



US009861802B2

(12) **United States Patent**
Mickelsen

(10) **Patent No.:** **US 9,861,802 B2**
(45) **Date of Patent:** **Jan. 9, 2018**

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

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

(71) Applicant: **UNIVERSITY OF IOWA RESEARCH FOUNDATION**, Iowa City, IA (US)

See application file for complete search history.

(72) Inventor: **Steven Richard Mickelsen**, Iowa City, IA (US)

(56) **References Cited**

(73) Assignee: **University of Iowa Research Foundation**, Iowa City, IA (US)

U.S. PATENT DOCUMENTS

5,281,213 A 1/1994 Milder et al.
5,334,193 A 8/1994 Nardella

(Continued)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 90 days.

FOREIGN PATENT DOCUMENTS

EP 1125549 8/2001
EP 0797956 6/2003

(Continued)

(21) Appl. No.: **14/400,455**

(22) PCT Filed: **Mar. 14, 2013**

OTHER PUBLICATIONS

(86) PCT No.: **PCT/US2013/031252**

Supplementary European Search Report for European Application No. 13827672.0, dated Jul. 11, 2016, 12 pages.

§ 371 (c)(1),

(2) Date: **Nov. 11, 2014**

(Continued)

(87) PCT Pub. No.: **WO2014/025394**

Primary Examiner — Aarti B Berdichevsky

PCT Pub. Date: **Feb. 13, 2014**

(74) *Attorney, Agent, or Firm* — Smith, Gambrell & Russell

(65) **Prior Publication Data**

US 2015/0182740 A1 Jul. 2, 2015

Related U.S. Application Data

(60) Provisional application No. 61/681,552, filed on Aug. 9, 2012.

(51) **Int. Cl.**

A61B 18/14 (2006.01)

A61M 39/10 (2006.01)

(Continued)

(57) **ABSTRACT**

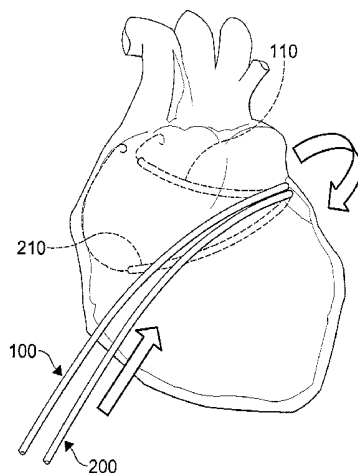
A percutaneous catheter system for use within the human body and an ablation catheter for ablating a selected tissue region within the body of a subject. The percutaneous catheter system can include two catheters that are operatively coupled to one another by magnetic coupling through a tissue structure. 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 struc-

(Continued)

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

CPC **A61M 39/10** (2013.01); **A61B 5/0538** (2013.01); **A61B 5/4833** (2013.01);

(Continued)



ture using the percutaneous catheter system are disclosed. Also disclosed are methods for ablating a selected tissue region using the ablation catheter.

17 Claims, 43 Drawing Sheets

(51) Int. Cl.

- A61B 5/053* (2006.01)
- A61B 5/00* (2006.01)
- A61B 18/12* (2006.01)
- A61M 25/01* (2006.01)
- A61M 25/06* (2006.01)
- A61M 25/00* (2006.01)
- A61B 17/22* (2006.01)
- A61B 18/02* (2006.01)
- A61B 18/18* (2006.01)
- A61B 18/20* (2006.01)
- A61B 18/00* (2006.01)
- A61B 34/00* (2016.01)
- A61B 17/00* (2006.01)

(52) U.S. Cl.

- CPC *A61B 18/1206* (2013.01); *A61B 18/1492* (2013.01); *A61M 25/0169* (2013.01); *A61M 25/0606* (2013.01); *A61B 17/22004* (2013.01); *A61B 18/02* (2013.01); *A61B 18/1815* (2013.01); *A61B 18/20* (2013.01); *A61B 2017/00876* (2013.01); *A61B 2018/00363* (2013.01); *A61B 2018/00375* (2013.01); *A61B 2018/00577* (2013.01); *A61B 2018/128* (2013.01); *A61B 2034/731* (2016.02); *A61B 2217/005* (2013.01); *A61B 2218/002* (2013.01); *A61M 2025/0089* (2013.01)

6,119,041	A	9/2000	Pomeranz et al.	
6,120,500	A	9/2000	Bednarek et al.	
6,146,381	A	11/2000	Bowe et al.	
6,164,283	A	12/2000	Lesh	
6,231,518	B1	5/2001	Grabek et al.	
6,251,107	B1	6/2001	Schaer	
6,251,128	B1	6/2001	Knopp et al.	
6,270,476	B1	8/2001	Santoianni et al.	
6,272,384	B1	8/2001	Simon et al.	
6,287,306	B1	9/2001	Kroll et al.	
6,314,963	B1	11/2001	Vaska et al.	
6,370,412	B1	4/2002	Armoundas et al.	
6,391,024	B1	5/2002	Sun et al.	
6,464,699	B1	10/2002	Swanson	
6,470,211	B1	10/2002	Ideker et al.	
6,502,576	B1	1/2003	Lesh	
6,503,247	B2	1/2003	Swartz et al.	
6,517,534	B1	2/2003	McGovern et al.	
6,527,724	B1	3/2003	Fenici	
6,527,767	B2	3/2003	Wang et al.	
6,592,581	B2	7/2003	Bowe	
6,595,991	B2	7/2003	Tollner et al.	
6,607,520	B2	8/2003	Keane	
6,623,480	B1	9/2003	Kuo et al.	
6,638,278	B2	10/2003	Falwell et al.	
6,666,863	B2	12/2003	Wentzel et al.	
6,669,693	B2	12/2003	Friedman	
6,702,811	B2 *	3/2004	Stewart	A61B 18/1492 606/41

(56)

References Cited

U.S. PATENT DOCUMENTS

5,341,807	A	8/1994	Nardella	
5,454,370	A	10/1995	Avitall	
5,545,161	A	8/1996	Imran	
5,617,854	A	4/1997	Munsif	
5,672,170	A	9/1997	Cho	
5,702,438	A	12/1997	Avitall	
5,706,823	A	1/1998	Wodlinger	
5,722,400	A	3/1998	Ockuly et al.	
5,779,699	A	7/1998	Lipson	
5,788,692	A	8/1998	Campbell et al.	
5,810,762	A	9/1998	Hofmann	
5,843,154	A	12/1998	Osypka	
5,849,028	A	12/1998	Chen	
5,871,523	A	2/1999	Fleischman et al.	
5,876,336	A	3/1999	Swanson et al.	
5,895,404	A	4/1999	Ruiz	
5,916,158	A	6/1999	Webster, Jr.	
5,921,924	A	7/1999	Avitall	
5,928,269	A	7/1999	Alt	
5,928,270	A	7/1999	Ramsey, III	
6,002,955	A	12/1999	Willems et al.	
6,006,131	A	12/1999	Cooper et al.	
6,009,351	A	12/1999	Flachman	
6,014,579	A	1/2000	Pomeranz et al.	
6,033,403	A	3/2000	Tu et al.	
6,045,550	A	4/2000	Simpson et al.	
6,068,653	A	5/2000	LaFontaine	
6,071,281	A	6/2000	Burnside et al.	
6,090,104	A	7/2000	Webster, Jr.	
6,096,036	A	8/2000	Bowe et al.	
6,113,595	A	9/2000	Muntermann	
7,001,383	B2	2/2006	Keidar	
7,171,263	B2	1/2007	Darvish et al.	
7,182,725	B2 *	2/2007	Bonan	A61N 5/1002 600/3
7,195,628	B2	3/2007	Falkenberg	
7,207,988	B2	4/2007	Leckrone et al.	
7,207,989	B2	4/2007	Pike, Jr. et al.	
7,229,402	B2	6/2007	Diaz et al.	
7,229,437	B2	6/2007	Johnson et al.	
7,250,049	B2	7/2007	Roop et al.	
7,285,116	B2	10/2007	de la Rama et al.	
7,285,119	B2 *	10/2007	Stewart	A61B 18/1492 606/41
7,326,208	B2 *	2/2008	Vanney	A61B 18/1492 606/41
7,346,379	B2	3/2008	Eng et al.	
7,387,629	B2 *	6/2008	Vanney	A61B 18/1492 606/41
7,387,630	B2	6/2008	Mest	
7,387,636	B2	6/2008	Cohn et al.	
7,416,552	B2	8/2008	Paul et al.	
7,419,477	B2	9/2008	Simpson et al.	
7,419,489	B2 *	9/2008	Vanney	A61B 18/1492 606/41
7,429,261	B2	9/2008	Kunis et al.	
7,435,248	B2	10/2008	Taimisto et al.	
7,527,625	B2 *	5/2009	Knight	A61B 18/1492 606/41
7,578,816	B2	8/2009	Boveja et al.	
7,588,567	B2	9/2009	Boveja et al.	

(56)		References Cited			
		U.S. PATENT DOCUMENTS			
				8,926,589 B2	1/2015 Govari
				8,932,287 B2	1/2015 Gelbart et al.
				8,945,117 B2	2/2015 Bencini
				8,979,841 B2 *	3/2015 Kunis A61B 18/1815
					606/41
7,623,899 B2	11/2009	Worley et al.		8,986,278 B2	3/2015 Fung et al.
7,678,108 B2 *	3/2010	Chrisitian A61B 18/1492		9,002,442 B2	4/2015 Harley et al.
				9,005,189 B2	4/2015 Davalos et al.
				9,005,194 B2	4/2015 Oral et al.
7,681,579 B2 *	3/2010	Schwartz A61B 8/12		9,011,425 B2	4/2015 Fischer et al.
				9,044,245 B2	6/2015 Condie et al.
				9,055,959 B2	6/2015 Vaska et al.
7,771,421 B2 *	8/2010	Stewart A61B 18/1492		9,072,518 B2	7/2015 Swanson
				9,078,667 B2	7/2015 Besser et al.
				9,101,374 B1	8/2015 Hoch et al.
				9,119,533 B2	9/2015 Ghaffari
				9,119,634 B2	9/2015 Gelbart et al.
7,805,182 B2	9/2010	Weese et al.		9,131,897 B2	9/2015 Harada et al.
7,850,642 B2	12/2010	Moll et al.		9,155,590 B2	10/2015 Mathur
7,850,685 B2 *	12/2010	Kunis A61B 18/1815		9,162,037 B2	10/2015 Belson et al.
				9,179,972 B2	11/2015 Olson
				9,186,481 B2	11/2015 Avitall et al.
7,857,808 B2	12/2010	Oral et al.		9,192,769 B2	11/2015 Donofrio et al.
7,857,809 B2	12/2010	Drysen		9,211,405 B2	12/2015 Mahapatra et al.
7,869,865 B2	1/2011	Govari et al.		9,216,055 B2	12/2015 Spence et al.
7,896,873 B2	3/2011	Hiller et al.		9,233,248 B2	1/2016 Luther et al.
7,917,211 B2	3/2011	Zacouto		9,237,926 B2	1/2016 Nollert et al.
7,918,819 B2	4/2011	Karmarkar et al.		9,262,252 B2	2/2016 Kirkpatrick et al.
7,918,850 B2	4/2011	Govari et al.		9,277,957 B2	3/2016 Long et al.
7,922,714 B2	4/2011	Stevens-Wright		9,282,910 B2	3/2016 Narayan et al.
7,955,827 B2	6/2011	Rubinsky et al.		9,289,258 B2	3/2016 Cohen
8,048,067 B2	11/2011	Davalos et al.		9,289,606 B2	3/2016 Paul et al.
8,048,072 B2	11/2011	Verin et al.		9,295,516 B2	3/2016 Pearson et al.
8,100,900 B2	1/2012	Prinz et al.		9,301,801 B2	4/2016 Scheib
8,108,069 B2	1/2012	Stahler et al.		9,375,268 B2	6/2016 Long
8,133,220 B2	3/2012	Lee et al.		9,414,881 B2	8/2016 Callas et al.
8,137,342 B2	3/2012	Crossman		9,468,495 B2 *	10/2016 Kunis A61B 18/1815
8,145,289 B2	3/2012	Calabro' et al.		9,474,574 B2	10/2016 Ibrahim et al.
8,147,486 B2	4/2012	Honour et al.		9,480,525 B2	11/2016 Lopes et al.
8,160,690 B2	4/2012	Wilfley et al.		9,486,272 B2	11/2016 Bonyak et al.
8,175,680 B2	5/2012	Panescu		9,486,273 B2	11/2016 Lopes et al.
8,206,384 B2	6/2012	Falwell et al.		9,492,227 B2	11/2016 Lopes et al.
8,206,385 B2	6/2012	Stangenes et al.		9,492,228 B2	11/2016 Lopes et al.
8,216,221 B2	7/2012	Ibrahim et al.		9,517,103 B2	12/2016 Panescu et al.
8,221,411 B2	7/2012	Francischelli et al.		9,526,573 B2	12/2016 Lopes et al.
8,226,648 B2	7/2012	Paul et al.		9,532,831 B2	1/2017 Reinders et al.
8,228,065 B2	7/2012	Wirtz et al.		9,539,010 B2	1/2017 Gagner et al.
8,235,986 B2	8/2012	Kulesa et al.		9,554,848 B2 *	1/2017 Stewart A61B 18/1492
8,235,988 B2	8/2012	Davis et al.		9,700,368 B2	7/2017 Callas et al.
8,251,986 B2	8/2012	Chornenky et al.		2001/0007070 A1 *	7/2001 Stewart A61B 18/1492
8,282,631 B2	10/2012	Davalos et al.			606/41
8,287,532 B2	10/2012	Carroll et al.		2001/0044624 A1	11/2001 Seraj et al.
8,414,508 B2	4/2013	Thapliyal et al.		2002/0111618 A1 *	8/2002 Stewart A61B 18/1492
8,430,875 B2	4/2013	Ibrahim et al.			606/41
8,433,394 B2	4/2013	Harlev et al.		2002/0169445 A1	11/2002 Jain et al.
8,449,535 B2	5/2013	Deno et al.		2002/0177765 A1	11/2002 Bowe et al.
8,454,594 B2	6/2013	Demarais et al.		2002/0183638 A1	12/2002 Swanson
8,463,368 B2	6/2013	Harlev et al.		2003/0204161 A1	10/2003 Ferek Petric
8,475,450 B2	7/2013	Govari et al.		2003/0229379 A1	12/2003 Ramsey
8,486,063 B2	7/2013	Werneth et al.		2004/0039382 A1	2/2004 Kroll et al.
8,500,733 B2	8/2013	Watson		2004/0049181 A1 *	3/2004 Stewart A61B 18/1492
8,535,304 B2	9/2013	Sklar et al.			606/41
8,538,501 B2	9/2013	Venkatachalam et al.		2004/0049182 A1	3/2004 Koblish et al.
8,562,588 B2	10/2013	Hobbs et al.		2004/0082948 A1 *	4/2004 Stewart A61B 18/1492
8,568,406 B2	10/2013	Harlev et al.			606/41
8,571,635 B2	10/2013	McGee		2004/0199157 A1	10/2004 Palanker et al.
8,571,647 B2	10/2013	Harlev et al.		2004/0236360 A1	11/2004 Cohn et al.
8,585,695 B2	11/2013	Shih		2004/0254607 A1	12/2004 Wittenberger et al.
8,597,288 B2	12/2013	Christian		2004/0267337 A1	12/2004 Hayzelden
8,608,735 B2	12/2013	Govari et al.		2005/0033282 A1	2/2005 Hooven
8,628,522 B2	1/2014	Ibrahim et al.		2005/0222632 A1	10/2005 Obino
8,632,534 B2	1/2014	Pearson et al.		2005/0261672 A1 *	11/2005 Deem A61B 18/1492
8,647,338 B2	2/2014	Chornenky et al.			606/41
8,708,952 B2	4/2014	Cohen et al.		2006/0009755 A1	1/2006 Sra
8,734,442 B2	5/2014	Cao et al.		2006/0217703 A1	9/2006 Chornenky et al.
8,771,267 B2 *	7/2014	Kunis A61B 18/1815		2006/0270900 A1	11/2006 Chin et al.
				2006/0287648 A1	12/2006 Schwartz
				2006/0293730 A1 *	12/2006 Rubinsky A61B 18/1492
					607/98
8,808,273 B2	8/2014	Caples et al.			
8,834,461 B2	9/2014	Werneth et al.			
8,834,464 B2 *	9/2014	Stewart A61B 18/1492			
8,868,169 B2	10/2014	Narayan et al.			
8,876,817 B2	11/2014	Avitall et al.			
8,886,309 B2	11/2014	Luther et al.			
8,903,488 B2	12/2014	Callas et al.			

(56)

References Cited

U.S. PATENT DOCUMENTS

2007/0005053	A1	1/2007	Dando	2013/0317385	A1	11/2013	Sklar et al.
2007/0021744	A1	1/2007	Creighton	2013/0331831	A1	12/2013	Werneth et al.
2007/0129760	A1	6/2007	Demarais et al.	2013/0338467	A1	12/2013	Grasse et al.
2007/0156135	A1	7/2007	Rubinsky et al.	2014/0005664	A1	1/2014	Govari et al.
2007/0167740	A1	7/2007	Grunewald et al.	2014/0024911	A1	1/2014	Harlev et al.
2007/0173878	A1	7/2007	Heuser	2014/0039288	A1	2/2014	Shih
2007/0208329	A1	9/2007	Ward et al.	2014/0051993	A1	2/2014	McGee
2007/0225589	A1	9/2007	Viswanathan	2014/0052118	A1	2/2014	Laske et al.
2007/0249923	A1	10/2007	Keenan	2014/0052126	A1	2/2014	Long et al.
2007/0270792	A1	11/2007	Hennemann et al.	2014/0052216	A1	2/2014	Long et al.
2008/0033426	A1	2/2008	Machell	2014/0081113	A1	3/2014	Cohen et al.
2008/0065061	A1	3/2008	Viswanathan	2014/0100563	A1	4/2014	Govari et al.
2008/0132885	A1	6/2008	Rubinsky et al.	2014/0107644	A1	4/2014	Falwell et al.
2008/0161789	A1	7/2008	Thao et al.	2014/0142408	A1	5/2014	De La Rama et al.
2008/0200913	A1	8/2008	Viswanathan	2014/0163480	A1	6/2014	Govari et al.
2008/0243214	A1	10/2008	Koblish	2014/0163546	A1	6/2014	Govari et al.
2008/0300574	A1	12/2008	Belson	2014/0171942	A1	6/2014	Werneth et al.
2008/0300588	A1	12/2008	Groth et al.	2014/0180035	A1	6/2014	Anderson
2009/0105654	A1	4/2009	Kurth et al.	2014/0194716	A1	7/2014	Diep et al.
2009/0149917	A1	6/2009	Whitehurst et al.	2014/0194867	A1	7/2014	Fish et al.
2009/0163905	A1	6/2009	Winkler et al.	2014/0200567	A1	7/2014	Cox et al.
2009/0228003	A1	9/2009	Sinelnikov	2014/0235986	A1	8/2014	Harlev et al.
2009/0240248	A1	9/2009	Deford et al.	2014/0235989	A1	8/2014	Wodlinger et al.
2009/0275827	A1	11/2009	Aiken et al.	2014/0243851	A1	8/2014	Cohen et al.
2009/0281477	A1	11/2009	Mikus et al.	2014/0276760	A1	9/2014	Bonyak et al.
2009/0306651	A1	12/2009	Schneider	2014/0276782	A1	9/2014	Paskar
2010/0023004	A1	1/2010	Francischelli et al.	2014/0288556	A1	9/2014	Ibrahim et al.
2010/0137861	A1	6/2010	Soroff et al.	2014/0303721	A1	10/2014	Fung et al.
2010/0191232	A1	7/2010	Boveda	2014/0343549	A1	11/2014	Spear et al.
2010/0241185	A1	9/2010	Mahapatra et al.	2014/0364845	A1	12/2014	Rashidi
2010/0261994	A1	10/2010	Davalos et al.	2014/0371613	A1	12/2014	Narayan et al.
2010/0292687	A1	11/2010	Kauphusman et al.	2015/0005767	A1	1/2015	Werneth et al.
2010/0312300	A1	12/2010	Ryu et al.	2015/0011995	A1	1/2015	Avitall et al.
2011/0028962	A1	2/2011	Werneth et al.	2015/0126840	A1	5/2015	Thakur et al.
2011/0098694	A1	4/2011	Long	2015/0133914	A1	5/2015	Koblish
2011/0106221	A1	5/2011	Neal, II et al.	2015/0138977	A1	5/2015	Dacosta
2011/0144524	A1	6/2011	Fish et al.	2015/0141978	A1	5/2015	Subramaniam et al.
2011/0144633	A1	6/2011	Govari	2015/0142041	A1	5/2015	Kendale et al.
2011/0160785	A1	6/2011	Mori et al.	2015/0148796	A1	5/2015	Bencini
2011/0276047	A1	11/2011	Sklar et al.	2015/0150472	A1	6/2015	Harlev et al.
2011/0276075	A1	11/2011	Fung et al.	2015/0157402	A1	6/2015	Kunis et al.
2011/0288544	A1	11/2011	Verin et al.	2015/0157412	A1	6/2015	Wallace et al.
2011/0313417	A1	12/2011	De La Rama et al.	2015/0173824	A1	6/2015	Davalos et al.
2012/0029512	A1	2/2012	Willard et al.	2015/0173828	A1	6/2015	Avitall
2012/0053581	A1	3/2012	Wittkamp et al.	2015/0174404	A1	6/2015	Rouso et al.
2012/0059255	A1	3/2012	Paul et al.	2015/0196217	A1	7/2015	Harlev et al.
2012/0071872	A1	3/2012	Rubinsky et al.	2015/0223726	A1	8/2015	Harlev et al.
2012/0095459	A1	4/2012	Callas et al.	2015/0230699	A1	8/2015	Berul et al.
2012/0101413	A1	4/2012	Beetel et al.	2015/0258344	A1	9/2015	Tandri et al.
2012/0165667	A1	6/2012	Altmann et al.	2015/0265342	A1	9/2015	Long et al.
2012/0172867	A1	7/2012	Ryu et al.	2015/0265344	A1	9/2015	Aktas et al.
2012/0197100	A1	8/2012	Razavi et al.	2015/0272656	A1	10/2015	Chen
2012/0209260	A1	8/2012	Lambert et al.	2015/0272664	A9	10/2015	Cohen
2012/0220998	A1	8/2012	Long et al.	2015/0272667	A1	10/2015	Govari et al.
2012/0265198	A1	10/2012	Crow et al.	2015/0282729	A1	10/2015	Harlev et al.
2012/0283582	A1	11/2012	Mahapatra et al.	2015/0304879	A1	10/2015	Dacosta
2012/0303019	A1	11/2012	Zhao et al.	2015/0320481	A1	11/2015	Cosman et al.
2012/0310052	A1	12/2012	Mahapatra et al.	2015/0321021	A1	11/2015	Tandri et al.
2012/0310230	A1	12/2012	Willis	2015/0342532	A1	12/2015	Basu et al.
2013/0030430	A1	1/2013	Stewart et al.	2015/0343212	A1	12/2015	Rouso et al.
2013/0060247	A1	3/2013	Sklar et al.	2015/0351836	A1	12/2015	Prutchi
2013/0060248	A1	3/2013	Sklar et al.	2015/0359583	A1	12/2015	Swanson
2013/0079768	A1	3/2013	De Luca et al.	2016/0008061	A1	1/2016	Fung et al.
2013/0090651	A1	4/2013	Smith	2016/0008065	A1	1/2016	Gliner et al.
2013/0103027	A1	4/2013	Sklar et al.	2016/0029960	A1	2/2016	Toth et al.
2013/0103064	A1	4/2013	Arenson et al.	2016/0038772	A1	2/2016	Thapliyal et al.
2013/0131662	A1	5/2013	Wittkamp	2016/0051204	A1	2/2016	Harlev et al.
2013/0158538	A1	6/2013	Govari	2016/0051324	A1	2/2016	Stewart et al.
2013/0172864	A1	7/2013	Ibrahim et al.	2016/0058506	A1	3/2016	Spence et al.
2013/0172875	A1	7/2013	Govari et al.	2016/0066993	A1	3/2016	Avitall et al.
2013/0226174	A1	8/2013	Ibrahim et al.	2016/0074679	A1	3/2016	Thapliyal et al.
2013/0237984	A1	9/2013	Sklar	2016/0095531	A1	4/2016	Narayan et al.
2013/0253415	A1	9/2013	Sano et al.	2016/0095642	A1	4/2016	Deno et al.
2013/0296679	A1	11/2013	Condie et al.	2016/0095653	A1	4/2016	Lambert et al.
2013/0310829	A1	11/2013	Cohen	2016/0100797	A1	4/2016	Mahapatra et al.
				2016/0100884	A1	4/2016	Fay et al.
				2016/0106498	A1	4/2016	Highsmith et al.
				2016/0106500	A1	4/2016	Olson
				2016/0113709	A1	4/2016	Maor

(56)

References Cited

OTHER PUBLICATIONS

U.S. PATENT DOCUMENTS

2016/0113712	A1	4/2016	Cheung et al.
2016/0120564	A1	5/2016	Kirkpatrick et al.
2016/0128770	A1	5/2016	Afonso et al.
2016/0166167	A1	6/2016	Narayan et al.
2016/0166311	A1	6/2016	Long et al.
2016/0174865	A1	6/2016	Stewart et al.
2016/0184003	A1	6/2016	Srimathveeravalli et al.
2016/0213282	A1	7/2016	Leo et al.
2016/0220307	A1	8/2016	Miller et al.
2016/0235470	A1	8/2016	Callas et al.
2016/0310211	A1	10/2016	Long
2016/0324564	A1	11/2016	Gerlach et al.
2016/0324573	A1	11/2016	Mickelson et al.
2016/0331441	A1	11/2016	Konings
2016/0354142	A1	12/2016	Pearson et al.
2016/0361109	A1	12/2016	Weaver et al.
2017/0035499	A1	2/2017	Stewart et al.
2017/0042449	A1	2/2017	Deno et al.
2017/0042615	A1	2/2017	Salahieh et al.
2017/0056648	A1	3/2017	Syed et al.
2017/0065330	A1	3/2017	Mickelsen et al.
2017/0065339	A1	3/2017	Mickelsen
2017/0065340	A1	3/2017	Long
2017/0065343	A1	3/2017	Mickelsen
2017/0151029	A1	6/2017	Mickelsen
2017/0181795	A1	6/2017	Debruyne
2017/0189097	A1	7/2017	Viswanathan et al.

FOREIGN PATENT DOCUMENTS

EP	1127552	6/2006
EP	1340469	3/2007
EP	1009303	6/2009
EP	2213729	8/2010
EP	2425871	3/2012
EP	1803411	8/2012
EP	2587275	5/2013
EP	2663227	11/2013
EP	1909678	1/2014
EP	2217165	3/2014
EP	2376193	3/2014
EP	2934307	10/2015
EP	2777585	6/2016
WO	WO 99/04851	2/1999
WO	WO 99/22659	5/1999
WO	WO 2004/086994	10/2004
WO	2007006055	1/2007
WO	WO 2007/079438	7/2007
WO	WO 2009/082710	7/2009
WO	WO 2009/137800	11/2009
WO	WO 2010/014480	2/2010
WO	WO 2011/154805	12/2011
WO	WO 2012/051433	4/2012
WO	WO 2013/019385	2/2013
WO	WO 2014/025394	2/2014
WO	WO 2014/031800	2/2014
WO	WO 2014/160832	10/2014
WO	WO 2015/066322	5/2015
WO	WO 2015/099786	7/2015
WO	WO 2015/103530	7/2015
WO	WO 2015/103574	7/2015
WO	WO 2015/130824	9/2015
WO	WO 2015/143327	9/2015
WO	WO 2015/171921	11/2015
WO	WO 2015/175944	11/2015
WO	WO 2015/192018	12/2015
WO	WO 2015/192027	12/2015
WO	WO 2016/059027	4/2016
WO	WO 2016/060983	4/2016
WO	WO 2016/081650	5/2016
WO	WO 2017/120169	7/2017

Notice of Reasons for Rejection for Japanese Application No. 2015-526522, dated Mar. 6, 2017, 3 pages.

Office Action for U.S. Appl. No. 15/201,997, dated Apr. 3, 2017, 6 pages.

Office Action for U.S. Appl. No. 15/334,646, dated Jul. 25, 2017, 19 pages.

International Search Report and Written Opinion for International Application No. PCT/US2016/057664, dated Feb. 24, 2017, 11 pages.

International Search Report and Written Opinion for International Application No. PCT/US2017/012099, dated May 18, 2017, 17 pages.

Partial Supplementary European Search Report for European Application No. 13827672.0, dated Mar. 23, 2016, 6 pages.

International Search Report and Written Opinion for International Application No. PCT/US2013/031252, dated Jul. 19, 2013, 12 pages.

International Search Report and Written Opinion for International Application No. PCT/US2015/010138, dated Mar. 26, 2015, 14 pages.

International Preliminary Report on Patentability for International Application No. PCT/US2015/010138, dated Jul. 12, 2016, 9 pages.

International Search Report and Written Opinion for International Application No. PCT/US2015/010223, dated Apr. 10, 2015, 19 pages.

International Preliminary Report on Patentability for International Application No. PCT/US2015/010223, dated Jul. 12, 2016, 12 pages.

International Search Report and Written Opinion for International Application No. PCT/US2015/029734, dated Nov. 24, 2015, 15 pages.

International Search Report and Written Opinion for International Application No. PCT/US2015/031086, dated Oct. 21, 2015, 16 pages.

International Search Report and Written Opinion for International Application No. PCT/US2015/055105, dated Mar. 1, 2016, 15 pages.

International Search Report and Written Opinion for International Application No. PCT/US2015/035582, dated Oct. 2, 2015, 17 pages.

International Search Report and Written Opinion for International Application No. PCT/US2015/035592, dated Oct. 2, 2015, 13 pages.

du Pre, B.C. et al., "Minimal coronary artery damage by myocardial electroporation ablation," *Europace*, 15(1):144-149 (2013).

Hobbs, E. P., "Investor Relations Update: Tissue Ablation via Irreversible Electroporation (IRE)," Powerpoint (2004), 16 pages.

Lavee, J. et al., "A Novel Nonthermal Energy Source for Surgical Epicardial Atrial Ablation: Irreversible Electroporation," *The Heart Surgery Forum* #2006-1202, 10(2), 2007 [Epub Mar. 2007].

Madhavan, M. et al., "Novel Percutaneous Epicardial Autonomic Modulation in the Canine for Atrial Fibrillation: Results of an Efficacy and Safety Study," *Pace*, 00:1-11 (2016).

Neven, K. et al., "Safety and Feasibility of Closed Chest Epicardial Catheter Ablation Using Electroporation," *Circ Arrhythm Electrophysiol.*, 7:913-919 (2014).

Neven, K. et al., "Myocardial Lesion Size After Epicardial Electroporation Catheter Ablation After Subxiphoid Puncture," *Circ Arrhythm Electrophysiol.*, 7(4):728-733 (2014).

Neven, K. et al., "Epicardial linear electroporation ablation and lesion size," *Heart Rhythm*, 11:1465-1470 (2014).

van Driel, V.J.H.M. et al., "Pulmonary Vein Stenosis After Catheter Ablation Electroporation Versus Radiofrequency," *Circ Arrhythm Electrophysiol.*, 7(4):734-738 (2014).

van Driel, V.J.H.M. et al., "Low vulnerability of the right phrenic nerve to electroporation ablation," *Heart Rhythm*, 12:1838-1844 (2015).

Wittkampf, F.H. et al., "Myocardial Lesion Depth With Circular Electroporation Ablation," *Circ. Arrhythm Electrophysiol.*, 5(3):581-586 (2012).

(56)

References Cited

OTHER PUBLICATIONS

Wittkamp, F.H. et al., "Feasibility of Electroporation for the Creation of Pulmonary Vein Ostial Lesions," J Cardiovasc Electrophysiol, 22(3):302-309 (Mar. 2011).

* cited by examiner

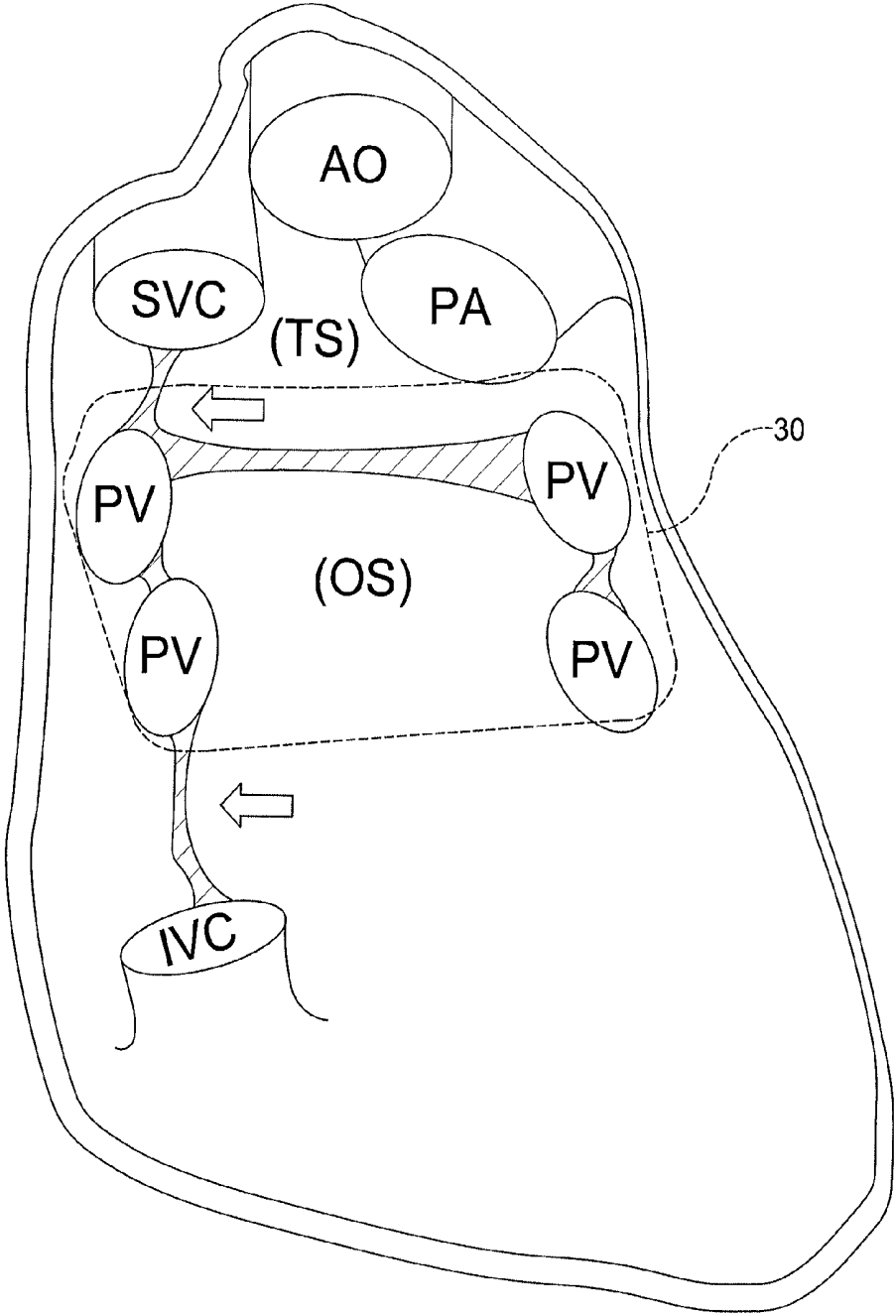


FIG. 1

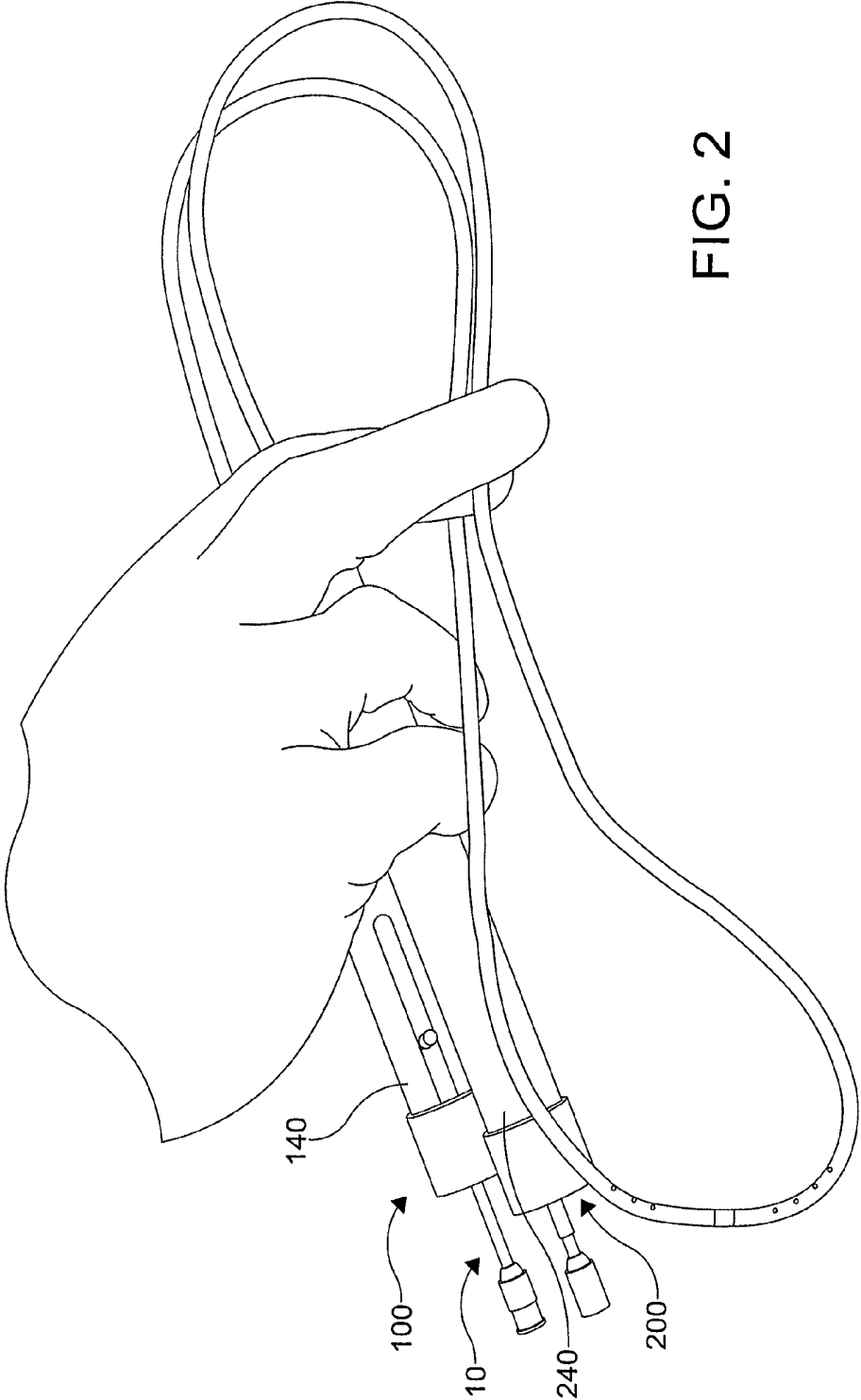


FIG. 2

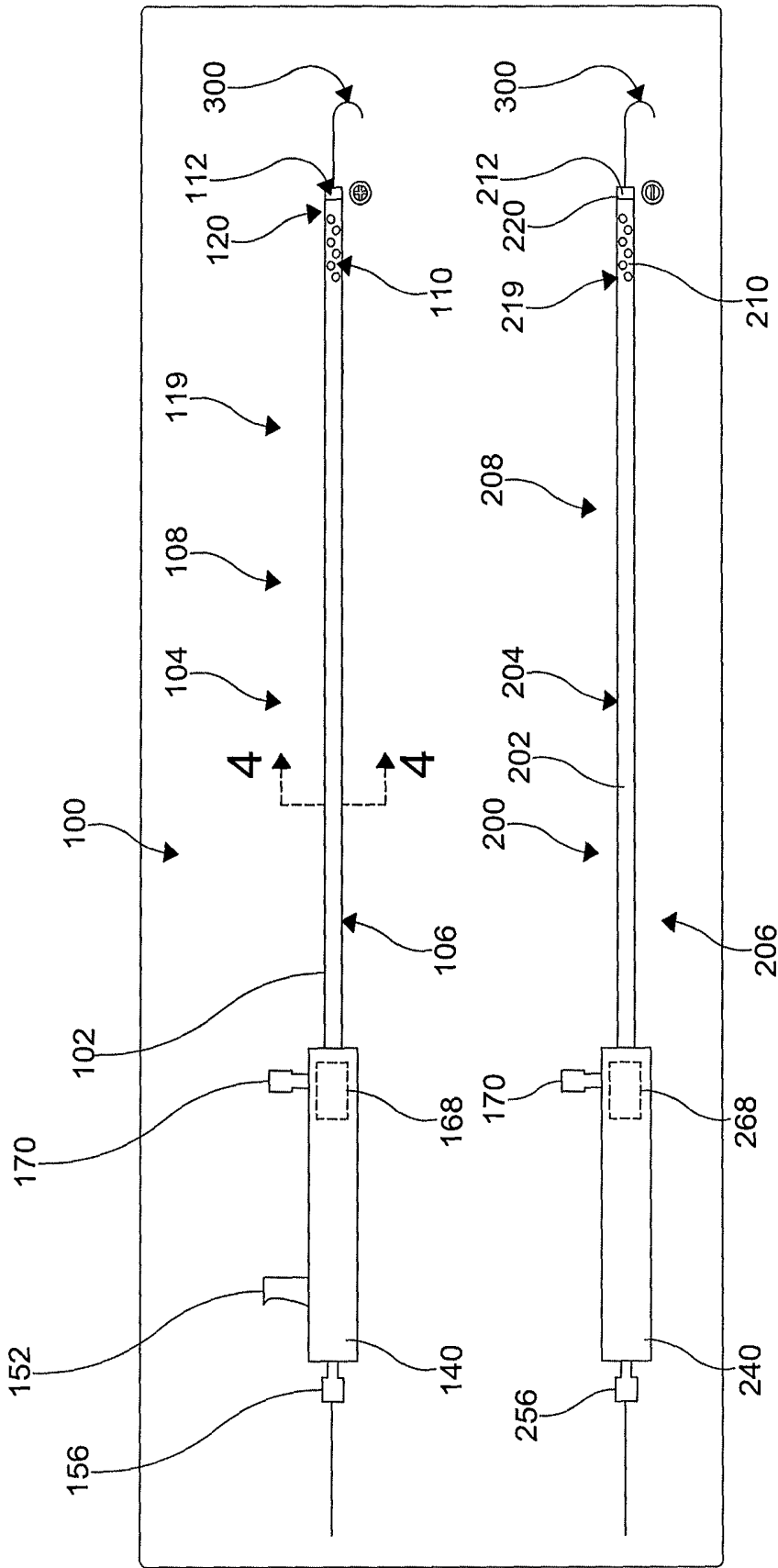


FIG. 3

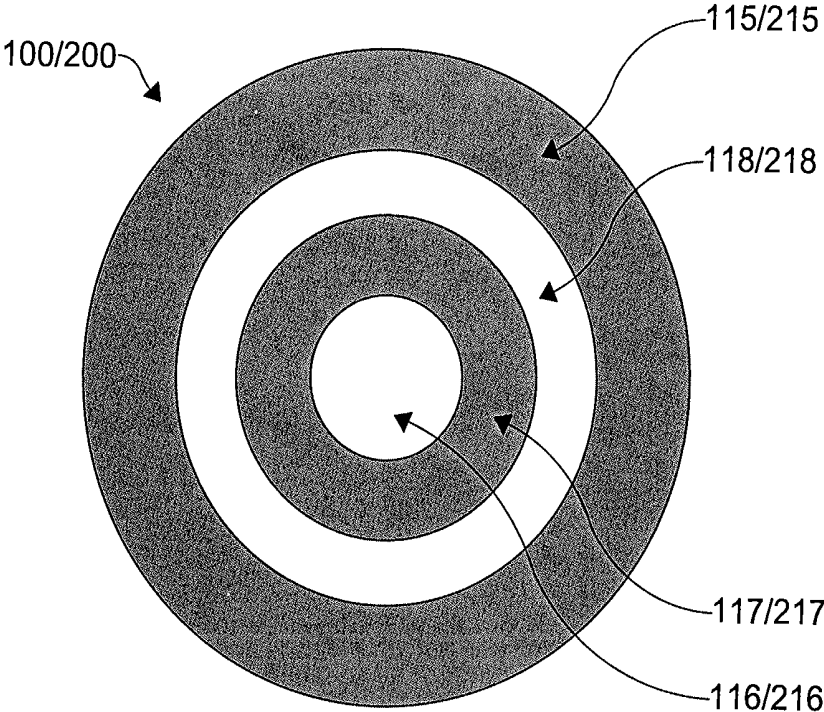


FIG. 4

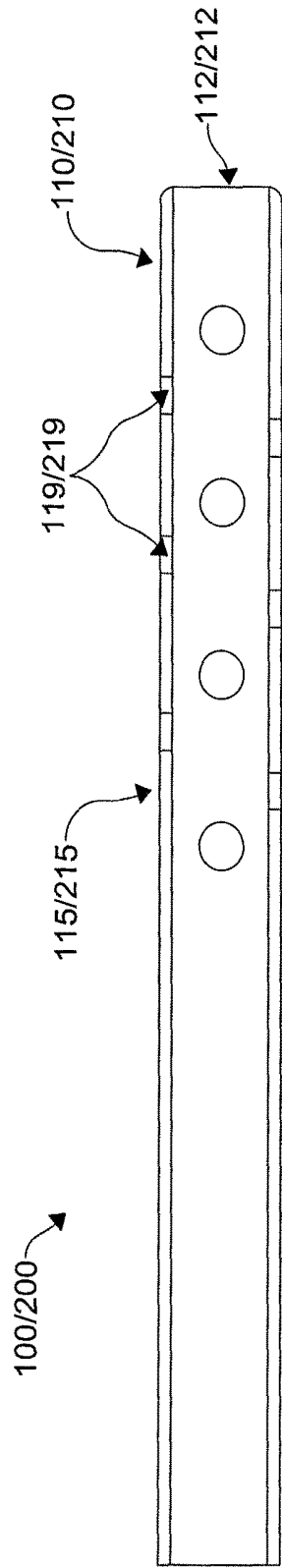


FIG. 5a

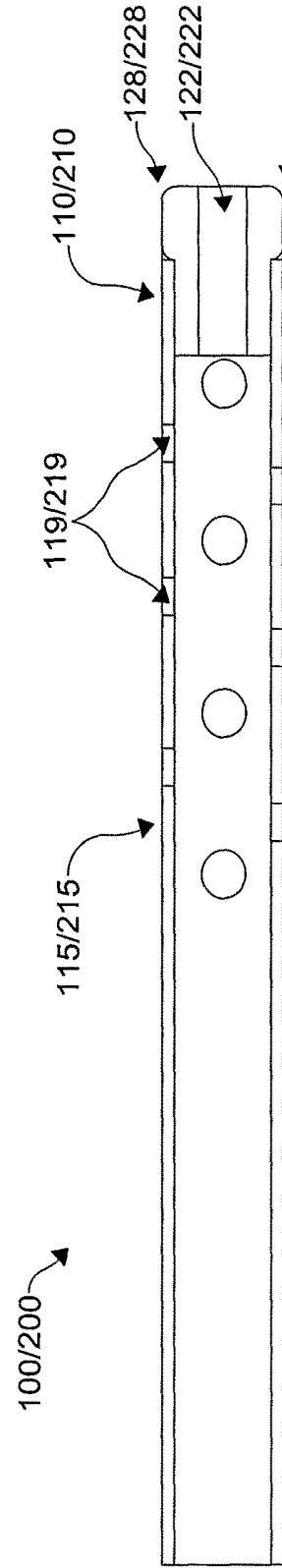


FIG. 5b

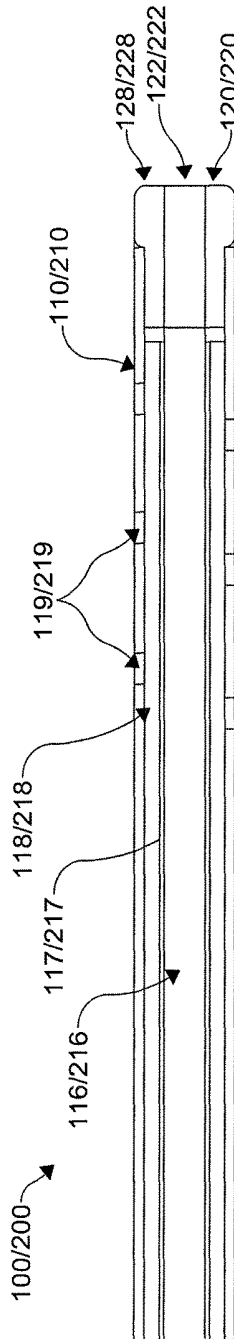


FIG. 5c

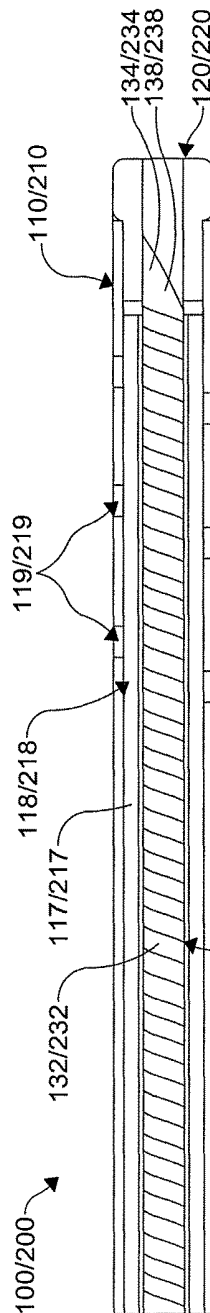


FIG. 5d

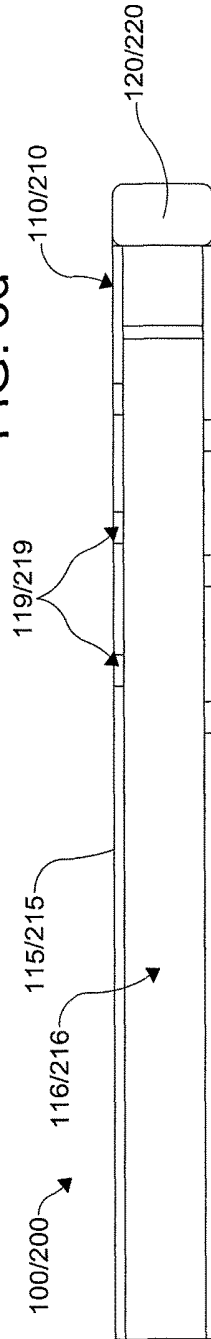


FIG. 5e

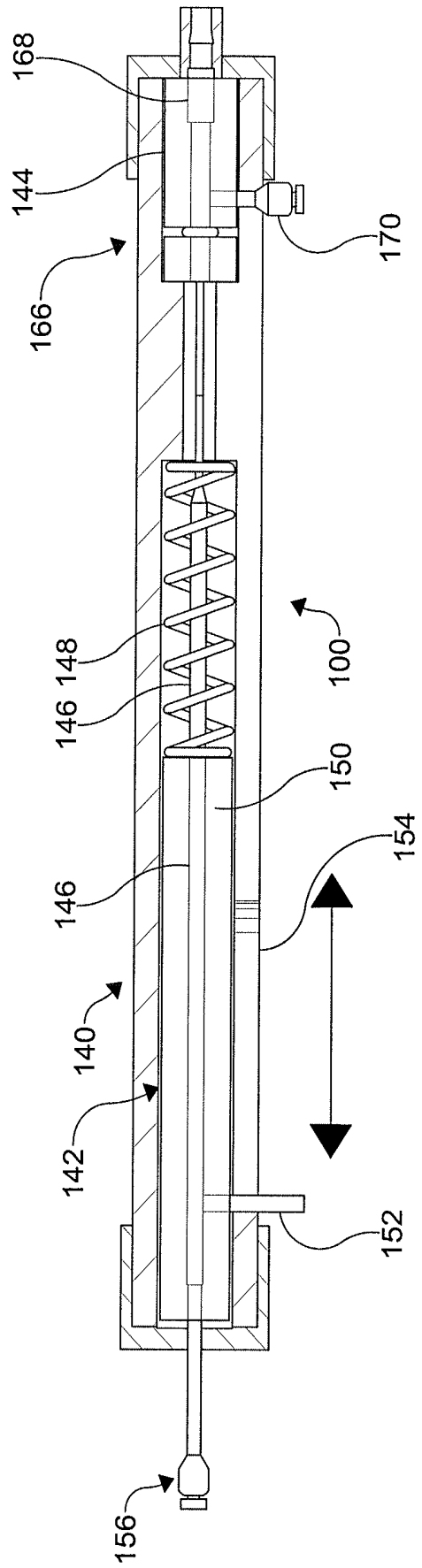


FIG. 6

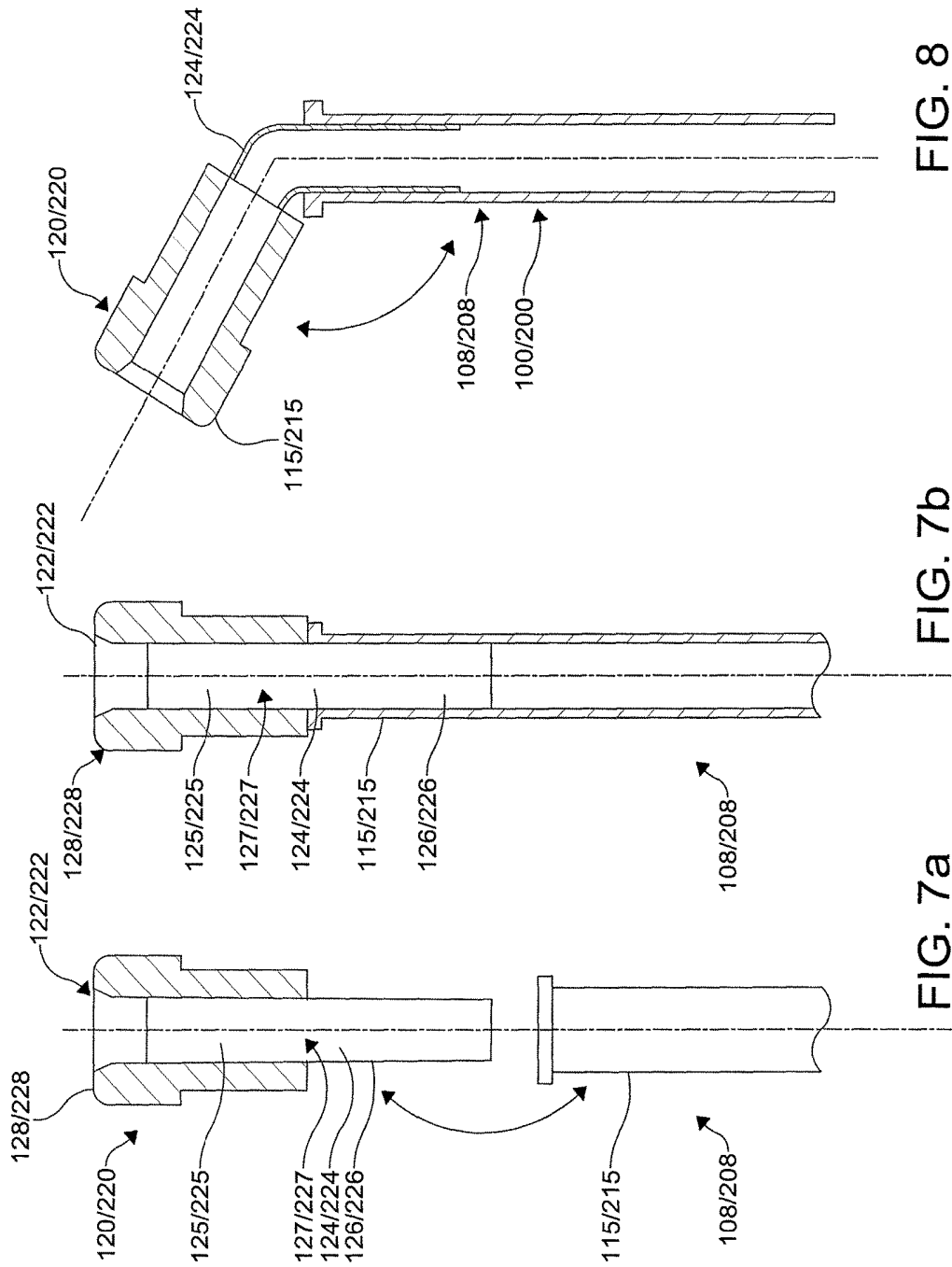


FIG. 8

FIG. 7b

FIG. 7a

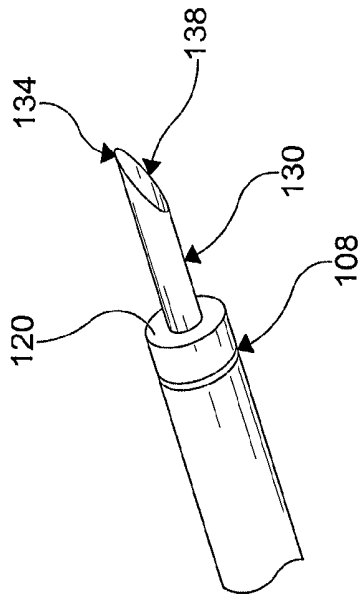


FIG. 9

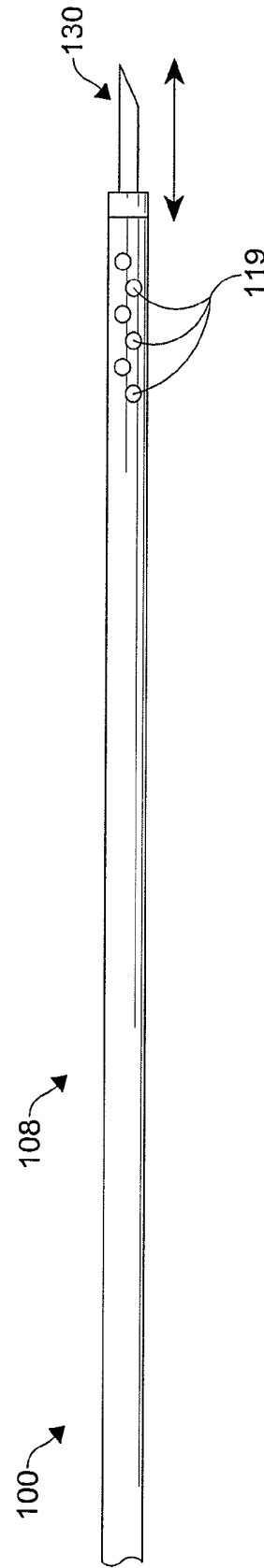


FIG. 10

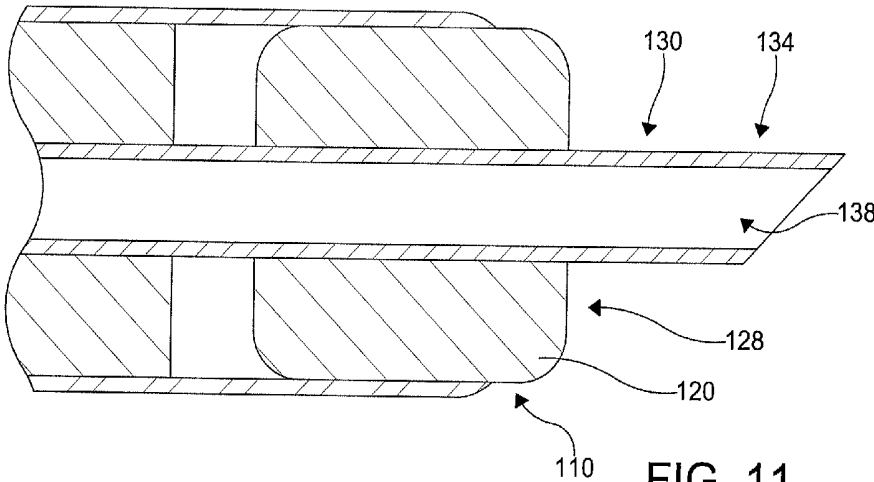


FIG. 11

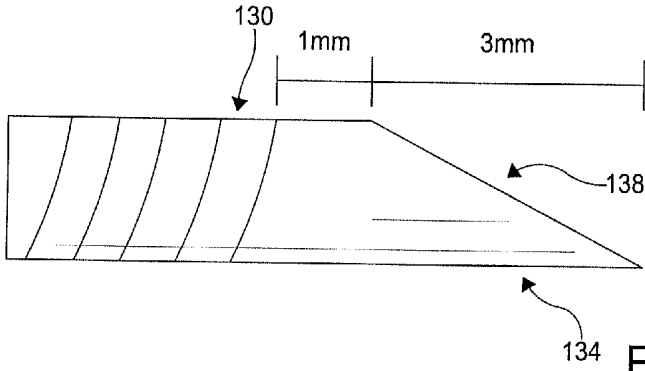


FIG. 12

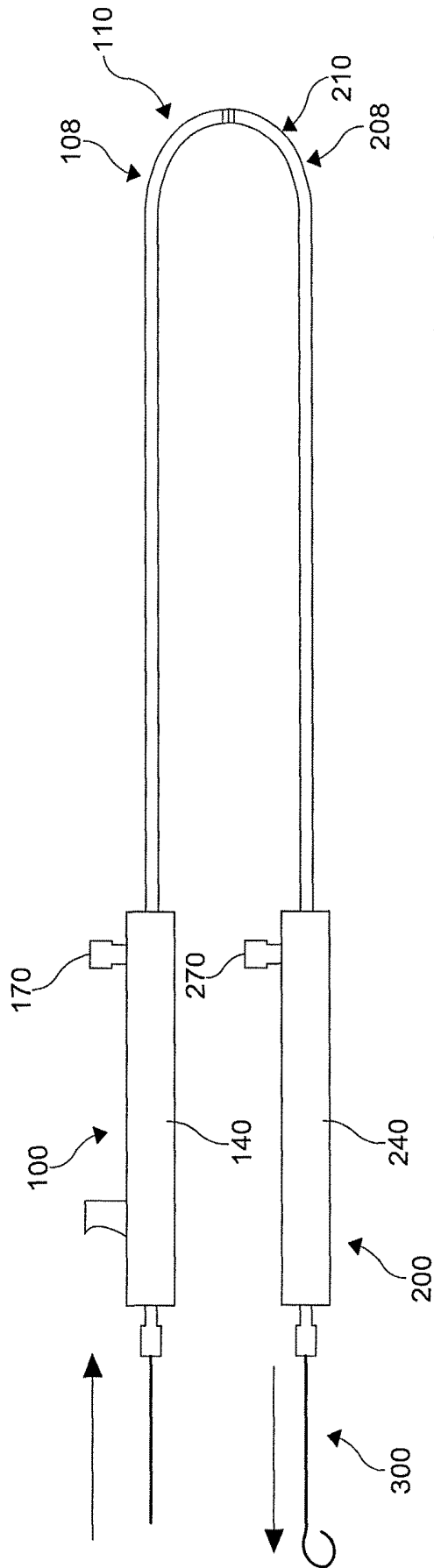


FIG. 13

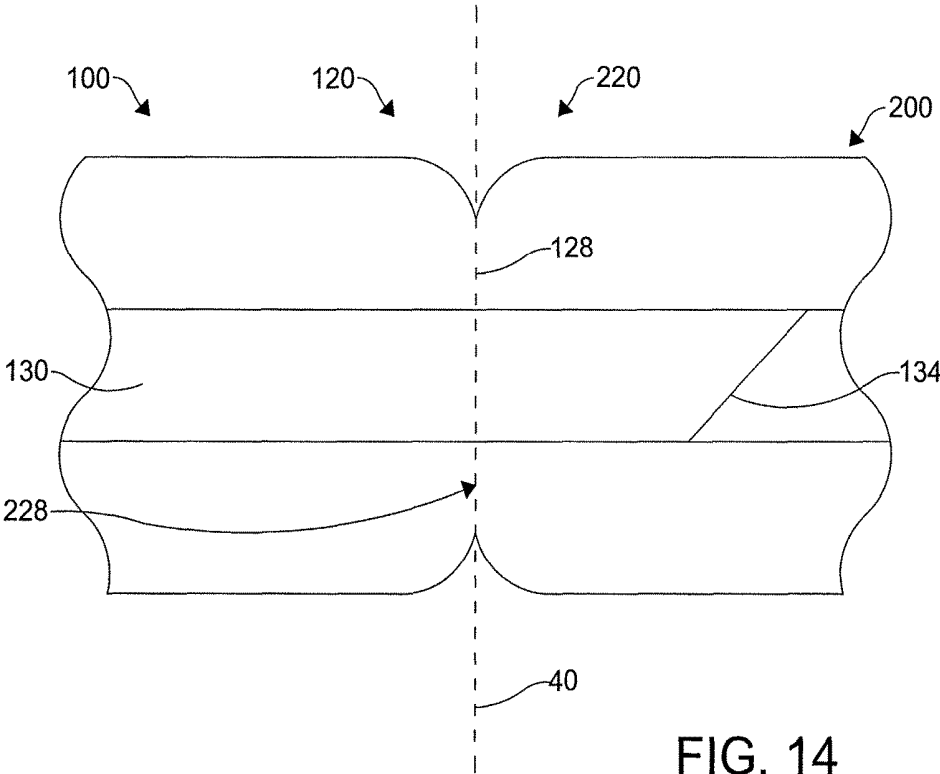


FIG. 14

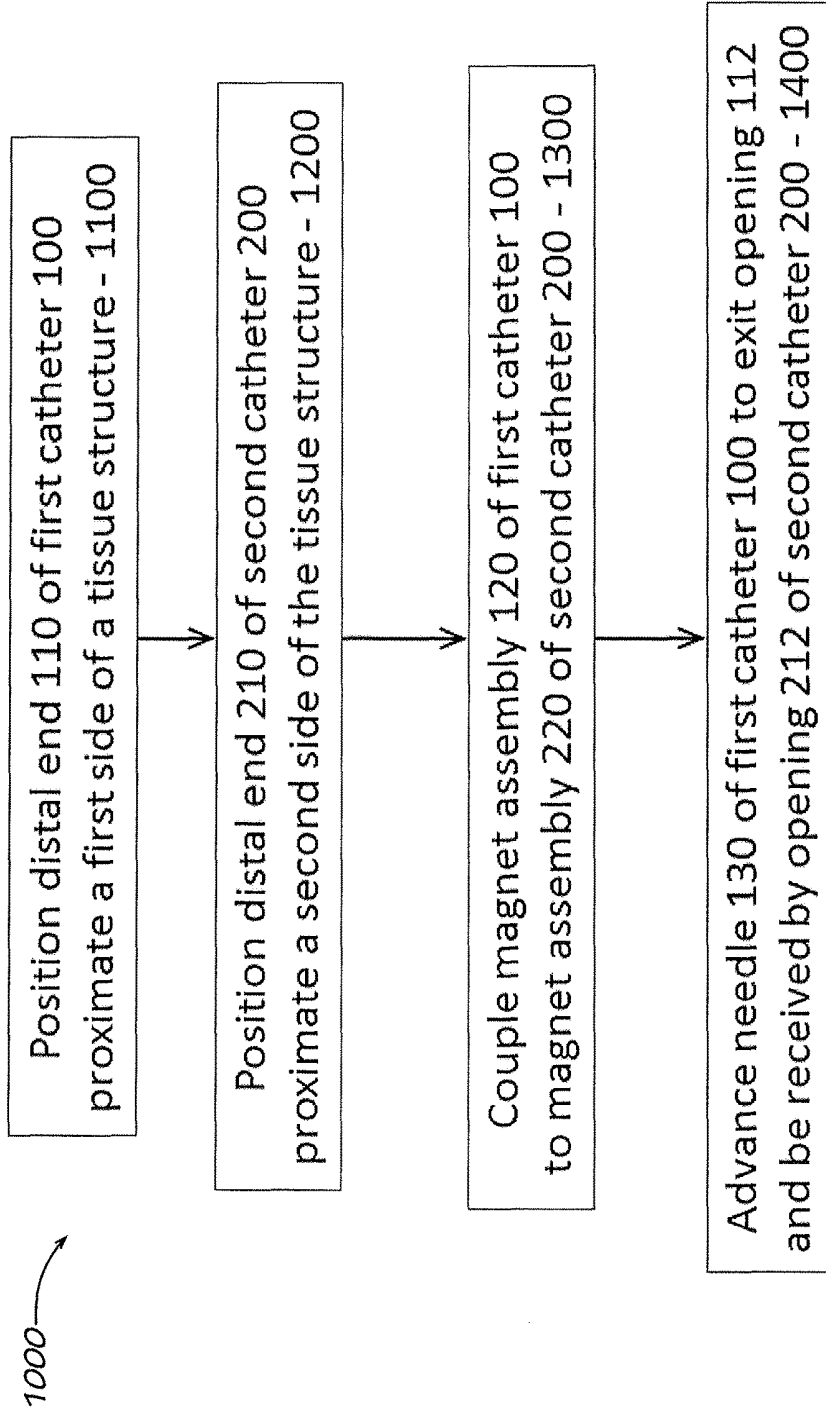


FIG. 15

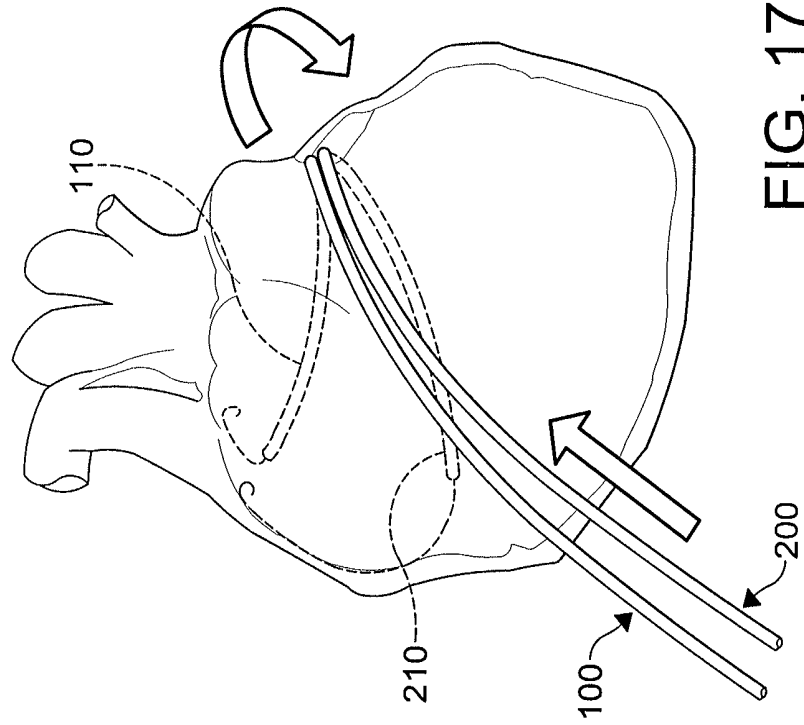


FIG. 17

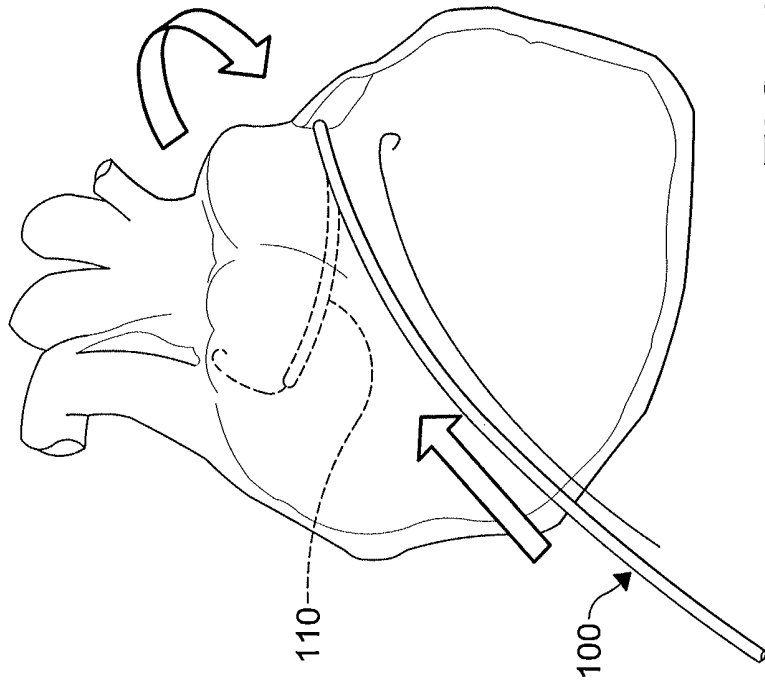


FIG. 16

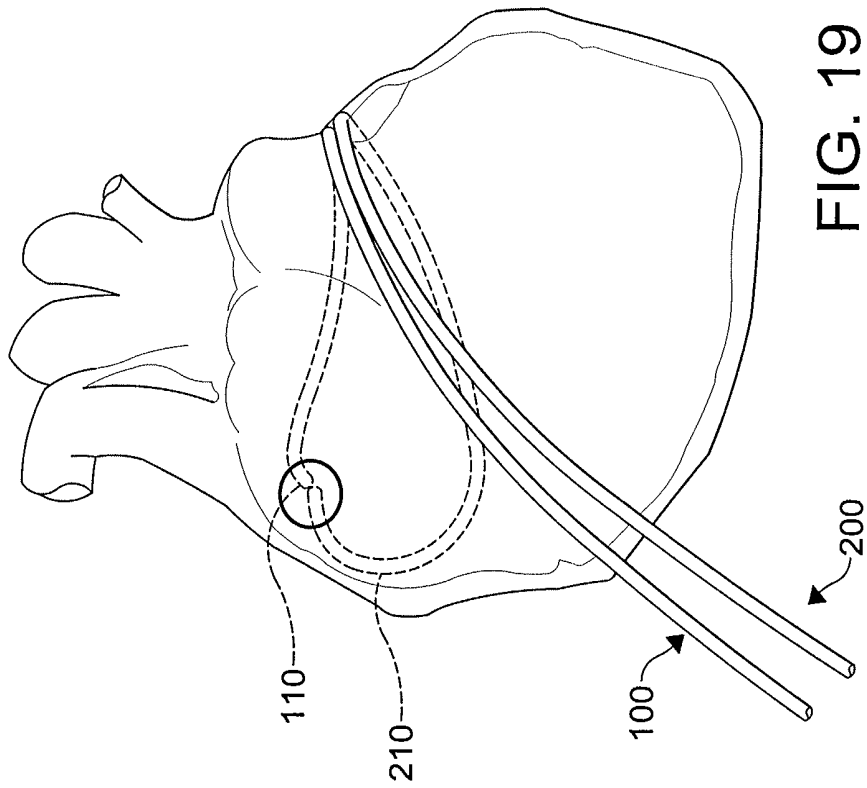


FIG. 19

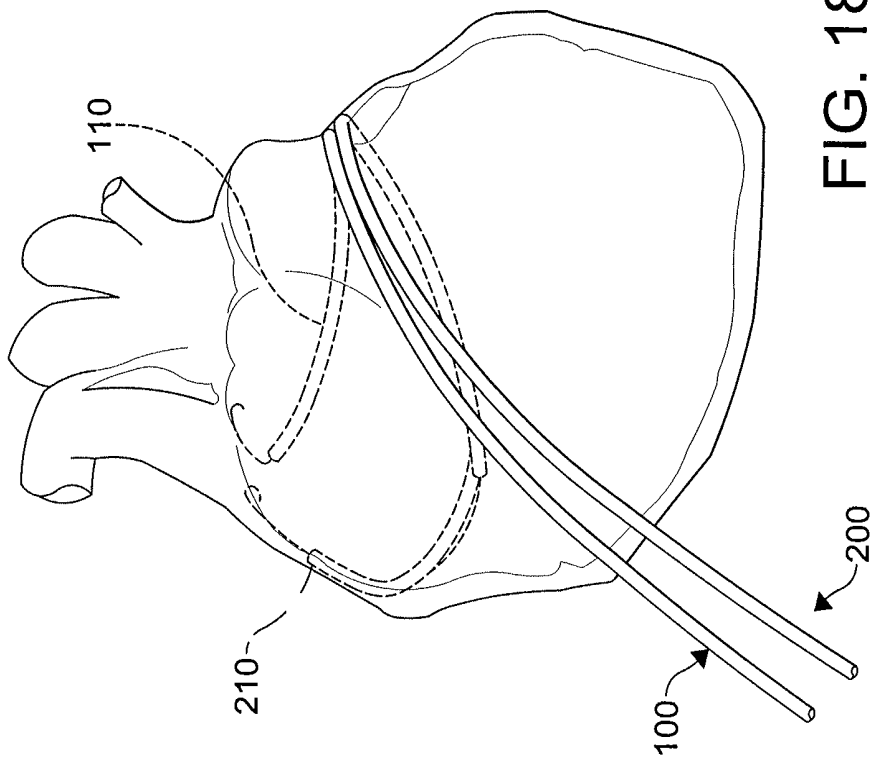
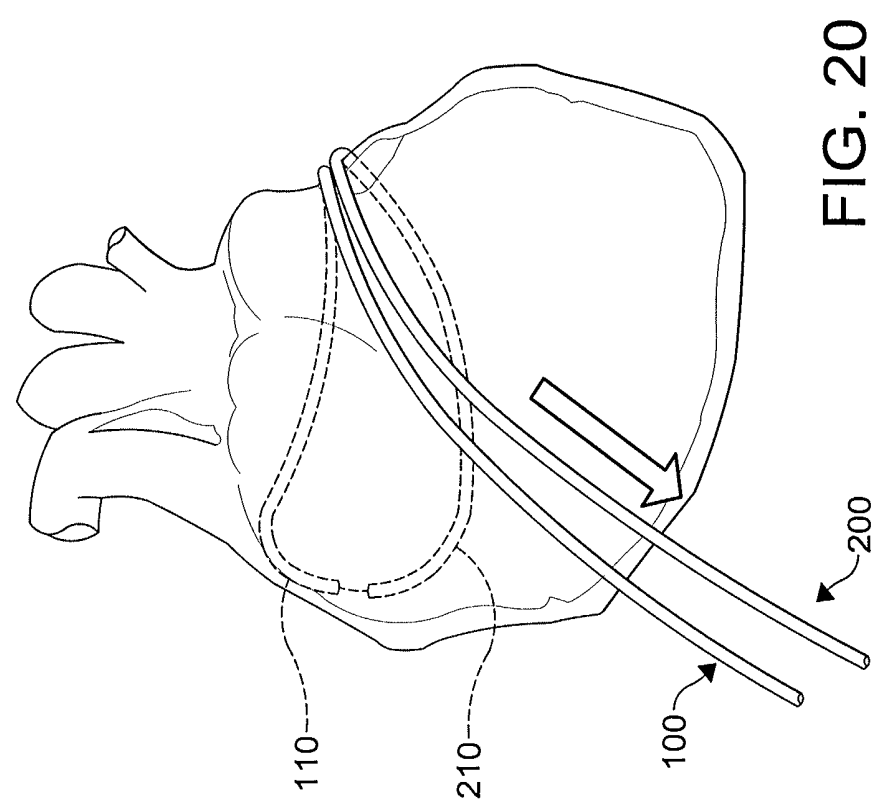
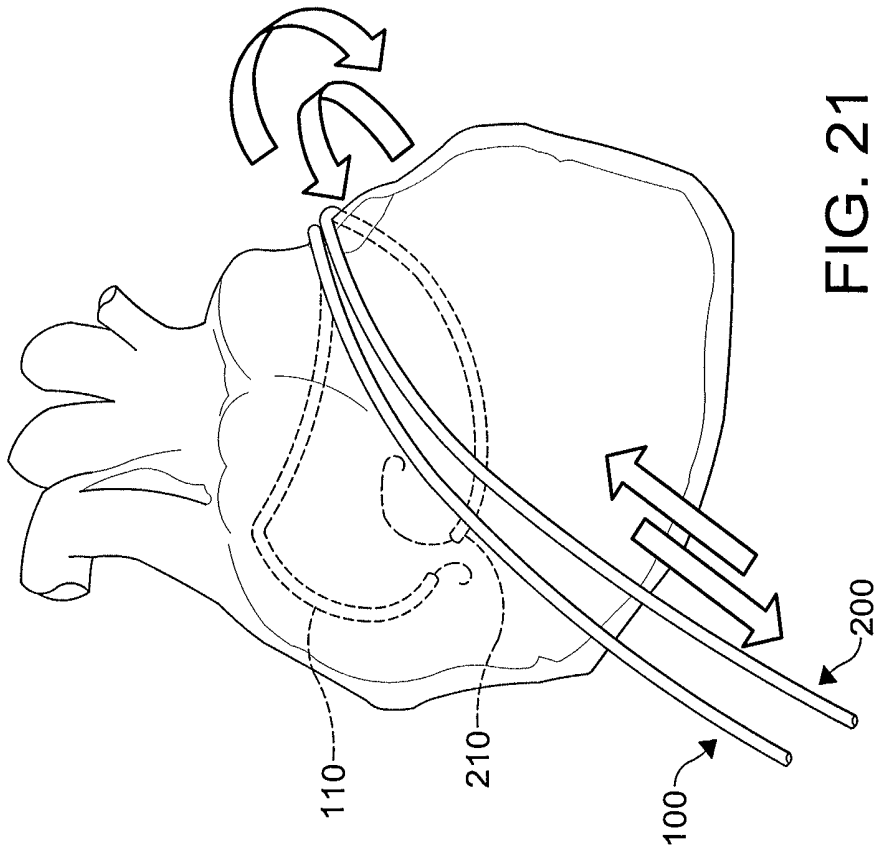


FIG. 18



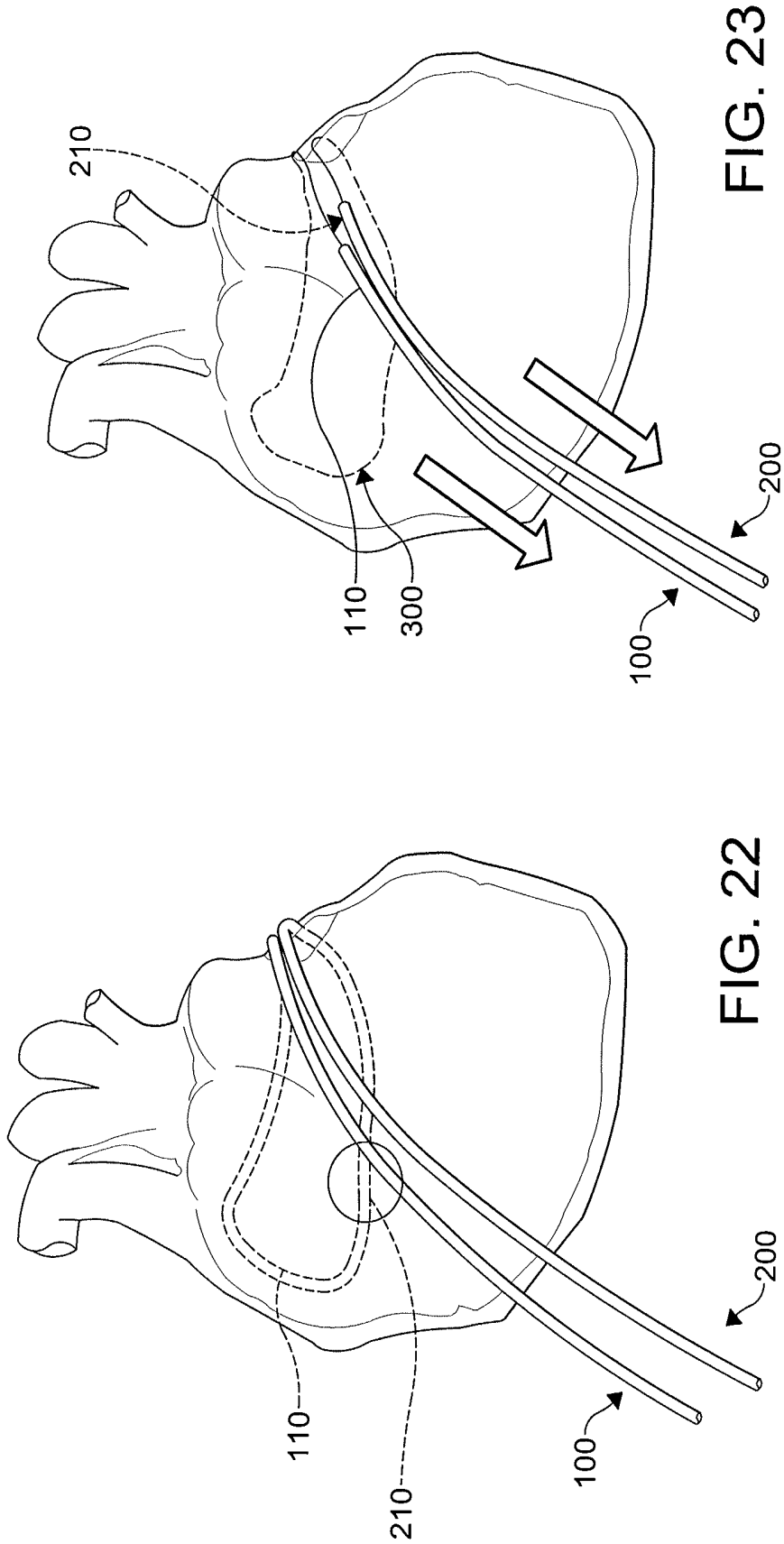


FIG. 23

FIG. 22

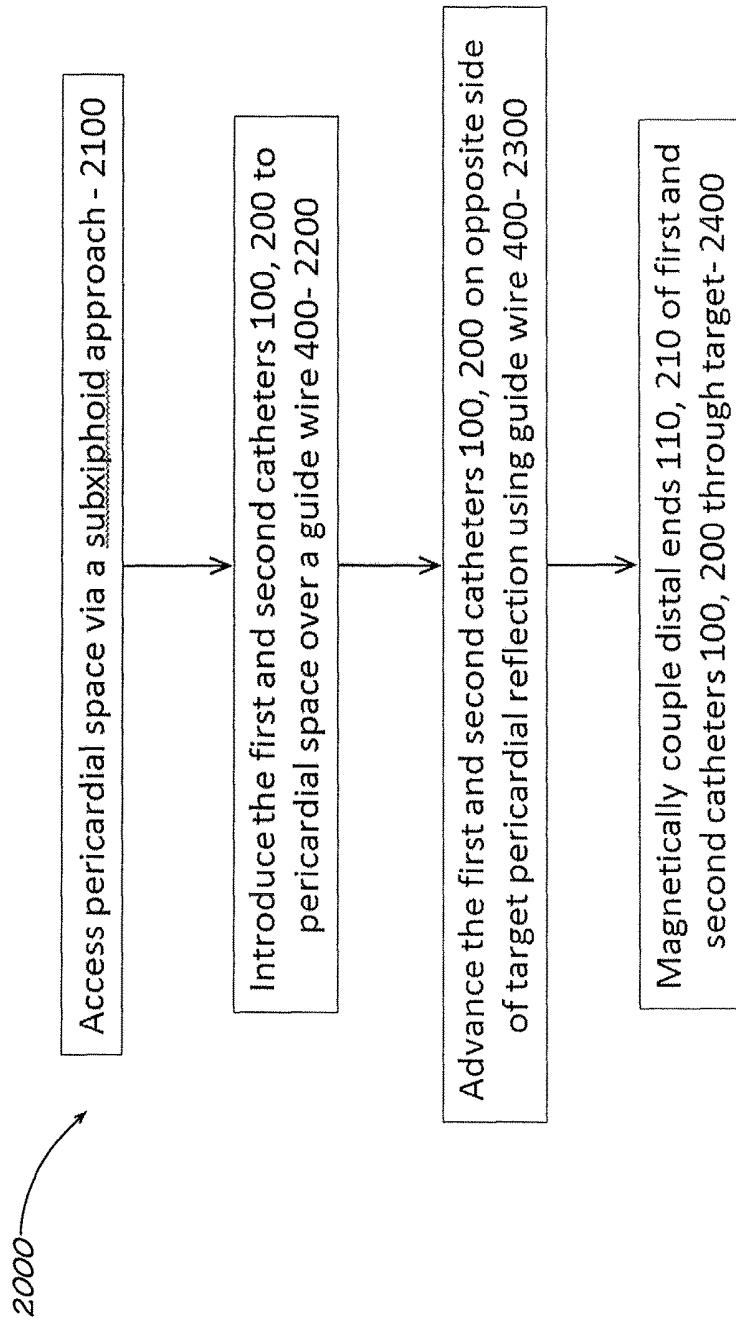


FIG. 24

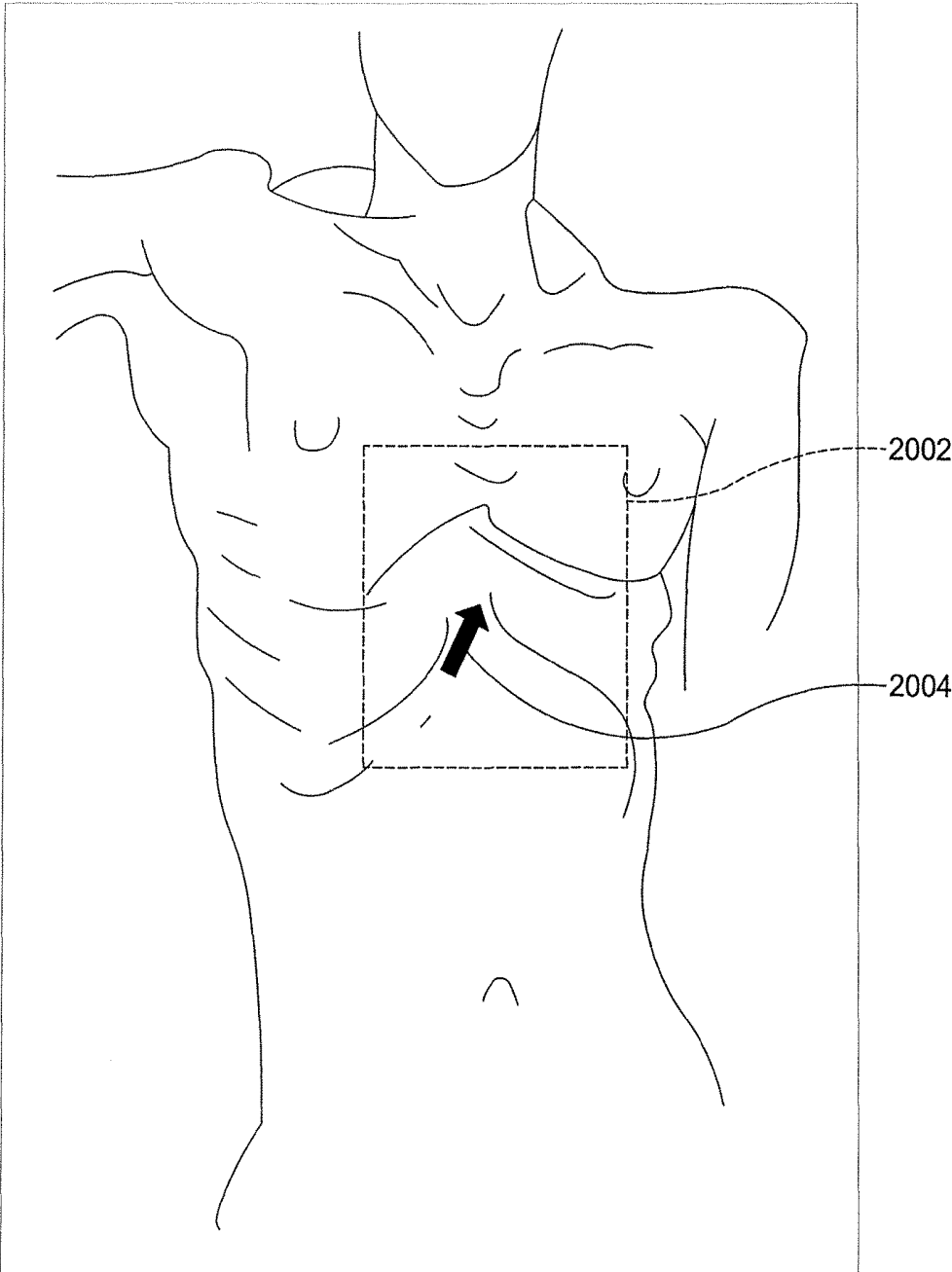


FIG. 25

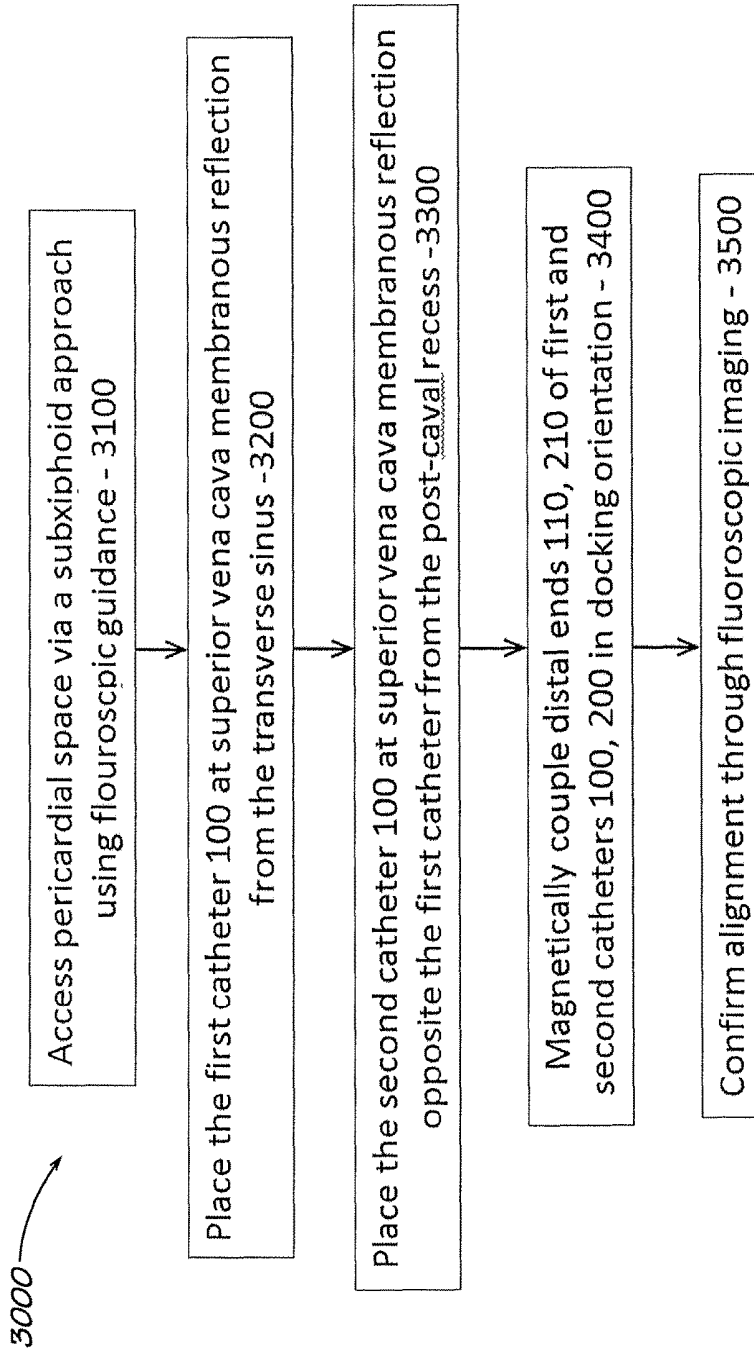


FIG. 26

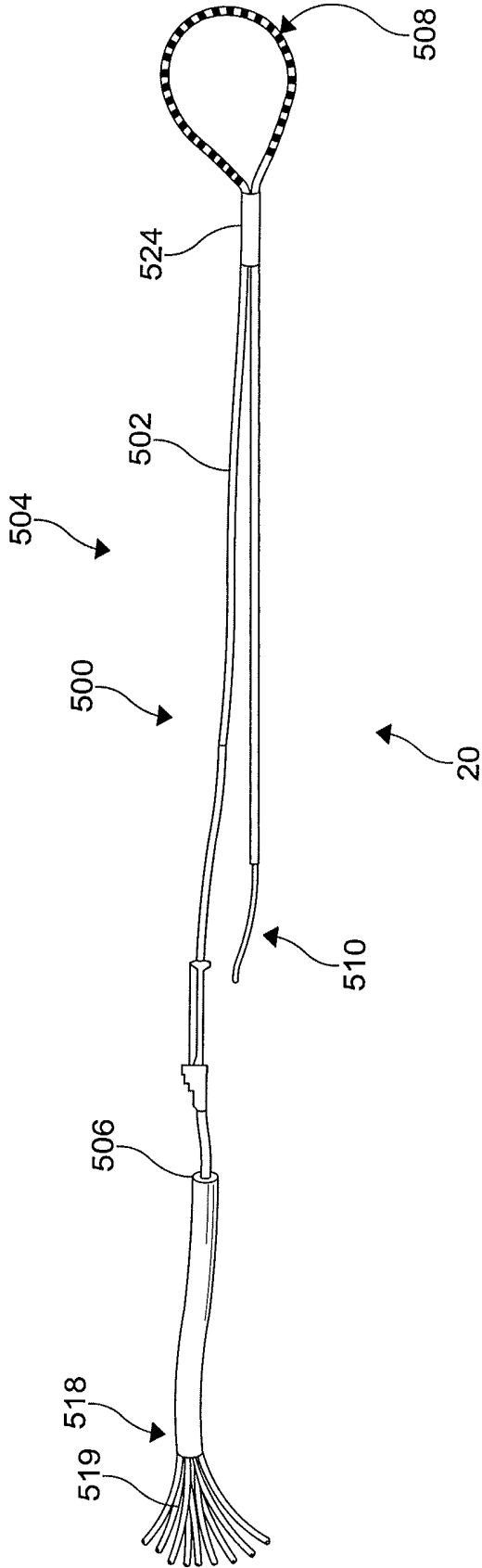


FIG. 27

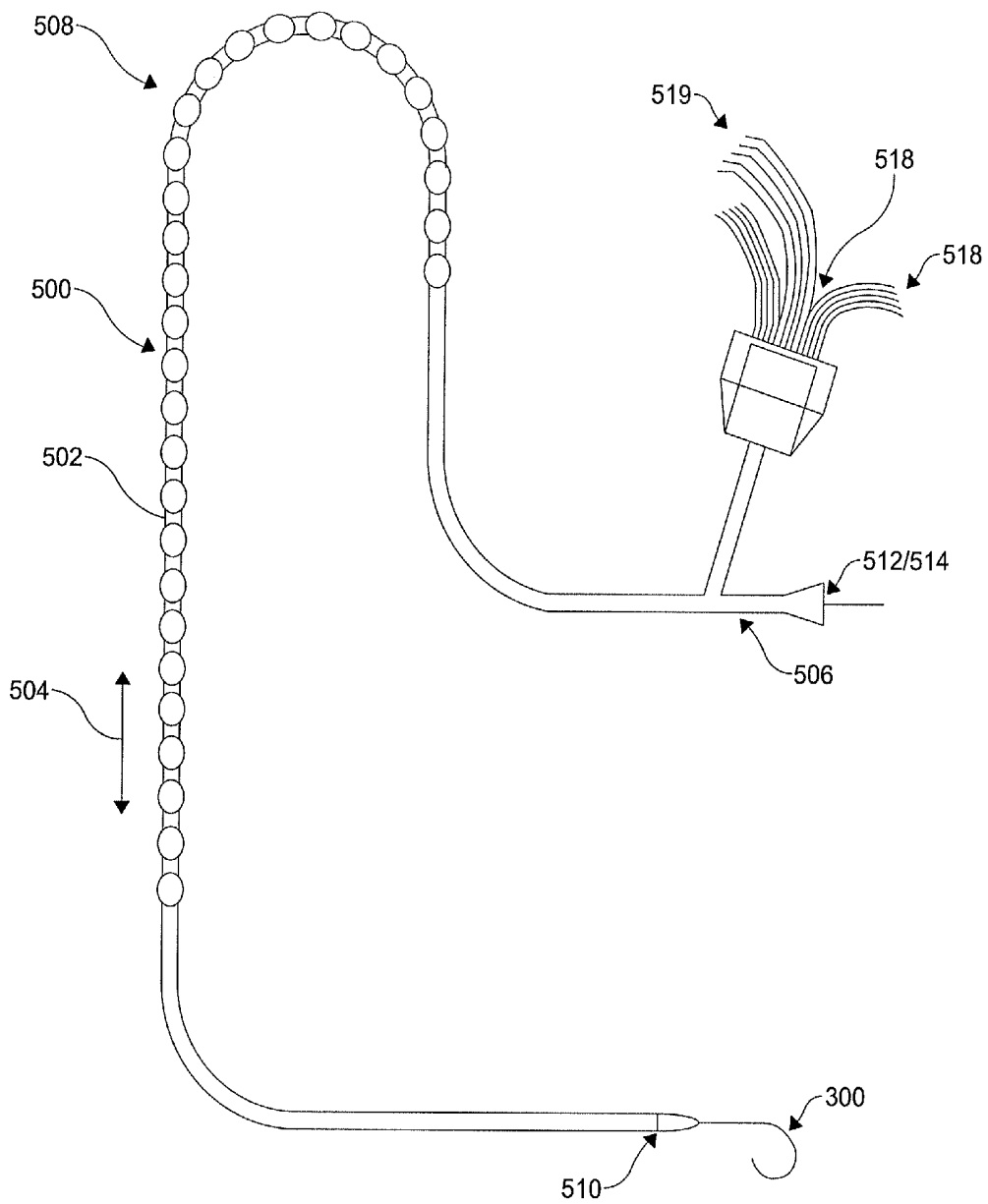


FIG. 28

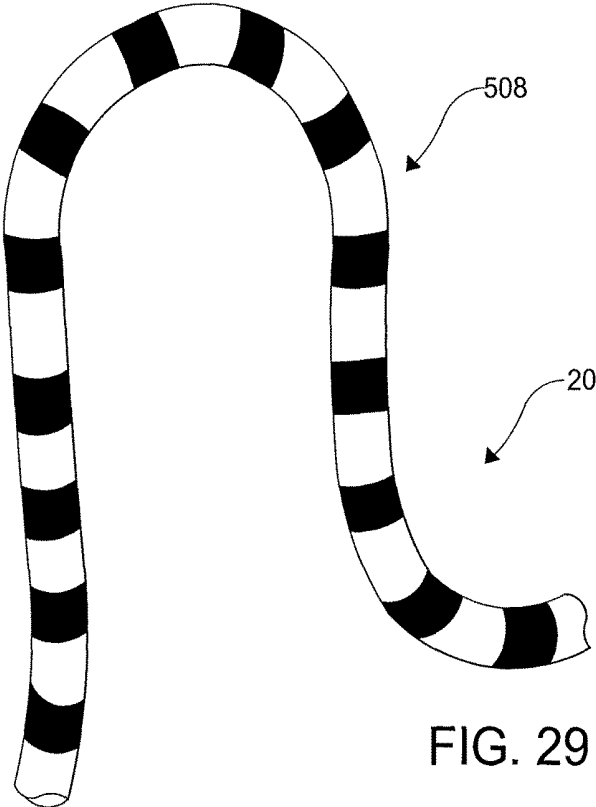


FIG. 29

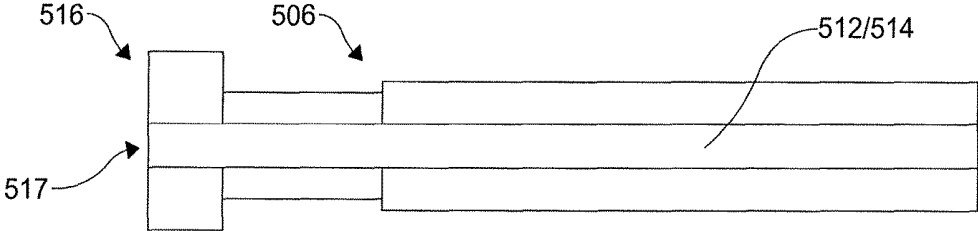


FIG. 30

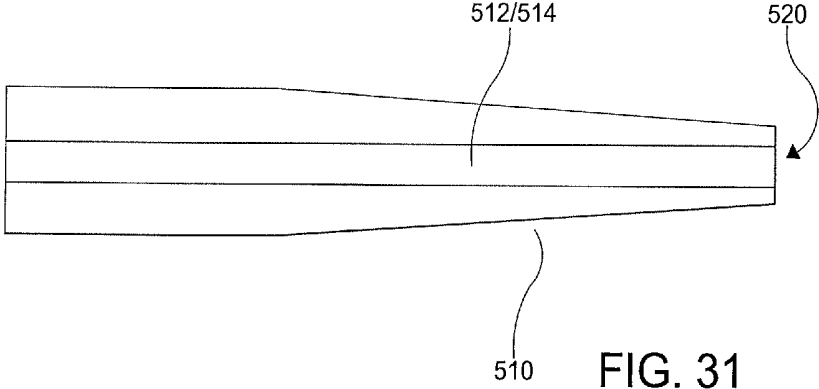


FIG. 31

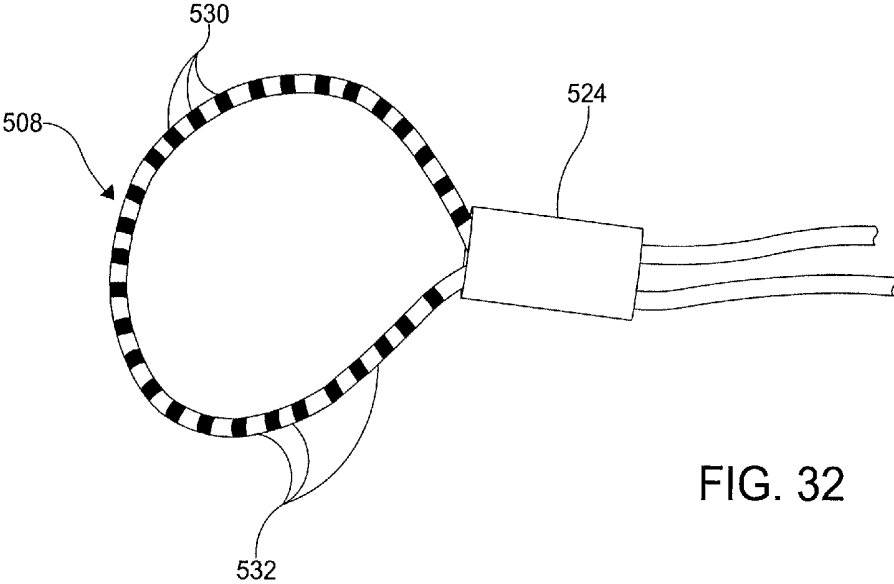
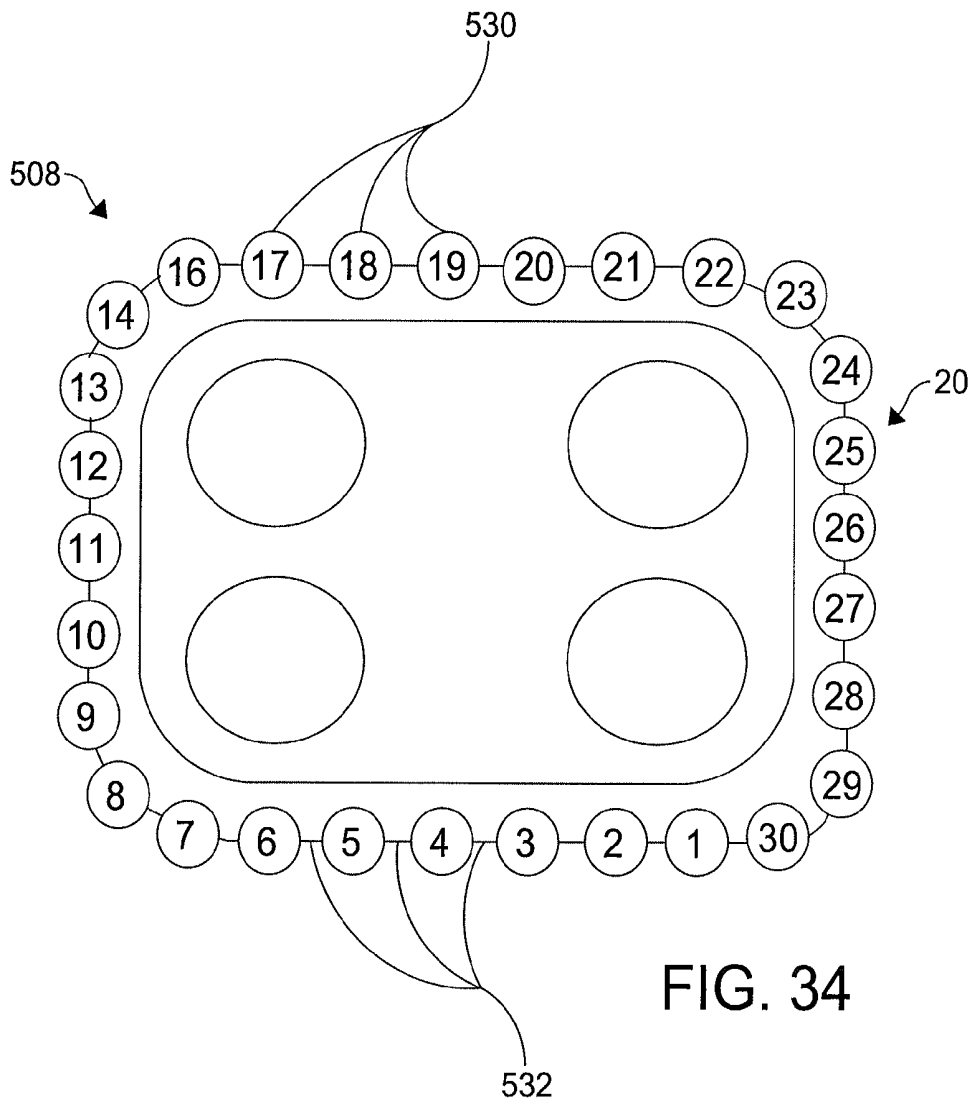


FIG. 32



FIG. 33



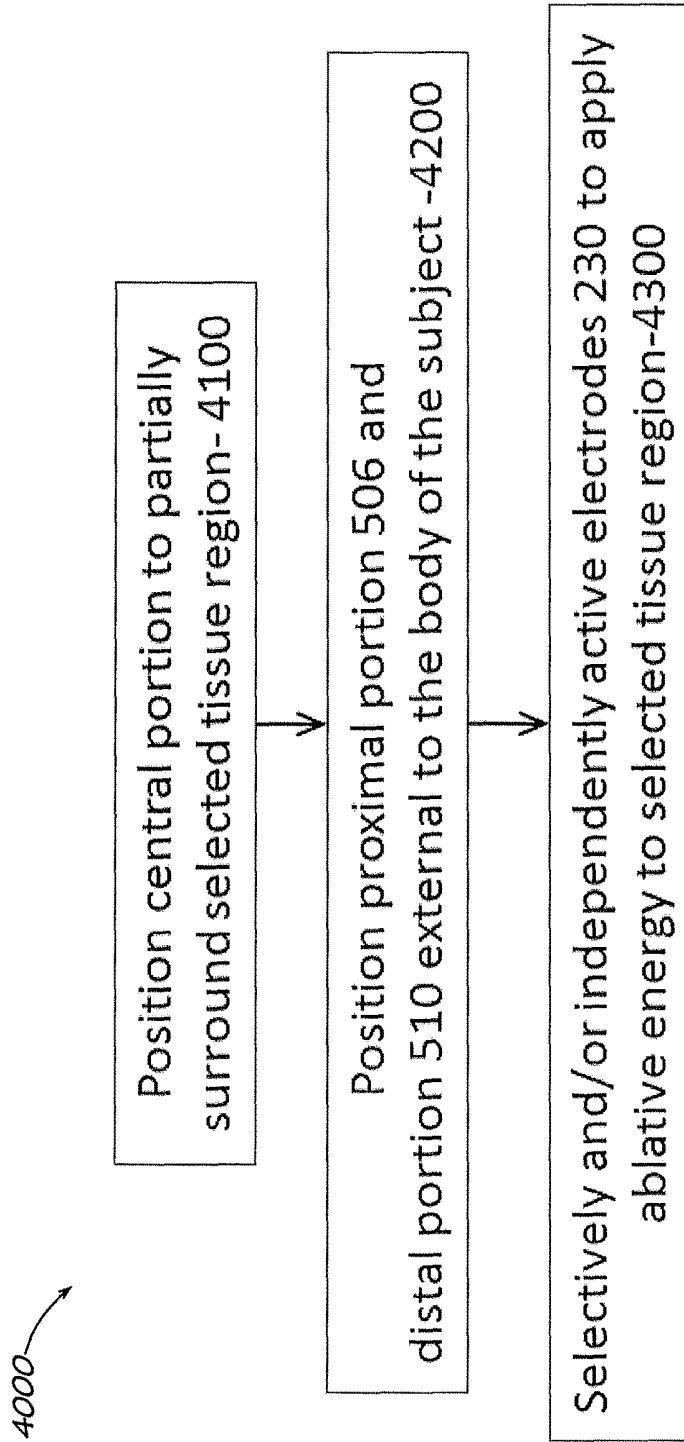


FIG. 35

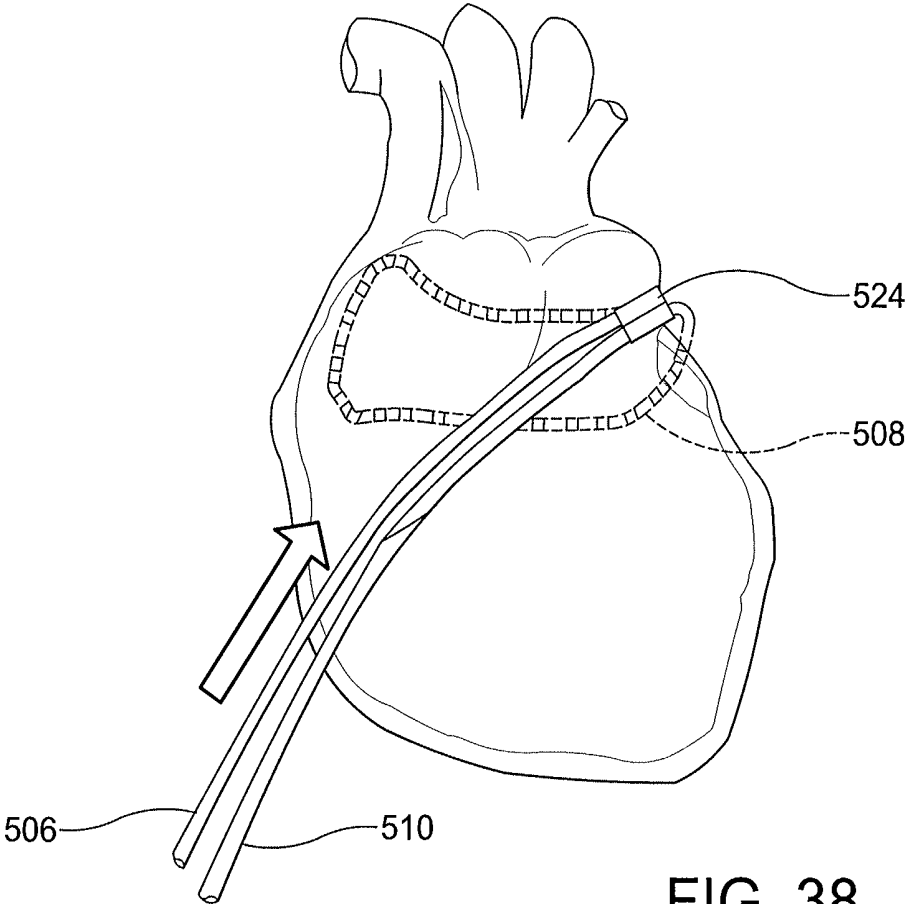


FIG. 38

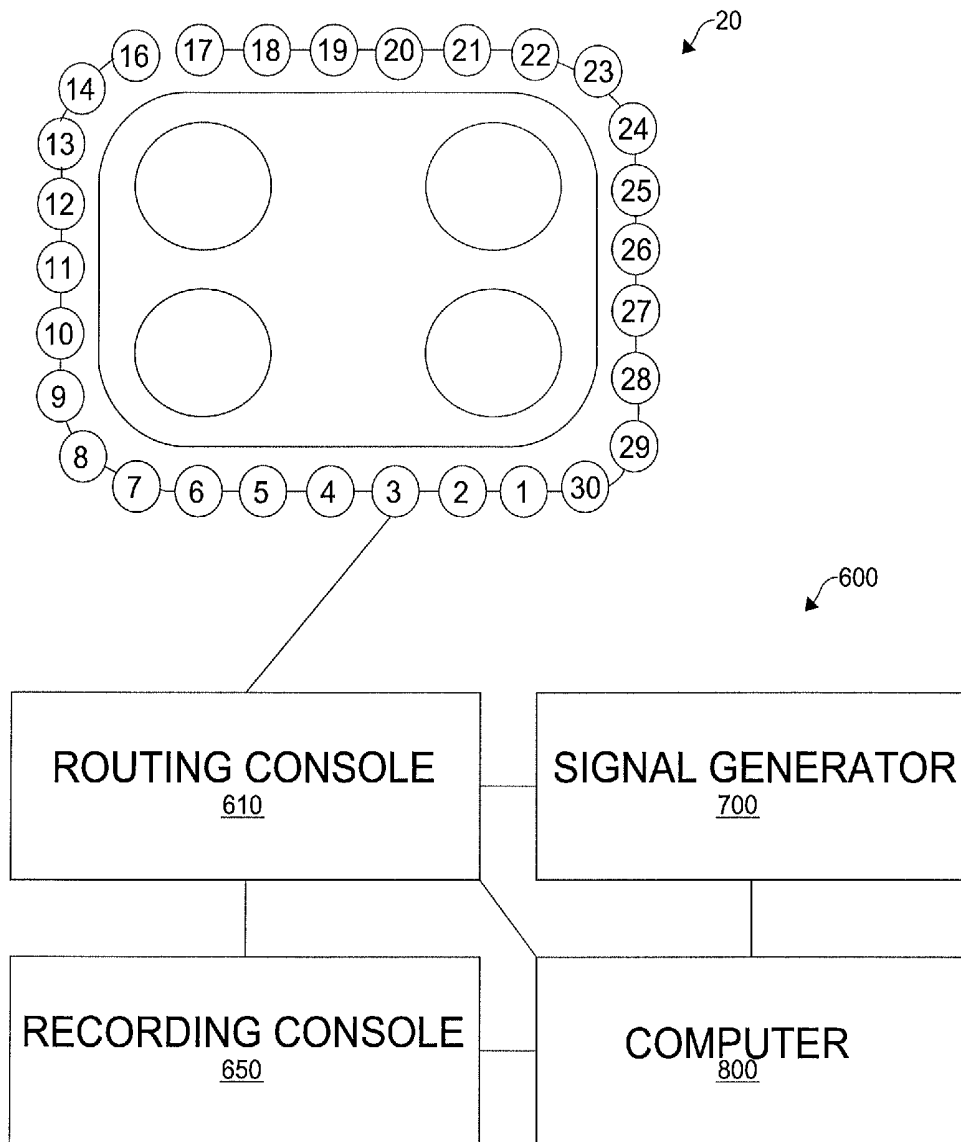


FIG. 39

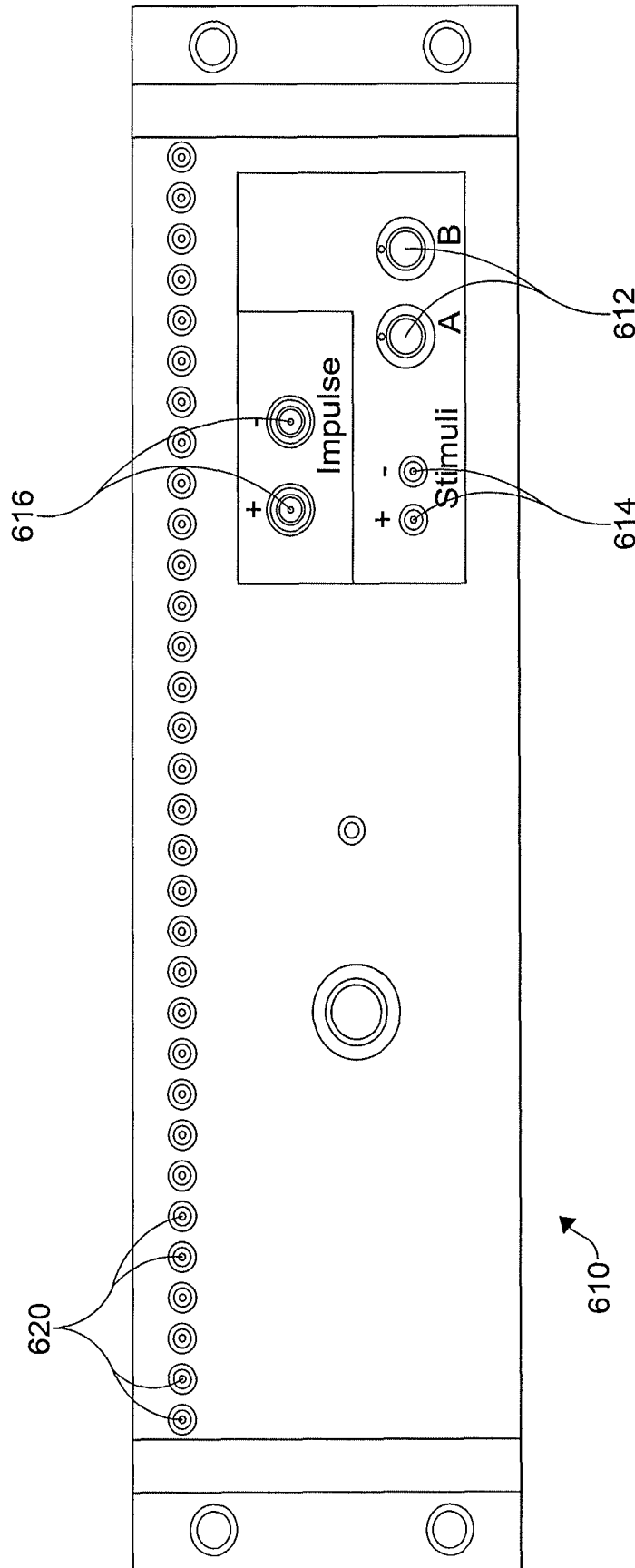


FIG. 40

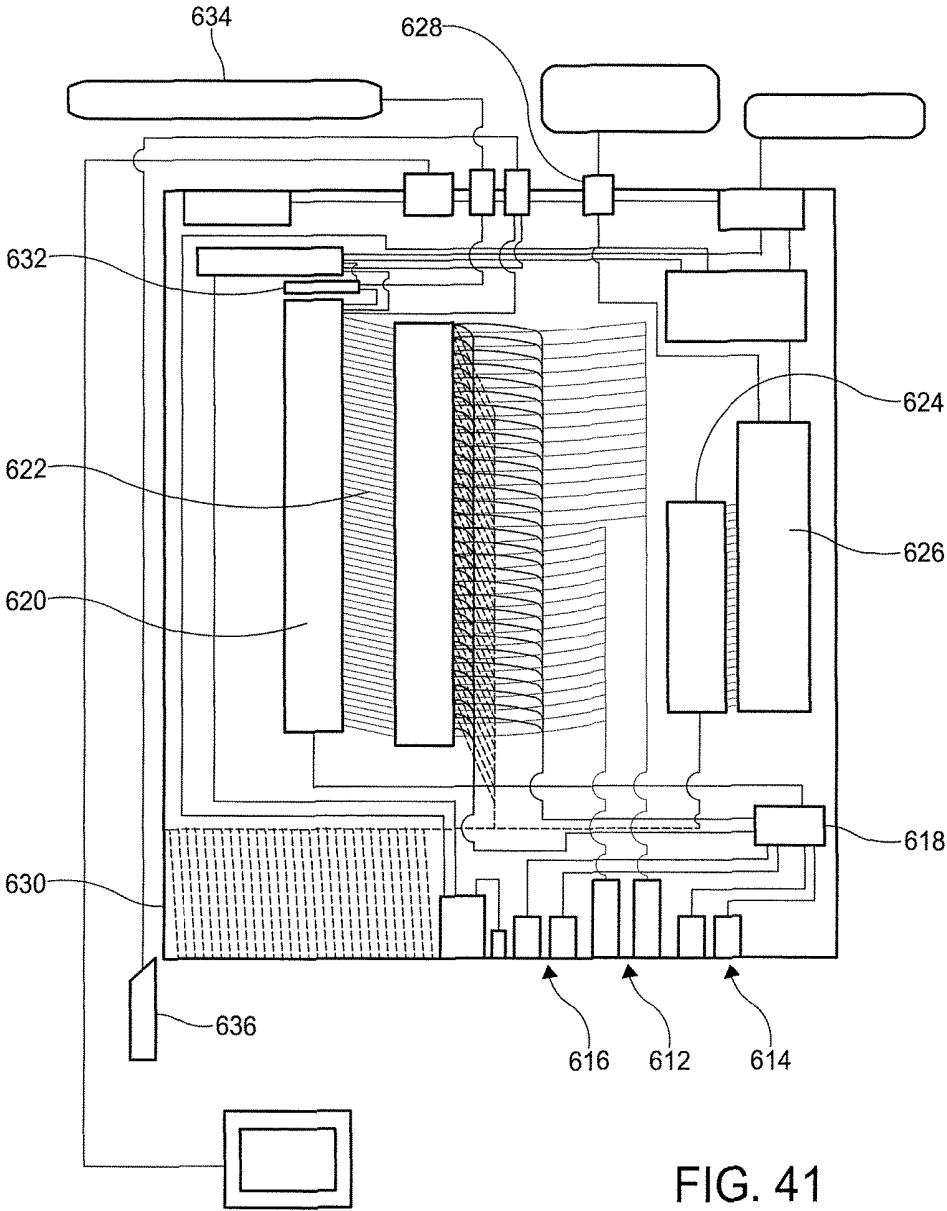


FIG. 41

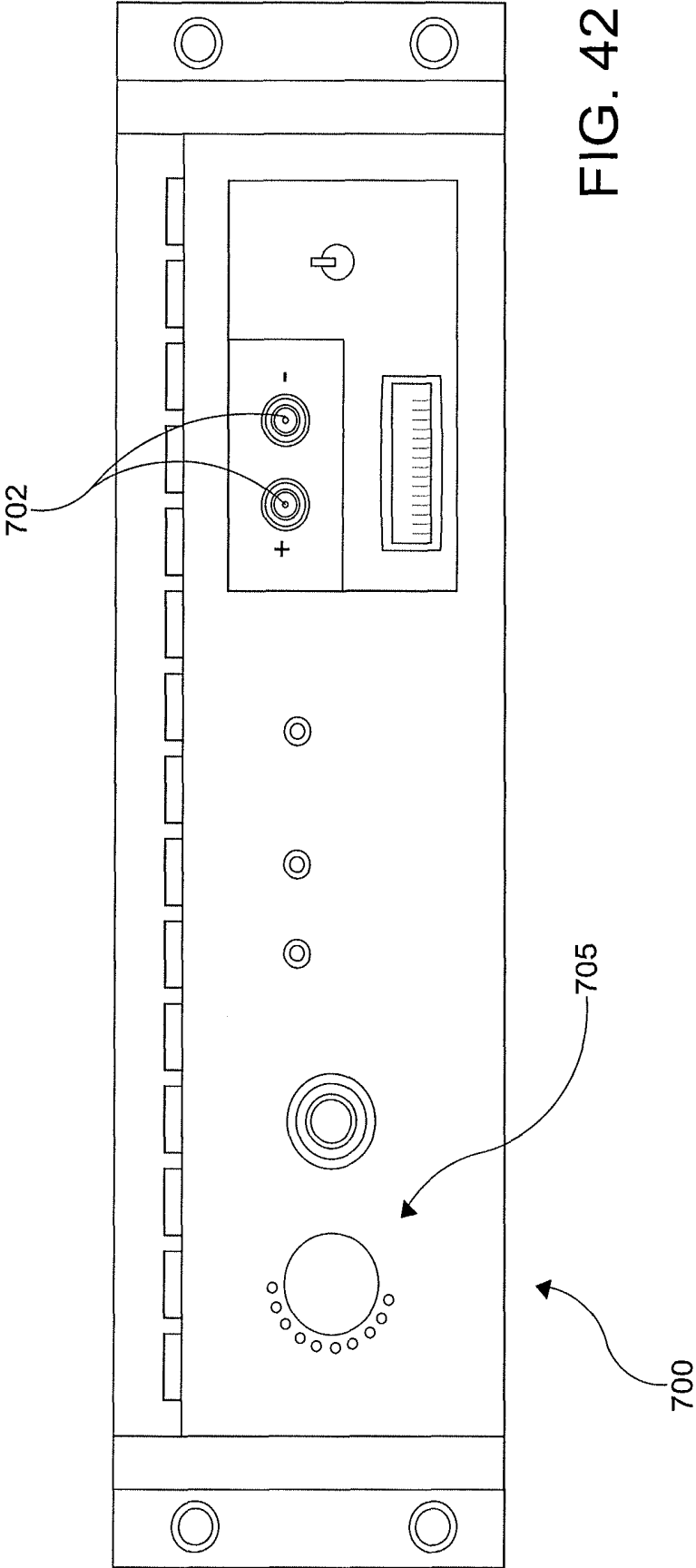
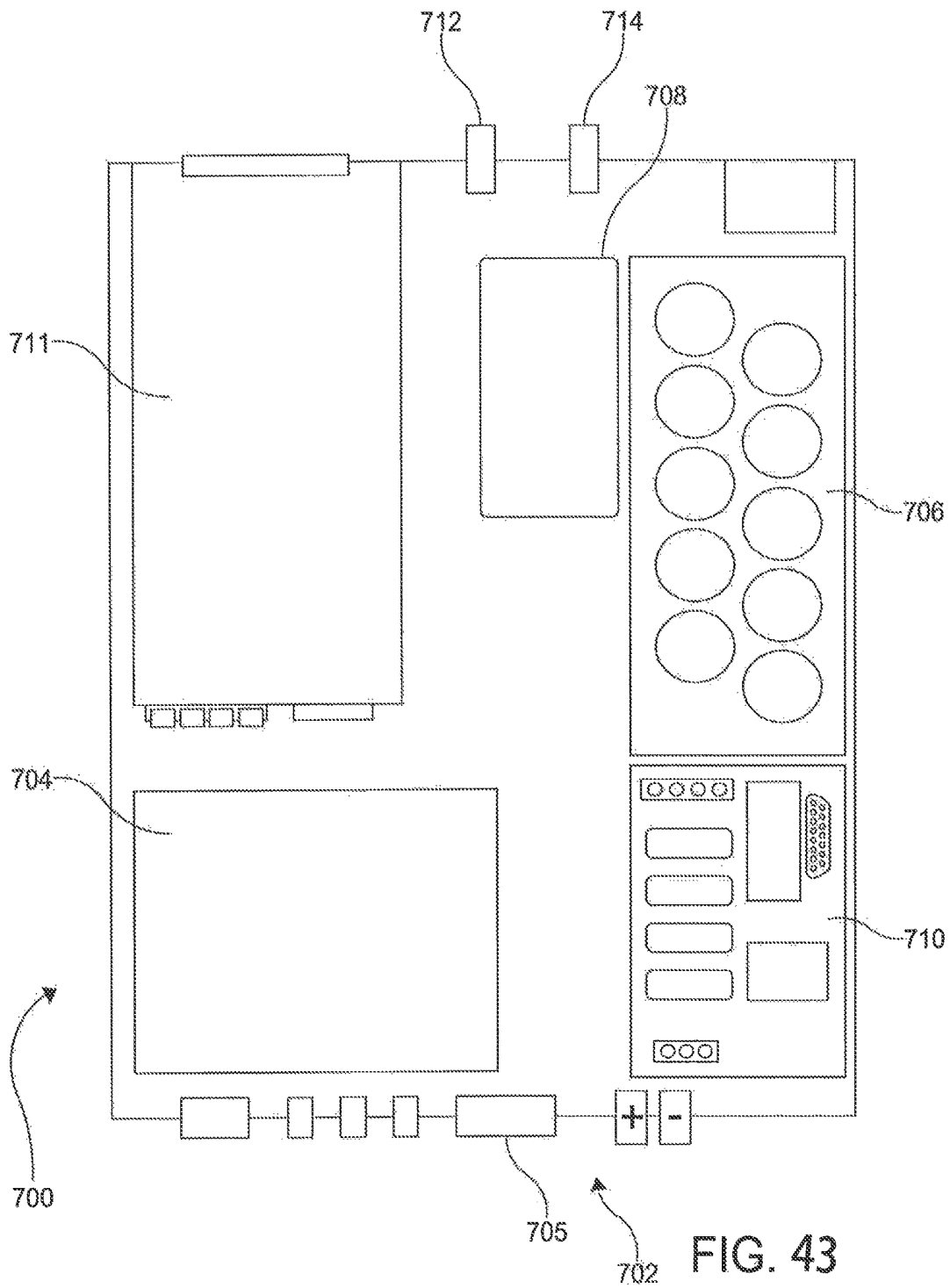


FIG. 42



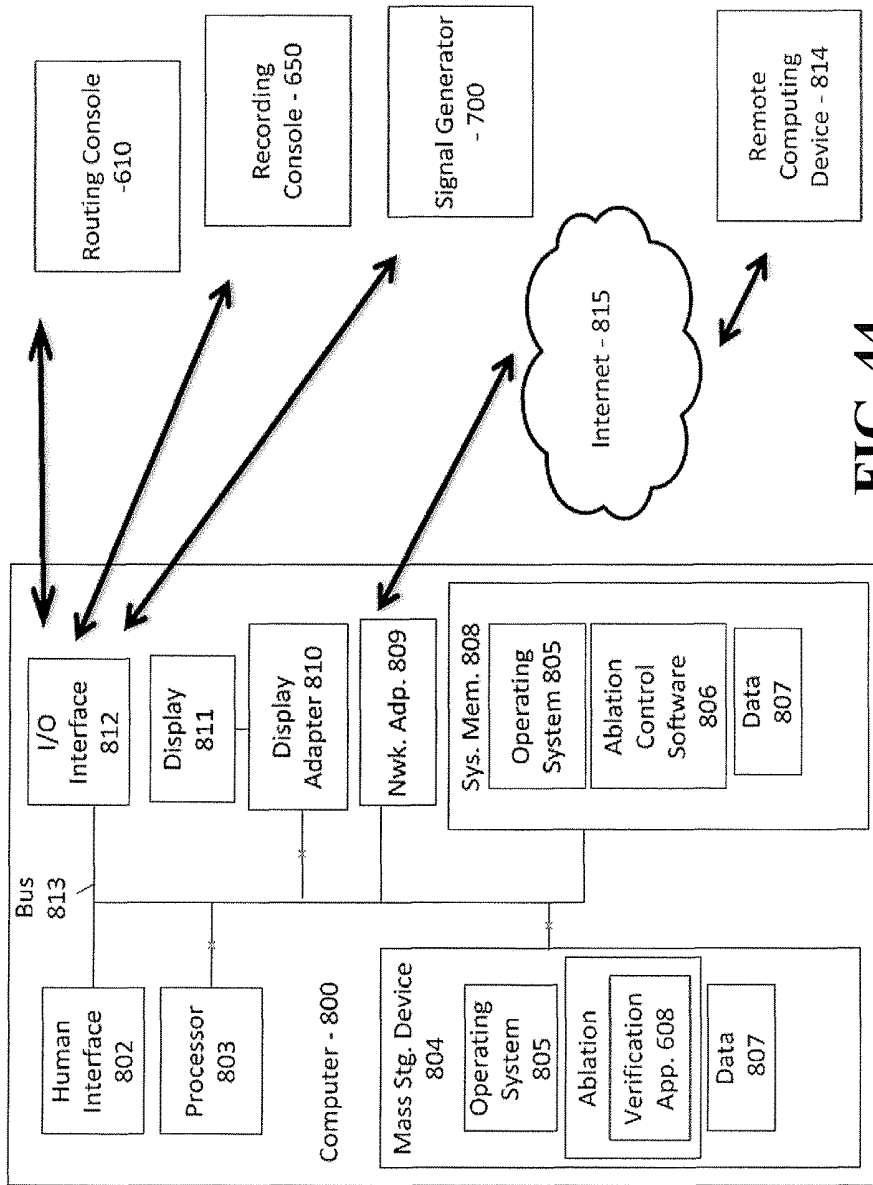


FIG. 44

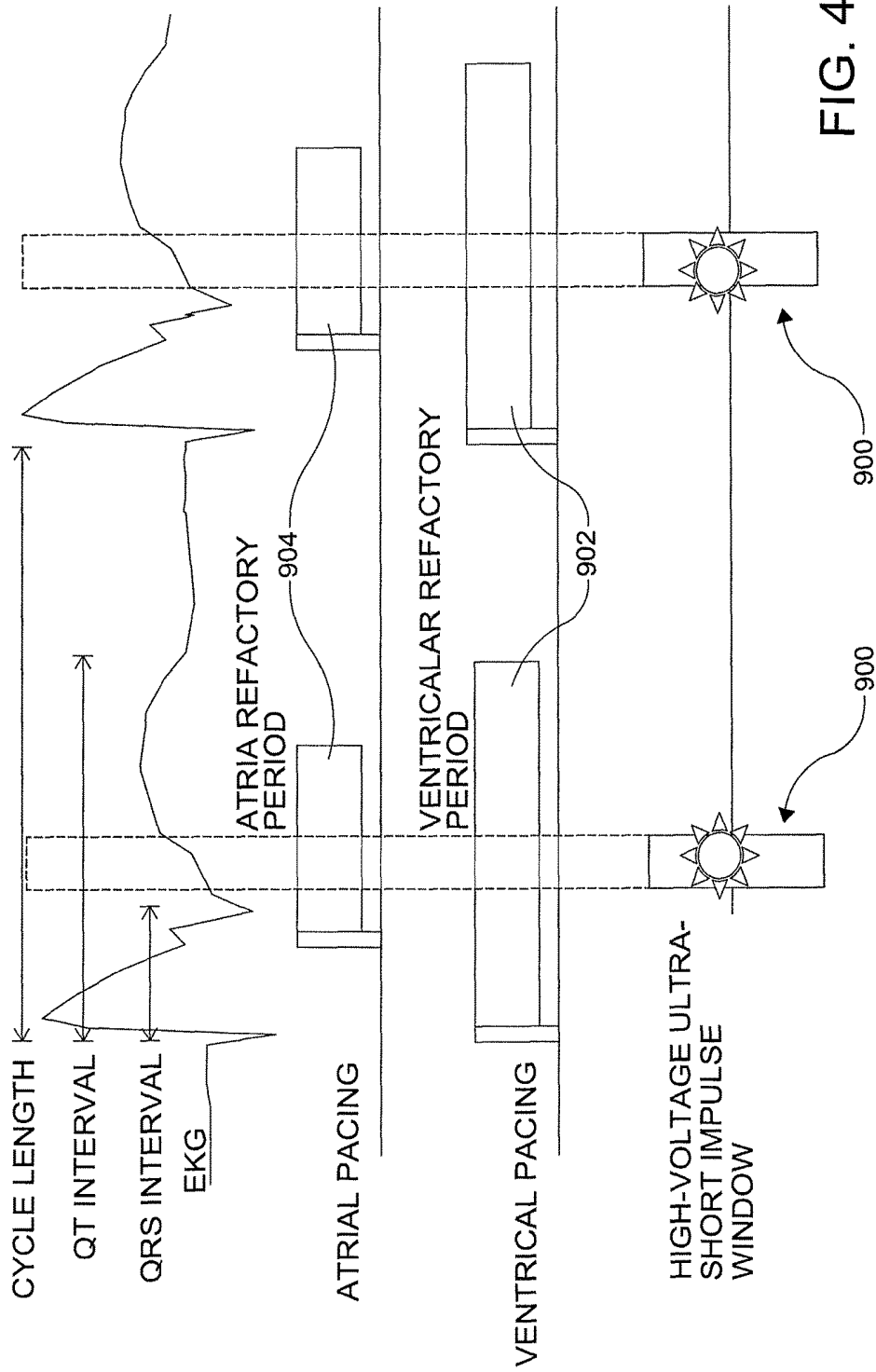


FIG. 45

5000 →

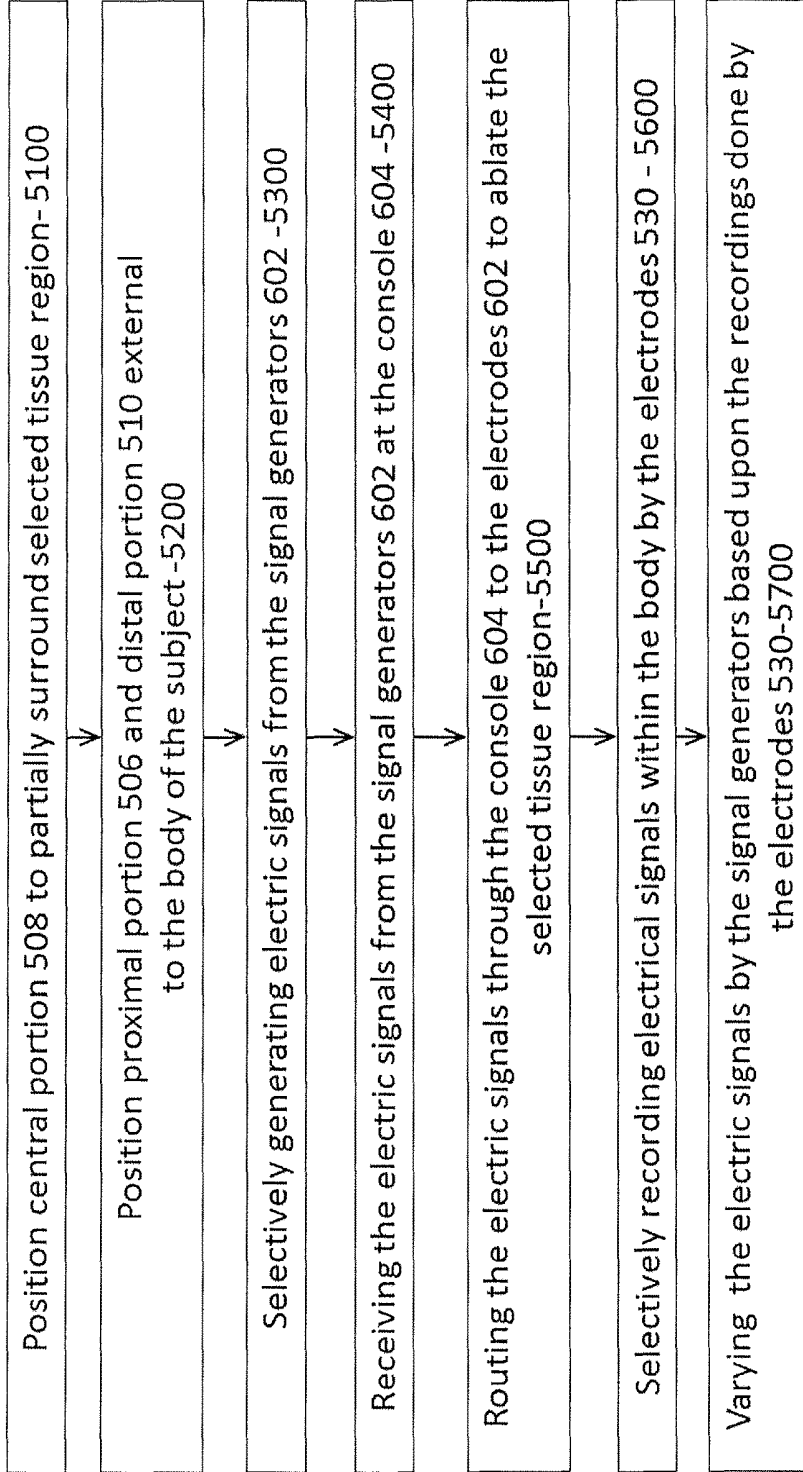


FIG. 46

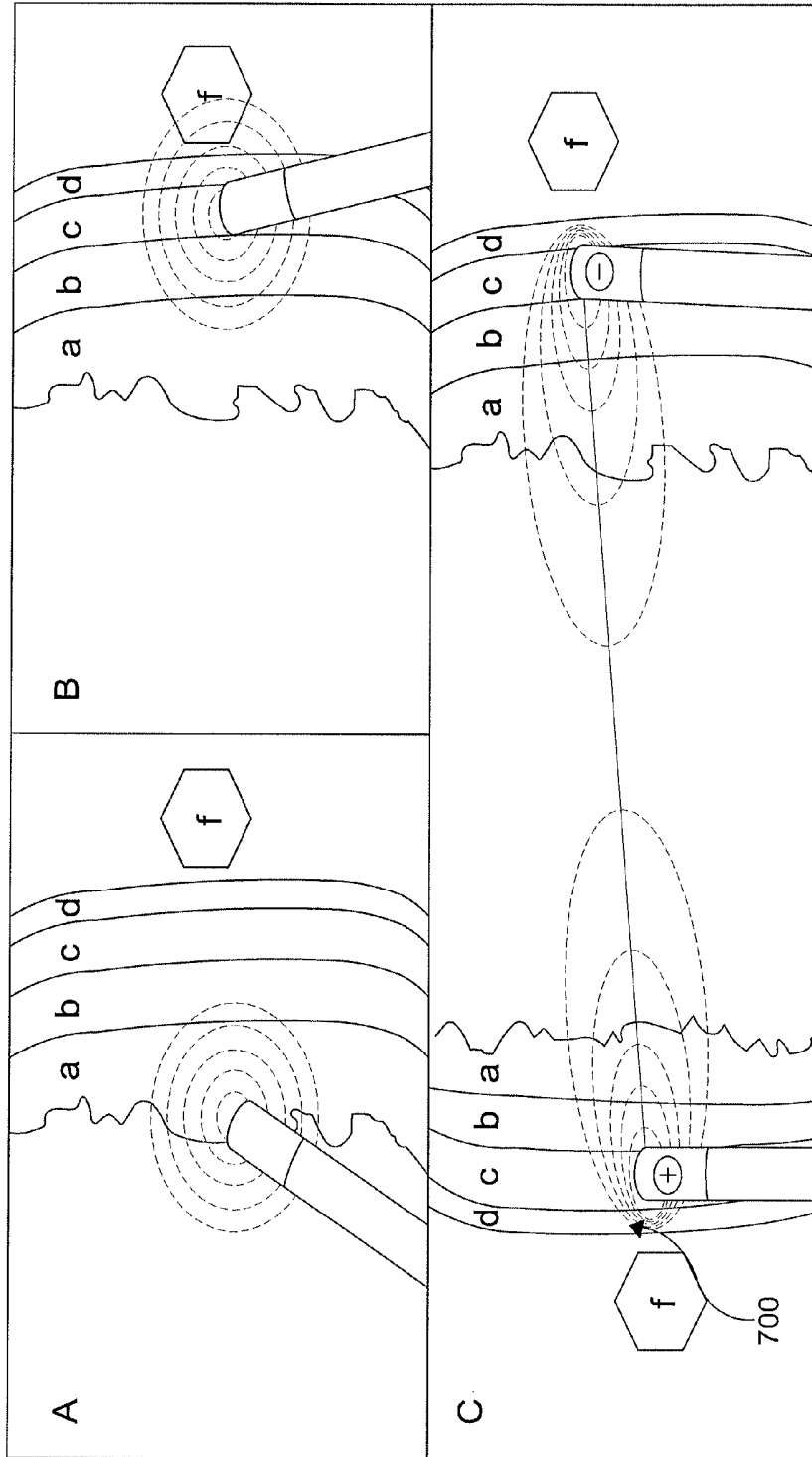


FIG. 47

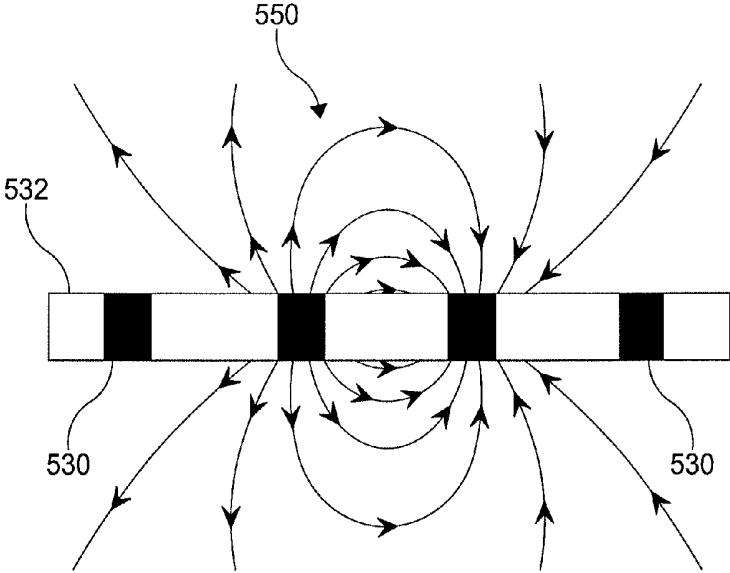


FIG. 48

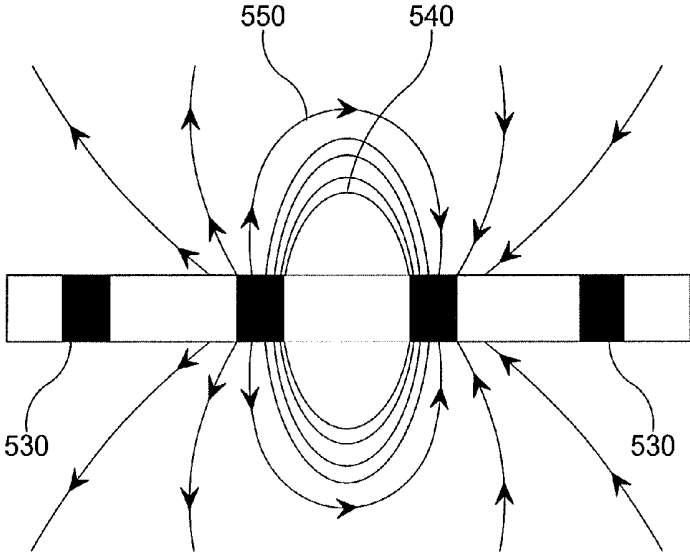


FIG. 49

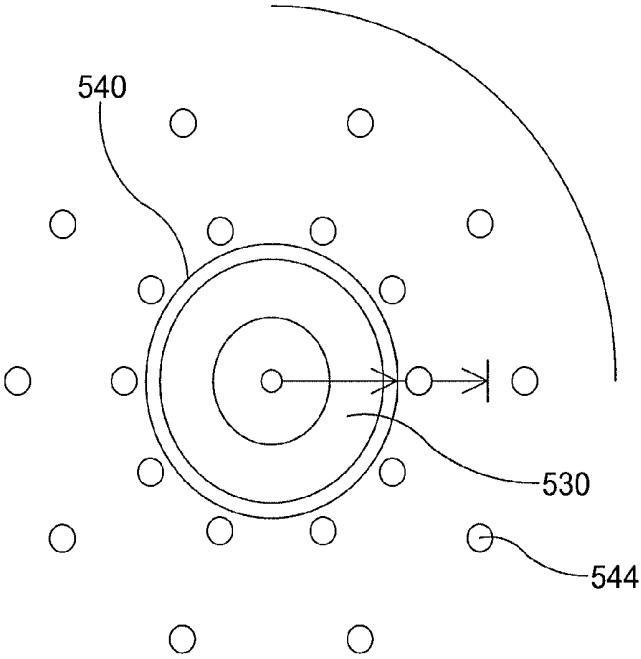


FIG. 50

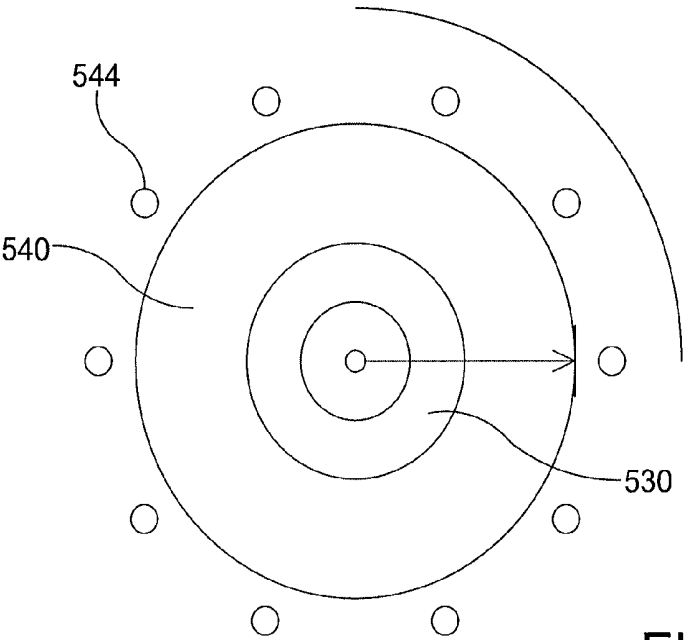


FIG. 51

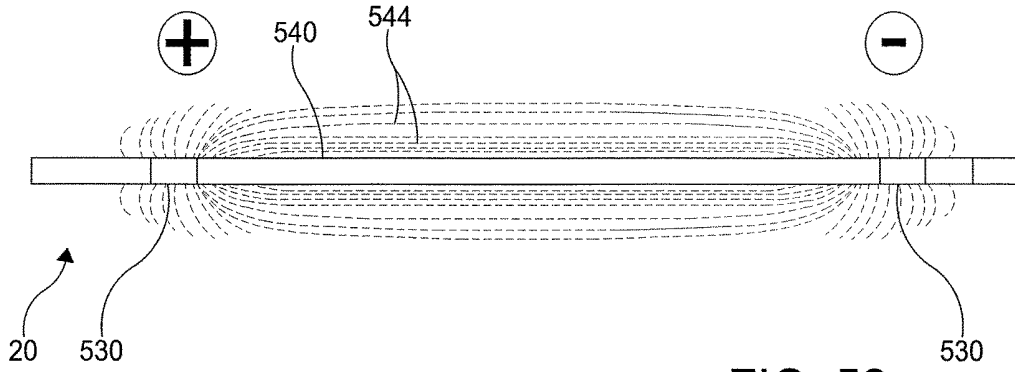


FIG. 52a

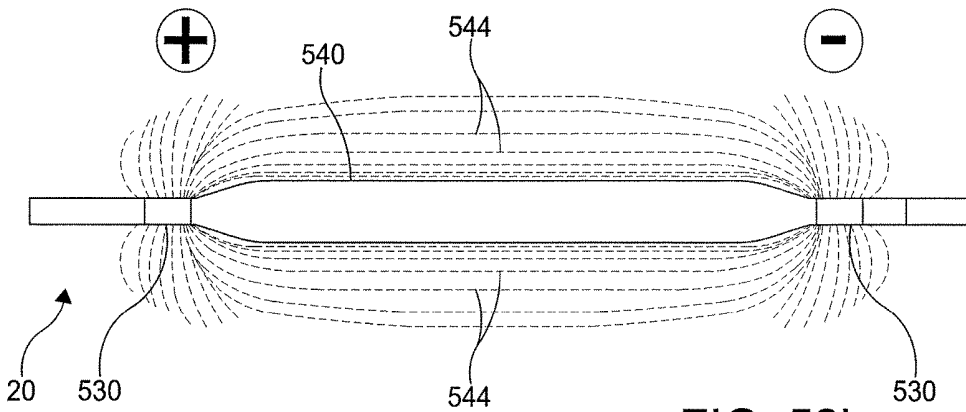


FIG. 52b

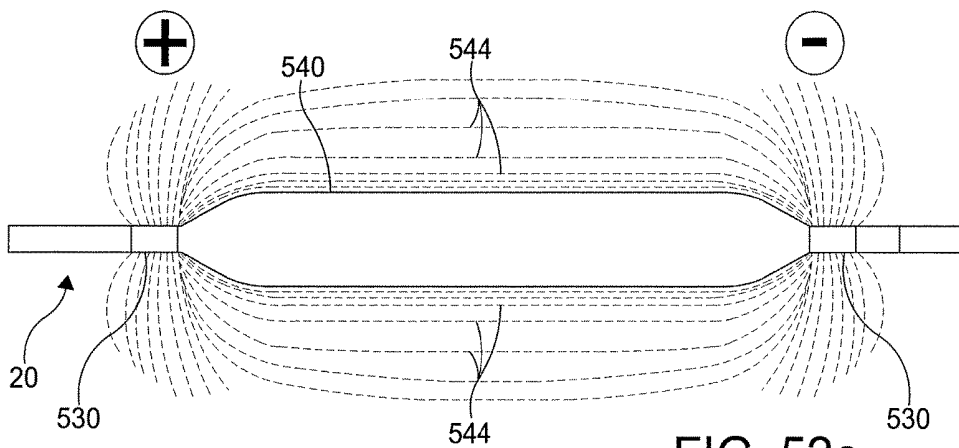
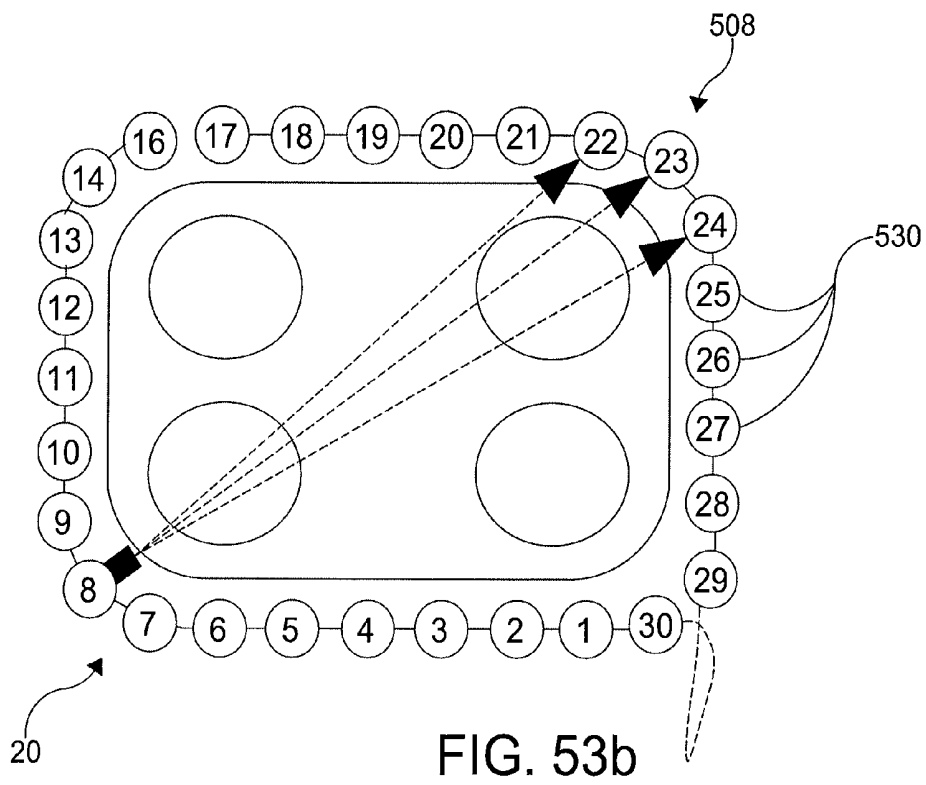
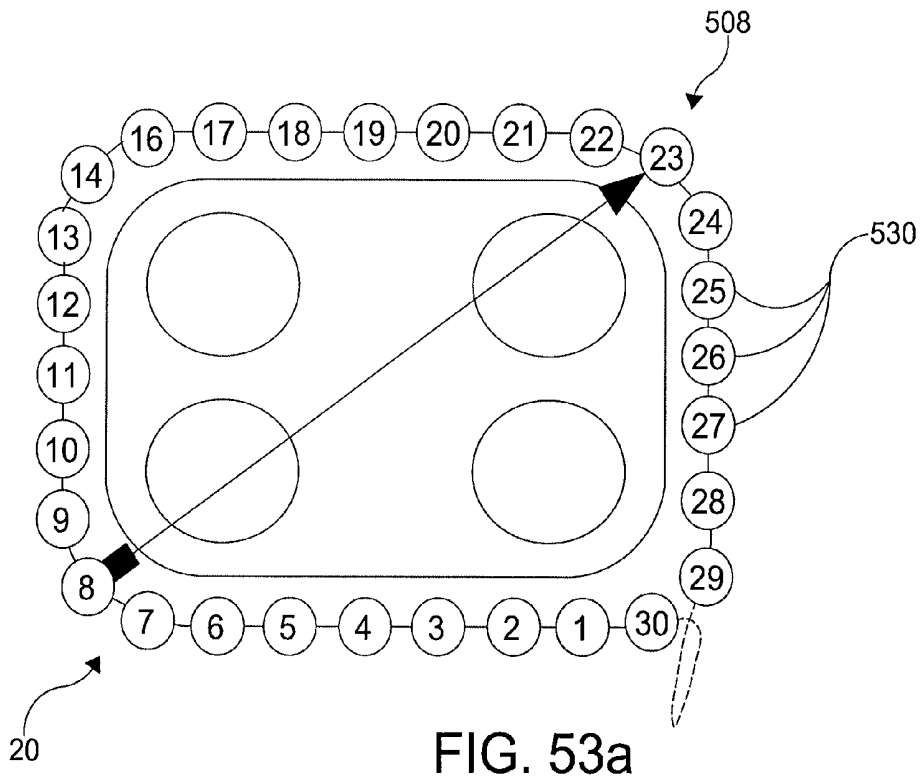
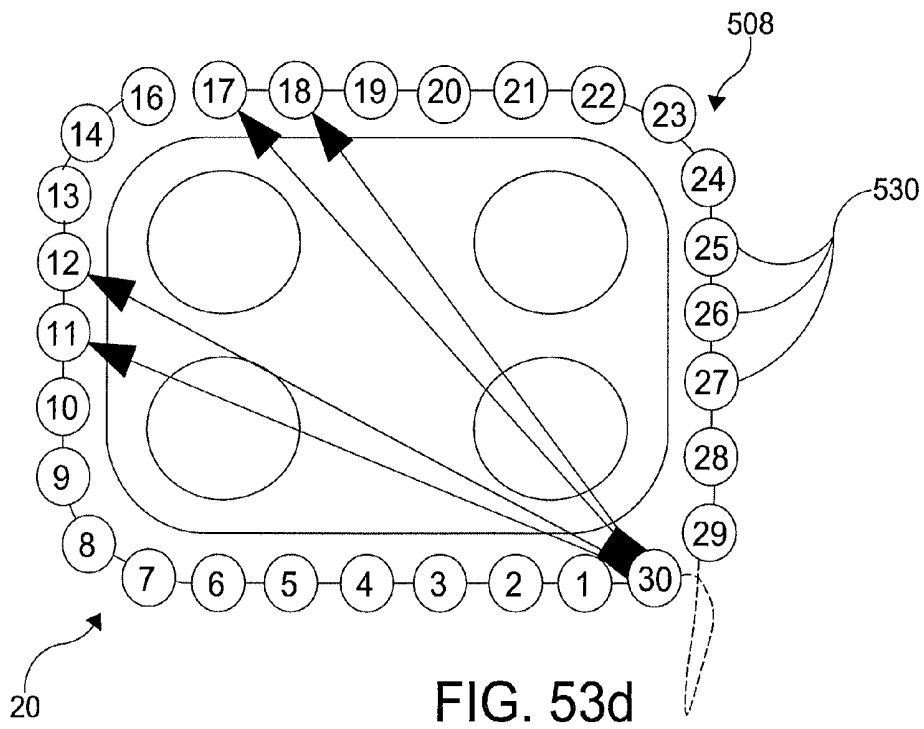
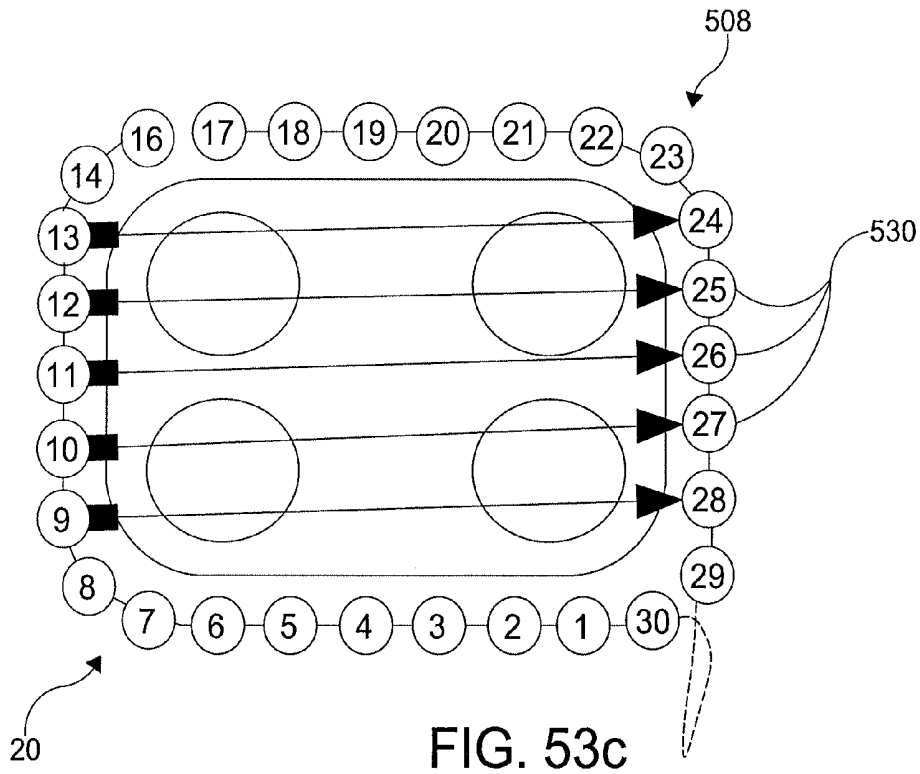


FIG. 52c





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

CLAIM OF PRIORITY

This application claims priority from U.S. Provisional Patent Application No. 61/681,552 filed on Aug. 9, 2012, which is relied upon and incorporated herein in its 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 0.02.5 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 to intersect the theoretical 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.

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.

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.

FIGS. 47a-c are 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 peri-

cardial 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 atraumatic 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 compounds. 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 lubri-

cious 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 boundaries. 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 adjuvant 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) 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 positioned 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 atrial 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 extending into the epicardial fat (b), pericardial space (c), and parietal pericardium (d). However, the field also extends to a bystander vulnerable structure (1). 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 530 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:

$$J = J_i \left(\frac{\Pi r^2 * I_i}{(\Pi r^2 * I_i)_2 - (\Pi r^2 * I_i)} \right)$$

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 to the electric field strength E). ($J = \text{Sigma} \cdot E$). 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 from 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 a laser light or ultrasound, the combination of electrodes **530** and high impedance structures **540** oriented on a 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 $\frac{1}{10}^{th}$ 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. Confirmation 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 RE 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 RE 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 dependency 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 comput-

ing 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 electrophysi-

ologically 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 of treating a heart of a subject from the pericardial space of the heart, comprising:

positioning within the body of the subject a catheter having a proximal portion, a distal portion, and a central portion disposed between the proximal portion and the distal portion, the central portion including a set of electrodes, such that the central portion of the catheter is disposed in the pericardial space of the heart of the subject, and such that the central portion of the 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 at least a portion of the left atrium of the heart of the subject using two or more electrodes of the set of electrodes, wherein the portion of the left atrium includes: a first wall portion of the left atrium; an endocardial space of the left atrium; and a second wall portion of the left atrium.

2. The method of claim 1, the positioning the catheter including passing the catheter through one or more pericardial reflections of the heart of the subject.

3. The method of claim 1, the positioning the catheter including inserting the catheter into the body of the subject using a subxiphoid pericardial access point.

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.

5. 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 catheter.

6. 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.

7. The method of claim 1, 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.

8. The method of claim 1, the delivering the electrical energy including delivering direct current impulses.

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

10. 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.

11. 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.

12. 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 heart beat of the subject.

13. The method of claim 1, wherein the catheter is an ablation catheter, further comprising creating a target path within the body of the subject using one or more percutaneous catheters, the positioning the ablation catheter including positioning the ablation catheter in the target path.

14. 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.

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

repositioning the catheter in a second position different from the first position, such that the central portion of the 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.

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

identifying one or more intended current vectors for delivering the electrical energy, the one or more intended current vectors passing through the endocardial space of the left atrium;

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.

17. The method of claim 1, the positioning the catheter including passing the catheter over a guidewire within the body of the subject.

* * * * *

专利名称(译)	导管，导管系统和穿刺组织结构的方法		
公开(公告)号	US9861802	公开(公告)日	2018-01-09
申请号	US14/400455	申请日	2013-03-14
[标]申请(专利权)人(译)	衣阿华大学研究基金会		
申请(专利权)人(译)	IOWA研究基金会大学		
当前申请(专利权)人(译)	IOWA研究基金会大学		
[标]发明人	MICKELSEN STEVEN RICHARD		
发明人	MICKELSEN, STEVEN RICHARD		
IPC分类号	A61B18/14 A61M39/10 A61M25/01 A61M25/06 A61B5/00 A61B18/12 A61B5/053 A61B34/00 A61B17/00 A61B18/20 A61B18/00 A61B18/18 A61B18/02 A61B17/22 A61M25/00		
CPC分类号	A61M39/10 A61B5/4833 A61B18/1206 A61B18/1492 A61M25/0169 A61M25/0606 A61B5/0538 A61M2025/0089 A61B17/22004 A61B18/02 A61B18/1815 A61B18/20 A61B2017/00876 A61B2018/00363 A61B2018/00375 A61B2018/00577 A61B2018/00613 A61B2018/128 A61B2034/731 A61B2217/005 A61B2218/002 A61B34/73 A61B2018/00839 A61B2018/1467 A61M25/0127 A61M25/09		
优先权	61/681552 2012-08-09 US		
其他公开文献	US20150182740A1		
外部链接	Espacenet USPTO		

摘要(译)

一种用于人体内的经皮导管系统和一种消融导管，用于消融受试者体内的选定组织区域。经皮导管系统可包括两个导管，这两个导管通过组织结构的磁耦合可操作地彼此连接。消融导管可包括位于中心部分内的电极。消融导管定位成使得柔性轴的中心部分至少部分地围绕所选择的组织区域。可以独立地激活消融导管的每个电极以将消融能量施加到所选择的组织区域。消融导管可以采用高阻抗结构来改变特定点处的电流密度。公开了使用经皮导管系统穿透组织结构的方法。还公开了使用消融导管消融选定组织区域的方法。

