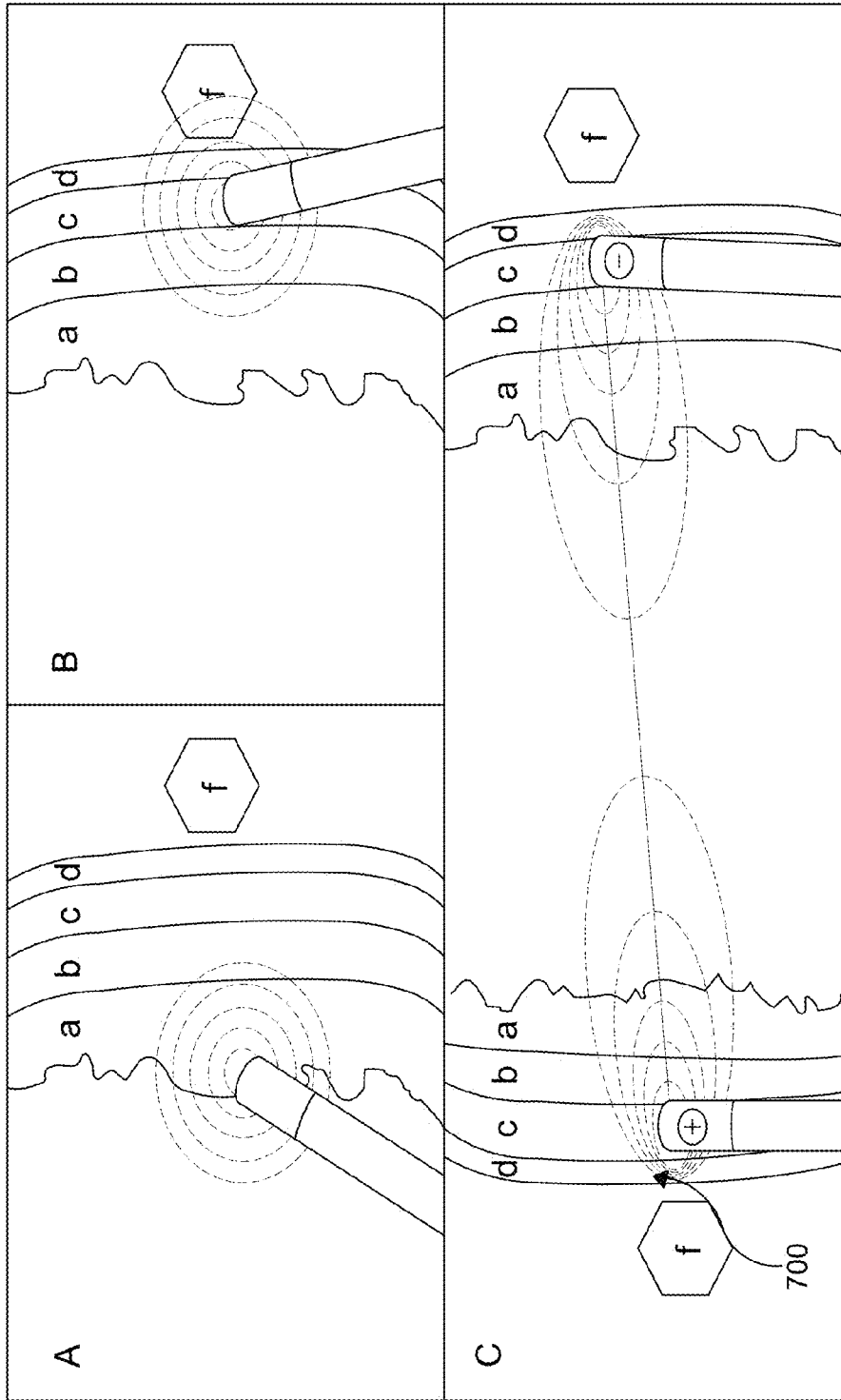


FIG. 47

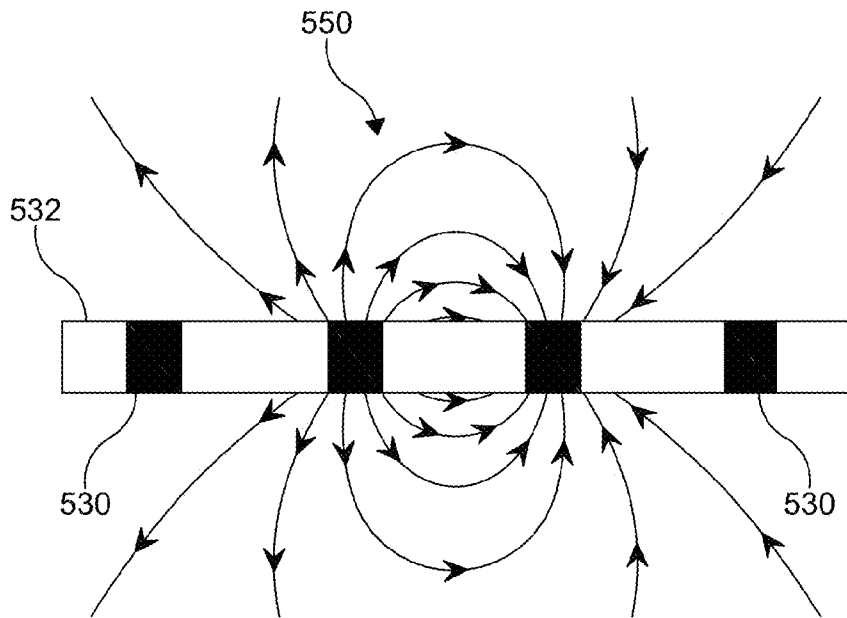


FIG. 48

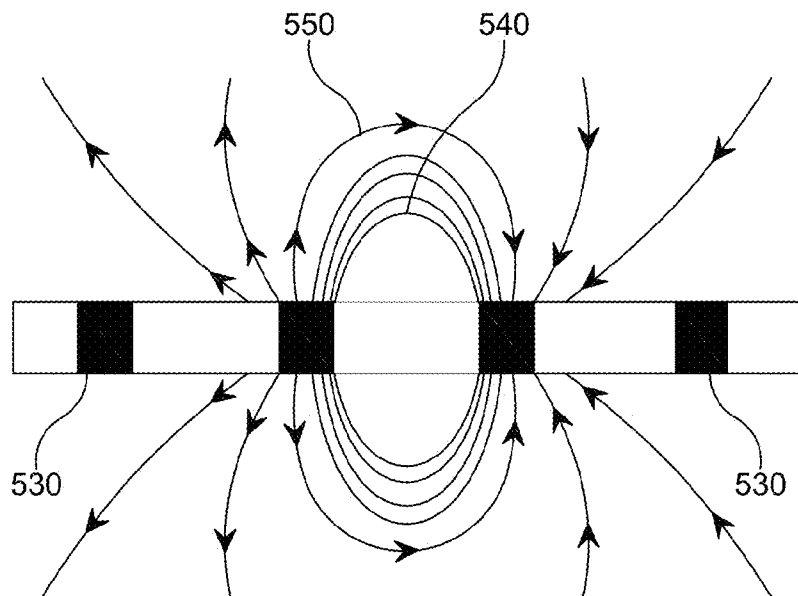


FIG. 49

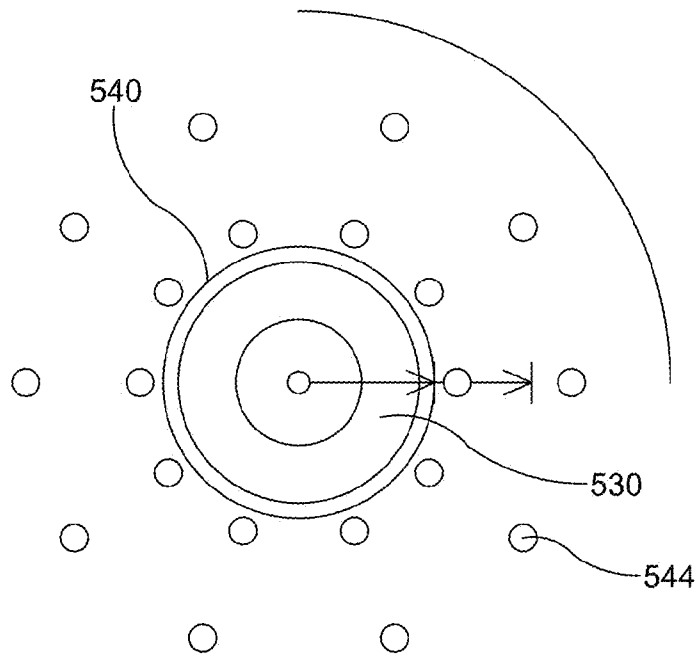


FIG. 50

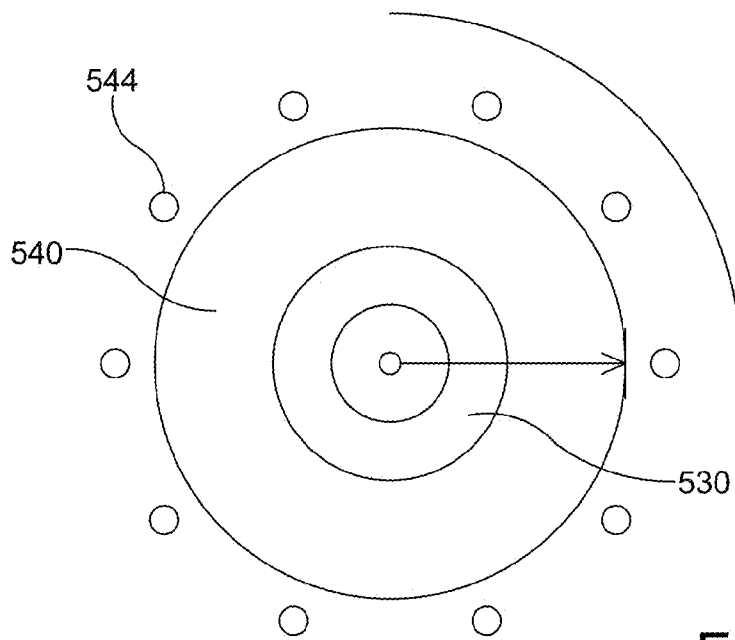
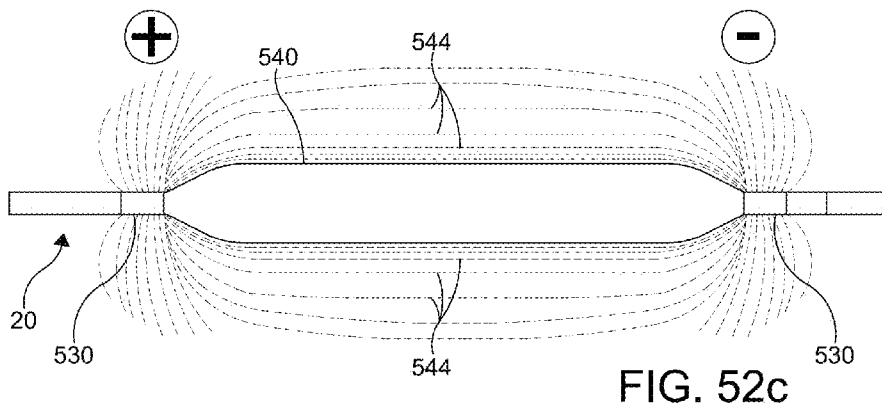
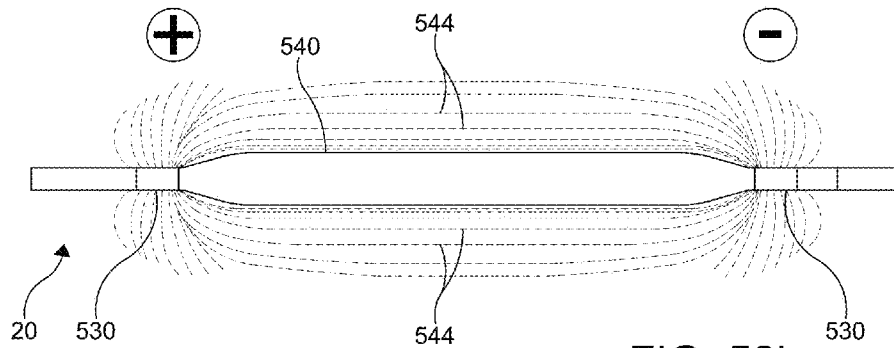
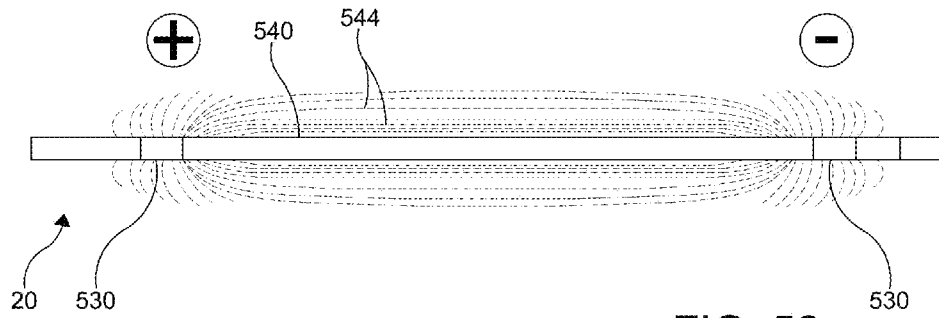
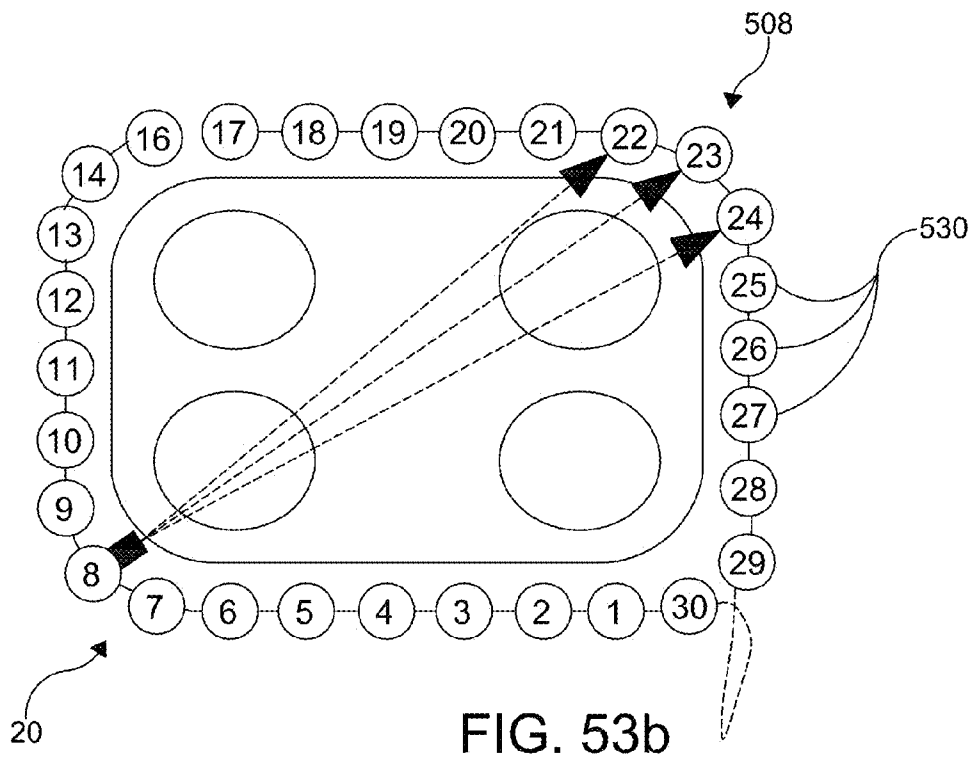
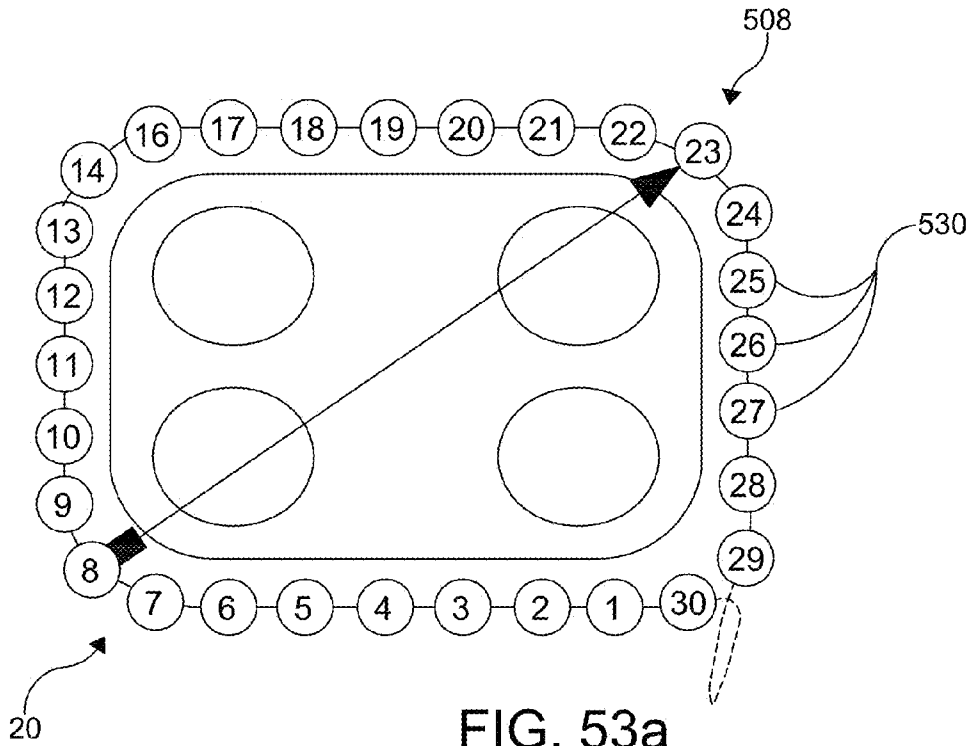


FIG. 51





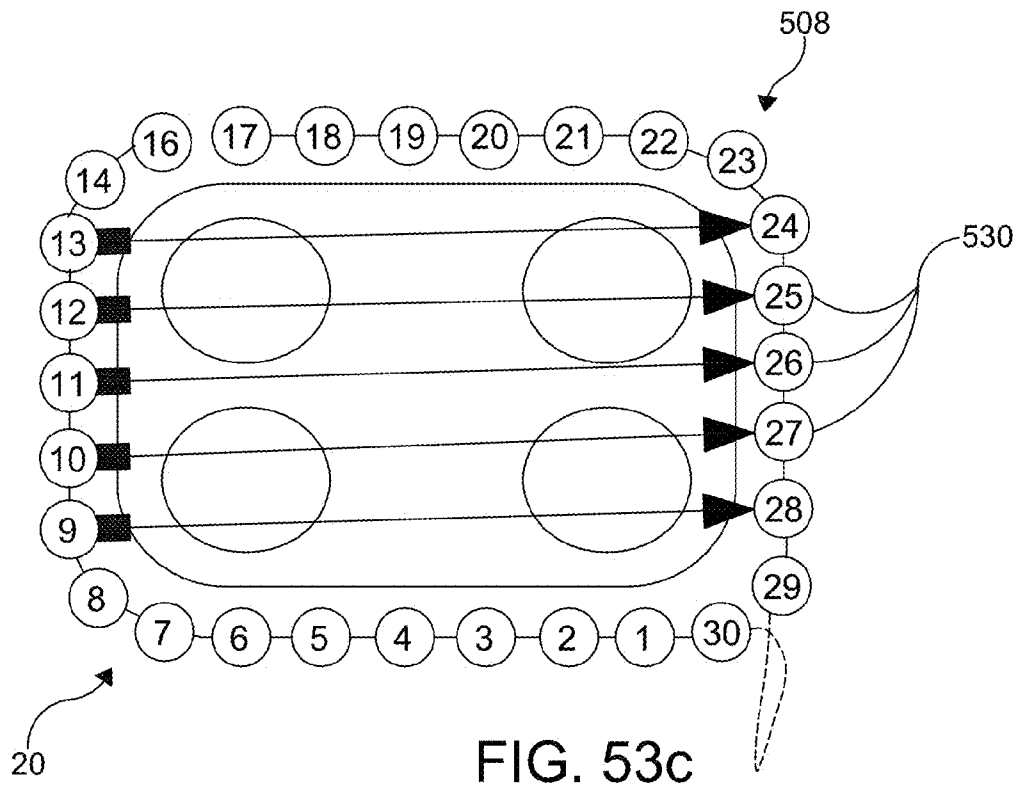


FIG. 53c

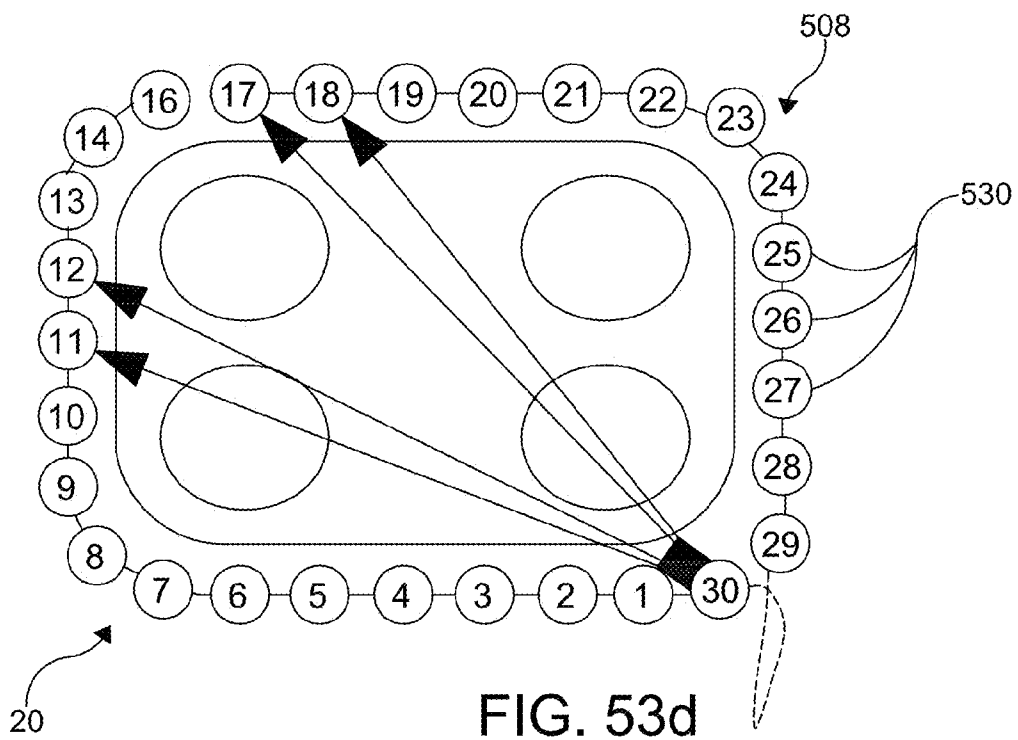


FIG. 53d

**CATHETERS, CATHETER SYSTEMS, AND
METHODS FOR PUNCTURING THROUGH A
TISSUE STRUCTURE AND ABLATING A
TISSUE REGION**

CLAIM OF PRIORITY

This application is a continuation of U.S. patent application Ser. No. 14/400,455, filed on Nov. 11, 2014, which is a National Phase of PCT/US2013/031252, filed on Mar. 14, 2013 and claims benefit of Provisional Patent Application No. 61/681,552, filed on Aug. 9, 2012, which are all relied upon and incorporated herein in their entirety by reference.

FIELD

This invention relates to percutaneous catheter systems and ablation catheters. More particularly, this invention relates to percutaneous catheter systems for puncturing through a tissue structure within the body of a subject and to ablation catheters for ablating a selected tissue region within the body of a subject.

BACKGROUND

Atrial fibrillation can be treated by isolating portions of the atria. Such isolation of the atria can be done by open-heart surgery (e.g., a modified Maze procedure) or, most commonly, by a trans-venous catheter technique. In the majority of cases, the doctor cauterizes the left atrial muscle tissues using radiofrequency ablation techniques, with the ablation lesion targeting and/or circumscribing the pulmonary veins. Isolation of these anatomic portions of atria prevents the electrical propagation of the arrhythmia into the remainder of the atria. The operator places electrophysiologic catheters into the right heart. Under fluoroscopic guidance, a catheter is advanced adjacent to the atrial septum. In most cases, a puncture of the atrial septum (right to left) is made with a specialized needle catheter. A guide-wire is then advanced into the left atrium.

The trans-septal catheter is removed and a guide catheter is delivered over the wire into the left atrium. An ablation catheter is then advanced into the left atrium under fluoroscopic guidance. Typically, electrophysiologists use additional imaging and mapping technology to improve safety and efficacy of the procedure, such as intercardiac ultrasound, cardiac CT, or non-contact mapping systems. Once the ablation/mapping catheters are in the left atrium, the operator delivers radiofrequency energy to the target sites. The operator moves the ablation catheter in a point-by-point fashion connecting the lesions to effectively electrically isolate the pulmonary veins from the rest of the atrium.

These known procedures typically take 3-6 hours to complete. The procedural success varies between operators and patient selection (success rate is between 50-85% for a single attempt). A substantial minority of patients requires subsequent ablation procedures to "touch up" the prior ablation site. The cost of these procedures is highly variable and increases substantially with duration of procedure and the addition of adjuvant imaging/mapping technology. The current procedures are associated with a 5-6% risk of procedural complications, including a $\frac{1}{200}$ risk of stroke due to the need to instrument (i.e., place one or more medical devices into) the left atrium. Other concerning complications include cardiac perforation, tamponade, pulmonary vein stenosis, and atrial-esophageal fistula. Despite attempts to simplify and streamline the procedure, the anatomic

variations of the left atrium and pulmonary veins have limited the utility of alternative ablation techniques.

Known epicardial techniques for atrial fibrillation also have various limitations. For example, most current epicardial ablation strategies require the operator to blindly navigate recesses of the pericardial space with an ablation catheter, and reflections of the pericardial anatomy pose an obstacle to delivery of a single contiguous lesion **30** using these techniques. (See the broken line in FIG. 1.) Thus, the pericardial anatomy greatly limits the efficacy and technical ease of current pericardial/epicardial catheter-based procedures.

Although the membranous reflections of the pericardial space that must be breached are very thin and relatively avascular, the angle, spatial limitations, and relative orientation of the surgical access point to the adjacent pericardial reflections do not facilitate simple puncture with a blunt catheter or a standard needle. Moreover, the large vessel and cardiac chambers adjacent to the pericardial reflections make the proposition of blind puncture with conventional catheters very risky.

Currently known cardiac ablation catheters typically require frequent repositioning and/or advanced noncontact mapping techniques to identify incomplete segments in the ablation lesion. For epicardial techniques performed from the pericardial space, such manipulation is fraught with danger and technical limitations. Standard unipolar applications require an externalized grounding pad that results in a diffuse or spherical virtual electrode. Current bipolar ablation techniques utilize electrode pairs that are in close proximity, require the use of cumbersome equipment, and often require entry into both the pericardium and the left atrial blood pool.

Accordingly, there is a need in the pertinent art for devices, systems, and methods for efficiently and reliably locating and puncturing pericardial reflections. There is a further need in the pertinent art for devices, systems, and methods for delivering a single contiguous lesion within the pericardial space without the need for repositioning of equipment.

SUMMARY

Described herein is a percutaneous catheter system including first and second catheters. Each catheter can include a longitudinal axis, a longitudinal length, a proximal portion, and a distal portion. The distal portion of each catheter defines a distal end of its respective catheter. Each catheter defines at least one lumen extending from an opening of the distal end of the catheter toward the proximal portion of the catheter along the longitudinal length of the catheter. Each catheter has a magnet assembly positioned proximate the distal end of the catheter and operatively coupled to the distal portion of the catheter. Optionally, the magnet assembly of each respective catheter can be permanently and/or fixedly attached to a flexible extension mounted within a lumen of the catheter. The magnet assembly of the first catheter is configured for magnetic coupling to the magnet assembly of the second catheter such that the longitudinal axis of the first catheter is substantially axially aligned with the longitudinal axis of the second catheter. The magnet assemblies of the first and second catheters can be configured for magnetic coupling to one another through a tissue structure, such as, for example, a pericardial reflection.

Methods of puncturing through a tissue structure are also described. In exemplary methods, the percutaneous catheter

system can permit an operator to deliver a guidewire around target structures, thereby facilitating the deployment of an over-the-wire ablation catheter system. The catheter systems provide means for delivering a single isolating lesion around the pulmonary veins using a subxiphoid pericardial access point. The circumscribing lesion can be produced by any currently known energy sources, including radiofrequency (RF), cryoablation, electroporation, microwave, laser, and ultrasound energy sources. However, the circumscribing lesion can also be produced by a non-energetic ablation.

In exemplary methods, extended bipolar application of high voltage ultra short direct current impulses (HVUS-DCI) are used. These impulses produce brief but extremely strong electric fields within the tissue leading to irreversible electroporation (IE), cell death, and injury. However, it should be noted that the total energy applied is relatively low averaging (estimated range 0.025 J to 45 J per pulse). At these energy levels there is very little tissue heating. Thus the mechanism of tissue injury is non-thermal; this is in contrast to RF ablation, which produces thermal tissue ablation through resistive heating.

Also described herein is an ablation catheter for ablating a selected tissue region. The ablation catheter can have a flexible elongate shaft and a plurality of electrodes spaced along a longitudinal length of the flexible elongate shaft. The flexible elongate shaft has a longitudinal axis, a longitudinal length, a proximal portion, a central portion, and a distal portion, with the central portion being positioned between the proximal portion and the distal portion along the longitudinal length of the flexible elongate shaft. The elongate shaft can also define a primary lumen (and, optionally, one or more secondary lumens) of the ablation catheter. The plurality of electrodes can be positioned exclusively within the central portion of the elongate shaft. The electrodes can be separated by high impedance structures. The flexible elongate shaft can be selectively positioned within the body of a subject such that the central portion of the elongate shaft at least partially surrounds the selected tissue region and the proximal and distal portions of the elongate shaft are positioned external to the body of the subject. Upon positioning of the elongate shaft in this manner, each electrode of the plurality of electrodes is configured for selective, independent activation to apply ablative energy to the selected tissue region. Each of the high impedance structures is configured for selective, independent activation to intersect the theoretic field lines created by surrounding electrodes. An ablation catheter system including an ablation catheter, one or more signal generators, and a routing console is also described.

Further described herein are methods of ablating the selected tissue region. In exemplary methods, the ablation catheter can be deployed into the pericardial space with both the proximal and distal portions of the catheter outside the body. The ablation catheter can be more flexible than other clinically available catheter-based ablation devices to thereby permit tissue contact around the left atrial structures. The electrodes of the ablation catheter can be capable of monitoring and/or delivering RF energy, electroporation impulses, and programmed cardiac pacing and/or neuro-stimulus. Unlike other known ablation catheters, the electrodes of the described ablation catheter also can have the capability of delivering extended bipolar high voltage, ultra-short impulses. The feature of individualizing the activation of each extended bipolar electrode can take advantage of the natural geometry inside the pericardial space to deliver energy to a series of electrodes arranged around the target structure and control the vector of the electrical current.

Once the ablation catheter is deployed, a linear lesion can be created without repositioning the catheter, thereby increasing efficiency and effectiveness (when compared to standard point-by-point techniques). This ablation catheter can provide a stable and contiguous array of electrodes along the target path that can deliver ablation and can also be used to confirm electrophysiologic block using an extended bipolar electrocardiographic technique. The ablation catheter takes advantage of the natural contours of the left atrial epicardial surface to provide reliable and stable electrode contact. Additionally, the high-voltage, ultra-short duration impulses used in electroporation techniques do not require that the electrode be in direct contact with the ablation target.

Moreover, the epicardial positioning can have mechanical advantages over endocardial multi-electrode arrays. Indeed, the positioning of the described ablation catheter can be varied with little effort to provide full circumferential coverage around a target structure. The flexibility of the ablation catheter provides a mechanism for ensuring secure tissue contact and/or tissue proximity around complex anatomic geometry. The natural spatial limitation of the pericardial space can provide a natural mechanism to assure electrode approximation. In addition, high impedance structures (e.g., insulators) found along the ablation catheter can change the contour of the current moving between electrodes. Such changes to the contour can lead to an increased current density at the farthest point along the flow of current and the electrodes.

The risks of performing ablation from the epicardial surface can place the electrodes of the ablation catheter closer to some important bystander structures. However, the electrodes of the ablation catheter can be configured to deliver ablative energy with programmed directional vectors. With RF energy, extended bipolar ablation can result in a 40-50% deeper lesion in the direction of the programmed vector. With electroporation, the potential for creating a preferential directional injury vector is greater. In exemplary methods, extended bipolar irreversible electroporation (which cause no thermal injury) can be delivered.

These and other objects and advantages of the invention will become apparent from the following detailed description of the preferred embodiment of the invention.

Both the foregoing general description and the following detailed description are exemplary and explanatory only and are intended to provide further explanation of the invention as claimed. The accompanying drawings are included to provide a further understanding of the invention and are incorporated in and constitute part of this specification, illustrate several embodiments of the invention, and together with the description serve to explain the principles of the invention.

BRIEF DESCRIPTION OF THE FIGURES

These and other features of the preferred embodiments of the invention will become more apparent in the detailed description in which reference is made to the appended drawings wherein:

FIG. 1 depicts the posterior pericardial anatomy with a membranous reflection illustrating a hypothetical lesion delivered to the left atria (note: heart is absent from the illustration).

FIG. 2 is a perspective view of a percutaneous catheter system according to an aspect.

FIG. 3 is a schematic plane view of a percutaneous catheter system according to an aspect.

5

FIG. 4 is a cross sectional view of a catheter of the system of FIG. 3 along line 4-4.

FIGS. 5a-5e are a series of cross sectional views of a portion of the catheter of the system of FIG. 3.

FIG. 6 is a cross sectional view of a portion of a catheter of the system of FIG. 3.

FIGS. 7a-b are cross sectional views of the assembly of a portion of a catheter of system FIG. 3.

FIG. 8 is a cross sectional view of the portion of the catheter of assembled in FIGS. 7a-b.

FIG. 9 is a perspective view of a needle of the percutaneous catheter system of FIG. 2.

FIG. 10 is a schematic plane view of a needle of the percutaneous catheter system of FIG. 3.

FIG. 11 is a cross sectional view of a portion of a catheter of the percutaneous catheter system of FIG. 3.

FIG. 12 is a schematic view of a needle of the percutaneous catheter system of FIG. 3.

FIG. 13 is a schematic plane view of docked catheters of percutaneous catheter system of FIG. 3.

FIG. 14 is a cross-sectional schematic view of the "docked" catheter system of FIG. 13.

FIG. 15 is a depiction of a process to puncture a tissue structure using a percutaneous catheter system according to an aspect.

FIGS. 16-23 are illustrations of the placement and use of a percutaneous catheter system according to an aspect.

FIG. 24 is a depiction of a process to puncture a tissue structure using percutaneous catheter system according to an aspect.

FIG. 25 is a schematic representation of the entry site for the process shown in FIG. 15.

FIG. 26 is a depiction of a process to position a percutaneous catheter system according to an aspect.

FIG. 27 depicts an exemplary ablation catheter according to an aspect.

FIG. 28 is a schematic representation of an ablation catheter according to an aspect.

FIG. 29 is a partial close-up view of a central portion the ablation catheter of FIG. 27.

FIG. 30 is a schematic cross-sectional view of a proximal end of an ablation catheter according to an aspect.

FIG. 31 is a schematic cross-sectional view of a distal end of an ablation catheter according to an aspect.

FIG. 32 is a partial close-up view of the central portion of the ablation catheter of FIG. 27.

FIG. 33 depicts the positioning of an ablation catheter during an exemplary ablation procedure as described herein.

FIG. 34 is a schematic representation of an ablation catheter positioned around the heart according to an aspect.

FIG. 35 is a depiction of a process to position and use an ablation catheter according to an aspect.

FIGS. 36-38 are illustrations of the placement and use of an ablation catheter according to an aspect.

FIG. 39 is a block diagram of an exemplary ablation catheter system according to an aspect.

FIG. 40 is a schematic front plane view of a routing console according to an aspect.

FIG. 41 is a block diagram of a routing console of FIG. 40.

FIG. 42 is a schematic front plane view of a signal generator according to an aspect.

FIG. 43 is a block diagram of a signal generator of FIG. 42.

FIG. 44 is a block diagram of an exemplary computer system according to an aspect.

6

FIG. 45 is an illustration of a graphic representation of a high-voltage impulse window according to an aspect.

FIG. 46 is a depiction of a process to position and use an ablation catheter according to an aspect.

FIG. 47 illustrates schematic representations of epicardial ablation techniques.

FIG. 48 is a schematic representation of an ablation catheter with electrodes according to an aspect.

FIG. 49 is a schematic representation of an ablation catheter with electrodes and a high impedance structure according to an aspect.

FIGS. 50-51 are schematic representations of a cross section of the ablation catheter according to an aspect.

FIGS. 52a-c is a schematic representation of an ablation catheter with electrodes and a high impedance structure according to an aspect.

FIGS. 53a-d display exemplary electrode assignments according to an embodiment.

DETAILED DESCRIPTION

The present invention can be understood more readily by reference to the following detailed description, examples, drawings, and claims, and their previous and following description. However, before the present devices, systems, and/or methods are disclosed and described, it is to be understood that this invention is not limited to the specific devices, systems, and/or methods disclosed unless otherwise specified, and, as such, can, of course, vary. It is also to be understood that the terminology used herein is for the purpose of describing particular aspects only and is not intended to be limiting.

The following description of the invention is provided as an enabling teaching of the invention in its best, currently known embodiment. To this end, those skilled in the relevant art will recognize and appreciate that many changes can be made to the various aspects of the invention described herein, while still obtaining the beneficial results of the present invention. It will also be apparent that some of the desired benefits of the present invention can be obtained by selecting some of the features of the present invention without utilizing other features. Accordingly, those who work in the art will recognize that many modifications and adaptations to the present invention are possible and can even be desirable in certain circumstances and are a part of the present invention. Thus, the following description is provided as illustrative of the principles of the present invention and not in limitation thereof.

As used throughout, the singular forms "a," "an" and "the" include plural referents unless the context clearly dictates otherwise. Thus, for example, reference to "a delivery conduit" can include two or more such delivery conduits unless the context indicates otherwise.

As used herein, the terms "optional" or "optionally" mean that the subsequently described event or circumstance may or may not occur, and that the description includes instances where said event or circumstance occurs and instances where it does not.

The word "or" as used herein means any one member of a particular list and also includes any combination of members of that list.

It is contemplated that the disclosed devices and systems can comprise elements of the devices and systems described in U.S. Pat. No. 6,314,963, the disclosure of which is incorporated herein by reference in their entirety.

It is contemplated that the percutaneous catheter system 10 and ablation catheter 20 of the present invention can

allow an operator to deliver a single isolating lesion around the pulmonary veins of a subject using a subxiphoid pericardial access point. The circumscribing lesion can be produced by any of the currently available energy sources, including, for example and without limitation, HVUS-DCI, RF, cryoablation, electroporation, microwave, laser, biologics, radiation, small molecule chemicals (e.g., ethanol ablation) and ultrasound. However, it is contemplated that the circumscribing lesion can be produced by any ablative energy source. In use, it is contemplated that, once an operator achieves a stable catheter position for the ablation catheter **20**, delivery of a single circumscribing lesion **30** around the pulmonary veins (as shown in FIG. **1**) of the subject can become much simpler. The atrial fibrillation ablation technique described herein can require fewer steps, catheters, time, and equipment than conventional atrial fibrillation ablation techniques. Further, it is contemplated that the described percutaneous catheter system **10** can minimize or avoid the need for expensive advanced mapping and imaging equipment; instead, the described percutaneous catheter system **10** can permit usage of a purely anatomic approach. Consequently, it is contemplated that the described percutaneous catheter system can minimize the expense of atrial fibrillation ablation, thereby making atrial fibrillation ablation to a larger population of patients.

Catheter System for Puncturing Through a Tissue Structure

With reference to FIGS. **2-24**, disclosed herein, is a percutaneous catheter system **10** for use within the body of a subject. In one aspect, the percutaneous catheter system **10** comprises a first catheter **100** and a second catheter **200**. The first catheter **100** can be referred to as the male catheter **100** and the second catheter **200** can be referred to as the female catheter **200**. In this aspect, the first catheter **100** and the second catheter **200** can each have respective longitudinal axes **102**, **202**, longitudinal lengths **104**, **204**, proximal portions **106**, **206**, and distal portions, **108**, **208**. In exemplary aspects, the first and second catheters **100**, **200** can each have a longitudinal length **104**, **204** ranging from about 20 cm to about 50 cm. In another exemplary aspect, the longitudinal length **104**, **204** of the first catheter **100** and the second catheter **200** are approximately the same. While the length of the catheters **100**, **200** in relation to one another is not critical in many aspects, it is important that the catheters **100**, **200** are configured to work as a pair. However, the lengths of the catheters **100**, **200** collectively need to have a combined length that is long enough to reach the key areas of the anatomy for which the catheter system **10** is being used. In these aspects, it is contemplated, following magnetic coupling between the first catheter **100** and the second catheter **200**, the total length of the first catheter **100** and the second catheter **200** can range from about 40 cm to about 100 cm.

In other exemplary aspects, at least one of the first catheter **100** and the second catheter **200** can be flexible. In other exemplary aspects, both the first catheter **100** and the second catheter **200** can be flexible. The catheters **100**, **200** should be comprised of a material that is also kink resistant. In an aspect, the catheters **100**, **200** can be comprised of kink resistant material such as expanded PTFE and/or more standard biocompatible materials (coil reinforced silicon, PFA, Pebax, and/or PVC). The construction can utilize expanded PTFE with progressively decreasing density distally, however other construction techniques could be employed. The stiffer proximal segment provides necessary column strength and transmission of torsional force for navigation. In an aspect, the distal portions **106** (which can range between 10-20 cm) are more flexible to permit a-tra-

umatic manipulation and navigation by over the wire techniques through tortuous anatomy. In some embodiments, in order to prevent kinking, braided reinforcement, as well as other types of reinforcement, can be utilized.

In an exemplary aspect, the first and second catheters **100**, **200** are configured to be flexible enough so that the catheters **100**, **200** can permit a 180° turn around a 1.5 cm obstacle. However, the catheters **100**, **200** can be made to perform to other standards (e.g., perform 180° turns around various sized obstacles) in other exemplary embodiments.

In another aspect, the distal portion **108** of the first catheter **100** can define a distal end **110** of the first catheter **100**. In an aspect, the distal end **110** can have a nominal outer diameter between 1 mm to 5 mm to accommodate a magnet assembly **120**. In this aspect, the distal end **110** of the first catheter **100** can define an opening **112**. In an aspect, the end of the proximal portion **106** is configured to be larger than the distal end **110** in order to facilitate the manipulation of the catheter **100** at the handle **140**, discussed in more detail below.

In an additional aspect, the first catheter **100** can define at least one lumen **116**, **118** extending from the opening **112** of the distal end **108** toward the proximal portion **106** of the first catheter **100** along at least a portion of the longitudinal length **104** of the first catheter **100**. The lumen can be defined by an outer shaft **115** of the catheter **100**. In a further aspect, the first catheter **100** can comprise a first magnet assembly **120** positioned proximate the distal end **110** of the first catheter **100** and operatively coupled to the distal portion **108** of the first catheter **100**.

In another aspect, the distal portion **208** of the second catheter **200** can define a distal end **210** of the second catheter **200**. In an aspect, the distal end **210** can have a nominal outer diameter between 1 mm to 5 mm to accommodate a magnet assembly **220**. In an aspect, the distal end **210** of the second catheter **200** can define an opening **212**. In an aspect, the end of the proximal portion **206** is configured to be larger than the distal end **210** in order to facilitate the manipulation of the second catheter **200** through the use of a handle **240**, discussed in more detail below.

In an additional aspect, the second catheter **200** can define at least one lumen **216**, **218** extending from the opening **212** of the distal end **210** toward the proximal portion **206** of the second catheter **200** along at least a portion of the longitudinal length **204** of the second catheter **200**. The lumen **216**, **218** can be defined by an outer shaft **215** of the second catheter **200**. In a further aspect, the second catheter **200** can comprise a second magnet assembly **220** positioned proximate the distal end **210** of the second catheter **200** and operatively coupled to the distal portion **208** of the second catheter **200**.

In an exemplary aspect, the first catheter **100** and the second catheter **200** can have a nominal outer diameter of 1 to 5 mm and in other respects the geometry of catheter **100** and **200** will be similar to provide a symmetric and complementary magnetic coupling surface for the magnet assemblies **120**, **220**. However, in other aspects, the outer diameter of the catheters **100**, **200** can vary. In an exemplary aspect, the first and second catheters **100**, **200** can have an inner diameter configured to accommodate a needle tube **130** discussed in more details below. In an exemplary aspect, inner diameter of the first and second catheters **100**, **200** can be configured to accommodate a needle tube **130** of approximately 1.473 mm in diameter. However, in other aspects, the inner diameter of the catheters **100**, **200**, as well as the diameter of the needle tube **130**, can vary. In other aspects,

when magnetic coupling and guide wire transfer are the only desired functions, the catheters **100/200** may not have a needle component.

In an exemplary aspect, the first magnetic assembly **120** of the first catheter **100** is configured for magnetic coupling to the second magnet assembly **220** of the second catheter **200**. In this aspect, it is contemplated that the first magnetic assembly **120** can be configured for magnetic coupling to the second magnet assembly **220** such that the longitudinal axis **102** of the first catheter **100** is substantially axially aligned with the longitudinal axis **202** of the second catheter **200**. It is further contemplated that the first magnet assembly **120** can be configured for magnetic coupling to the second magnet assembly **220** through a tissue structure within the body of the subject, discussed further below.

It is contemplated that the at least one lumen of the first catheter **100** can comprise a primary lumen **116**. Similarly, it is contemplated that the at least one lumen of the second catheter **200** can comprise a primary lumen **216**. Optionally, in another exemplary aspect, the at least one lumen of the first catheter **100** can further comprise one or more auxiliary lumens **118**. Similarly, it is contemplated that the at least one lumen of the second catheter **200** optionally can further comprise one or more auxiliary lumens **218**. In an aspect, the primary lumen **116**, **216** and the auxiliary lumen **118**, **218** can be separate by an inner shaft **117**, **217** in each catheter **100**, **200**, with the primary lumen **116**, **216** being contained within the inner shaft **117**, **217**, and the auxiliary lumen **118**, **218** being contained between the inner shaft **117**, **217** and the outer shaft **115**, **215**. The primary lumen **116**, **216** can be configured to receive the needle tube **130**. In some aspects, the inner shaft **117**, **217** can move up and down the outer shaft **115**, **215** of the catheters **100**, **200** respectively.

Optionally, it is contemplated that the one or more auxiliary lumens **118** of the first catheter **100** can be configured for delivery of one or more fluids to the opening **112** of the distal end **110** of the first catheter **100**, while the one or more auxiliary lumens **218** of the second catheter **200** can be configured for delivery of one or more fluids to the opening **212** of the distal end **210** of the second catheter **200**. Optionally, it is further contemplated that the one or more auxiliary lumens **118** of the first catheter **100** can be configured for application of suction to the opening **112** of the distal end **110** of the first catheter **100**, while the one or more auxiliary lumens **218** of the second catheter **200** can be configured for application of suction to the opening **212** of the distal end **210** of the second catheter **200**.

In another aspect, the auxiliary lumens **118**, **218** can perform the delivery of fluids and the application of suction through irrigation ports/side openings/side holes **119**, **219** approximate the openings **112**, **212** of the distal ends **110**, **210** of the catheters **100**, **200**. In one optional exemplary aspect, the at least one lumen of the first catheter **100** and/or second catheter **200** can comprise a primary lumen **116**, **216** and an auxiliary lumen **118**, **218**, with the auxiliary lumen **118**, **218** radially surrounding the primary lumen **116**, **216**.

In one aspect, the first catheter **100** can further comprise a needle **130** operatively positioned within the primary lumen **116** of the first catheter **100**, as shown in FIGS. **5e**, **9-12** and **14**. The needle **130** can further comprise a flexible tubular needle **130**. In an exemplary aspect, the flexible tubular needle **130** can comprise a modified hypodermic needle spirally cut circumferentially around a shaft **132** of the needle **130**. The needle **130** can have a progressive pitch to the coil providing increasing flexibility at a distal tip **134**. The needle **130** can be made of materials that include, but are not limited to, metal, plastic, or other suitable com-

pounds. In an aspect, the needle **130** can be a composite with a coating to improve mechanical and/or functional characteristics (examples include, but are not limited to, a lubricious polymer, insulator, electrical components, and/or bio-compatible metals). A proximal portion of the needle **130** can connect to a mounting hub, the inner shaft **117**, and/or other elements to provide a method of fixation within the catheter **100** and/or a deployment mechanism **146** in the catheter handle **140**. In an exemplary aspect, the needle **130** is mounted to the inner shaft **117** of the first catheter **100**. In other aspects, the needle **130** can extend the length of the catheter **100**. In additional aspects, the needle **130** can be connected to the inner wall of the outer shaft **115** of the catheter **100**.

In an exemplary embodiment, the tubular needle **130** can have a flexibility to accommodate a 1.5 cm turn radius. However, in other aspects, the flexibility of the needle **130** can vary depending on the needs of the application. In one exemplary aspect, it is contemplated that the needle **130** of the first catheter **100** can have a distal puncturing surface **134** and be configured for selective axial movement relative to the longitudinal axis **102** of the first catheter **100**.

In an aspect, the distal tip **134** is configured to serve as a puncturing surfaces **134**. In an exemplary aspect, the puncturing surface **134** can be flared at a 45° angle and OD 2.5 mm. However, in other aspects, the puncturing surface **134** can be configured differently. It is still further contemplated that the distal puncturing surface **134** of the needle **130** of the first catheter **100** can be configured to puncture through a tissue structure within the body of the subject positioned between the distal ends **110**, **210** of the first and second catheters **100**, **200** respectively when the ends **110**, **210** are magnetically coupled, discussed below.

Optionally, in one aspect, the needle **130** of the first catheter **100** can be retractably secured within the primary lumen **116** of the first catheter **100**. In this aspect, the needle **130** of the first catheter **100** can define a delivery lumen **138**. In this aspect, the delivery lumen **138** of the needle **130** of the first catheter **100** can be configured to receive a guide wire **300** (shown in FIG. **3**). The guide wire **300** can be utilized before and after the placement of the catheters **100**, **200**. In this aspect, upon receipt of at least a portion **134** of the needle **130** of the first catheter **100** within the opening **212** of the distal end **210** of the second catheter **200** (as shown in FIG. **14**), the delivery lumen **138** of the needle **130** of the first catheter **100** can be configured to permit transfer of a guide wire **300** from the first catheter **100** to the second catheter **200**. **116**

In an aspect, as illustrated in FIGS. **3** and **6**, the handles **140**, **240** are found approximate the proximal ends **106**, **206** of the catheters **100**, **200**. The handles **140**, **240** can be made of a rigid material, such as, but not limited to, machined aluminum, carbon fiber, and the like. The handles **140**, **240** provide the means of manual manipulation of the catheters **100**, **200** when in use. The handles **140**, **240** provide a place to apply force to advance, withdrawal, and apply rotational torsion to catheters **100**, **200**.

As shown in FIG. **6**, the handle **140** of the male catheter **100** (i.e., the catheter **100** operating the needle **130**) can include a proximal chamber **142** and a distal chamber **144**. In an aspect, the proximal chamber **142** can contain a stylus/integrated lever **146** that is connected to the inner shaft **117** of the catheter **100**. The stylus/integrated lever **146** allows for the independent manipulation of the needle **130** within the outer shaft **115** of the catheter **100**. In an aspect, the stylus **146** allows for the independent manipulation of the inner catheter **117** to manipulate the needle **130** within

the outer shaft **115** of the catheter **100**. In a further aspect the control of the inner shaft **117** by the integrated lever **146** provides a means to transmit force distally and deploy the needle **130** through the central bore **122** of the magnetic assembly **120**. The stylus/integrated lever **146** can include a compression spring **148** that ensures that the needle **130** is not deployed until actually called on by the user. In an aspect, the spring **148** prevents the stylus/integrated lever **146** from the inner shaft **117** from deploying the needle until called upon.

In an aspect, the integrated lever **146** includes a rigid tube **150** connected to the proximal end of the spring **148**. The rigid tube **150** is hollow, and allows passage of the guidewire **300** and other components to the distal end **110** of the catheter **100**. A projection **152** extends from the rigid tube **150** through a slot **154** found on the outer portion of the handle **140**. The projection **152** allows the user to activate the integrated lever/stylus **146**, compressing the spring **148** and pushing the needle **130** distally along the catheter **100**. Lastly, the handle **140** can include a guidewire entry point **156**. In an aspect, the inner shaft **117** passes through a fluid hub **168** found in the distal chamber **144**.

In an aspect, the handle **240** of the female catheter **200** can include all of the same components of as described above for the male catheter **100**, but it is not necessary. For example, when a female catheter **200** is used that does not employ a needle **230**, the handle **240** does not need to have an integrated lever and the associated components to control the needle and inner shaft **217**. In another aspect, the catheter pair **100/200** can be constructed without an inner needle **130/230**, and be equipped to form magnetic coupling with central lumen for the passage of a guide wire. In other aspects, the female catheter **200** can have a proximal chamber **242** and a distal chamber **244**, with the proximal chamber **242** providing a guidewire entry point **256** to receive a guide wire **300** to pass through to the primary lumen **216** and the distal chamber **244** including a fluid hub **268**.

In an aspect, the handles **140, 240** can include a hemostasis/fluid management system. The fluid management systems include proximal valves (not shown) that prevent unwanted fluid leakage through the primary lumens **116, 216** of the respective male catheter **100** and female catheter **200**. In addition, the proximal valves prevent the introduction of unwanted air through the centers lumen **116, 216**. In an aspect, a second fluid valve (**166** in FIG. **6**) can be used to provide a seal of the auxiliary lumens **118, 218**. Both the first and second fluid valves can include silicon o-rings and various other seal-creating mechanisms.

Fluid hubs **168, 268** can be found within the handles **140, 240** near the proximal ends **106, 206** of the male catheter **100** and female catheter **200** respectively. The fluid hub **168, 268** of each catheter **100, 200** can be in communication with their respective auxiliary lumen **118, 218**. Fluid ports **170, 270** provide access to the fluid hubs **168, 268**. In an aspect, the combination of the fluid ports **170, 270**, fluid hubs **168, 268**, auxiliary lumen **118, 218** and side openings **119, 219** create the fluid management system. The fluid management system provides for the delivery of radio contrast agents for intra-pericardial navigation under x-ray fluoroscopic guidance. In addition, the fluid management systems provide a means to inject and suck moderate volumes of fluid through the lumen **118, 218** quickly. This is specifically used to inject and withdraw radio contrast agents and/or other fluids (including but not limited to saline, medications, etc.) within the pericardial space; thus accentuating anatomic boundar-

ies. The system, through the side openings **119, 219** can also be used to manage and/or drain a pericardial effusion.

In another aspect, it is contemplated that the first magnet assembly **120** of the first catheter **100** can be positioned within the primary lumen **116** of the first catheter **100**, as shown in FIG. **5**. In this aspect, it is further contemplated that the second magnet assembly **220** can be positioned within the primary lumen **216** of the second catheter **200**. It is still further contemplated that the first magnet assembly **120** of the first catheter **100** can define a central bore **122** configured to receive the needle **130** of the first catheter **100**. Similarly, it is contemplated that the second magnet assembly **220** of the second catheter **200** can define a central bore **222** configured to receive the needle **130** of the first catheter **100**.

In an aspect, as shown in FIGS. **7a-b** and **8**, the magnet assemblies **120, 220** can be coupled to the distal ends **110, 210** of respective catheters **100, 200** through the use of a flexible needle guide **124, 224**. The flexible needle guides **124, 224** include a distal portion **125, 225** and a proximal portion **126, 226**. The flexible needle guides **124, 224** can include central lumen **127, 227** that extend the length of the guides **124, 224** and are configured to receive the needle **130, 230**. The distal portions **125, 225** of the needle guides **124, 224** are secured within central bores **122, 222** of the magnet assemblies **120, 220**, with the proximal portions being secured within the primary lumens **116, 216** at the distal portions **108, 208** of the catheters **100, 200**. The needle guide **124, 224** can be attached coaxially through adhesive or by mounting over a thin walled rigid tube that has been affixed to the magnetic assembly and extends proximally from the magnet **120, 220**.

The needle guides **124, 224** provide a means to maintain central alignment of the inner and outer shafts of the catheters **100, 200** while allowing independent degrees of lengthwise movement. In an aspect, the flexible needle guides **124, 224** can provide a way to introduce a fixed and/or adjustable angle at the distal ends **110, 210** of the catheters **100, 200**. In the cases where the distal portions **110, 210** and magnet assemblies **120, 220** of the catheters **100, 200** meet curved portions, the flexible needle guide **124, 224** provides a flexible curved angle between the most distal portion **125, 225** and proximal portions **126, 226**, as shown in FIG. **8**. Further, the guides **124, 224** prevent the needle **130** from exiting the opening **112, 212** when the distal end **110, 210** encounters a curve, preventing accidental punctures. In an aspect, a rigid tube guide **124, 224** can be utilized. In such an aspect, the segment of the needle guide **124, 224** extending proximally from the magnet may be aligned with the long axis **102, 202** of the inner lumen **116, 216** or the rigid component may bend providing a means to introduce a fixed curve into the tip of the assembled catheter. The variations in performance requirements and mounting techniques will influence magnet assembly **120, 220** and needle guide **124, 224** dimensions and shape.

It is still contemplated that the first magnet assembly **120** can have a distal surface **128** substantially flush with the distal end **110** of the first catheter **100**. Similarly, it is contemplated that the second magnet assembly **220** of the second catheter **200** can have a distal surface **228** substantially flush with the distal end **210** of the second catheter **200**. In exemplary aspects, the first magnet assembly **120** can be permanently fixedly secured to the first catheter **100**. Similarly, it is contemplated that the second magnet assembly **220** can be permanently fixedly secured to the second catheter **200**. However, in other aspects, the first and second

magnet assemblies can be removably coupled to the first and second catheters **100**, **200** respectively.

In an aspect, the magnet assembly **120** of the first catheter **100** and the magnet assembly **220** of the second catheter **200** are configured to be magnetically attracted to one another. In an exemplary aspect, it is desired that the magnet assemblies **120**, **220** are strong enough to automatically magnetically couple to one another when the magnet assemblies **120**, **220** come within approximately 1 cm of each other. In the exemplary catheter we found magnetic field strength between 0.5 kG to 1.5 kG was ample to provide the desired coupling characteristics. However, in all aspects, the strength of the magnetic attraction has to be strong enough to magnetically couple the magnet assemblies **120**, **220** and hold them together magnetically on opposite sides of human tissue. In an aspect, the magnetic attraction can occur automatically. In another aspect, the magnetic attraction between the two magnet assemblies **120**, **220** can be manually controlled.

It is contemplated that, upon magnetic coupling between the first magnet assembly **120** of the first catheter **100** and the second magnet **220** assembly of the second catheter **200** such that the longitudinal axis **102** of the first catheter **100** is substantially axially aligned with the longitudinal axis **202** of the second catheter **200**, the needle **130** can be configured for axial movement relative to the longitudinal axis **102** of the first catheter **100** such that at least a portion **134** of the needle **130** exits the opening **112** of the distal end **110** of the first catheter **100** and is received within the opening **212** of the distal end **210** of the second catheter **200**.

Similarly, in another optional aspect, the second catheter **200** can further comprise a needle **230** operatively positioned within the primary lumen **216** of the second catheter **200**. In this aspect, the needle **230** of the second catheter **200** can be configured for selective axial movement relative to the longitudinal axis **202** of the second catheter **200**. It is further contemplated that, upon magnetic coupling between the magnet assemblies **120**, **220** of the first and second catheters **100**, **200** such that the longitudinal axis **102** of the first catheter **100** is substantially axially aligned with the longitudinal axis **202** of the second catheter **200**, the needle **230** of the second catheter **200** can be configured for axial movement relative to the longitudinal axis **202** of the second catheter **200** such that at least a portion **232** of the needle **230** exits the opening **212** of the distal end **210** of the second catheter **200** and is received within the opening **212** of the distal end **210** of the first catheter **100**. The needle **230** can also include a delivery lumen **238**.

In use, the disclosed percutaneous catheter system **10** can be incorporated into methods of puncturing through a tissue structure within the body of a subject (method **1000**), as shown in FIG. **15**. In one aspect, an exemplary method of puncturing through a tissue structure within the body of a subject can comprise positioning the distal end **110** of the first catheter **100** proximate a first side of the tissue structure (step **1100**). In another aspect, the exemplary method can comprise positioning a distal end **210** of a second catheter **200** proximate a second side of the tissue structure (step **1200**). In an additional aspect, the exemplary method can comprise magnetically coupling the first magnet assembly **120** of the first catheter **100** to the second magnet assembly **220** of the second catheter **200** through the tissue structure such that the longitudinal axis **102** of the first catheter **100** is substantially axially aligned with the longitudinal axis **202** of the second catheter **202** (step **1300**). In a further aspect, the exemplary method can comprise selectively advancing a needle **130** through the at least one lumen **114** (e.g., the

primary lumen **116** in the exemplary aspect) of the first catheter **100** such that at least a portion **132** of the needle **130** exits the opening **112** of the distal end **110** of the first catheter **100** and is received within the opening **212** of the distal end **210** of the second catheter **200**, piercing the tissue structure **40** (step **1400**), as shown in FIG. **14**. In exemplary aspects, the tissue structure can comprise an anatomical pericardial reflection adjacent to the heart of the subject. In these aspects and others, both catheters **100**, **200** can employ a guide wire **300** to reach their positions incrementally, with the operator using standard over-the-wire maneuvering techniques to advance the catheters **100**, **200**.

In an exemplary aspect of the method (**1000**) discussed above, the distal end **110** of the first catheter **100** being positioned in the transverse sinus (step **1100**), as illustrated in FIG. **16**. The distal end **210** of the female catheter **200** can be introduced over the anterior/superior aspect of the ventricle (FIG. **17**), and then advanced toward the right pericardial “gutter” by way of the posterior/inferior cardiac border (FIG. **18**) to be proximate the first catheter **100** (step **1200**). When in place, the magnet assemblies **120**, **220** of the male and female catheters **100**, **200** can then be magnetically coupled (Step **1300**), as illustrated in FIG. **19**. The needle **130** can then exit the distal end **110** of the male catheter **100** to be received within the bore **222** of the magnet assembly **220** of the female catheter (step **1400**), as shown in FIG. **14**.

In addition, steps of the method as discussed above can be repeated during certain procedures. Referring back to the exemplary aspect discuss above, after step **1400** has been completed, the second catheter **200** can be withdrawn into the obtuse sinus (step **1100**), as shown in FIG. **20**. The male catheter **100** can be positioned adjacent the second catheter **200** (step **1200**) (FIG. **21**) and couple the targeted pericardial reflection sandwiched in between (steps **1300**), as shown in FIG. **22**. The needle **130** can then puncture the tissue (step **1400**). After the needle **130** has punctured the tissue, the guidewire **300** can be advanced from the proximal male catheter across the magnetic coupled ends and out the proximal end of the female catheter **200**. The catheters **100**, **200** can be removed, leaving the guidewire **300** in place, as shown in FIG. **23**. In additional aspects, it is contemplated that the percutaneous catheter system **10** can be used to cross and/or puncture through other anatomic boundaries within the body of a subject. For example, it is contemplated that the percutaneous catheter system **10** can be used to cross and/or puncture through the pericardium and plural space (to create a pericardial window). In another exemplary aspect, it is contemplated that the percutaneous catheter system **10** can be used to create access between various organ structures in a controlled manner (e.g., between the bladder and the perineum or between ventricles in a brain (for drainage or placement of electrodes)). In yet another exemplary aspect, it is contemplated that the percutaneous catheter system can be used intravascularly to create an AV fistula in a dialysis patient. In still another exemplary aspect, it is contemplated that the percutaneous catheter system **10** can be used to accomplish trans-venous delivery of electrodes, such as electrodes used in pacemakers and/or nerve stimulators, when an electrical generator is positioned remotely from an electrode target and surgical tunneling is not a desirable option.

In exemplary applications, it is contemplated that the percutaneous catheter system **10** can safely perform punctures across membranous pericardial reflections. The catheter system **10** can be introduced into the pericardium by one of several common transcatheter techniques.

The following exemplary method (2000) can be employed following access to the pericardial space via a subxiphoid approach (step 2100) as shown in FIG. 24; however, it is understood that the method described below can also be employed following other conventional approaches. FIG. 25 illustrates the sterile field 2002 for percutaneous access into the pericardial space. The entry site 2004 is also shown. It is contemplated that the respective longitudinal lengths 104, 204 of the first and second catheters 100, 200 of the percutaneous catheter system 10 can be sufficiently long to permit advancement of the first and second catheters 100, 200 into the transverse sinus of the pericardium from the subxiphoid approach. Thus, it is contemplated that the longitudinal length 102, 202 of each respective catheter 100, 200 can range from about 20 cm to about 50 cm.

In exemplary aspects, the first and second catheters 100, 200 can be introduced into the pericardial space over a guide wire 300 (step 2200). The catheters 100, 200 can then be directed to opposite sides of the target pericardial reflection using standard over-the-wire steering techniques and/or fluoroscopic guidance (step 2300). When the distal ends 110, 210 of the catheters 100, 200 respectively are within close proximity, the magnet assemblies 120, 130 of the catheters will be drawn together magnetically, magnetically coupling the distal ends 110, 210 of the first and second catheters 100, 200 together (step 2400). Under conditions where there is a thin intervening tissue membrane, it is contemplated that the distal ends 110, 210 of the catheters 100, 200 can “sandwich” the membrane orthogonally to the primary lumens 116, 216, of the two catheters 100, 200. It is further contemplated that the magnetic field created by the magnet assemblies 120, 220 of the catheters 100, 200 can align the primary lumen 116 of the first catheter 100 with the corresponding primary lumen 216 of the second catheter 200, thereby facilitating longitudinal continuity. It is still further contemplated that the strength of the magnet assemblies 120, 220 and the size and flexibility of the catheters 100, 200 can allow the distal ends 110, 210 of the catheters 100, 200 to align when in close proximity.

Using fluoroscopic guidance, the operator can position the two complementary catheters 100, 200 on opposite sides of a target pericardial reflection (method 3000), as shown in FIG. 26. Visualization of key pericardial and cardiac landmarks can be facilitated by varying concentrations of radiopaque contrast injected and withdrawn through the irrigation ports 119, 219 of the catheters 100, 200. The catheters 100, 200 can access the pericardial space via a subxiphoid approach (step 3100). Referring to the exemplary pericardial reflection depicted in FIG. 1, it is contemplated that the male catheter 100 (i.e., the catheter of the two in which the needle is advanced) can be placed at the membranous reflection of the superior vena cava from the transverse sinus (step 3200), while the female catheter 200 (i.e., the catheter receiving the needle) can be advanced to the same membranous reflection via the post-caval recess (step 3300). Fluoroscopic navigation can be facilitated by delivery of 5-10 cc of one or more known radio-contrast agents that are injected into the pericardial space. It is contemplated that the first and second catheters 100, 200 can have a plurality of irrigation ports/side openings 119, 219 located at their distal ends 110, 210 to permit injection and suction of fluids, including, for example and without limitation, radio-contrast agents, saline, medications, and body fluids. It is further contemplated that the membranous reflection at this location can have a thickness ranging from about 0.25 mm to about 1 mm. After the catheters 100, 200 are positioned in near proximity (e.g., within about 1-2 cm of

one another), the magnet assemblies attract and align the distal ends of the catheters in a “docking” orientation (step 3400). Proper “docking” orientation can be confirmed by fluoroscopic imaging (step 3500).

In exemplary aspects, both male and female catheters 100, 200 can have a central lumen 116, 216 to accommodate a standard guide wire 300. In these aspects, it is contemplated that the standard guide wire 400 can be withdrawn once the catheters 100, 200 are positioned at a desired site and orientation. It is further contemplated that, through the use of fluoroscopic guidance, the position of the male and female catheters 100, 200 can be confirmed by injection and/or suction of one or more radio-contrast agents into or from the pericardial space. It is still further contemplated that the male catheter 100 can have a retractable puncture needle 130 that can extend and “dock” with the female catheter 200 when the two distal ends 110, 210 are aligned.

Once the catheters 100, 200 are magnetically attached and aligned, with the target membrane sandwiched in between the distal ends 110, 210 of the catheters 100, 200, the operator can advance a stylus 146 (i.e., the elongate member) of the male catheter 100 until the needle 130 punctures through the target membrane and “docks” with the female catheter 200. The operator can then advance the guide wire 300 from the male catheter 100 into the primary lumen 216 of the female catheter 200. The needle 130 can then be retracted, and the catheters 100, 200 can be withdrawn, leaving the guide wire 300 in place. It is contemplated that the previously described steps can be repeated as necessary to create a path for circumnavigating the left atrial target structures. For example, it is contemplated that the above-described method can be used to create a puncture across the pericardial reflection between the superior vena cava and the right superior pulmonary vein located at the rightward terminus of the transverse sinus and a second pericardial reflection puncture located between the inferior vena cava and the right inferior pulmonary vein traversing from the rightward aspect of the pericardial space into the oblique sinus. Following removal of the catheters 100, 200 from the body of the subject, one or more ablation catheters 20 can be delivered and positioned over the guide wire 300.

It is contemplated that the percutaneous catheter system 10 can perform the puncture methods described herein without the need for direct visualization and/or mechanically advantageous positioning, as is required for more conventional puncture techniques. Typically, the restrictions of space and geometric boundaries of the pericardial space constrain over-the-wire catheter design. However, the disclosed catheters 100, 200 of the percutaneous catheter system 10 can be flexible enough to navigate multiple turns while maintaining rotational rigidity for “steer-ability” and direct of the guide wire. Additionally, the distal ends 110, 210 of the catheters 100, 200 can be blunt and/or rounded, thereby reducing the risk of inadvertent puncture of surrounding vascular structures. With the magnetic “docking” capabilities of the catheters 100, 200 through their respective magnetic assemblies 120, 220 it is contemplated that the needle 130 can be deployed when the target membrane is the only structure in jeopardy; otherwise, the needle 130 will be housed within a lumen 116 of the catheter system 10 such that there is no risk of inadvertent puncture. While the exemplary aspects of the percutaneous catheter system 10 have been disclosed in relation to first catheter 100 as being the male catheter, and the second catheter 200 being the female catheter, either assignments can differ based upon which ever catheter is configured to control the advancement of the needle. For example, in an exemplary aspect, the

second catheter **200** can include a need **230** with a lumen **238** and a sharp edge **234** that is longitudinally controlled along the primary lumen **216** by a stylus **246**.

In additional exemplary applications, it is contemplated that the percutaneous catheter system **10** can be applied anywhere precision catheter-based puncture between two adjacent anatomic spaces (as described above) is desired. For example, it is contemplated that a dialysis fistula can be performed by advancing opposing catheters of a percutaneous catheter system **10** to a site of adjacent artery and vein to make a controlled perforation and shunt. In another exemplary application, it is contemplated that a controlled trans-cardiac puncture can be performed across the atrial wall into the pericardial space of a subject to accomplish epicardial pacemaker lead implantation. Where a trans-vascular puncture site is remote, it is contemplated that other biosensor and/or stimulator lead placement could be performed using the disclosed percutaneous catheter system **10**. In still further exemplary aspects, it is contemplated that the percutaneous catheter system **10** can be used for shunt placement between internal cavities, such as the plural space and parental space, for chronic plural effusions, or for creating a fistula between the bladder and a drain. It is further contemplated that the disclosed percutaneous catheter system **10** can be modified as necessary to permit usage of the catheter system in percutaneous procedures where special and anatomic restrictions do not facilitate precise puncture of a tissue structure and/or guide-wire manipulation.

Ablation Catheter

With reference to FIGS. **27-34**, described herein is an ablation catheter **20** for ablating a selected tissue region within the body of a subject. In exemplary aspects, the ablation catheter **20** is an over-the-wire multi-electrode ablation catheter **20** that can create a linear circumferential ablation lesion using one or more of radiofrequency (RF) energy, irreversible electroporation (IE) impulses, and other hybrid electro cautery techniques. The ablation catheter **20** is designed to apply high-voltage, ultra-short direct current pulses to tissue that causes tissue injury, cell death, and in some instances, only cell function disruption.

However, it is contemplated that other ablative techniques such as cooling, microwave, ultrasound, light, and/or chemical ablation techniques could also be used as alternative and/or as adjunct to the ablation approaches described herein. For example, aspects of the ablation catheter **20** can apply HVUS-DCI, RF, cryoablation, electroporation, microwave, laser, biologics, radiation, and small molecule chemicals. These impulses produce brief but extremely strong electric fields within the tissue leading to irreversible electroporation (IE), cell death, and injury. However, in an aspect, the total energy applied is relatively low averaging (estimated range 0.025 J to 45 J per pulse).

In additional exemplary aspects, the ablation catheter **20** can be used in conjunction with the percutaneous catheter system **10** described above. In these aspects, the percutaneous catheter system **10** can be used to place a guide wire **300** within the heart of a subject, and the ablation catheter **20** can be advanced within the heart over the guide wire. Following placement of the ablation catheter **20**, ablative energy can be selectively applied within the heart of the subject. In exemplary aspects, the entire ablation procedure can be performed without administration of anesthesia.

In one aspect, as illustrated in FIGS. **27-32**, the ablation catheter **20** comprises a flexible elongate shaft **500** having a longitudinal axis **502**, a longitudinal length **504**, a proximal portion **506**, a central portion **508**, and a distal portion **510**.

In this aspect, the elongate shaft **500** can define a primary lumen **512**. In this aspect, it is contemplated that the primary lumen **512** can be configured to receive the guide wire **300**. While the ablation catheter **20** can be comprised of many different materials, the material should be flexible. In exemplary aspects, the ablation catheter **20** can be highly flexible such that, upon deployment, the flexible elongate shaft **500** of the catheter **20** can conform to the natural contours of the anatomy. In these aspects, the flexibility of the ablation catheter **20** can facilitate positioning of electrodes **530** around the outside of asymmetric and/or complex contours.

In another aspect, the ablation catheter **20** further comprises a plurality of electrodes **530** spaced along the longitudinal length **504** of the central portion **508** of the flexible elongate shaft **500**. In this aspect, it is contemplated that the plurality of electrodes **530** can be integrally formed with the elongate shaft **500**. Each of the electrodes **530** is configured to be connected to a signal source through an independent wire **518** (shown in FIG. **28**) that is connected by pins **519** to the signal source. The electrodes **530** are configured to apply a signal to the targeted area to perform an ablation. Individual electrodes **530** can be assigned polarity and function in real time to optimize direction of current vectors during ablation. In an aspect, the electrodes **530** can be capable of monitoring and/or delivering RF energy, electroporation impulses, and programmed cardiac pacing and/or neuro-stimulus. Unlike other known ablation catheters, the electrodes **530** of the described ablation catheter **20** also can have the capability of delivering extended bipolar high voltage, ultra-short impulses.

In an aspect, in addition to being configured to apply a signal, the electrodes **530** are configured to be capable of selectively record signals. In this aspect, the signals can be described by an impulse strength, a duration, a duty cycle, and a timing. When the electrode **530** is configured to record the signals, the electrode **530** can record the above described characteristics of the signal(s) applied. The electrode **530** can capture this information, and send it to a console, described in more detail below. In an aspect, an electrode **530** that is not applying a signal can act as a recording electrode **530**. In another aspect, the electrodes **530** of the ablation catheter **20** can be configured to act as a recording electrode and signal delivering electrode **530** at the same time.

In another aspect, the electrodes **530** can be configured to monitor the vital signals of the subject. For example, the electrodes **530** can receive the electronic signals produced by the subject's heart to which the electrode **530** is in contact. In an aspect, the electrode **530** can act like an EKG. In another aspect, the electrode **530** can monitor the atrial pacing (including the atria refractory period), the ventricular pacing (including the ventricular refractory period), the cycle length, the QT interval, and the QRS interval of the subject's heart. The information can be passed along to other components discussed in more detail below.

In exemplary aspects, the plurality of electrodes **530** can be spaced to provide adequate coverage for creating a contiguous linear ablation lesion **40**. In these aspects, it is contemplated that the ratio of the spacing **532** between consecutive electrodes **530** to the longitudinal length of each electrode can be less than about 3:1 and, more preferably, less than about 2:1. In additional exemplary aspects, it is contemplated that the plurality of electrodes **530** can comprise between about 20 to about 40 independent electrodes **530**. In an example, the ablation catheter **200** can have 30 independent electrodes (e.g., FIG. **34**). In further exemplary aspects, it is contemplated that the plurality of electrodes

530 can be spaced along a sufficient length of the elongate shaft **500** (e.g., ranging from about 15 cm to about 30 cm) to create a circumscribing lesion **30** around a left atrial target and pulmonary veins. It is contemplated that the plurality of electrodes **530** can be positioned centrally along the longitudinal length **504** of the ablation catheter **20** so that the proximal portion **504** and distal portion **510** of the elongate shaft **500** are of sufficient length such that at least a portion of the proximal portion **504** and the distal portion **510** are positioned external to the body when the central portion **506** of the elongate shaft **500** (including the plurality of electrodes **530**) is deployed around the left atrial target structures. It is contemplated that the ratio between the longitudinal length of the proximal portion **506** to the longitudinal length of the central portion **508** and the ratio between the longitudinal length of the distal portion **510** and the longitudinal length of the central portion **508** can each range from about 1.5:1 to about 2:1. It is further contemplated that the proximal portion **506** and the distal portion **510** of the elongate shaft **500** can each have a longitudinal length ranging from about 40 cm to about 60 cm.

In exemplary aspects, the flexible elongate shaft **500** can be configured for selective positioning within the body of the subject such that the central portion **508** of the elongate shaft **500** at least partially surrounds the selected tissue region (shown in FIGS. 33-34) and the proximal **506** and distal portions **510** of the elongate shaft **500** are positioned external to the body of the subject. In these aspects, it is contemplated that, upon positioning of the elongate shaft **500** such that the central portion **508** of the elongate shaft **500** at least partially surrounds the selected tissue region, each electrode **530** of the plurality of electrodes **530** is configured for selective, independent activation to apply ablative energy to the selected tissue region. **518**

Optionally, in one aspect, the flexible elongate shaft **500** can further comprise one or more secondary lumens **514** defined by the flexible elongate shaft **500** and/or positioned within the primary lumen **512**. In an aspect, at least one secondary lumen **514** of the one or more secondary lumens **514** or the primary lumen **512** of the flexible elongate shaft **500** can be configured to receive the guide wire **300**. In such an aspect, the other lumen **512**, **514** that are not for use with the guide wire **300** can be configured to receive a flexible stylus and/or other mechanical support. Further, such lumens can be configured to carry and/or deliver a cooling fluid, an irrigation fluid, small molecules, peptides, and/or DNA/RNA to improve ablation characteristics. It is further contemplated that the elongate shaft **500** can be configured for deployment within the body of the subject over the guide wire **300**. However, it is contemplated that the ablation catheter **20** can optionally be deployed within the body of a subject in a manual fashion (without a guide wire).

In an aspect, the proximal end **506** of the catheter **20** can include a luer lock **516** and opening **518** to receive a guidewire **300** in the primary lumen **512** or secondary lumen **514**, as shown in FIG. 30. The distal end **510** can include an opening **520** that continues to the secondary lumen **514**, allowing a guidewire **300** to exit, as shown in FIG. 31. Further, the distal end **510** can have a tapered shape as well.

In an aspect, the ablation catheter **20** can include a catheter noose **524**, as shown in FIGS. 27 and 32. The catheter noose **524** is configured to apply tension to the elongated body **500** of the catheter **20** when the catheter **20** is positioned around the targeted sight. In an aspect, and discussed in further details below, the central portion **508** of the catheter **20** is positioned around the targeted area within the body, with the proximal **506** and distal **510** ends posi-

tioned outside of the body. The catheter noose **524** is then used to tighten the loop formed by the center portion **508** of the catheter **20** around the targeted area. In an aspect, the catheter noose **524** can include two lumens (not shown). The first lumen can be configured to receive the proximal end **506** of the catheter **20**. The second lumen can be configured to receive the distal end **510** of the catheter **20** after the catheter **20**, and more specifically the central portion **508**, has been positioned around the targeted area within the body and the distal end **510** and proximal end **506** are positioned outside the body. The catheter noose **524** can then be advanced along the proximal and distal portions **506**, **510** until the central portion **508** is fully secured, as shown in FIGS. 33-34.

In use, the ablation catheter **20** can be employed in a method for ablating a selected tissue region within the body of a subject. In one aspect, the method for ablating the selected tissue region (**4000**), as shown in FIG. 35, can comprise selectively positioning the flexible elongate shaft of the ablation catheter within the body of the subject such that the central portion of the elongate shaft at least partially surrounds the selected tissue region (step **4100**). In this aspect, the proximal portion **506** and the distal portion **510** of the elongate shaft **500** of the ablation catheter **20** can optionally be positioned external to the body of the subject (step **4200**). In another aspect, the method for ablating the selected tissue region can comprise selectively, independently activating each electrode **530** of the plurality of electrodes **530** of the ablation catheter **20** to apply ablative energy to the selected tissue region (step **4300**).

In an exemplary aspect of the method **4000** described above, the distal end **510** of the catheter **20** can be advanced along the guidewire **300** to be positioned around the left arterial target structures, with the distal end **510** being deployed to cross the pericardial reflection into the transverse sinus and through until the central portion **508** is positioned correctly (step **4100**), as shown in FIGS. 36-37. The proximal portion **506** and distal portion **510** can be placed outside of body (step **4200**), as shown in FIG. 37. Once in place, the catheter noose **524** can be advanced to cinch the loop, as shown in FIG. 38. In cases where the circumference is less than the length **504** of the catheter **20** along the central portion **508** (i.e., the multi-electrode **530** array), excess proximal electrodes **530** are deactivated and pulled proximally into the catheter noose **524** before applying ablative energy (step **4300**). If the circumference of the targeted area is greater than the length **504** along the central portion **508**, the central portion **508** will require an additional repositioning after applying the ablative energy (step **4300**).

In exemplary aspects, it is contemplated that the ablation catheter **20** can be included in an ablation catheter system **600** for ablating a selected tissue region within the body of a subject, as shown in FIGS. 39-44. In an aspect, the ablation catheter system **600** can include a routing console **610**, a recording console **650**, a signal generator **700**, and a computer **800**. The routing console **610** is electrically coupled to the plurality of electrodes **530** of the ablation catheter **20**. More specifically, the routing console **610** is connected to each pin **519** of each independent wire **518** from each electrode **530**. The routing console **610** can carry signals from the signal generator **700** to the electrodes **530**, as well as assign polarity and function in real time to optimize the direction of current vectors during ablation, discussed in more detail below.

As shown in FIGS. 40-41, the routing console **610** includes catheter connectors **612** to receive the pins **519** of

the ablation catheter 20. An exemplary routing console 610 can include two 16 pin connectors used to accommodate thirty (30) independent electrodes 230 on the exemplary ablation catheter 200. However, the total number of catheter connectors can be adjusted to accommodate any range of electrode arrays. The routing console also includes pacing inputs 614, which can receive monitoring information from devices (EKG, etc.) used to monitor the function of the subjects' vital parts, including the heart. The routing console 610 can include signal inputs 616. The signal inputs 616 receive the signal(s) from the signal generator 700. In an aspect, the signal inputs 616 can include high voltage inputs 616. In other aspects, the signal inputs can accept RF and/or any electrical ablation energy source generated by the signal generator 700. The pacing inputs 614 and signal inputs 616 feed into the input signal relay 618, which passes along all the information and signals to the various other components of the ablation catheter system 600, including the signal generator 700, recording console 650 and computer 800, as well as other components of the routing console 610.

The input signal relay 618 is connected to logic controllers 620 and a relay bank 622. The logic controllers 620 and relay bank 622 work in tandem to send signals to a specific electrode 530 based upon the information and commands received from other components, including the signal generator 700, the computer 800, and the pacing inputs 612. The relay bank 622 can pass signal information, as well as other information, to another relay bank 624 which is connected to an I/O interface 626. The I/O interface 626 can be in communication with the signal generator 700 through a signal generator output 628. The first relay bank 622 can also pass along any information related to the signals that are being monitored by an electrode 530 to sensing outputs 630, which can be connected to the recording console 650. The routing console can also include a timing relay 632 which works with the controllers 620 to control the delivery of the signals to the electrodes 530. The timing relay 632 is connected to a synchronization trigger 634, which is in communication with the signal generator 700.

In an aspect, the synchronization trigger 634 ensures that when signals are sent to the electrodes 530 for ablation, the signals are applied in synchronization with the cardiac cycle, discussed in more detail below. The synchronization trigger 634 can receive monitoring information monitoring devices through the pacing inputs 614 or through electrodes 530 that are assigned to a monitoring function. The synchronization trigger 634 can monitor the EKG results, the atrial pacing (including the atria refractory period), the ventricular pacing (including the ventricular refractory period), the cycle length, the QT interval, and the QRS interval of the subject's heart to indicate when a signal should be delivered to the electrodes 530. For example, as shown in FIG. 45, the synchronization trigger 634 can determine the impulse window 900 (i.e., when to apply the signal) by identifying when the ventricular refractory period 902 and the atria refractory period 904 overlap. The synchronization trigger 634 can then alert the routing console 610 and the signal generator 700 of the window 900 to apply the signal.

The routing console 610 includes a fire button 636. The fire button activates the signal generator 700 to generate a signal to deliver a signal to the routing console 610. The routing console 610 will then deliver the signal to the desired electrodes 530. The computer 800 can direct the routing console 610 as to which electrodes 230 to deliver the signal.

The routing console 610 is electrically coupled to the signal generator 700. In an aspect, the signal generator 700 can comprise one or more signal generators 700. It is

contemplated that each signal generator 700 of the one or more signal generators 700 can be configured to selectively generate one or more electrical signals. The signal generator 700 can create several types of signals, including, but not limited to, radio-frequency (RF), high voltage ultra-short direct current (DC) impulses (as used in electroporation), stimulus range impulses, and/or hybrid electrical impulses. In addition, the signal generator 700 can vary at least one of the impulse strength, duration, duty cycle, and timing of the signals that the signal generator 700 generates.

In an aspect, as illustrated in FIGS. 42-43, the signal generator 700 includes pulse/high voltage outputs 702 that are configured to connect with the pulse/high voltage inputs 616 of the routing console 610. The outputs 702 deliver the signal to the routing console 610. The signal generator 700 can include a control circuit 704 that controls the characteristics of the signal that it generates, discussed in more detail below. The control circuit 704 can also be connected to a voltage level controller 705. The pulse outputs 702 receive the signal from a capacitor 706. In an aspect, the capacitor 706 can comprise a bank of capacitors 706. A power supply 708 can provide the power needed to the capacitor(s) 706 to generate a signal. In an aspect, the capacitor 706 can pass along the signal to a transistor 710. In an aspect the transistor 708 can include an insulated-gate bi-polar transistor 710. The signal generator 700 also includes a commercially available pulse capacitor charger 711 which provides a high voltage source for the capacitor bank and a feedback control to adjust peak voltage charge.

In an aspect, the signal generator 700 can also include various inputs to reference information and commands. For example, the signal generator 700 can be connected to the computer 800 and the routing console 610 through an input/output connection 712. The input/output connection can comprise a plurality of input/output connections 712. In addition, the signal generator can be connected to the fire button through a separate input 714. Parameters/commands from the computer 800 and information from the routing console 610, including the synchronization trigger 634 and activation of the fire button 636, are received by the control circuit 704. Based upon the information received, the control circuit 704 controls the generation of the signal. For example, the control circuit 704 can control the pulse duration, the number of pulses within a burst, the burst pulse spacing, the voltage of the signal, and other signal parameters. In another aspect, the control circuit 704 can initiate the signal upon receiving a response from the fire button. In another aspect, the control circuit 704 can control when the signal is generated based upon information received from the synchronization trigger 634 in order to deliver a signal within the pulse window 900.

In an aspect, the recording console 650 can receive and record all the information that is collected by the various other components of the system 600. For example, the recording console 650 can record the pacing information that the routing console 610 receives from monitoring devices associated with the subject. In addition, the recording console 650 can receive monitoring information from the electrodes 530 monitoring the subject. In an aspect, the recording console 650 can also receive the signal information from the recording electrodes 530. In another aspect, the recording console 650 can receive other information from the signal generator 700 regarding the timing and strength of the signals generated, as well as other information. In an aspect, the recording console 650 can be a separate component from the computer 800 and routing console 610. It can be a display device that immediately displays conditions to

the users of the system 600. In other aspects, the recording console 650 can be an application within the computer 800. The physical characteristics of the recording console 650 are not important, nor whether it is a separate entity from the other components of the ablation system 600.

In an aspect, the computer (shown in FIG. 44) can include ablation control software 806 that controls the overall function of the ablation system 600. The ablation control software 806 can use the other components of the system 600 to retrieve information (gathering signal information from the signal generator 700/electrodes 230, and pacing information from the routing console 610/electrodes 530) in order to initiate and maintain the ablation treatment. In other aspects, the ablation control software 806 can also control the synchronization trigger 634, or supply the synchronization trigger 634 with the needed information to apply the signal during the window 900,

In these aspects, the routing console 610 can be configured to receive the one or more electrical signals from the one or more signal generators 700. It is contemplated that the routing console 610 can be further configured to selectively activate the plurality of electrodes 530 by delivery of the one or more electrical signals from the signal generators 700. In an aspect, the routing console 610 can be configured to selectively activate at least one electrode 530 of the plurality of electrodes 530 of the ablation catheter 20 such that the at least one electrode 530 has a first polarity that is different from a polarity of at least one other electrode 530 of the plurality of electrodes 530, which, in turn, can provide means for customizing the ablation vector for each electrode 530 individually and/or delivering pacing and/or ablation impulses in quick succession.

In exemplary aspects, the ablation catheter system 600 can be employed in a method for ablating a selected tissue region within the body of a subject 5000, as shown in FIG. 46. In one aspect, the method 5000 for ablating a selected tissue region can comprise selectively positioning the flexible elongate shaft 500 of the ablation catheter 20 within the body of the subject such that a central portion 508 of the elongate shaft 500 at least partially surrounds the selected tissue region (step S100) and a proximal portion 506 and a distal portion 510 of the elongate shaft 500 are positioned external to the body of the subject (step S200). In another aspect, the method for ablating the selected tissue region can comprise selectively generating one or more electrical signals using the one or more signal generators 610 (step S300). In an additional aspect, the method for ablating the selected tissue region can comprise, through the routing console 620, receiving the one or more electrical signals from the one or more signal generators 610 (step S400). In a further aspect, the method for ablating the selected tissue region can comprise, through the routing console 620, delivering the one or more electrical signals to the plurality of electrodes 530 of the ablation catheter 20 such that each electrode 530 of the plurality of electrodes 530 is selectively, independently activated to apply ablative energy to the selected tissue region (step S500). In an exemplary aspect, the method for ablating the selected tissue region can further comprise, through the plurality of electrodes 530, selectively recording one or more electrical signals within the body of the subject (step S600). In another exemplary aspect, the method for ablating the selected tissue region can further comprise, through the one or more signal generators 610, selectively varying at least one of the impulse strength, the duration, the duty cycle, and the timing of the one or more electrical signals generated by the one or more signal generators 610 based upon the one or more electrical signals

recorded by the plurality of electrodes 530 (step S700). In a further exemplary aspect, it is contemplated that the step of, through the routing console, delivering the one or more electrical signals to the plurality of electrodes 530 can comprise selectively activating at least one electrode 530 of the plurality of electrodes 530 such that the at least one electrode 530 has a first polarity that is different from a polarity of at least one other electrode of the plurality of electrodes 530, as discussed above.

In exemplary aspects, the ablation catheter 20 can be highly flexible such that, upon deployment, the flexible elongate shaft 500 of the catheter 20 can conform to the natural contours of the anatomy. In these aspects, the flexibility of the ablation catheter 20 can facilitate positioning of electrodes 530 around the outside of asymmetric and/or complex contours.

It is contemplated that the ablation catheter 20 can be configured to deliver both radio frequency (RF) and/or high intensity ultra short duration electrical impulses/irreversible electroporation (IE) to ablate adjacent tissue. RF ablation in the closed pericardial space has some important limitations. First, RF ablation can produce tissue injury through resistive heating. The lesion depth resulting from RF ablation can be limited by the energy and thermodynamics of the tissue environment. For example, a unipolar RF lesion created from the epicardium can require greater energy to create a transmural lesion than the same lesion delivered from an endocardial approach; this is because the endocardium is cooled by the blood pool and there is often a layer of epicardial fat that adds thickness. (See FIG. 47.) Using an extended bipolar electrode arrangement, it is contemplated that approximately 50% more directional penetration can be achieved (using RF techniques).

FIG. 47 shows the potential advantages of an extended bipolar ablation arrangement for epicardial ablation techniques. Panel (A) depicts a virtual electrode from a standard unipolar RF ablation on an endocardial surface. As shown, the field of the unipolar signal extends substantially only along the myocardium (a) and epicardial fat (b). Panel (B) shows unipolar RF ablation from an epicardial approach, with the field of the unipolar signal extends into the epicardial fat (b), pericardial space (c), and parietal pericardium (d). However, the field also extends to a bystander vulnerable structure (f). Panel (C) illustrates the distortion of the virtual electrode by using an extended bipolar orientation. As shown, the bipolar orientation leads the field to extend into the ventricular myocardium (a), epicardial fat (b), pericardial space (c), and parietal pericardium (d) without impacting the bystander vulnerable structure (f).

It is contemplated that the use of high-voltage, ultra-short impulses (irreversible electroporation) can substantially increase the directionality of the ablation vector. In a closed pericardial space, the thermal conduction can continue to be problematic, causing undesirable collateral damage and/or accumulation of proteinaceous material on the electrodes, which can require device removal, cleaning, and/or reinsertion. However, despite these limitations, it is contemplated that RF techniques may be preferred for ablation targets that are epicardial structures, such as autonomic ganglia.

The selected polarity of each electrode 530 of the plurality of electrodes 530 can be assigned based upon the geometric orientation of each respective electrode 530 toward the ablation target. Optionally, the assignment of polarity to each respective electrode 530 can be performed in real time using the routing console 610 attached to the catheter 20 outside the body. In an aspect, the polarity assignment for each respective electrode 530 can be adjusted to tailor the

intended vectors of ablation current. In another aspect, the polarity assignment can optionally be performed in connection with a remote electrode located within or external to the body. In these aspects, the vector of current between any two electrodes of the plurality of electrodes can be directed toward the intended ablation target by choosing an electrode 530 combination that optimizes the intended vector and away from bystander structures (see FIG. 47). In another aspect, the electrode combination can comprise two or more electrodes 530 of the central portion 508 of the ablation catheter 50.

In another aspect, a high impedance structure 540 can be positioned between the electrodes 530. The high impedance structure 540 is configured to change and/or direct the current path between selected electrodes 530, as illustrated in FIGS. 48-52(a-c). In an aspect, the ablation catheter 20 can use a plurality of high impedance structures 540. The high impedance structures 540 are configured to intersect the theoretic field lines 550 (see FIGS. 48-49) created by two bipolar electrodes 530 by creating an obstacle to a baseline current flow. For example, in a homogeneous conductor such as seawater or blood plasma, the predicted current path will follow the shortest path (i.e., the current will follow the path of least resistance), as shown in FIG. 48. By placing a high impedance structure 540 between adjacent electrodes 530, the current contour 550, as shown in FIG. 49, can be distorted by the contours of the high impedance structure 540, with the current density decreasing linearly between the electrodes 530 but increasing orthogonally along the surface of the high impedance structure 540. FIGS. 50-51 show an axial perspective of the change of the location of the current density 544 of a coaxially cylindrical insulator 540 relative to the insulator circumference. As shown in FIG. 50, when the circumference of the insulator/high impedance structure 540 is small, the current density 544 is approximate the surface of the electrode 530. However, as the high impedance structure 540 expands, the current density 544 becomes located further from the surface of the electrode 530.

In exemplary aspects, the shape, and more specifically the height of the high impedance structure 540 relative to the axis 502 of the ablation catheter 20, is adjustable. For example, the high impedance structure 540 comprises an inflatable balloon 540 made of a suitable nonporous material with high dielectric constant (i.e., effectively an electric insulator). The inflatable balloon 540 is coaxially situated between two electrodes 530, as shown in FIG. 52a-c. As the inflatable balloon 540 is inflated, the current density 544 along the surface of the balloon will decrease linearly while the relative current density 544 at an arbitrary point between the electrodes 530 and orthogonally remote from the axis 502 of the ablation catheter 20 increases. The adjustment of the inflatable balloon 540 provides a way to project and/or direct the electric field along an orthogonal/radial vector to increase the current density 544. While the exemplary aspect utilizes a balloon 540 to provide low profile delivery, other articulated, fixed and/or mechanical high impedance structures 540, including a wide variety of insulators, can be employed. Further, it is preferable that the high impedance structures 540 are controllably adjustable, for the reasons discussed below.

The current density at the surface of the cylindrical insulator symmetrical positioned between two ring electrodes is geometrically related to the radius of the cylinder. In such an exemplary aspect can be determined by the following formula:

$$J = J_i (\Pi r^2 * I_i) / ((\Pi r^2 * I_i)_2 - (\Pi r^2 * I_i))$$

where J is the resulting density, J_i is the initial density, $(\Pi r^2 * I_i)$ is the initial area of the high impedance structure before activation, and $(\Pi r^2 * I_i)_2$ is the area of the high impedance structure after activation.

In our exemplary aspect, the electrical conductivity ranges 50-100 S/M (conductivity σ is defined as the ratio of the current density J to the electric field strength E). ($J = \text{Sigma.E}$). The predicted electric field strength at the surface of the insulator balloon 540 (represented by A in FIGS. 48-49) will be related to the current density/conductivity of the environment.

Positioning of the high impedance structure or insulator 540 between the dipole formed from adjacent electrodes 530 will change the contour of the current path and increase the relative electric field strength at point A, as shown in FIGS. 48-49. The shape of the high impedance structure 540 can be varied to project/amplify the relative the current orthogonal to the axis 502 of the ablation catheter 20. Other shapes and materials can be used as high in combination with high impedance structures/insulators 540 to focus the current asymmetrically or to isolate the current source form the target tissue. In an aspect, the high impedance structure or insulator 540 can comprise an insulator balloon 540 configured to expand off center to provide a preferential path for current ipsilateral to the shorter axis's.

In other aspects, the high impedance structure or insulator 540 can be constructed to geometrically isolate current from one source electrode 530 from untargeted nearby structures but allow the current to travel through a fenestration or other geometrically oriented opening, there by changing the current density. In a simple example, a balloon when expanded would partially cover the electrodes 530 while creating a prescribed tunnel for the current to travel through. In an aspect, an asymmetrical balloon 530 can focus current along the path of least resistance (generally the shortest linear distance). In another aspect, an expanding mesh high impedance structure 530 can surround the electrode 530 to safely increase current at that electrode 530 with less risk of unwanted collateral damage by simply maintaining a prescribed distance from soft tissues. Such a high impedance structure allows an increase current density at one end of a bipole near an ablation target while protecting structures at the counterpoint. The use of geometric high impedance structures or insulators 540 to contour the current path of a current created between dipole electrodes 530 within a conductive media such as tissue could be used to precisely deliver electrical ablation or stimulus energy to targeted tissues adjacent to the high impedance structure 540.

While the combination of the electrodes 530 and the high impedance structures 540 are directed to deliver high voltage ultra short ablation impulses in the pericardial/epicardial space for the purpose of treating cardiac arrhythmia, there is an immediate implication for other ablation procedures using the electrode 530/high impedance structures 540 for contouring ablation energy to vascular walls (in stent restenosis) and/or contour the virtual electrode 530 in ablation procedures targeting solid tumors and/or prostatic hypertrophy. While balloon catheters are known in the art for the purpose of providing mechanical force, geometric stabilization, and or the delivery of ablation energy such as laser light or ultrasound, the combination of electrodes 530 and high impedance structures 540 oriented on an ablation catheter 20 is fundamentally distinct as the ablation catheter 20 uses the high impedance structures 540 to shape the electric current used in an in vivo therapy.

It is contemplated that the independent electrodes 530 can be assigned polarity individually or in groups. Depending on

these polarity assignments, it is contemplated that the relative orientation of the electrical impulses and the virtual electrode properties (e.g., the surface area and thus control current density) of the electrodes **530** can be selectively adjusted. In exemplary aspects, the plurality of electrodes **530** of the ablation catheter **20** can be connected to a routing console/switchboard **610** outside the body where the electrodes **530** can be assigned a role as a recording electrode, an active pacing, and/or an ablation electrode, as discussed above. The console **610**, in turn, can be operatively coupled to a computer-controlled signal generator **700** and recording console **650**. In an aspect, the electrode polarity assignments can be changed as needed to achieve one or more desired effects. By changing the relative polarity assignments of the electrodes **530**, at least one of the virtual electrode shape and the current density can be selectively varied.

In another aspect, the ablation energy can be delivered to a single electrode **530** or to multiple electrodes **530** simultaneously. In an aspect, FIGS. **53a-d** display an array of exemplary electrode **530** assignments. FIG. **53a** illustrates an extended bipolar arrangement with equal current density between electrodes **8** and **23**. The selected electrodes **8** and **23** can deliver an ablation impulse for every cardiac cycle, changing the active bipoles with every cardiac cycle in a step-wise manner. In an example, if the heart is paced at a 500 ms cycle length the circumferential linear lesion will be delivered in 7.5 seconds.

FIG. **53b** illustrates an extended bipolar arrangement with asymmetric current density, wherein electrode **8** is assigned a different polarity than electrodes **22**, **23**, and **24**. This assignment decreases the current density at one of the bipoles to reduce injury to bystander structures near the pole.

FIG. **53c** illustrates an extended bipolar arrangement with equal current density but activated as a simultaneous array. As illustrated, electrodes **9-13** are assigned one polarity, whereas electrodes **24-28** are assigned another. The electrodes **530** are activated simultaneously to form complementary arrays. This could be employed in cases where sub straight accommodated more rapid ablation sequencing (2-3 cycle lengths).

An extended bipolar arrangement with asymmetric current density is illustrated in FIG. **53d**. As shown, electrodes **11**, **12**, **17**, and **18** are assigned a polarity different from electrode **30**, which creates an extended bipolar arrangement with a gap in the complementary electrode array. Such an arrangement can be used to avoid inadvertent ablation of a vulnerable bystander structures, including the phrenic nerve.

It is still further contemplated that the impulses can be delivered in a programmed manner, triggered by feedback from a bio-potential or physiologic signal (such as respirations, nerve impulses, fluctuations in blood pressure, and/or the cardiac action potential) or an outside event.

In exemplary applications, as described above, the ablation catheter **20** can be deployed such that both the proximal portion **506** and distal portion **510** of the elongate shaft **500** are external to the body (the central portion **508** of the catheter with the multi-electrode array remains internal). However, in additional applications, it is contemplated that the ablation catheter **20** can be customized to take advantage of target anatomy; in some cases, the distal portion **510** of the ablation catheter **20** can remain in the body, and a remote electrode can be used to complete the ablation procedure.

In exemplary applications, the ablation catheter **20** can be employed in a catheter-based epicardial atrial fibrillation ablation procedure performed in a closed pericardium. In this atrial fibrillation ablation procedure, the ablation catheter **20** can be advanced over a guide wire **300** that has

already been positioned around the epicardial left atrial structures. Thus, the ablation catheter **20** can be deployed into the pericardial space from a subxiphoid or apical percutaneous approach, as discussed above.

It is contemplated that the guide wire **300** can be delivered around the left atrium by using the percutaneous catheter system **10** described herein to puncture through two key anatomic obstacles (pericardial reflections near the vena cava and the right pulmonary veins). Using this method, the guide wire **300** can enter the pericardium and then travel under the inferior-lateral left ventricle, along the lateral left atria, into the transverse sinus, along the roof of the left atria, between the right superior pulmonary vein and superior vena cava (SVC) through a pericardial puncture site. Then, the guide wire **300** can travel along the right lateral aspect of the left ventricle, between the right inferior pulmonary vein and inferior vena cava (IVC), traveling through the second pericardial puncture into the obtuse sinus under the posterior left atria. The guide wire **300** can then extend under the ventricle and out of the pericardium such that both ends of the guide wire **300** are outside the body. Once the guide wire **300** has been positioned, the ablation catheter **20** can be advanced along the guide wire **300**. From this advantageous position, the ablation energy can be delivered directly to the key left atrial ablation targets, thereby creating a circumferential lesion without the need for repositioning the ablation catheter **20** or entering the left atrial blood pool. However, the ablation catheter **20** can be repositioned to perform other targeted epicardial ablations, including, for example and without limitation, ablation of autonomic ganglia or creation of additional linear ablation lesions.

In an aspect a goal of the disclosed ablation procedure can be the electrophysiological isolation/decoupling of key segments of the heart (e.g., the left atrium and the ostia of the pulmonary veins) that are thought to be involved in the genesis and/or maintenance of atrial fibrillation. The disclosed percutaneous catheter system **10** and ablation catheter **20**, and the associated ablation catheter system **600**, can provide means for creating a "box" lesion around ostia of the pulmonary veins without the need to enter the arterial blood pool. In use, after the ablation catheter **20** is deployed over the guide wire **300**, one or more electrodes **530** of the plurality of electrodes **530** of the ablation catheter **20** can be used to measure local electrograms and/or deliver mapping stimuli. Using an extended bipolar arrangement of the electrodes **530**, the directional electrograms adjacent to the electrodes **530** can be assessed to permit identification of changes in the substrate and local conduction block. As further described herein, the ablation catheter **20** can be connected to one or more impulse generators **700** and a routing console **610**. It is contemplated that the operator can select an electrode configuration to optimize the vector of current for each segment of the lesion. In exemplary aspects, the procedure can be at least partially computer-automated while requiring at least some input from the operator to identify a preferred current vector. The impulse generator **700** can then deliver ablative energy to the electrodes **530** of the ablation catheter.

In exemplary applications, the ablation catheter **20** can be configured to deliver high intensity ultra-short duration impulses/IE to produce a transmural lesion. In an aspect, the IE impulses can be delivered by the electrodes **530** in synchrony with the cardiac cycle (e.g., from about 200 ms to about 300 ms after detection of a QRS complex) to reduce the chance of inducing arrhythmias. In an aspect, the impulse strength, duration, duty cycle and timing of the IE impulses can be selectively adjusted to tailor the ablation

characteristics in real time. In such an aspect, the real-time adjustments can be required to address changes in tissue conductance as the lesion evolves. In exemplary aspects, the power can be adjusted to maintain a constant current density in the virtual electrode, thereby reducing the tissue conductance. In such aspects, the tissue conductance can be measured between impulses and integrated into an automated feedback circuit. In such aspects, the impulse strength can be adjusted to electroporation impulses using a standard unipolar configuration or an extended bipolar configuration.

Irreversible electroporation (IE) is a non-thermal ablation technique that can be advantageously used within the pericardial space. IE works by delivery on ultra-short (nanoseconds) high voltage (100-10,000V) impulses that cause very brief disruption in the membrane of cells. The disruption in the lipid bilayer leads to cell death through necrosis or apoptosis, depending on the field strength involved. In exemplary aspects, the ablation catheter 20 can permit customization of the direction of ablation energy within the pericardium. When compared to RF ablation, IE ablation can produce a lesion that follows a geometric pattern more closely approximating the contours of the virtual electrode 530. In such an aspect, the ablation catheter 20 can take advantage of these electrophysiologic properties to create a more focal lesion that directs the vector of current toward the target and also reduces the risk of unintended collateral injury. Although RF ablation using the same extended bipolar technique shows directionality, local tissue heating can reduce the current vector effect. (See FIG. 26). Additionally, the IE ablation can leave the intracellular matrix of tissue relatively undistorted, thereby reducing the risk of structural tissue instability, rupture, and fistula formation; there is typically limited or no opportunity for "char" formation on the electrode, so it generally will not need to be removed, cleaned, or redeployed. Because nerve fibers are particularly resistant to injury from IE techniques, IE ablation can reduce the risk of damage to nearby phrenic nerves. IE ablation can produce effective lesions in a fraction of the time required to create a transmural lesion by RF techniques. In exemplary aspects, IE impulses can be delivered via the ablation catheter 20 through the electrodes 530 in an automated fashion in a variety of extended bipolar orientations to create the complete linear circumscribing lesion in less than $\frac{1}{10}$ the time it would take to produce the same lesion set using RF ablation techniques. IE ablation techniques are not dependent on tissue thermodynamics, thereby improving the chance of creating a full thickness lesion. Thermal techniques such as resistive heating from RF energy can be less effective because conductive cooling properties of the blood pool can protect the endocardium. In an aspect, IE ablation techniques can be selectively tuned to create lesions by apoptosis (as opposed to necrosis), leaving a very clean scar with less local inflammation.

In exemplary configurations, the ablation catheter system 600 can comprise the ablation catheter 20 and a routing console 610 that is linked to a commercially available signal generator 700 which is capable of arbitrary electrical waveform generation, including simple DC stimulus, radiofrequency monophasic and biphasic impulse generation, and high voltage ultra short impulse generation

In use, after an operator has positioned a guide wire 300 around the left atrium, the ablation catheter 20 can be advanced over the guide wire 300 so that the array of electrodes 530 (located at the central portion 508 of the elongate shaft 500) now surrounds the left atrium. The distal portion 510 of the ablation catheter 20 can extend outside the body of the subject and be passed through the means for

applying tension 524 (e.g., a loop tensioner 524), as further described herein. The loop tensioner 524 can then be advanced over the proximate portion 506 and distal portion 510 of the ablation catheter 500 to provide lateral tension and create a closed loop around the left atrial target structures. The guide wire 300 can then be removed to provide more flexibility and improved tissue contact along the left atrial contours. Small adjustments can be made using the loop tensioner 524 and/or a variety of custom styluses 524 that can be inserted into the catheter wire lumen 512/514. Once a desired position of the electrodes 530 of the ablation catheter 20 around the targeted tissue region is achieved, it is contemplated that the ablation catheter 20 will not need to be repositioned.

The operator can then conduct a limited electrophysiologic study, checking left atrial pacing thresholds and local electrocardiograms. The operator can then evaluate the radiographic orientation of the electrodes 530 around the left atrium and assign a polarity to the each respective electrode 530. Optionally, this assignment procedure can be partially automated to reduce the total steps needed to create and optimal extended bipolar vector. The tissue conductance and impedance can be measured at each electrode 530 at baseline. In an aspect, these measurements can be performed in an automated procedure performed by an automated recorder and potentially integrated into the control algorithm to make voltage adjustments, and/or can be performed manually by the operator. These baseline measurements can be periodically re-measured to assess local ablation effects. The data can be used to adjust the applied ablation energy in an automated fashion when such automated functions are available. It is contemplated that each electrode 530 of the plurality of electrodes 530 of the ablation catheter 20 can be used to monitor, pace and/or deliver energy for ablation. In exemplary aspects, the ablation energy can be delivered to the plurality of electrodes 530 using a programmed computerized protocol synchronized with the cardiac cycle of the subject. In exemplary applications, the operator can selectively initiate a sequence activating each electrode 530 individually and/or in series.

It is contemplated that the linear ablation should be completed in less than about 60 seconds (depending on the baseline heart rate and total length of the linear lesion being created). In the exemplary system we will overdrive pace the heart at a rate between 100 and 120 beats per minute. In order to deliver ablation pulses or train of pulses to each electrode we will discharge the device $n^{*1/2}$ times the number of electrodes in the array. In our example we use 30 electrodes therefor a completed cycle will take 7.5 seconds. Conceivably the entire procedure could be performed in 7.5 milliseconds with commercially available solid-state high voltage relays.

In an aspect, an electrophysiologic study of conduction block can be performed without any repositioning of the ablation catheter 20. The operator can perform a programmed stimulus protocol to identify gaps in the linear lesion. In the example the operator would perform an electrophysiologic study prior to the ablation. The principal maneuver would be to measure the pacing threshold at each point along the ablation catheter 20. The electrodes 530 of the ablation catheter 20 can be used for measuring the pacing threshold, or other pacing measuring devices can be used. After the ablation is delivered the operator could retest the capture threshold. The anticipated results would be an increase in the local pacing threshold. Furthermore a more standard electrophysiologic study can be performed using pacing electrodes in the pericardial space and/or standard

diagnostic electrophysiologic catheters in the right atria, coronary sinus and right ventricle. Conformation that the pulmonary veins are electrically uncoupled from the rest of the left atria is a standard clinical practice. Atrial pacing form inside the lesion boundary can be performed using a remote stimulus electrode, which can optionally be a part of the loop tensioner 524. When there is evidence of conduction outside the lesion (as evidenced by capture of the atria), the operator can evaluate the local electrograms to identify potential gaps in the lesion. It is contemplated that the extended bipolar arrangement of the electrodes 530 can be useful in determining timing and direction of local depolarization. Electrodes overlaying these potential incomplete ablation sites can be identified and additional energy can be delivered as needed.

Once complete electrophysiologic block around the pulmonary veins is verified, it is further contemplated that the ablation catheter 20 can also be used to evaluate autonomic ganglia that are common along this path. These potential targets can be identified with neuro-stimulus techniques and evaluation of epicardial signals. The operator can choose to deliver RF ablation to these select sites, if desired. After the ablation is complete, it is contemplated that the ablation catheter 20 can be removed or repositioned to create lesions at additional ablation target sites.

As described herein, the ablation catheter 20 is an over-the-wire ablation catheter with an array of multiple electrodes 530 located on its mid (central) portion 508. The ablation catheter 20 can be more flexible than other clinically available catheter-based ablation devices to permit tissue contact around the left atrial structures. The electrodes 530 can be capable of monitoring and/or delivering RF energy, electroporation impulses, and programmed cardiac pacing and/or neuro-stimulus. The ability of the disclosed ablation catheter 20 to individualize the as-extended bipolar electrode 530 can take advantage of the natural geometry inside the pericardial space to deliver energy to a series of electrodes arranged around the target structure.

In use, once the ablation catheter 20 is deployed, it is contemplated that a linear lesion can be created without need to reposition the catheter 20. It is further contemplated that the ablation catheter 20 can provide a stable and contiguous array of electrodes 530 along the target path that can deliver ablation energy and can also be used to confirm electrophysiologic block using an extended bipolar electrocardiographic technique. It is contemplated that the use of high impedance structures 540 positioned along the bipolarly aligned electrodes can further adjust the density of the current applied. It is contemplated that the ability to perform the entire procedure without repositioning of the ablation catheter 20 can save time and potentially make this approach more effective than standard point-by-point techniques, which often require frequent repositioning and/or advanced noncontact mapping techniques to identify incomplete segments in the ablation lesion. For epicardial techniques performed from the pericardial space, such manipulation is fraught with danger and technical limitations. The disclosed ablation catheter 20 takes advantage of the natural contours of the left atrial epicardial surface to provide reliable and stable electrode contact.

As will be appreciated by one skilled in the art, the methods and systems described above in relation to the ablation catheter system 600 may take the form of an entirely hardware embodiment, an entirely software embodiment, or an embodiment combining software and hardware aspects. Furthermore, the methods and systems may take the form of a computer program product on a computer-readable

storage medium having computer-readable program instructions (e.g., computer software) embodied in the storage medium. More particularly, the present methods and systems may take the form of web-implemented computer software. Any suitable computer-readable storage medium may be utilized including hard disks, CD-ROMs, optical storage devices, or magnetic storage devices.

Some embodiments of the methods and systems discussed above and below can be described with reference to block diagrams and flowchart illustrations of methods, systems, apparatuses and computer program products. It will be understood that each block of the block diagrams and flowchart illustrations, and combinations of blocks in the block diagrams and flowchart illustrations, respectively, can be implemented by computer program instructions. These computer program instructions may be loaded onto a general purpose computer, special purpose computer, or other programmable data processing apparatus to produce a machine, such that the instructions which execute on the computer or other programmable data processing apparatus create a means for implementing the functions specified in the flowchart block or blocks.

These computer program instructions may also be stored in a computer-readable memory that can direct a computer or other programmable data processing apparatus to function in a particular manner, such that the instructions stored in the computer-readable memory produce an article of manufacture including computer-readable instructions for implementing the function specified in the flowchart block or blocks. The computer program instructions may also be loaded onto a computer or other programmable data processing apparatus to cause a series of operational steps to be performed on the computer or other programmable apparatus to produce a computer-implemented process such that the instructions that execute on the computer or other programmable apparatus provide steps for implementing the functions specified in the flowchart block or blocks.

Accordingly, blocks of the block diagrams and flowchart illustrations support combinations of means for performing the specified functions, combinations of steps for performing the specified functions and program instruction means for performing the specified functions. It will also be understood that each block of the block diagrams and flowchart illustrations, and combinations of blocks in the block diagrams and flowchart illustrations, can be implemented by special purpose hardware-based computer systems that perform the specified functions or steps, or combinations of special purpose hardware and computer instructions.

The methods and systems that have been introduced above, and discussed in further detail below, have been and will be described as comprised of units. One skilled in the art will appreciate that this is a functional description and that the respective functions can be performed by software, hardware, or a combination of software and hardware. A unit can be software, hardware, or a combination of software and hardware. The units can comprise the ablation control software 806 as illustrated in FIG. 44 and described below. In one exemplary aspect, the units can comprise a computer 800 as illustrated in FIG. 44 and described below.

FIG. 44 is a block diagram illustrating an exemplary operating environment for performing the disclosed methods. This exemplary operating environment is only an example of an operating environment and is not intended to suggest any limitation as to the scope of use or functionality of operating environment architecture. Neither should the operating environment be interpreted as having any depen-

gency or requirement relating to any one or combination of components illustrated in the exemplary operating environment.

The present methods and systems can be operational with numerous other general purpose or special purpose computing system environments or configurations. Examples of well known computing systems, environments, and/or configurations that can be suitable for use with the systems and methods comprise, but are not limited to, personal computers, server computers, laptop devices, and multiprocessor systems. Additional examples comprise set top boxes, programmable consumer electronics, network PCs, minicomputers, mainframe computers, distributed computing environments that comprise any of the above systems or devices, and the like.

The processing of the disclosed methods and systems can be performed by software components. The disclosed systems and methods can be described in the general context of computer-executable instructions, such as program modules, being executed by one or more computers or other devices. Generally, program modules comprise computer code, routines, programs, objects, components, data structures, etc. that perform particular tasks or implement particular abstract data types. The disclosed methods can also be practiced in grid-based and distributed computing environments where tasks are performed by remote processing devices that are linked through a communications network. In a distributed computing environment, program modules can be located in both local and remote computer storage media including memory storage devices.

Further, one skilled in the art will appreciate that the systems and methods disclosed herein can be implemented via a general-purpose computing device in the form of a computer **800**. The components of the computer **800** can comprise, but are not limited to, one or more processors or processing units **803**, a system memory **808**, and a system bus **813** that couples various system components including the processor **803** to the system memory **808**. In the case of multiple processing units **803**, the system can utilize parallel computing.

The system bus **813** represents one or more of several possible types of bus structures, including a memory bus or memory controller, a peripheral bus, an accelerated graphics port, and a processor or local bus using any of a variety of bus architectures. By way of example, such architectures can comprise an Industry Standard Architecture (ISA) bus, a Micro Channel Architecture (MCA) bus, an Enhanced ISA (EISA) bus, a Video Electronics Standards Association (VESA) local bus, an Accelerated Graphics Port (AGP) bus, and a Peripheral Component Interconnects (PCI), a PCI-Express bus, a Personal Computer Memory Card Industry Association (PCMCIA), Universal Serial Bus (USB) and the like. The bus **813**, and all buses specified in this description can also be implemented over a wired or wireless network connection and each of the subsystems, including the processor **803**, a mass storage device **804**, an operating system **805**, ablation control software **806**, data **807**, a network adapter **809**, system memory **808**, an Input/Output Interface **812**, a display adapter **810**, a display device **811**, and a human machine interface **802**, can be contained within one or more remote computing devices **814** at physically separate locations, connected through buses of this form, in effect implementing a fully distributed system.

The computer **800** typically comprises a variety of computer readable media. Exemplary readable media can be any available media that is accessible by the computer **800** and comprises, for example and not meant to be limiting, both

volatile and non-volatile media, removable and non-removable media. The system memory **808** comprises computer readable media in the form of volatile memory, such as random access memory (RAM), and/or non-volatile memory, such as read only memory (ROM). The system memory **808** typically contains data such as data **807** and/or program modules such as operating system **805** and ablation control software **806** that are immediately accessible to and/or are presently operated on by the processing unit **803**.

In another aspect, the computer **800** can also comprise other removable/non-removable, volatile/non-volatile computer storage media. By way of example, FIG. 1 illustrates a mass storage device **804** which can provide non-volatile storage of computer code, computer readable instructions, data structures, program modules, and other data for the computer **800**. For example and not meant to be limiting, a mass storage device **804** can be a hard disk, a removable magnetic disk, a removable optical disk, magnetic cassettes or other magnetic storage devices, flash memory cards, CD-ROM, digital versatile disks (DVD) or other optical storage, random access memories (RAM), read only memories (ROM), electrically erasable programmable read-only memory (EEPROM), and the like.

Optionally, any number of program modules can be stored on the mass storage device **804**, including by way of example, an operating system **805** and ablation control software **806**. Each of the operating system **805** and ablation control software **806** (or some combination thereof) can comprise elements of the programming and the ablation control software **806**. Data **807** can also be stored on the mass storage device **804**. Data **807** can be stored in any of one or more databases known in the art. Examples of such databases comprise, DB2®, Microsoft® Access, Microsoft® SQL Server, Oracle®, mySQL, PostgreSQL, and the like. The databases can be centralized or distributed across multiple systems.

In another aspect, the user can enter commands and information into the computer **800** via an input device (not shown). Examples of such input devices comprise, but are not limited to, a keyboard, pointing device (e.g., a "mouse"), a microphone, a joystick, a scanner, tactile input devices such as gloves, and other body coverings, and the like. These and other input devices can be connected to the processing unit **803** via a human machine interface **802** that is coupled to the system bus **813**, but can be connected by other interface and bus structures, such as a parallel port, game port, an IEEE 1394 Port (also known as a Firewire port), a serial port, or a universal serial bus (USB).

In yet another aspect, a display device **811** can also be connected to the system bus **813** via an interface, such as a display adapter **810**. It is contemplated that the computer **800** can have more than one display adapter **810** and the computer **800** can have more than one display device **811**. For example, a display device can be a monitor, an LCD (Liquid Crystal Display), or a projector. In addition to the display device **811**, other output peripheral devices can comprise components such as speakers (not shown) and a printer (not shown) which can be connected to the computer **800** via Input/Output Interface **812**. Any step and/or result of the methods can be output in any form to an output device. Such output can be any form of visual representation, including, but not limited to, textual, graphical, animation, audio, tactile, and the like. Likewise, the routing console **610**, recording console **650**, and signal generator **700** can communicate with the computer **800** and its components through the Input/Output Interface **812**.

The computer **800** can operate in a networked environment using logical connections to the routing console **610**, recording console **650**, and signal generator **700** and/or to one or more remote computing devices **814**. By way of example, a remote computing device can be a personal computer, portable computer, a server, a router, a network computer, a wireless connected tablet or mobile device, a peer device or other common network node, and so on. Logical connections between the computer **800** and a remote computing device **814** can be made via a local area network (LAN) and a general wide area network (WAN). Such network connections can be through a network adapter **809**. A network adapter **809** can be implemented in both wired and wireless environments. Such networking environments are conventional and commonplace in offices, enterprise-wide computer networks, intranets, cellular networks and the Internet **815**.

For purposes of illustration, application programs and other executable program components such as the operating system **805** are illustrated herein as discrete blocks, although it is recognized that such programs and components reside at various times in different storage components of the computing device **800**, and are executed by the data processor(s) of the computer. An implementation of ablation control software **806** can be stored on or transmitted across some form of computer readable media. Any of the disclosed methods can be performed by computer readable instructions embodied on computer readable media. Computer readable media can be any available media that can be accessed by a computer. By way of example and not meant to be limiting, computer readable media can comprise "computer storage media" and "communications media." "Computer storage media" comprise volatile and non-volatile, removable and non-removable media implemented in any methods or technology for storage of information such as computer readable instructions, data structures, program modules, or other data. Exemplary computer storage media comprises, but is not limited to, RAM, ROM, EEPROM, flash memory or other memory technology, CD-ROM, digital versatile disks (DVD) or other optical storage, magnetic cassettes, magnetic tape, magnetic disk storage or other magnetic storage devices, or any other medium which can be used to store the desired information and which can be accessed by a computer.

The methods and systems can employ Artificial Intelligence techniques such as machine learning and iterative learning. Examples of such techniques include, but are not limited to, expert systems, case based reasoning, Bayesian networks, behavior based AI, neural networks, fuzzy systems, evolutionary computation (e.g. genetic algorithms), swarm intelligence (e.g. ant algorithms), and hybrid intelligent systems (e.g. Expert inference rules generated through a neural network or production rules from statistical learning).

The proposed procedures are performed under conscious sedation and local anesthesia in a standard cardiac catheterization laboratory. The patient is prepped in the typical manner for an electrophysiologic study with an additional sterile field exposing the anterior chest and upper abdomen. Stimulus and mapping catheters are positioned in the RA, RV, and CS position. Percutaneous access to the pericardial space is achieved using a modified Seldinger technique or clinically available pericardial access tool. A small volume of iodinated contrast is injected into the pericardial space for visualization of key cardiac landmarks. The percutaneous track is expanded to accommodate catheter insertion. The clinical goal of the procedure will be to position a multi-

electrode ablation catheter within the pericardial space for the purpose of ablation. The catheter will follow a course that circumferentially divides the more anterior left atrial structures from the pulmonary veins. Once in a stable position, the catheter's multi-electrode array will be used to deliver a single linear ablation lesion that can electrophysiologically isolate arrhythmogenic substrate of pulmonary veins from the greater left atrium.

As further described herein, it is contemplated that epicardial positioning the ablation catheter **20** can have mechanical advantages over endocardial multi-electrode arrays. The ablation catheter **20** can tailor the circumference of the loop formed by the elongate shaft **500** of the catheter **20** with little effort to provide full coverage. The flexibility of the ablation catheter **20** can provide a mechanism for secure tissue contact around complex anatomic geometry. It is further contemplated that the natural spatial limitation of the pericardial space provides a natural mechanism to assure electrode approximation. Furthermore, the risks of performing ablation from the epicardial surface place the ablation electrode **530** closer to some important bystander structures that necessitate the delivery of ablative energy with programmed directional vectors. (See FIG. **23**). With RF energy ablation, extended bipolar ablation can result in 40-50% deeper lesion in the direction of the programmed vector. With IE ablation, the potential for creating a preferential directional injury vector can be greater because there is limited or no thermal energy. Typically, unipolar applications utilize an externalized grounding pad that results in a diffuse or spherical virtual electrode, while currently known bipolar ablation techniques typically utilize electrode pairs that are in very close proximity, require equipment is cumbersome, and require entry into both the pericardium and the left atrial blood pool.

In exemplary aspects, it is contemplated that the ablation catheter **20** can be modified to deliver gene therapy. In these aspects, it is contemplated that the elongate shaft **500** of the ablation catheter **20** can be modified to have irrigation side ports. It is further contemplated that a DNA or RNA vector can be delivered via the catheter using a tailored electroporation impulse.

In other exemplary aspects, it is contemplated that the ablation catheter **20** can be employed in a method for prostate ablation. In these aspects, it is contemplated that, in patients with benign prostatic hypertrophy and urinary obstruction, the ablation catheter **20** can be positioned to deliver irreversible electroporation impulses in an extended bipolar or unipolar configuration. High impedance structures **540** can be further utilized by the ablation catheter **20** in an extended bipolar configuration to increase the density current at targeted areas. In use, the ablation catheter can be advanced over a guide wire **300** that has been delivered into the bladder non-traumatically. It is contemplated that this technique can provide substantial advantages over current procedures, which are typically traumatic to the transitional endothelium of the urethra. With irreversible electroporation, it is contemplated that the impulse can be tailored to minimize inflammation and damage to the greater tissue architecture.

In other exemplary aspects, it is contemplated that the ablation catheter **20** can be used to preserve erectile function. In these aspects, the ablation catheter **20** can be used to ablate selected nerve axons.

In further exemplary aspects, it is contemplated that the ablation catheter **20** can be configured for therapy for solid tumors. Typically, current electroporation devices are created to place a pair of needle electrodes into the tumor using

open and minimally-invasive surgical techniques. However, it is contemplated that the ablation catheter **20**, with its over-the-wire electrode array, can be used in treating tumors which can be accessed through the vascular space (e.g., palliative therapy for renal cell carcinoma that is extending into the vena cava).

In still further exemplary aspects, it is contemplated that the ablation catheter **20** can be used to treat pulmonary hypertension where there is substantial endothelial remodeling and hypertrophy of the pulmonary vascular structures. In these aspects, the ablation catheter **20** can be used to “prune” the smooth muscle mass in these hypertrophied vessels and potentially lead to a favorable remodeling. It is contemplated that the electrodes of the ablation catheter **20** can be advanced around the hilum of the kidneys (using laparoscopic techniques) for purposes performing renal denervation and managing malignant refractory hypertension.

Although several embodiments of the invention have been disclosed in the foregoing specification, it is understood by those skilled in the art that many modifications and other embodiments of the invention will come to mind to which the invention pertains, having the benefit of the teaching presented in the foregoing description and associated drawings. It is thus understood that the invention is not limited to the specific embodiments disclosed hereinabove, and that many modifications and other embodiments are intended to be included within the scope of the appended claims. Moreover, although specific terms are employed herein, as well as in the claims which follow, they are used only in a generic and descriptive sense, and not for the purposes of limiting the described invention, nor the claims which follow.

What is claimed is:

1. A method, comprising:

positioning a distal end of a first catheter proximate to a first side of a first pericardial reflection of a subject, the first catheter including a proximal portion, the distal end of the first catheter defining a first opening, the first catheter further including a first lumen extending there-through, the first lumen formed between the opening of the distal end of the first catheter and the proximal portion of the first catheter, the first catheter including a first magnet assembly coupled to the distal end of the first catheter; and

positioning a distal end of a second catheter proximate to a second side of the first pericardial reflection, the second catheter including a proximal portion, the distal end of the second catheter defining a second opening, the second catheter further including a second lumen extending therethrough, the second lumen formed between the opening of the distal end of the second catheter and the proximal portion of the second catheter, the second catheter including a second magnet assembly coupled to the distal end of the second catheter, such that the first magnet assembly and the second magnet assembly are magnetically coupled across the first pericardial reflection, and such that the lumen of the first catheter is substantially axially aligned with the lumen of the second catheter;

piercing the first pericardial reflection by advancing a needle through the first lumen of the first catheter, through the first pericardial reflection, and into the second lumen of the second catheter to generate a pierced portion in the first pericardial reflection;

withdrawing the needle from the second lumen of the second catheter and from the first pericardial reflection;

positioning the distal end of the first catheter proximate to a first side of a second pericardial reflection of the subject;

positioning the distal end of the second catheter proximate to a second side of the second pericardial reflection;

piercing the second pericardial reflection by advancing the needle through the first lumen of the first catheter, through the second pericardial reflection, and into the second lumen of the second catheter to generate a pierced portion in the second pericardial reflection;

withdrawing the needle from the second lumen of the second catheter and from the second pericardial reflection;

delivering a guidewire through the first lumen of the first catheter, the pierced portion in the first pericardial reflection, the pierced portion in the second pericardial reflection, and the second lumen of the second catheter;

withdrawing the first catheter and the second catheter from the body of the subject while leaving the guidewire in place;

positioning a third catheter in the pericardial space of the heart of the subject by passing the third catheter over the guidewire, through the pierced portion of the first pericardial reflection and through the pierced portion of the second pericardial reflection, such that a central portion of the third catheter at least partially encircles the left pulmonary veins and the right pulmonary veins, the central portion of the third catheter including a set of electrodes; and

delivering electrical energy to at least a portion of the left atrium of the heart of the subject using two or more electrodes of the set of electrodes.

2. The method of claim **1**, the delivering the electrical energy including ablating tissue in the portion of the left atrium to create a lesion.

3. The method of claim **1**, the delivering the electrical energy including ablating tissue in the portion of the left atrium to create a circumferential lesion without repositioning the third catheter.

4. The method of claim **1**, the delivering the electrical energy including ablating tissue in the portion of the left atrium to create a lesion via irreversible electroporation.

5. The method of claim **1**, the delivering the electrical energy including delivering high voltage ultra short direct current impulses (HVUS-DCI).

6. The method of claim **1**, the delivering the electrical energy including delivering the electrical energy in synchronization with the cardiac cycle of the subject.

7. The method of claim **1**, further comprising monitoring the cardiac cycle of the subject, the delivering the electrical energy including delivering the electrical energy in synchronization with the cardiac cycle of the subject.

8. The method of claim **1**, further comprising monitoring the cardiac cycle of the subject using one or more electrodes of the set of electrodes, the delivering the electrical energy including delivering the electrical energy in synchronization with the cardiac cycle of the subject.

9. The method of claim **1**, the electrical energy including one or more current impulses, the delivering the electrical energy including delivering one or more current impulses for every heartbeat of the subject.

10. The method of claim **1**, wherein the portion of the left atrium is a first portion, further comprising delivering electrical energy to a second portion of the left atrium of the heart of the subject using two or more electrodes of the set of electrodes.

11. The method of claim 1, wherein the portion of the left atrium is a first portion, the positioning the third catheter including positioning the third catheter in a first position, further comprising:

repositioning the third catheter in a second position different from the first position, such that the central portion of the third catheter at least partially encircles the left pulmonary veins and the right pulmonary veins in the pericardial space of the heart of the subject; and delivering electrical energy to a second portion of the left atrium of the heart of the subject using two or more electrodes of the set of electrodes.

12. The method of claim 1, wherein the delivering includes:

identifying one or more intended current vectors for delivering the electrical energy; selecting, based on the one or more intended current vectors, the two or more electrodes; and delivering the electrical energy using the two or more electrodes and along the one or more current vectors.

13. The method of claim 1, the positioning the distal end of the first catheter including inserting the first catheter into the body of the subject using a subxiphoid pericardial access point, and the positioning the distal end of the second catheter including inserting the second catheter into the body of the subject using the subxiphoid pericardial access point.

14. The method of claim 13, the delivering the electrical energy including ablating tissue in the portion of the left atrium to create a lesion around an ostium or ostia of one or more of the left pulmonary veins and the right pulmonary veins formed in the left atrium.

15. An apparatus, comprising:

a first catheter including a proximal portion and a distal end, the distal end of the first catheter defining a first opening, the first catheter further including a first lumen extending therethrough, the first lumen formed between the opening of the distal end of the first catheter and the proximal portion of the first catheter, the first catheter including a first magnet assembly coupled to the distal end of the first catheter, the distal end of the first catheter configured to be disposed proximate to a first side of a first pericardial reflection of a subject;

a second catheter including a proximal portion and a distal end, the distal end of the second catheter defining a second opening, the second catheter further including a second lumen extending therethrough, the second lumen formed between the opening of the distal end of the second catheter and the proximal portion of the second catheter, the second catheter including a second magnet assembly coupled to the distal end of the second catheter, such that during use:

the distal end of the second catheter is configured to be disposed proximate to a second side of the first pericardial reflection; and

the first magnet assembly and the second magnet assembly are configured to magnetically couple across the first pericardial reflection, such that the lumen of the first catheter is substantially axially aligned with the lumen of the second catheter;

a needle configured to be positioned within the first lumen and configured for movement within the first lumen, the needle including a distal puncturing surface, such

that during use, the distal puncturing surface of the needle is configured to puncture the first pericardial reflection and to be disposed within the second lumen of the second catheter to generate a pierced portion in the first pericardial reflection, and to be subsequently withdrawn from the second lumen of the second catheter and from the first pericardial reflection,

the first catheter, second catheter, and needle further configured to be repositioned during use to puncture a second pericardial reflection of the subject to generate a pierced portion in the second pericardial reflection; and

a third catheter including a flexible shaft having a central portion, the third catheter configured to be positioned during use in the pierced portion of the first pericardial reflection and in the pierced portion of the second pericardial reflection, the third catheter further including a set of electrodes disposed along the central portion, the central portion configured to at least partially encircle the left pulmonary veins and the right pulmonary veins, the set of electrodes configured to deliver electrical energy to at least a portion of the left atrium of the heart of the subject using two or more electrodes of the set of electrodes.

16. The apparatus of claim 15, the electrical energy including one or more high voltage ultra short direct current impulses (HVUS-DCI).

17. The apparatus of claim 15, further comprising:

a routing console including one or more connectors operably coupled to the set of electrodes; and

a synchronization trigger operably coupled to the routing console, the synchronization trigger configured for identifying timing information associated with a cardiac cycle of the subject, the synchronization trigger further configured for communicating an indication of the timing information to the routing console,

the routing console further configured for selectively activating the two or more electrodes of the set of electrodes based on the indication of timing information.

18. The apparatus of claim 15, the set of electrodes including one or more pacing electrodes, further comprising:

a routing console including one or more connectors operably coupled to the set of electrodes; and

a synchronization trigger operably coupled to the one or more pacing electrodes and the routing console, the synchronization trigger configured for:

receiving cardiac cycle information associated with the subject from the one or more pacing electrodes;

identifying timing information based on the cardiac cycle information; and

communicating an indication of the timing information to the routing console,

the routing console further configured for selectively activating the two or more electrodes of the set of electrodes based on the indication of timing information.

19. The apparatus of claim 15, the electrical energy including one or more current impulses, the routing console further configured for selectively activating the two or more electrodes for delivering one or more current impulses for every heartbeat of the subject.