



US009220427B2

(12) **United States Patent**
Narayan et al.

(10) **Patent No.:** **US 9,220,427 B2**
(45) **Date of Patent:** ***Dec. 29, 2015**

(54) **SYSTEM AND METHOD FOR
RECONSTRUCTING CARDIAC ACTIVATION
INFORMATION**

(71) Applicants: **The Regents of the University of
California**, Oakland, CA (US); **Topera,
Inc.**, Menlo Park, CA (US)

(72) Inventors: **Sanjiv Narayan**, La Jolla, CA (US);
Carey Robert Briggs, La Jolla, CA
(US); **Ruchir Sehra**, Scottsdale, AZ
(US)

(73) Assignees: **The Regents of the University of
California**, Oakland, CA (US); **Topera,
Inc.**, Menlo Park, CA (US); **The United
States of America as Represented by
the Department of Veterans Affairs,
Office of the General Counsel**,
Washington, DC (US)

(*) Notice: Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 0 days.

This patent is subject to a terminal dis-
claimer.

(21) Appl. No.: **14/670,300**

(22) Filed: **Mar. 26, 2015**

(65) **Prior Publication Data**

US 2015/0196212 A1 Jul. 16, 2015

Related U.S. Application Data

(63) Continuation of application No. 13/840,334, filed on
Mar. 15, 2013, which is a continuation-in-part of
application No. 13/438,534, filed on Apr. 3, 2012, now
Pat. No. 8,594,777, which is a continuation of

(Continued)

(51) **Int. Cl.**
A61B 5/04 (2006.01)
A61B 5/042 (2006.01)

(Continued)

(52) **U.S. Cl.**
CPC **A61B 5/04012** (2013.01); **A61B 5/0205**
(2013.01); **A61B 5/046** (2013.01);

(Continued)

(58) **Field of Classification Search**

CPC .. A61B 5/0452; A61B 5/4362; A61B 5/0245;
A61B 5/0464; A61N 1/3962; A61N 1/3956;
A61N 1/368; A61N 1/3627; A61N 1/36507
USPC 607/4-5, 9, 25, 26
See application file for complete search history.

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,029,082 A 7/1991 Shen et al.
5,121,750 A 6/1992 Katims

(Continued)

FOREIGN PATENT DOCUMENTS

WO 9421168 A1 9/1994
WO 9625096 A1 8/1996

(Continued)

OTHER PUBLICATIONS

Holm, M. et al., "A New Method for Analysis of Atrial Activation
During Chronic Atrial Fibrillation in Man", IEEE Transactions on
Biomedical Engineering, vol. 43, No. 2, Feb. 1, 1996, pp. 198-210.

(Continued)

Primary Examiner — Christopher D Koharski

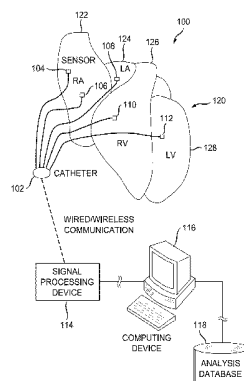
Assistant Examiner — Roland Dinga

(74) *Attorney, Agent, or Firm* — Eleanor Musick; Greer,
Burns & Crain, Ltd.

(57) **ABSTRACT**

In a system and method for reconstructing cardiac activation
information, an analysis cardiac signal and a reference car-
diac signal are processed to determine a first point of change
in a derivative of the analysis cardiac signal with respect to a
derivative of the reference cardiac signal. The analysis car-
diac signal and the reference cardiac signal are processed to
determine a second point of change in a different deriva-
tive of the analysis cardiac signal with respect to a different deriva-
tive of the reference cardiac signal. An activation onset time is
assigned in the analysis cardiac signal at a point based on a
mathematical association of the first point of change and the
second point of change to define cardiac activation indicating
a beat in the analysis cardiac signal.

37 Claims, 8 Drawing Sheets



Related U.S. Application Data

application No. 13/217,123, filed on Aug. 24, 2011,
now Pat. No. 8,165,666.

- (60) Provisional application No. 61/481,607, filed on May 2, 2011.

(51) **Int. Cl.**

A61B 5/0452 (2006.01)

A61B 5/00 (2006.01)

A61B 5/0205 (2006.01)

A61B 5/046 (2006.01)

A61B 5/0464 (2006.01)

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

CPC *A61B5/04011* (2013.01); *A61B 5/0422*
(2013.01); *A61B 5/0452* (2013.01); *A61B*
5/0464 (2013.01); *A61B 5/7203* (2013.01);
A61B 5/7239 (2013.01); *A61B 2562/046*
(2013.01)

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,172,699	A	12/1992	Svenson et al.
5,427,112	A	6/1995	Noren et al.
5,433,198	A	7/1995	Desai
5,450,846	A	9/1995	Goldreyer
5,458,621	A	10/1995	White et al.
5,480,422	A	1/1996	Ben-Haim
5,487,391	A	1/1996	Panescu
5,582,173	A	12/1996	Li
5,645,070	A	7/1997	Turcott
5,657,755	A	8/1997	Desai
5,662,108	A	9/1997	Budd et al.
5,687,737	A	11/1997	Branham et al.
5,718,241	A	2/1998	Ben-Haim et al.
5,795,303	A	8/1998	Swanson et al.
5,810,740	A	9/1998	Paisner
5,817,134	A	10/1998	Greenhut et al.
5,840,025	A	11/1998	Ben-Haim
5,848,972	A	12/1998	Triedman et al.
5,868,680	A	2/1999	Steiner et al.
5,954,665	A	9/1999	Ben-Haim
6,052,618	A	4/2000	Dahlke et al.
6,066,094	A	5/2000	Ben-Haim
6,115,628	A	9/2000	Stadler et al.
6,188,924	B1	2/2001	Swanson et al.
6,208,888	B1	3/2001	Yonce
6,236,883	B1	5/2001	Ciaccio et al.
6,256,540	B1	7/2001	Panescu et al.
6,301,496	B1	10/2001	Reisfeld
6,360,121	B1	3/2002	Shoda et al.
6,438,406	B2	8/2002	Yonce
6,438,409	B1	8/2002	Malik et al.
6,449,503	B1	9/2002	Hsu
6,522,905	B2	2/2003	Desai
6,584,345	B2	6/2003	Govari
6,847,839	B2	1/2005	Ciaccio et al.
6,856,830	B2	2/2005	He
6,889,081	B2	5/2005	Hsu
6,892,091	B1	5/2005	Ben-Haim et al.
6,920,350	B2	7/2005	Xue et al.
6,941,166	B2	9/2005	MacAdam et al.
6,950,696	B2	9/2005	Bjorling et al.
6,959,212	B2	10/2005	Hsu et al.
6,975,900	B2	12/2005	Rudy et al.
6,978,168	B2	12/2005	Beatty et al.
7,016,719	B2	3/2006	Rudy et al.
7,043,292	B2	5/2006	Tarjan et al.
7,117,030	B2	10/2006	Berenfeld et al.
7,123,954	B2	10/2006	Narayan et al.
7,206,630	B1	4/2007	Tarler
7,245,962	B2	7/2007	Ciaccio et al.
7,263,397	B2	8/2007	Hauck et al.
7,283,865	B2	10/2007	Noren
7,289,843	B2	10/2007	Beatty et al.
7,328,063	B2	2/2008	Zhang et al.
7,505,810	B2	3/2009	Harlev et al.
7,515,954	B2	4/2009	Harlev et al.
7,567,835	B2	7/2009	Gunderson et al.
7,580,744	B2	8/2009	Hsu
7,734,333	B2	6/2010	Ghanem et al.
7,742,812	B2	6/2010	Ghanem et al.
7,751,882	B1	7/2010	Helland
7,761,142	B2	7/2010	Ghanem et al.
7,761,150	B2	7/2010	Ghanem et al.
7,769,452	B2	8/2010	Ghanem et al.
7,907,993	B2	3/2011	Ghanem et al.
7,907,994	B2	3/2011	Stolarski et al.
7,930,018	B2	4/2011	Harlev et al.
7,930,020	B2	4/2011	Zhang et al.
7,953,475	B2	5/2011	Harlev et al.
7,962,202	B2	6/2011	Bhunia
8,050,732	B2	11/2011	Desai
8,095,205	B2	1/2012	Bhunia
8,095,206	B2	1/2012	Ghanem et al.
8,160,684	B2	4/2012	Ghanem et al.
8,165,666	B1	4/2012	Briggs et al.
8,165,671	B2	4/2012	Freeman et al.
8,175,702	B2	5/2012	Efimov et al.
8,306,618	B2	11/2012	Ghanem et al.
8,315,697	B2	11/2012	Hsu
8,340,766	B2	12/2012	Ryu et al.
8,386,024	B2	2/2013	Gunderson et al.
8,435,185	B2	5/2013	Ghanem et al.
8,521,266	B2	8/2013	Narayan et al.
8,594,777	B2	11/2013	Briggs et al.
8,639,325	B2	1/2014	Efimov et al.
8,676,303	B2	3/2014	Narayan
8,700,140	B2	4/2014	Narayan et al.
8,838,222	B2	9/2014	Narayan et al.
8,838,223	B2	9/2014	Narayan et al.
8,868,169	B2	10/2014	Narayan et al.
9,050,006	B2	6/2015	Narayan et al.
9,055,876	B2	6/2015	Narayan et al.
9,055,877	B2	6/2015	Narayan et al.
9,055,878	B2	6/2015	Narayan et al.
2002/0010392	A1	1/2002	Desai
2003/0018277	A1	1/2003	He
2003/0083587	A1	5/2003	Ferek-Petric
2003/0236466	A1	12/2003	Tarjan
2004/0073262	A1	4/2004	Lovett
2005/0148892	A1	7/2005	Desai
2005/0203502	A1	9/2005	Boveja et al.
2006/0084970	A1	4/2006	Beatty et al.
2006/0161206	A1	7/2006	Efimov et al.
2007/0055167	A1	3/2007	Bullinga
2007/0208260	A1	9/2007	Afonso
2007/0232948	A1	10/2007	Stadler et al.
2007/0299351	A1	12/2007	Harley et al.
2008/0097539	A1	4/2008	Belalcazar et al.
2008/0109041	A1	5/2008	de Voir
2008/0114258	A1	5/2008	Zhang et al.
2008/0188765	A1	8/2008	Stolarski et al.
2009/0069704	A1	3/2009	MacAdam et al.
2009/0099468	A1	4/2009	Thiagalingam et al.
2009/0112106	A1	4/2009	Zhang
2009/0112110	A1	4/2009	Zhang et al.
2009/0112199	A1	4/2009	Zhang et al.
2009/0177071	A1	7/2009	Harlev et al.
2009/0177072	A1	7/2009	Harlev et al.
2009/0299424	A1	12/2009	Narayan
2010/0094274	A1	4/2010	Narayan
2010/0204592	A1	8/2010	Hatib et al.
2010/0217143	A1	8/2010	Whittington et al.
2010/0239627	A1	9/2010	Zhang et al.
2010/0298729	A1	11/2010	Zhang et al.
2010/0305456	A1	12/2010	Brainard, II
2011/0087121	A1	4/2011	Zhang et al.
2011/0112425	A1	5/2011	Muhlsteff et al.
2011/0130801	A1	6/2011	Maskara et al.
2011/0196249	A1	8/2011	Staeuber et al.
2011/0251505	A1	10/2011	Narayan

(56)

References Cited**U.S. PATENT DOCUMENTS**

2011/0257547	A1	10/2011	Zhang et al.
2011/0282227	A1	11/2011	Zhang
2012/0184858	A1	7/2012	Harlev et al.
2012/0232417	A1	9/2012	Zhang et al.
2012/0283579	A1	11/2012	Briggs et al.
2013/0006131	A1	1/2013	Narayan et al.
2013/0150740	A1	6/2013	Narayan et al.
2013/0150742	A1	6/2013	Briggs et al.
2013/0324871	A1	12/2013	Dubois et al.
2013/0331718	A1	12/2013	Narayan et al.
2014/0228696	A1	8/2014	Narayan et al.
2014/0276152	A1	9/2014	Narayan et al.

FOREIGN PATENT DOCUMENTS

WO	9632885	A1	10/1996
WO	9632897	A1	10/1996
WO	9639929	A1	12/1996
WO	9724983	A2	7/1997
WO	0045700	A1	8/2000
WO	2008035070	A3	3/2008

OTHER PUBLICATIONS

Kadish, A., et al., "Characterization of fibrillatory rhythms by ensemble vector directional analysis", *Am J Physiol.—Heart Circ. Physiol.*, vol. 285, Oct. 2003, pp. H1705-H1719.

Masse, S., et al., "Wave similarity of human ventricular fibrillation from bipolar electrograms", *Eurospace* (2007) vol. 9, pp. 10-19.

Ulphani, J.S., et al., "Frequency gradients during two different forms of fibrillation in canine atria", *Heart Rhythm*, vol. 4, No. 10, Oct. 2007, pp. 1315-1323.

EP12855266.8: Supplementary European Search Report & European Search Opinion, Jun. 2, 2015, 9 pages.

EP12855738.6: Supplementary European Search Report & European Search Opinion, Jun. 5, 2015, 9 pages.

Houben, R.P.M., et al., "Automatic mapping of human atrial fibrillation by template matching", *Heart Rhythm*, vol. 3, No. 10, Oct. 1, 2006, pp. 1221-1228.

Nademanee, Koonlawee, et al., "A new approach for catheter ablation of atrial fibrillation: mapping of the electrophysiologic substrate", *J. Amer.Coll.Cardiol.*, vol. 43, No. 11, Jun. 2, 2004, pp. 2044-2053.

Narayan, S.M., et al., "Dynamics factors preceding the initiation of atrial fibrillation in humans", *Heart Rhythm*, vol. 5, No. 6, Jun. 1, 2008, pp. S22-S25.

Ciaccio, Edward J. et al., "Development of Gradient Descent Adaptive Algorithms to Remove Common Mode Artifact for Improvement of Cardiovascular Signal Quality", *Annals of Biomedical Engineering*, vol. 35, No. 7, Jul. 2007, pp. 1146-1155.

Sornborger, Andrew, et al., "Extraction of Periodic Multivariate Signals: Mapping of Voltage-Dependent Dye Fluorescence in the Mouse Heart", *IEEE Transactions on Medical Imaging*, vol. 22, No. 12, Dec. 2003, pp. 1537-1549.

Sun, Yan, et al., "Characteristic wave detection in ECG signal using morphological transform", *BMC Cardiovascular Disorders*, vol. 5, No. 28, 2005.

Tai, Dean C.S., et al., "Correction of motion artifact in transmembrane voltage-sensitive fluorescent dye emission in hearts", *Am. J. Physiol. Heart Circ. Physiol.*, vol. 287, 2004, pp. H985-H993.

Lin, Y-J, et al., "Electrophysiological Characteristics and Catheter Ablation in Patients With Paroxysmal Right Atrial Fibrillation", *Circulation*, Sep. 20, 2005; 112(12): 1692-1700, EPub Sep. 12, 2005.

Houben, R.P.M., et al., "Processing of Intracardiac Electrograms in Atrial Fibrillation", *IEEE Engineering in Medicine and Biology Magazine*, Nov./Dec. 2006, pp. 40-51.

Saksena, S., et al., "Regional Endocardial Mapping of Spontaneous and Induced Atrial Fibrillation in Patients With Heart Disease and Refractory Atrial Fibrillation", *Am J Cardiol*, 1999; 84:880-889.

PCT/US2011/031468: International Preliminary Report on Patentability, Oct. 9, 2012, 8 pages.

PCT/US2009/060178: International Preliminary Report on Patentability, Apr. 12, 2011, 10 pages.

PCT/US2011/031470: International Preliminary Report on Patentability, Oct. 9, 2012, 7 pages.

PCT/US2012/029935: International Search Report and Written Opinion, Nov. 8, 2012, 9 pages.

PCT/US2012/036157: International Preliminary Report on Patentability, Nov. 5, 2013, 8 pages.

PCT/US2012/068640: International Preliminary Report on Patentability, Jun. 10, 2014, 5 pages.

PCT/US2012/068639: International Preliminary Report on Patentability, Jun. 10, 2014, 6 pages.

EP09819953: Supplementary European Search Report and European Search Opinion, Feb. 7, 2012, 5 pages.

EP12711553: Supplementary European Search Report and European Search Opinion, Sep. 11, 2013, 7 pages.

Umapathy, K, et al. "Spatiotemporal Frequency Analysis of Ventricular Fibrillation in Implanted Human Hearts," *IEEE Transactions in Biomedical Engineering*, IEEE Service Center, Piscataway, NJ USA, vol. 56, No. 2, Feb. 1, 2009, pp. 328-335.

Kalifa, J, et al. "Mechanisms of wave fractionation at boundaries of high-frequency excitation in the posterior left atrium of the isolated sheep heart during atrial fibrillation," *Circulation*, vol. 113, No. 5, Feb. 7, 2006, pp. 626-633.

Yenn-Jiang L, et al. "Electrophysiological Mechanisms and Catheter Ablation of Complex Atrial Arrhythmias from Crista Terminalis: Insight from Three-Dimensional Noncontact Mapping," *Pacing and Clinical Electrophysiology*, vol. 27, No. 9, Sep. 1, 2004, pp. 1231-1239.

EP12779506.0: Supplementary European Search Report & European Search Opinion, Nov. 18, 2014, 8 pages.

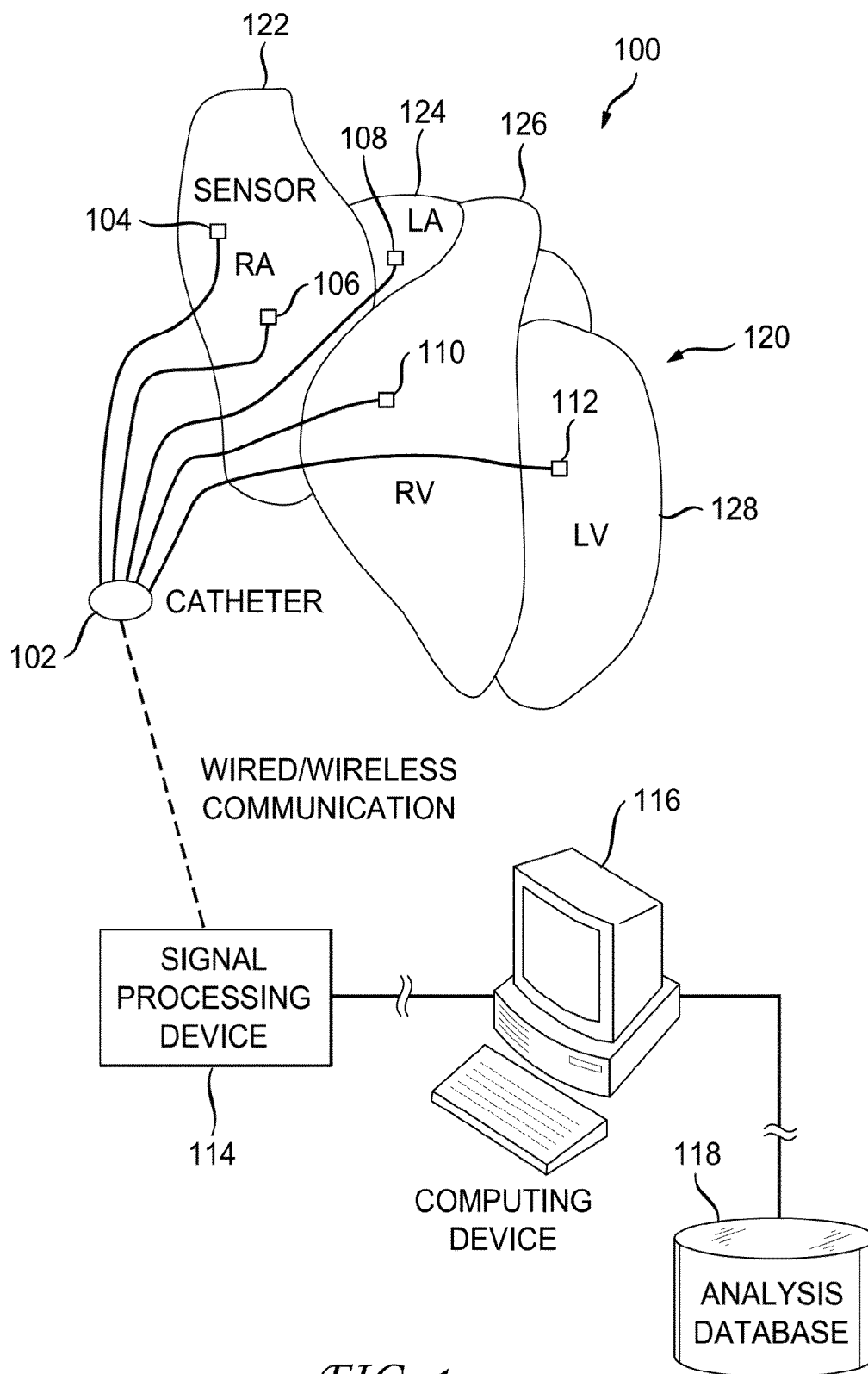
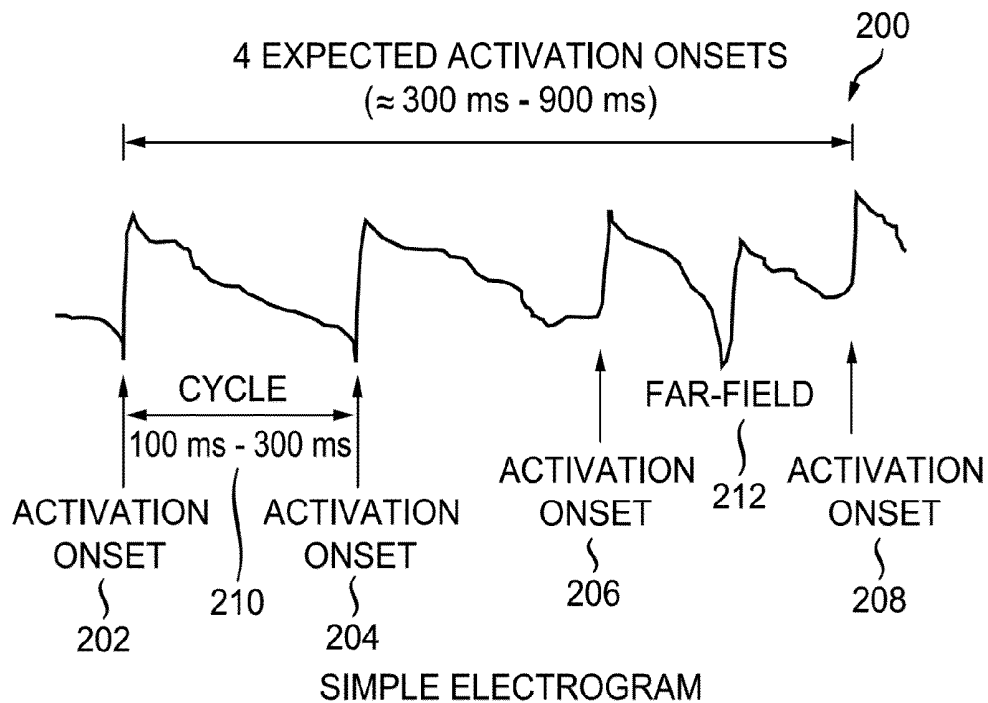
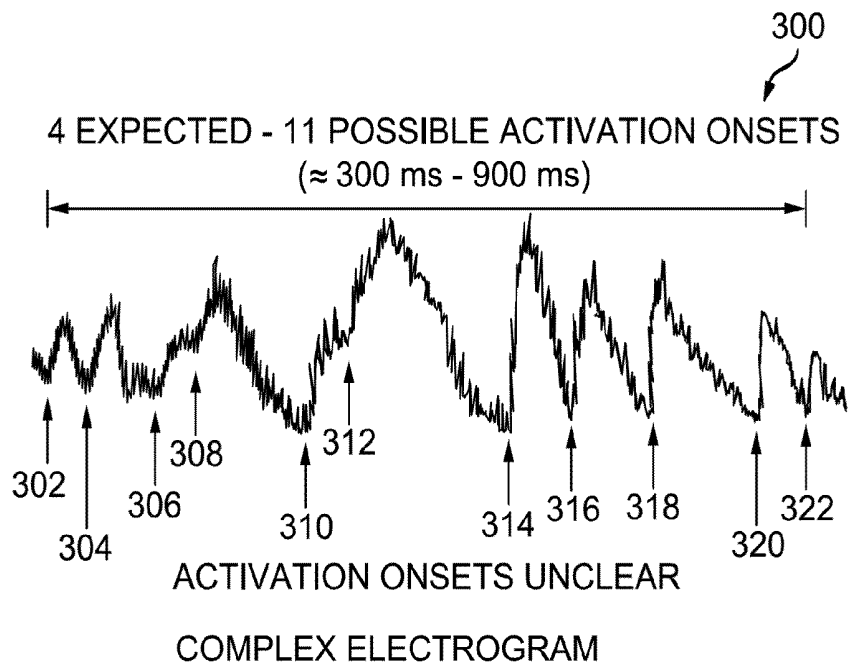


FIG. 1

*FIG. 2**FIG. 3*

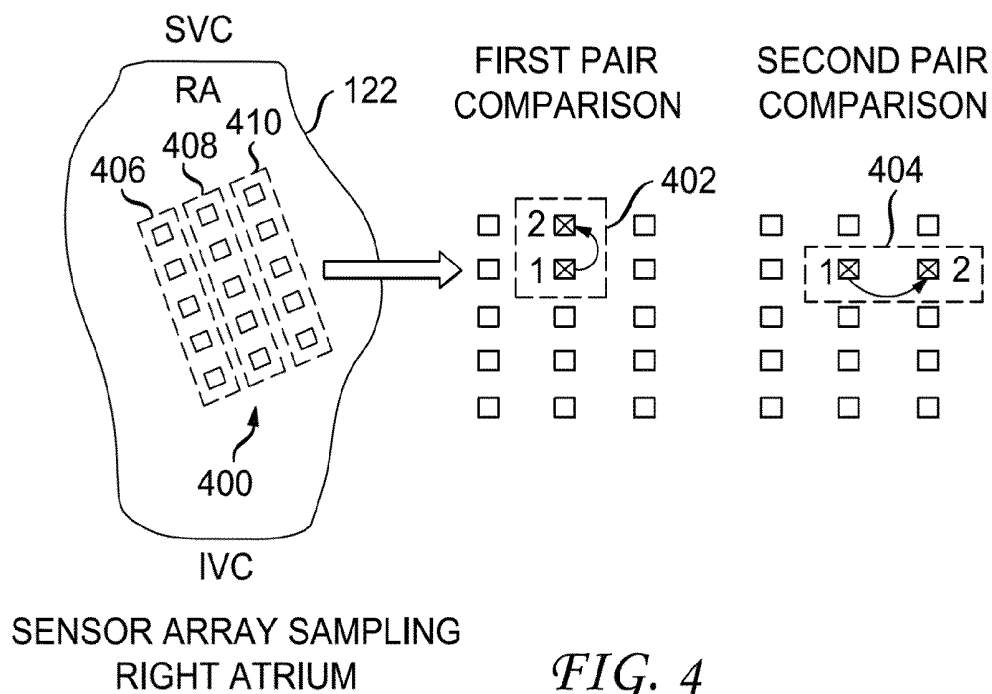


FIG. 4

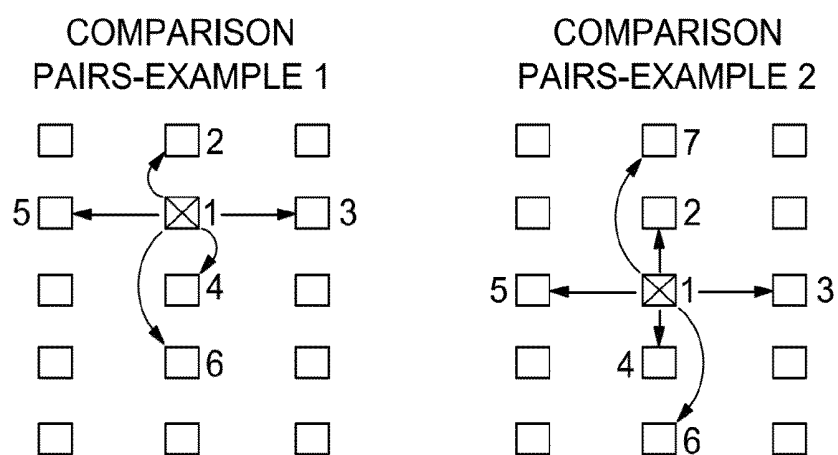


FIG. 5

FAR-FIELD ACTIVATION
(NOT SIGNIFICANTLY LOCAL)

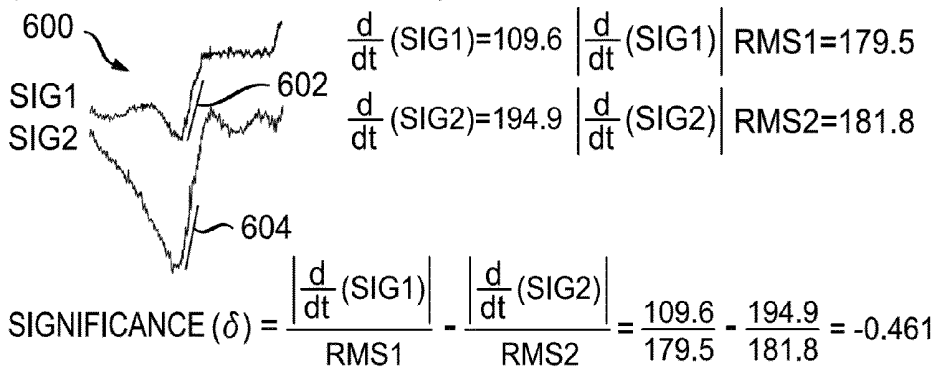


FIG. 6

BACKGROUND NOISE

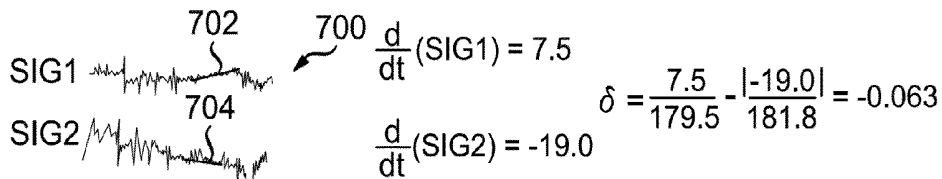
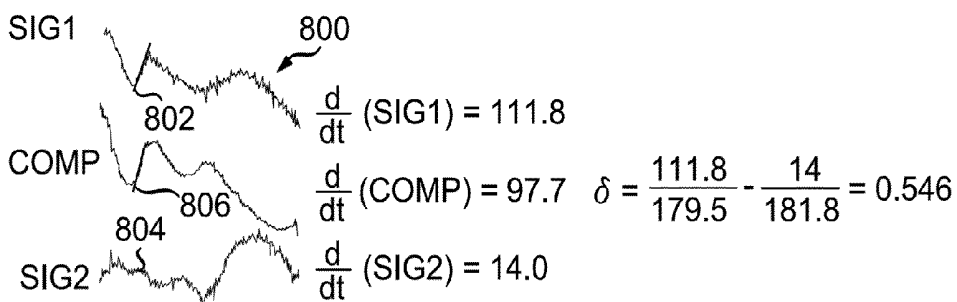


FIG. 7

LOCAL ACTIVATION ONSET



USING THE COMPOSITE:

$$\delta = \frac{\left| \frac{d}{dt}(\text{SIG2}) \right| - \left| \frac{d}{dt}(\text{COMP}) \right|}{\ln \left[\left| \frac{d}{dt}(\text{SIG1}) - \frac{d}{dt}(\text{COMP}) \right| \right]} = \frac{|14 - 97.7|}{\ln [|111.8 - 97.7|]} = 31.63$$

FIG. 8

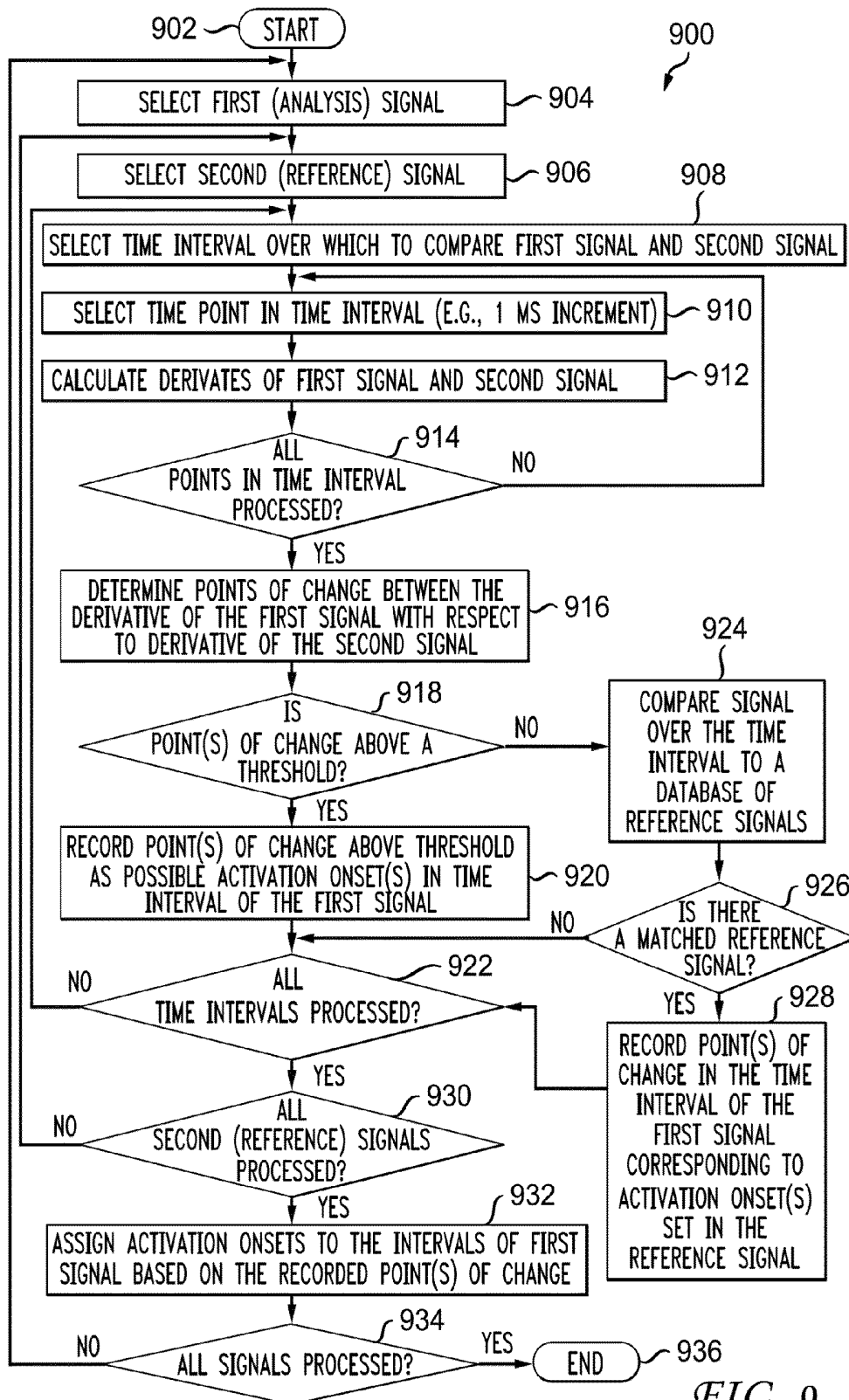
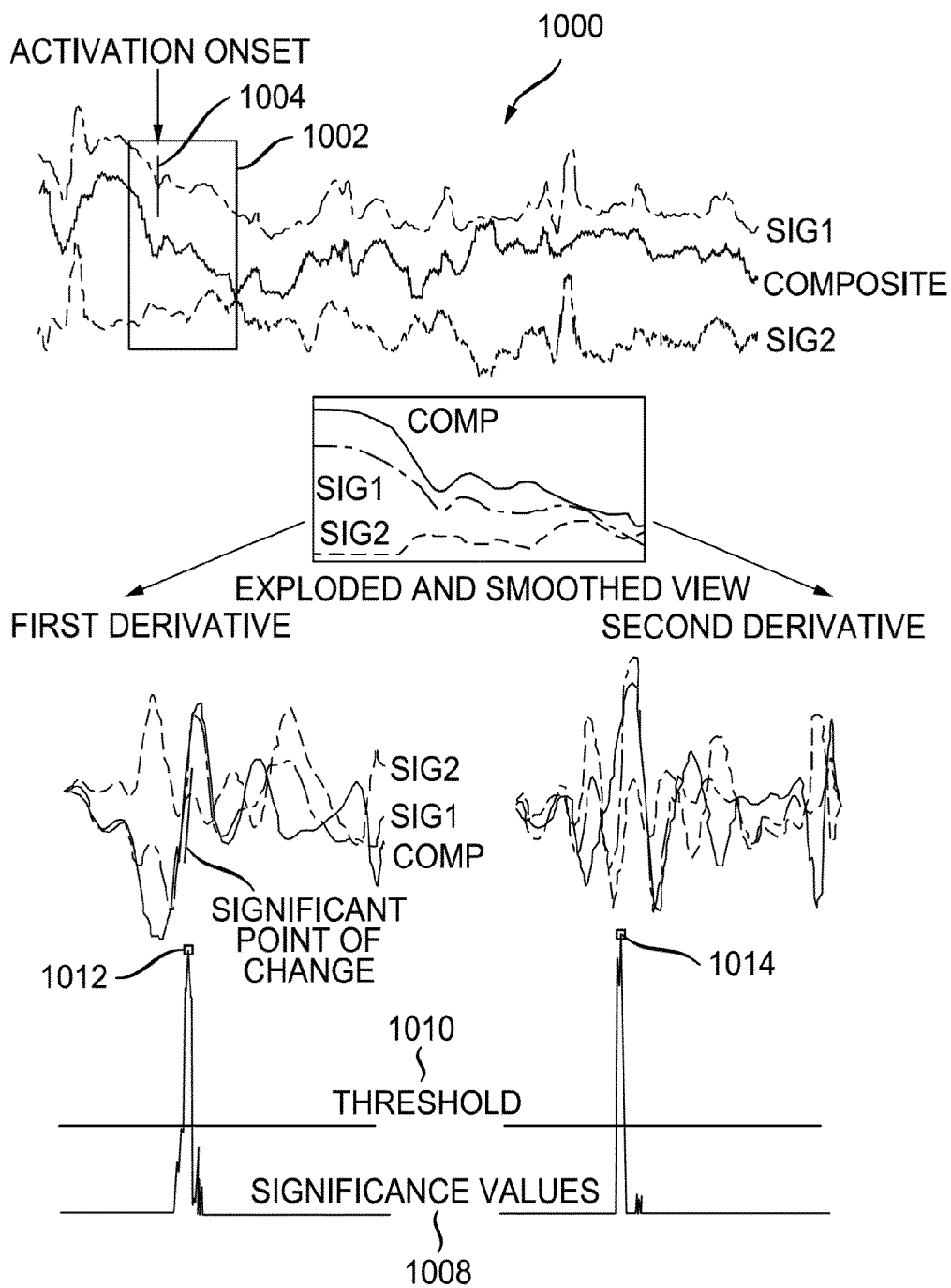


FIG. 9

*FIG. 10*

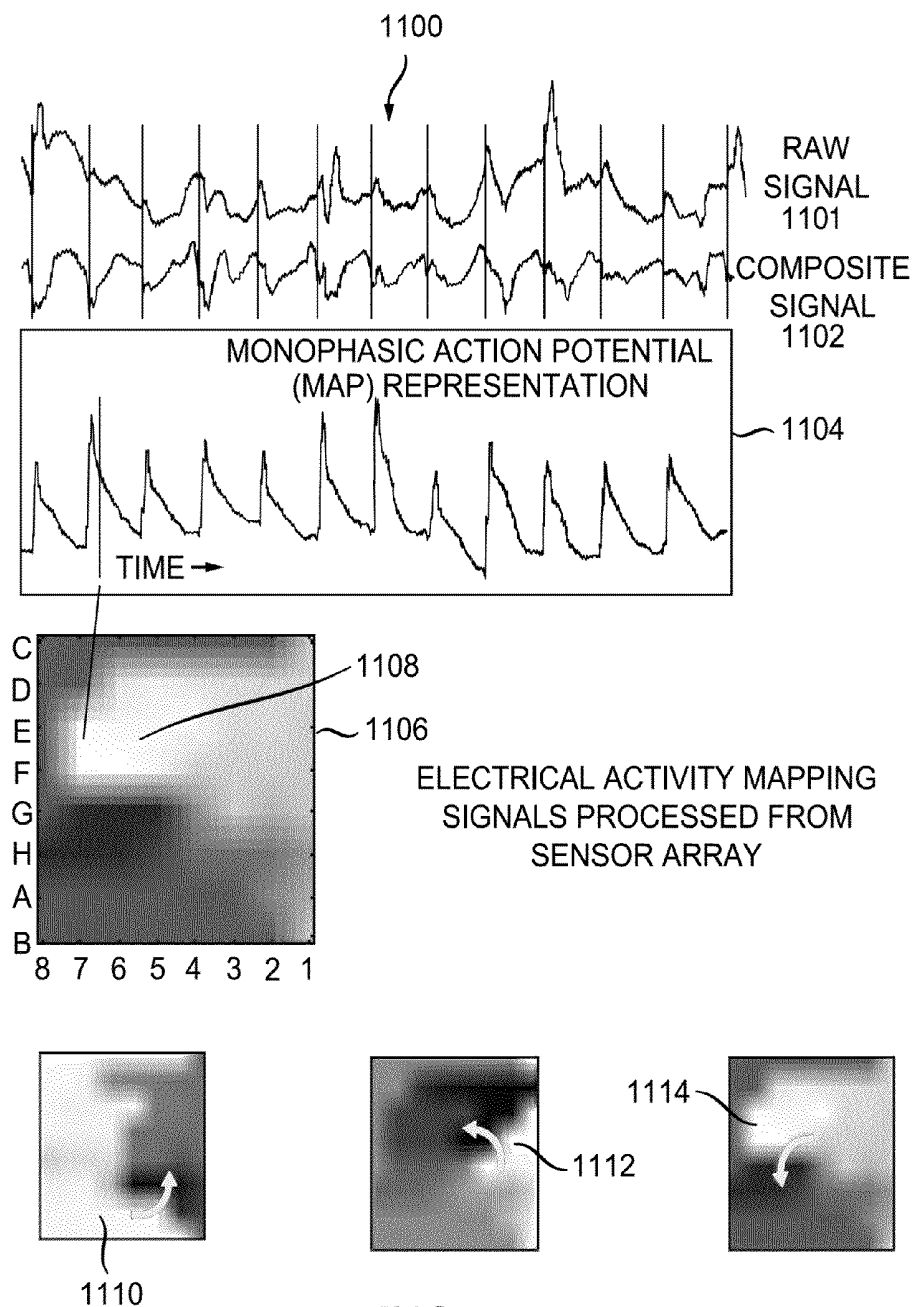
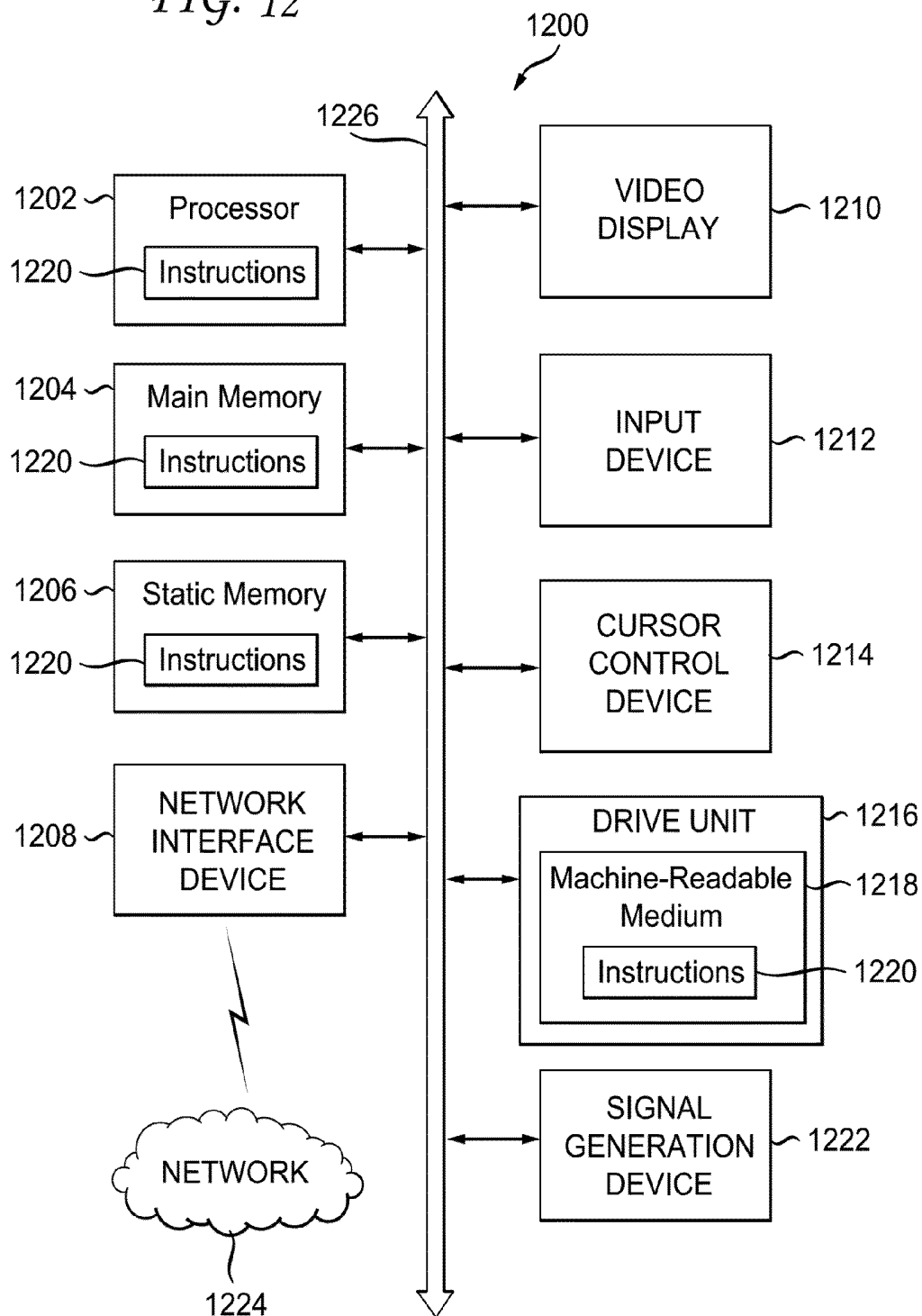


FIG. 11

FIG. 12



SYSTEM AND METHOD FOR RECONSTRUCTING CARDIAC ACTIVATION INFORMATION

RELATED APPLICATIONS

This is a continuation of U.S. application Ser. No. 13/840,334, filed Mar. 15, 2013, now issued as U.S. Pat. No. 9,050,006, which is a continuation-in-part of U.S. application Ser. No. 13/438,534, filed Apr. 3, 2012, now issued as U.S. Pat. No. 8,594,777, which is continuation of U.S. application Ser. No. 13/217,123, filed Aug. 24, 2011, now issued as U.S. Pat. No. 8,165,666, which claims the benefit of the priority of U.S. Provisional Application No. 61/481,607, filed May 2, 2011. Each of the listed applications is incorporated herein by reference in its entirety.

GOVERNMENT RIGHTS

This invention was made with government support under Grants R01 HL83359, HL83359-S1 and HL103800 awarded by the National Institutes of Health. The U.S. Government has certain rights in the invention.

BACKGROUND

1. Field

The present application relates generally to heart rhythm disorders. More specifically, the present application is directed to a system and method for reconstructing cardiac activation information (activation onset) associated with heart rhythm disorders.

2. Brief Discussion of Related Art

Heart (cardiac) rhythm disorders are common and represent significant causes of morbidity and death throughout the world. Malfunction of the electrical system in the heart represents a proximate cause of heart rhythm disorders. Heart rhythm disorders exist in many forms, of which the most complex and difficult to treat are atrial fibrillation (AF), ventricular tachycardia (VT) and ventricular fibrillation (VF). Other rhythm disorders are more simple to treat, but may also be clinically significant including atrial tachycardia (AT), supraventricular tachycardia (SVT), atrial flutter (AFL), supraventricular ectopic complexes/beats (SVE) and premature ventricular complexes/beats (PVC). While under normal conditions the sinus node keeps the heart in sinus rhythm, under certain conditions rapid activation of the normal sinus node can cause inappropriate sinus tachycardia or sinus node reentry, both of which also represent heart rhythm disorders.

Treatment of heart rhythm disorders—particularly complex rhythm disorders of AF, VF and polymorphic VT—can be very difficult. Pharmacologic therapy for complex rhythm disorder is not optimal, with poor efficacy and significant side effects. Ablation has been used increasingly in connection with heart rhythm disorders by maneuvering a sensor/probe to the heart through the blood vessels, or directly at surgery, and delivering energy to a location of the heart that harbors a cause of the heart rhythm disorder to mitigate and in some cases to eliminate the heart rhythm disorder. However, in complex rhythm disorders ablation is often difficult and ineffectual because tools that identify and locate a cause of the heart rhythm disorder are poor and hinder attempts to deliver energy to the correct region of the heart to eliminate the disorder.

Certain systems and methods are known for treating simple heart rhythm disorders. In a simple heart rhythm disorder (e.g., atrial tachycardia), consistent activation onset patterns

from beat to beat can generally be traced back to an earliest location, which can be ablated to mitigate and in some cases to eliminate the disorder. Even in simple heart rhythm disorders, such ablation of the cause of a heart rhythm disorder is challenging and experienced practitioners often require hours to ablate simple rhythm disorders with consistent beat-to-beat activation patterns, such as atrial tachycardia.

There are no known systems and methods that have been successful with respect to identifying causes for the complex rhythm disorders such as AF, VF or polymorphic VT. In a complex rhythm disorder, an earliest location of activation onsets cannot be identified because activation onset patterns change from beat to beat, and are “continuous” such that there is no identifiable earliest point (or start) or latest point (or end).

Diagnosing and treating heart rhythm disorders often involves the introduction of a catheter having a plurality of sensors/probes into the heart through the blood vessels of a patient. The sensors detect electric activity of the heart at sensor locations in the heart. The electric activity is generally processed into electrogram signals that represent the activation of the heart at the sensor locations.

In a simple heart rhythm disorder, the signal at each sensor location is generally consistent from beat to beat in timing and often in shape and number of its deflections, enabling identification of activation onsets at each sensor location. However, in a complex rhythm disorder, the signal at each sensor location from beat to beat may transition between one, several, and multiple deflections of various shapes. For instance, when a signal for a sensor location in AF includes 5, 7, 11 or more deflections, it is difficult if not impossible to identify which deflections in the signal are at or near the sensor location in the heart (i.e., local activation) versus a further removed location still sensed by the sensor in the heart (i.e., far-field activation) or simply noise from another part of the patient's heart, other anatomic structures, movement or motion of the sensor relative to the heart or external electronic systems.

There are no known systems and methods that have been able to reconstruct cardiac activation information (onsets) in variously shaped signals associated with heart rhythm disorders, especially in complex rhythm disorders, to facilitate identification of a cause of the heart rhythm disorders and their elimination.

SUMMARY

The present invention is applicable to reconstructing activation information of various rhythm disorders, including heart rhythm disorders, as well as other biological rhythm disorders, such as neurological seizures, esophageal spasms, bladder instability, irritable bowel syndrome, and other biological disorders for which biological activation information can be reconstructed to permit determination, diagnosis, and/or treatment of the cause or source of the disorders. It is particularly useful, however, in complex rhythm disorders which result in complex activation patterns, and especially useful in complex rhythm disorders of the heart, in order to find the cause(s) or source(s) of the disorders such that they can be treated with expediency.

Complex heart rhythm disorders typically result in activation patterns that are extremely difficult to decipher and the ability to determine accurate activation information of heart beats in complex disorders has previously not been possible. Among the advantages of the present invention is the ability to reconstruct cardiac activation information such that a determination of the cause and/or source of the disorder can

be determined and treated. Another advantage is that the present invention provides a system and method which can be carried out rapidly while a sensing device—such as a catheter having sensors thereon—is used in or near the patient and can be followed by treatment of cardiac tissue to ameliorate the disorder and in many cases cure the disorder. Treatment may thus occur immediately upon computing the reconstructed cardiac information, since it will provide the location(s) of the cause or source of the disorder.

Prior systems and methods suffered from the inability to determine the source of heart rhythm disorders and consequently provided no means of targeting the source for meaningful and curative treatment. Additionally, prior systems and methods required numerous and complex steps of treatment and yet still failed to provide a means of reconstructing cardiac activation information sufficient to identify the cause(s) or source(s) of the heart rhythm disorder.

In contrast to prior systems and methods, the present invention provides a relatively few number of steps to reconstruct the activation information in order to determine the activation onset times at various sensor locations for a heartbeat amidst the virtually indiscernible activation patterns.

As used herein, reconstruction is a process of identifying activation onset time in a cardiac or biological signal at a sensor location distinct from nearby or adjacent sensor locations for one or more beats of a biological or cardiac rhythm disorder.

As used herein, activation onset time is a time point at which activation commences in a cell or tissue, as opposed to other time points during activation.

As used herein, activation is a process whereby a cell commences its operation from a quiescent (diastolic) state to an active (electrical) state.

In accordance with an embodiment or aspect, a system to reconstruct biological activation information is disclosed. The system includes at least one computing device. The computing device is configured to process an analysis cardiac signal and a reference cardiac signal to determine a first point of change in a derivative of the analysis cardiac signal with respect to a derivative of the reference cardiac signal. The derivative of the analysis cardiac signal and the derivative of the reference cardiac signal are selected from zero, first and second order derivative. The computing device is also configured to process the analysis the analysis cardiac signal and the reference cardiac signal to determine a second point of change in a different derivative of the analysis cardiac signal with respect to a different derivative of the reference cardiac signal. The different derivative of the analysis cardiac signal and the different derivative of the reference cardiac signal are selected from zero, first and second order derivative. The computing device is further configured assign an activation onset time in the analysis cardiac signal at a point based on a mathematical association of the first point of change and the second point of change to define cardiac activation indicating a beat in the analysis cardiac signal.

In accordance with another embodiment or aspect, a method of reconstructing biological activation information is disclosed. The method includes processing an analysis cardiac signal and a reference cardiac signal to determine a first point of change in a derivative of the analysis cardiac signal with respect to a derivative of the reference cardiac signal. The derivative of the analysis cardiac signal and the derivative of the reference cardiac signal are selected from zero, first and second order derivative. The method also includes processing the analysis cardiac signal and the reference cardiac signal to determine a second point of change in a different derivative of the analysis cardiac signal with respect to different derivative

of the reference cardiac signal. The different derivative of the analysis cardiac signal and the different derivative of the reference cardiac signal are selected from zero, first and second order derivative. The method further includes assigning an activation onset time in the analysis cardiac signal at a point based on a mathematical association of the first point of change and the second point of change to define cardiac activation indicating a beat in the analysis cardiac signal.

In accordance with a further embodiment or aspect, a method of treating a cardiac rhythm disorder is disclosed. The method includes iteratively accessing an analysis cardiac signal and a reference cardiac signal from a plurality of cardiac signals. The analysis cardiac signal and the reference cardiac signal are processed to determine a first point of change in a derivative of the analysis cardiac signal with respect to a derivative of the reference cardiac signal. The derivative of the analysis cardiac signal and the derivative of the reference cardiac signal are selected from zero, first and second order derivative. The analysis cardiac signal and the reference cardiac signal are processed to determine a second point of change in a different derivative of the analysis cardiac signal with respect to a different derivative of the reference cardiac signal. The different derivative of the analysis cardiac signal and the different derivative of the reference cardiac signal are selected from zero, first and second order derivative. Activation onset times are assigned in the analysis cardiac signal at points based on a mathematical association of first points of change and second points of change to define cardiac activations indicating beats in the analysis cardiac signal. The method further includes reconstructing a cardiac activation pattern based on the assigned activation onset times to indicate a source of the cardiac rhythm disorder. Still further, the method includes treating cardiac tissue at the source to suppress or eliminate the cardiac rhythm disorder.

These and other purposes, goals and advantages of the present application will become apparent from the following detailed description when read in connection with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

Some embodiments or aspects are illustrated by way of example and not limitation in the figures of the accompanying drawings in which:

FIG. 1 illustrates an example cardiac activation reconstruction system;

FIG. 2 illustrates an example simple electrogram signal of a heart rhythm disorder from a sensor positioned at a sensor location in a heart illustrated in FIG. 1;

FIG. 3 illustrates an example complex electrogram signal of a heart rhythm disorder from a sensor positioned at a sensor location in a heart illustrated in FIG. 1;

FIG. 4 illustrates an example array of sensors of a catheter illustrated in FIG. 1 and an example selection of signals from the sensors to reconstruct cardiac activation information;

FIG. 5 illustrates example comparison pairs of signals from the sensors of the array illustrated in FIG. 4;

FIG. 6 is an illustration of an example signal pair comparison of analysis signal (SIG1) and reference signal (SIG2);

FIG. 7 is an illustration of another example signal pair comparison of analysis signal (SIG1) and reference signal (SIG2);

FIG. 8 is an illustration of a further example signal pair comparison of analysis signal (SIG1) and reference signal (SIG2) utilizing a composite signal;

5

FIG. 9 is a flowchart that illustrates an example method of reconstructing cardiac activation information associated with heart rhythm disorders;

FIG. 10 is an illustration of an example signal pair comparison of analysis signal (SIG1) and reference signal (SIG2) that can be processed in accordance with the method of FIG. 9 to reconstruct cardiac activation information;

FIG. 11 is an illustration of an example mapping of processed signals in accordance with FIGS. 1-10; and

FIG. 12 is a block diagram of an illustrative embodiment of a general computer system.

DETAILED DESCRIPTION

A system and method for reconstructing cardiac activation information associated with heart rhythm disorders are disclosed herein. In the following description, for the purposes of explanation, numerous specific details are set forth in order to provide a thorough understanding of example embodiments or aspects. It will be evident, however, to one skilled in the art that an example embodiment may be practiced without all of the disclosed specific details.

FIG. 1 illustrates an example cardiac activation reconstruction system 100. The example system 100 is configured to detect and reconstruct cardiac activation information collected/detected from a patient's heart in connection with a heart rhythm disorder. The heart includes a right atrium 122, left atrium 124, right ventricle 126 and left ventricle 128.

The example system 100 includes a catheter 102, signal processing device 114, computing device 116 and analysis database 118.

The catheter 102 is configured to detect cardiac activation information in the heart and to transmit the detected cardiac activation information to the signal processing device 114, either via a wireless or wired connection. The catheter includes a plurality of probes/sensors 104-112, which can be inserted into the heart through the patient's blood vessels.

In some embodiments or aspects, one or more of the sensors 104-112 are not inserted into the patient's heart. For example, some sensors may detect cardiac activation via the patient's surface (e.g., electrocardiogram) or remotely without contact with the patient (e.g., magnetocardiogram). As another example, some sensors may also derive cardiac activation information from cardiac motion of a non-electrical sensing device (e.g., echocardiogram). In various embodiments or aspects, these sensors can be used separately or in different combinations, and further these separate or different combinations can also be used in combination with sensors inserted into the patient's heart.

The sensors 104-112, which are positioned at sensor locations in the heart under consideration, can detect cardiac activation information at the sensor locations and can further deliver energy to ablate the heart at the sensor locations. It is noted that the sensors 104-112 can also detect cardiac activation information from overlapping regions of the heart (e.g., right atrium 122 and left atrium 124).

The signal processing device 114 is configured to process (e.g., clarify and amplify) the cardiac activation information detected by the sensors 104-112 at the sensor locations into electrogram signals and to provide the processed cardiac signals to the computing device 116 for analysis or processing in accordance with methods disclosed herein. In processing the cardiac activation information from the sensors 104-112, the signal processing device 114 can subtract cardiac activation information from overlapping regions of the heart 120 to provide processed cardiac signals to the computing device 116 for analysis. While in some embodiments or aspects, the

6

signal processing device 114 is configured to provide unipolar signals, in other embodiments or aspects, the signal processing device 114 can provide bipolar signals.

The computing device 116 is configured to receive (or access) cardiac signals from the signal processing device 114 and further configured to analyze or process the cardiac signals in accordance with methods, functions or logic disclosed herein to reconstruct cardiac activation information in the cardiac signals such that it is possible to locate a cause of the heart rhythm disorder and to eliminate the cause.

For example, the computing device 116 can process a first cardiac signal and a second cardiac signal from the received cardiac signals to determine whether there is a point of change in a derivative of the first cardiac signal with respect to a derivative of the second cardiac signal above a threshold. The computing device 116 can then assign an activation onset time in the first signal at the point of change to define cardiac activation indicating a beat in the first signal if it is determined that the point of change is above the threshold.

As another example, the computing device 116 can iteratively select pairs of cardiac signals from the received cardiac signals, each pair having a first cardiac signal and second cardiac signal. The computing device 116 can process and assign for each of the pairs in order to define multiple cardiac activations indicating beats for the first cardiac signal in each of the pairs. For example, the computing device 116 is configured to perform processing and assigning to define multiple cardiac activations indicating beats in the first cardiac signal. The computing device 116 can then reconstruct a cardiac activation pattern based on assigned activation onset times of cardiac activations from the received cardiac signals to indicate a source of a rhythm disorder. In some embodiments or aspects, the computing device 116 can also display the reconstructed cardiac activation pattern to facilitate treatment of cardiac tissue at the source to suppress, lessen or eliminate the cardiac rhythm disorder.

The analysis database 118 is configured to support or aid in the analysis of the signals by the computing device 116. In some embodiments or aspects, the analysis database 118 can store a catalog of reference signals and associated activations to enable the computing device 116 to determine an activation onset associated with a signal being considered (e.g., when point of change is below threshold during a time window), as will be described in greater detail herein.

FIG. 2 illustrates an example simple electrogram signal 200 of a heart rhythm disorder from a sensor positioned at a sensor location in the heart 120. For example, sensor 104 of catheter 102 can be positioned at a sensor location in the right atrium 122, as shown in FIG. 1. As an example, the heart rhythm disorder can be a complex rhythm disorder AF, VF and polymorphic VT, or another heart rhythm disorder.

The example signal 200 is for a time period between about 300 ms and about 900 ms. During this time period, the signal 200 is expected to have four (4) local activation onsets 202-208, e.g., those activation onsets that originate at or near (locally to) the sensor location in the heart 120 of sensor 104. Specifically, based on established observations in heart rhythm disorders, cycle length between activation onsets of about 100 ms to about 300 ms can be expected for AF, and cycle length between activation onsets of about 180 ms to about 240 ms can be expected for complex ventricular arrhythmias. As an example, cycle length 210 of about 100 ms to about 300 is expected between activation onset 202 and activation onset 204. In the example signal 200, the activation onsets 202-208 are generally identifiable as having a small degree of baseline wander superposed in the local signal with few far-field artifacts that could be mistaken as local activity.

Local activity in this example can be characterized by an activation onset with a sharp inflection point and high slope, followed by a period of gentle, low-deviation slope representing repolarization, typically lasting between about 100 ms and 250 ms.

In the example signal **200**, an example far-field deflection **212** is illustrated between location activation onset **206** and local activation onset **208**, e.g., an activation onset that originates at a location in the heart **120** that is different than the sensor location associated with the sensor **104**. Specifically, the heart **120** at the sensor location associated with sensor **104** cannot physiologically activate again after activation onset **206** in a shorter cycle than about 100 ms to about 300 ms because local tissue must undergo repolarization. Moreover, the deflection **212** cannot be local to the sensor location associated with the sensor **104** when the deflection **212** is also significantly present in signals collected by neighbor sensors in multiple directions to sensor **104**. For example, the far-field deflection **212** detected by sensor **104** can be associated with activation onset at a sensor location associated with sensor **106**.

FIG. 3 illustrates an example complex electrogram signal **300** of a heart rhythm disorder from a sensor positioned at a sensor location in the heart **120**. For example, sensor **106** of catheter **102** can be positioned at a sensor location in the right atrium **122**, as shown in FIG. 1. As an example, the heart rhythm disorder can be a complex rhythm disorder AF, VF and polymorphic VT, or another heart rhythm disorder.

Similarly to example signal **200**, example signal **300** is for a time period between about 300 ms and about 900 ms. During this time period, the signal **300** is expected to have four (4) local activation onsets, e.g., activation onsets that originate locally to the sensor location in the heart **120** of sensor **106**. However, in the example signal **300** there are eleven (11) possible activation onsets **302-322**. Multiple deflections of short duration (shorter than shortest cycle length of about 100 ms) caused by the heart rhythm disorder makes the discernment of local activation onsets at the sensor location of sensor **104** as opposed to far-field activations or simply noise prohibitively difficult.

FIG. 4 illustrates an example array of sensors **400** of catheter **102** and an example selection of signals from the sensors to reconstruct cardiac activation information (e.g., activation onsets). The array **400** includes fifteen (15) example sensors for simplicity and clarity of the description. It is to be understood that the array **400** can include fewer or more sensors to as may be determined to cover different portions of the heart **120**. In some embodiments or aspects, the array **400** can include 160 or more sensors.

The sensors of the array **400** are shown in example spatial arrangement with respect to the right atrium **122** of the heart **120**. Similarly, the array **400** can be spatially arranged in other chambers of the heart, e.g., left atrium, right ventricle, left ventricle, or for combinations of chambers including the endocardial or epicardial surfaces. In FIG. 4, the spatial arrangement of electrodes in the array **400** is shown to be uniform and planar for simplicity and clarity of the description. However, the heart **120** is not a uniform or planar structure. Accordingly, the spatial arrangement of electrodes in the array **400** can be varied with respect to the shape of the heart **120** to improve detection of electric activity in the heart **120**.

In one example embodiment or aspect, catheter **102** of FIG. 1 can be a basket catheter with the example sensors of the array **400** disposed in spatial arrangements along splines **406-408** of the basket catheter **102**. Different catheters with vari-

ous spatial arrangements of the sensors in the sensor array **400** can be used, such as spiral, radial spokes or other spatial arrangements.

Pairs of sensors (signals of sensors) in the array **400** are iteratively selected for processing as will be described in greater detail herein in order to reconstruct cardiac activation information (activation onsets) of the heart **120** in the right atrium **122**, or another chamber in which the array **400** may be disposed.

As illustrated at **402**, an analysis signal (1) is selected for processing. A reference signal (2)—a neighbor to the analysis signal (1)—is then selected to form a first pair that is processed to determine activation onsets in the analysis signal (1). Similarly, as illustrated at **404**, an analysis signal (1) is selected for processing. A reference signal (2)—another neighbor to the analysis signal (1)—is then selected to form a second pair that is processed to determine activation onsets in the analysis signal (1). The activation onsets from the first pair and the second pair of signals can be stored in memory of computing device **116** or database **118** of FIG. 1. The neighboring sensors (signals) can but do not have to be adjacent, as will be described in greater detail below.

The selections and processing are repeated for the sensors of the array **400** (signals) that neighbor the analysis signal (1). The activation onsets in the analysis signal (1) for all pairs of signals can also be stored in memory of computing device **116** or database **118**. Thereafter, another analysis signal is selected and the selections and processing are repeated for that analysis signal. In this fashion, each of the plurality of analysis signals in array **400** is processed against its neighboring signals. The number of neighboring signals for a given analysis signal can be fewer or greater depending on the spatial arrangement of the sensors in the array **400**, the chamber of the heart analyzed and the heart rhythm disorder treated.

FIG. 5 illustrates example comparison pairs of signals from the sensors of the array **400** illustrated in FIG. 4. Neighbor signals can include not only those signals that are immediately adjacent to the analysis signal but also those signals not adjacent to the analysis signal. Spatially separating the paired sensors can have the effect of spatially extending the area over which deflections are considered to be local activity. Local activity is therefore approximately defined by the separation of the paired sensors. As illustrated in example 1 of FIG. 5, selected analysis signal (1) is processed against adjacent signals (2)-(5) and also against a non-adjacent signal (6). As further illustrated in example 2 of FIG. 5, selected analysis signal (1) is processed against adjacent signals (2)-(5) and also against a non-adjacent signals (6) and (7). While closest neighbor signals are preferred, neighbor signals in various spatial orientations with respect to the analysis signal can be used.

For each analysis signal, there could be a plurality of reference signals (e.g., four (4) reference signals or greater). A final activation onset in the analysis signal is determined with reference to or based on the combination of the reference signals' possible activation onsets. Specifically, the activation onsets determined from each pair can be referenced against each other to check for correspondence or association of activations in the analysis signal. An activation onset for the analysis signal is finalized based on the possible activation onsets of the referenced pairs of signals.

The final activation onset for the analysis signal can be determined in various ways. In one embodiment or aspect, the final activation onset for the analysis signal can be determined based on an average of the possible activation onsets from the various pairs of referenced signals. In another embodiment or

aspect, the final activation onset for the analysis signal can be determined based on an average of the possible activation onsets from those pairs of signals in which a majority of the possible activation onsets are within a predetermined time interval of each other (e.g., ± 5 ms). Similarly, the final activation onset for the analysis signal can be determined based on an average of the possible activation onsets from those pairs of signals in which a plurality of the possible activation onsets are within a predetermined time interval of each other (e.g., ± 5 ms). The time interval used can be chosen to be lower or higher. Alternatively, the final activation can also be determined by performing a "center-of-mass" calculation weighted by the significance value of each of the possible activation onsets in the majority, or by analysis of a predominant direction of activation onsets relative to sensor locations.

With reference to example 1 in FIG. 5, if an analysis signal has been determined to have possible activation onsets of 170 ms, 190 ms, 193 ms, 165 ms and 172 ms in connection with the five (5) reference signal pairs, respectively, then the final activation onset for the analysis signal can be determined based on a majority of the possible activation onsets within the predetermined time interval (e.g., ± 5 ms) to be $(170+165+172)/3=169$ ms. The activation onsets of 190 ms and 193 ms that are outside the time interval can be discounted from the determination of the final activation onset for the analysis signal. The final activation onset determined for each signal can be saved in the database 118 of FIG. 1.

With reference to example 2 in FIG. 5, if an analysis signal has been determined to have possible activation onsets of 170 ms, 176 ms, 165 ms, 200 ms, 202 ms and 204 ms, in connection with the six (6) reference signal pairs, respectively, then the final activation onset for the analysis signal can be determined based on a plurality of the possible activation onsets within the predetermined time interval (e.g., ± 5 ms) to be $(200+202+204)/3=202$ ms. The activation onset of 176 ms is outside the time interval (e.g., ± 5 ms) and is discounted from the determination of the final activation onset for the analysis signal. Activation onsets 165 ms and 170 ms do not form a plurality of possible activation onsets within the predetermined time interval (e.g., ± 5 ms). The final activation onset determined for each signal can be saved in the database 118 of FIG. 1.

While in the forgoing examples for the sake of brevity and clarity, only one activation onset was determined for the analysis signal in connection with each reference signal, it should be understood that each signal (from a sensor of array 400) can represent multiple successive analysis intervals (e.g., activation cycles) as illustrated in FIG. 2, each of which can have an activation onset as determined based on the same time interval of multiple reference signals (neighboring sensors of array 400).

FIG. 6 is an illustration of an example signal pair comparison 600 of example analysis signal (SIG1) and example reference signal (SIG2). For example, the signals can be from comparison pair 402 (or comparison pair 404) illustrated in FIG. 4, or from any comparison pair illustrated in FIG. 5. It is noted that the signals are illustrative and occur during the same analysis interval. As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles), as illustrated in FIG. 2.

The signals are processed at one or more successive time points (e.g., every millisecond, two milliseconds, or other time points) to determine whether there is a point of change in a derivative of the analysis signal with respect to a derivative of the reference signal above a threshold. The point of change can be determined from one or more of slope, amplitude, timing and shape for the first cardiac signal and the second

cardiac signal. It is noted that in some embodiments or aspects, processing of some time points can be omitted (e.g., every other time point or two of three time points). While the slope can be determined by a first-order derivative and/or a second-order derivative for each of the time points in the signals, this example shows calculations using the first-order derivative. A root mean squared is determined for each of the signals. For example, RMS1 and RMS2 are determined by taking a root mean squared of the derivatives (first-order and/or second-order) for the entire signal of each of the signals (e.g., all activation cycles). RMS can be used to normalize the amplitude of the signals with respect to one another, such that amplitudes (e.g., voltage) of the deflections in the signals do not affect the processing of the signals as described below.

A time point (same time point or about the same time point) is successively selected from each of the signals (SIG1, SIG2) for consideration and processing. For each time point under consideration, a time increment 602, 604 in each signal starting at that time point can be considered. For example, a time increment of 10 ms can be used. Different time increments can be selected. A line which is pinned to the point under consideration in each signal and which provides the best fit to the time points in the time increment of each signal is determined. The determined lines represent the slopes (e.g., volts/per second) of the signals for the selected time point. As described, the slopes of the lines can be determined by first-order and/or second order derivatives. More specifically, the determined lines represent slopes of the signals at the selected time point for the same time increment (e.g., 10 ms). A significance value (δ) is determined with respect to the slopes.

The significance value (δ) can be determined by taking an absolute value of the first slope over its associated root mean squared value and subtracting an absolute value of the second slope over its associated root mean squared value. A determination is made as to whether the resulting (δ)= -0.461 is above a significance threshold (e.g., 0.25). The significance threshold indicates that there is a potentially significant point of change (based on slopes) for the time point in the signals under consideration, e.g., that the derivatives diverge sufficiently from each other. In the example signal pair comparison 600, the significance value (δ)= -0.461 is below the significance threshold of 0.25. The low significance value indicates that the deflection in SIG1 is far-field and not sufficiently local to a sensor location from which the signal originated, e.g., a sensor shown in FIG. 4. Accordingly, there is no potentially significant point of change in the example signal pair comparison 600.

While the foregoing significance value (δ) calculations are shown and described with reference to the slopes using first-order derivatives, it should be noted that second-order derivatives can be computed instead of, or in addition to, the first-order derivatives in a similar fashion. A second significance value (δ) calculated based on slopes using second order derivatives can then be compared to a second significance threshold defined for the second-order derivative. Significance thresholds can be different for the first and second calculated significance values (δ).

As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles), as illustrated in FIG. 2. In each analysis interval, it is possible to have zero, one or more potentially significant points of change as described above. The time point under consideration and the potentially significant point(s) of change can be recorded, such as in database 118.

FIG. 7 is an illustration of an example signal pair comparison 700 of example analysis signal (SIG1) and example ref-

erence signal (SIG2). Similarly, the signals can be from comparison pair **402** (or comparison pair **404**) illustrated in FIG. **4**, or from any comparison pair illustrated in FIG. **5**. The signals are illustrative and occur during the same analysis interval. As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles), as illustrated in FIG. **2**.

The signals are processed at one or more successive time points to determine whether there is a point of change in a derivative of the analysis signal with respect to a derivative of the reference signal above a threshold. In some embodiments or aspects, processing of some time points can be omitted (e.g., every other time point or two of three time points). A first derivative (or second derivative) is determined for each of the time points in the signals. A root mean squared is further determined for each of the signals. A time point (same time point or about the same time point) is successively selected from each of the signals (SIG1, SIG2) for consideration and processing. For each time point under consideration, a time increment **702, 704** (e.g., 10 ms) in each signal starting at that time point can be considered. A line which is pinned to the point under consideration in each signal and which provides the best fit to the time points in the time increment of each signal is determined. The determined lines represent the slopes (e.g., volts/per second) of the signals for the selected time point. More specifically, the determined lines represent the slopes at the selected time point for the same time increment. A significance value (δ) is determined with respect to the slopes.

The significance value (δ) can be determined by taking an absolute value of the first slope over its associated root mean squared value and subtracting an absolute value of the second slope over its associated root mean squared value. A determination is made as to whether the resulting (δ)= -0.063 is above a significance threshold (e.g., 0.25). In the example signal pair comparison **700**, the significance value (δ)= -0.063 is well below the significance threshold of 0.25. The low significance value indicates low amplitude noise. Accordingly, there is no potentially significant point of change in the example signal pair comparison **700**.

A noise level can be defined as fraction of the significance threshold (first, second or third) or can be defined programmatically in various ways. For example, noise level can be one-tenth (0.025) of the significance threshold (0.25). A different fraction level can be selected. As another example, the noise level can be defined as a Gaussian standard deviation of a plurality of significance values. Other ways of defining the noise level are contemplated. It is noted that the significance threshold (e.g., 0.25) is higher than the noise level that can be associated with the analysis signal and reference signal in the example signal pair comparison **700**. Accordingly, a point of change at or below noise level can be associated with one or more signals from other regions of a heart, respiratory system, gastrointestinal tract, neurological system as well as electronic interference.

While the foregoing significance value (δ) calculations are shown and described with reference to the slopes using first-order derivatives, it should be noted that second-order derivatives can be computed instead of, or in addition to, the first-order derivatives in a similar fashion. A second significance value (δ) calculated based on slopes using second order derivatives can then be compared to a second significance threshold defined for the second-order derivative. Significance thresholds can be different for the first and second calculated significance values (δ).

As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles) and in each analysis

interval, it is possible to have zero, one or more potentially significant points of change as described above. The time point under consideration and the potentially significant point(s) of change can be recorded, such as in database **118**.

FIG. **8** is an illustration of an example signal pair comparison **800** of example analysis signal (SIG1) and example reference signal (SIG2) utilizing a composite signal. As in the other examples, the signals can be from comparison pair **402** (or comparison pair **404**) illustrated in FIG. **4**, or from any comparison pair illustrated in FIG. **5**. The signals are illustrative and occur during the same analysis interval. As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles), as illustrated in FIG. **2**.

The signals are processed at one or more successive time points to determine whether there is a point of change in a derivative of the analysis signal with respect to a derivative of the reference signal above a threshold. In some embodiments or aspects, processing of some time points can be omitted (e.g., every other time point or two of three time points). A first derivative (zero order derivative or second derivative) is determined for each of the time points in the signals. A root mean squared is further determined for each of the signals. A time point (same time point or about the same time point) is successively selected from each of the signals (SIG1, SIG2) for consideration and processing. For each time point under consideration, a time increment **802, 804** (e.g., 10 ms) in each signal starting at that time point can be used. A line which is pinned to the point under consideration in each signal and which provides the best fit to the time points in the time increment of each signal is determined. The determined lines represent the slopes (e.g., volts/per second) of the signals for the selected time point. More specifically, the determined lines represent the slopes of the signals at the selected time point for same time increment. A significance value (δ) is determined with respect to the slopes.

In some embodiments or aspects, the significance value can be determined by taking an absolute value of the first slope over its associated root mean squared value and subtracting an absolute value of the second slope over its associated root mean squared value. A determination is made as to whether the resulting (δ)= 0.546 is above a significance threshold (e.g., 0.25). In the example signal pair comparison **800**, the significance value (δ)= 0.546 is determined to be above the significance threshold of 0.25.

Accordingly, there is a potentially significant point of change in the example signal pair comparison **800** at the time point under consideration. As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles) and in each analysis interval, it is possible to have zero, one or more potentially significant points of change as described above. The time point under consideration and the potentially significant point(s) of change can be recorded, such as in database **118**.

While the foregoing significance value (δ) calculations are shown and described with reference to the slopes using first-order derivatives, it should be noted that second-order derivatives can be computed instead of, or in addition to, the first-order derivatives in a similar fashion. A second significance value (δ) calculated based on slopes using second order derivatives can then be compared to a second significance threshold defined for the second-order derivative. Significance thresholds can be different for the first and second calculated significance values (δ).

In other embodiments or aspects, the significance value (δ) can be determined with respect to a composite signal. Specifically, a composite signal (COMP) is computed by subtracting SIG2 (reference signal) from SIG1 (analysis signal),

e.g., $COMP = SIG2 - SIG1$. The composite signal can represent a bipolar signal (COMP) of constituent unipolar signals (SIG1, SIG2). In alternate embodiments or aspects, the composite signal COMP can also be computed by adding signals SIG1 and SIG2. The signals in the signal pair comparison **800** are illustrative and occur during the same analysis interval. As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles), as illustrated in FIG. 2.

The signals SIG1, SIG2 are processed at one or more successive time points with respect to the composite signal COMP to determine whether there is a point of change in a derivative of the analysis signal with respect to a derivative of the reference signal above a threshold. A first derivative (or second derivative) is determined for each of the time points in the signals, SIG1, SIG2, COMP. A time point (same time point or about the same time point) is successively selected from each of the signals (SIG1, SIG2, COMP) for consideration and processing. For each time point under consideration, a time increment **802**, **804**, **806** (e.g., 10 ms) in each signal starting at that time point can be considered. A line which is pinned to the point under consideration in each signal and which provides the best fit to the time points in the time increment of each signal is determined. The determined lines represent the slopes (e.g., volts/per second) of the signals for the selected time point. More specifically, the determined lines represent the slopes of the signals at the selected time point for the same time increment. A significance value (δ) is determined with respect to the slopes.

In the embodiments or aspects employing the composite signal, the significance value (δ) can be determined by a ratio taking an absolute value of the second slope and subtracting an absolute value of the composite slope, and dividing by a logarithm of a result of an absolute value the first slope subtracting an absolute value of the composite slope. The resulting significance value for the time point under consideration is (δ)=31.63. Significance values can be computed for all points under consideration. A significance threshold can be determined to be an average of the computed significance values (δ) plus a standard deviation. Thereafter, only those significance values (δ) that are above the significance threshold can be considered to be potentially significant points of change for the comparison pair **800**. For the example signals in the signal pair comparison **800** of FIG. 8, the determined significance threshold can be 10. It is noted that the significance value(s) that is above the significance threshold generally extends substantially above the significance threshold. For example, a significance value (δ)—having the largest ratio—can therefore be selected.

While the foregoing significance value (δ) calculations using the composite signal are shown and described with reference to the slopes using first-order derivatives, it should be noted that second-order derivatives can be computed instead of, or in addition to, the first-order derivatives in a similar fashion. In embodiments using the composite signals, a third significance value (δ) calculated based on slopes using second order derivatives can then be compared to a third significance threshold defined for the second-order derivative. Significance thresholds can be different for the first, second and third calculated significance values (δ).

Accordingly, there is a potentially significant point of change in the example signal pair comparison **800** at the time point under consideration. As noted herein, the signals can have multiple successive analysis intervals (e.g., activation cycles) and in each analysis interval, it is possible to have zero, one or more potentially significant points of change as

described above. The time point under consideration and the potentially significant point(s) of change can be recorded, such as in database **118**.

FIG. 9 is a flowchart that illustrates an example method **900** of reconstructing cardiac activation information (activation onset) associated with heart rhythm disorders. The example method **900** can be performed by the computing device **116** illustrated in FIG. 1. More specifically, the example method **900** starts at operation **902** at which signals are received by the computing device **116** via signal processing device **114** from sensors disposed in the heart **120**. For example, signals can be received from sensors of the sensor array **400** disposed in the right atrium **122** of the heart **120**, as shown in FIGS. 1 and 4. In some embodiments or aspects, at least a portion of the signals from the sensors can be recorded by signal processing device **114** and then provided to computing device **116**.

At operation **904**, a first signal (analysis signal) is selected. At operation **906**, a second signal (reference signal) is selected. Selection of the analysis signal and the reference signal can be performed as described in greater detail with reference to FIGS. 4 and 5. In some embodiments or aspects, a root mean squared (RMS) can be determined for the first signal and for the second signal. At operation **908**, a time interval over which the first signal and the second signal are to be compared is selected. The time interval can be selected to be an activation cycle (e.g., 100 ms to 300 ms) as described in FIG. 2. In some embodiments or aspects, the time interval can be determined by a dominant frequency analysis or other analysis of the average cycle length of the first (analysis) signal. A default time interval of 200 ms can be used if the time interval cannot be determined computationally. In other embodiments or aspects, the time interval can be selected manually, computationally by a different analysis method, from a database that catalogs such time intervals for patients of a certain age, gender and type of heart rhythm disorder, or defaulted to a value between about 100 ms and about 300 ms.

In some embodiments or aspects, a composite signal can be determined based on the selected first signal and the second signal, such as by subtracting or adding the signals as described with reference to FIG. 8.

At operation **910**, a time point is selected for consideration in the selected time interval. The same or about the same time point is selected for consideration in each signal (e.g., first signal and second signal). At operation **912**, derivatives are calculated for a time increment (e.g., 10 ms) extending from the point of consideration in each signal. It is noted that first-order derivatives and/or second-order derivatives are calculated in each signal. In those embodiments or aspects that use a composite signal, a derivative is also calculated for a time increment (e.g., 10 ms) extending from a time point of consideration in the composite signal. Similarly, in those embodiments or aspects using the composite signal, first-order derivatives and/or second-order derivatives are calculated in each signal. The time point of consideration in the composite signal is the same or about the same as in the other signals (e.g., first signal and second signal).

At operation **914**, a determination is made as to whether all points in the selected time interval have been processed. If it is determined that all point in the selected time interval were processed, the method **900** continues at operation **916**. Alternatively, the method **900** performs operations **910**, **912** until all points in the selected time interval are determined to be processed at operation **914**.

At operation **916**, points of change between the first-order derivatives of the first signal with respect to the first-order derivatives of the second signal are determined in the time interval under consideration. Alternatively or additionally to

the points of changes between first-order derivatives, points of change between the second-order derivatives of the first signal with respect to the second-order derivatives of the second signal are determined in the time interval under consideration. For example, a first significance value (δ) can be determined at each point of change using the first-order derivatives, and/or a second significance value (δ) can be determined at each point of change using the second-order derivatives, as described with reference to FIGS. 6-8.

At operation 918, a determination is made as to whether there is a point(s) of change in the derivative of the first cardiac signal with respect to the derivative of the second cardiac signal above a threshold. As described hereinabove with reference to FIGS. 6-8, the point of change for the first-order derivatives can be determined with reference to a first threshold, while the point of change for second-order derivatives can be determined with reference to a second threshold, and the point of change for composite signals can be determined with reference to a third threshold.

For example, it can be determined whether the significance value (δ) at the point of change is above the first threshold for first-order derivatives and above the second threshold for second-order derivatives. In some embodiments or aspects that do not use a composite signal, the first threshold can be 0.25 (or another value) as described with reference to FIGS. 6-8 and the second threshold can be the same or a different value, while in those embodiments or aspects that use a composite signal, a third threshold can be computed as an average value plus a standard deviation of all points of change as described with reference to FIG. 8.

If it is determined that there is a point(s) of change above the threshold (e.g., first threshold, second threshold, third threshold), the method 900 continues at operation 920 where the significant point(s) of change is recorded (selected) as a possible activation onset(s) for the time interval under consideration in the first (analysis) signal. If however, it is determined that there is no point of change above the threshold (no significant point of change), the method 900 continues at operation 924 where the first signal is compared over the time interval to a catalog of reference signals. For example, the catalog of reference signals for heart rhythm disorders can be maintained in database 118. At operation 926, a determination is made as to whether there is a match to a reference signal in the database. The comparison can be based on at least one characteristic of the first signal to at least one characteristic of the reference signal, such as shape, slope, amplitude, frequency and/or timing. Other characteristics can be used together with or instead of the enumerated characteristics.

If there is no match to a reference signal at operation 926, the method 900 continues at operation 922. Alternatively, the method 900 continues at operation 928 where the point(s) of change in the time interval under consideration is recorded (selected), which would correspond to activation onset(s) in the reference signal that was matched.

At operation 922, a determination is made as to whether all time intervals in the signals have been processed. If it is determined that all time intervals have not been processed, the method 900 continues to perform operations 908-922 to process subsequent time intervals until it is determined that all time intervals have been processed. The subsequent time interval can be determined from the point(s) of change that represents the possible activation onset at 920. Specifically, if only one point of change (above the threshold) is recorded at 920, then the next time interval (e.g., 100 ms to 300 ms) can start at the onset time associated with the point of change plus a half of a cycle length (e.g., 50 ms to 150 ms). If there are

multiple points of change, then the onset time associated with the largest point of change (significance value) is used to determine the next time interval for operations 908-922. It is noted that the determination of the next time interval can be extended to consider significant points of change from all second (reference) signals for the same time interval under consideration. However, if it is determined that all time intervals have been processed at operation 922, the method 900 continues at operation 930.

At operation 930, a determination is made as to whether all second (reference) signals have been processed in association with the selected first (analysis signal). If it is determined that all second signals have not been processed, the method 900 continues to perform operations 906-930 until it is determined that all second (reference) signals have been processed for the first (analysis) signal. However, if it is determined that all second signals have been processed, the method 900 continues to operation 932.

At operation 932, an activation onset(s) is assigned in the first signal at the point(s) of change to define cardiac activation(s) indicating a beat(s) in the first signal if it is determined (at operation 918) that the point(s) of change is above the threshold (e.g., first threshold, second threshold, third threshold). Contention among points of change based on the different thresholds (first, second and third thresholds) can be resolved as set forth below in greater detail. Similarly, at operation 932 an activation onset(s) can be assigned in the first signal at the point(s) of change to define cardiac activation(s) indicating a beat(s) in the first signal based on a matched reference signal (at operation 928). More specifically, activation onsets are assigned to the time intervals of the first signal based on the recorded (or significant) point(s) of change of the first signal with reference to the second signal(s). That is, an activation onset is assigned to each time interval in the first (analysis) signal based possible activation onset(s) associated with the significant point(s) of change in the same time interval of the second (reference) signal(s).

As described with reference to FIG. 5, the activation onset for the time interval of the first (analysis) signal can be determined based on an average of the activation onsets with reference to the second (reference) signals. In another embodiment or aspect, the activation onset for the time interval of the first signal can be determined based on an average of activation onsets with reference to those second signals in which a majority of activation onsets are within a predetermined time interval of each other (e.g., ± 5 ms). Moreover, the activation onset for the time interval can be determined based on an average of activation onsets with reference to those second signals in which a plurality of activation onsets are within a predetermined time interval of each other (e.g., ± 5 ms). The assigned onset can be recorded for each interval in the first (analysis) signal such as in database 118.

At operation 934, a determination is made as to whether all signals have been processed or analyzed as first (analysis) signals against second (reference) signals. If it is determined that all signals have not been processed, then the method 900 continues to perform operations 904-932 until all signals have been processed. Alternatively, if it is determined that all signals have been processed, the method 900 ends at operation 936.

At the conclusion of the method 900, signals collected from the heart 120 have been reconstructed with cardiac activation information (activation onsets) such that a cause of the heart rhythm disorder can be determined. More specifically, unipolar electrograms or monophasic action potentials (MAPs) can be mapped to the reconstructed activation onsets of the signals to show unipolar or MAP sequences or repre-

sentations for the signals. An activation map or pattern can be constructed from these unipolar voltage or MAP voltage representations of the signals to locate the cause of the heart rhythm disorder. An example MAP representation and example activation map are illustrated in FIG. 11.

FIG. 10 is an illustration of an example signal pair comparison 1000 of analysis signal (SIG1) and reference signal (SIG2) that can be processed in accordance with method 900 of FIG. 9 to assign an activation onset 1004. As illustrated in comparison 1000, a time interval 1002 (e.g., 100 ms-300 ms) is selected for comparison and processing. In some example embodiments or aspects, the signals in the time interval (SIG1, SIG2, COMP) are smoothed, such as via median filter. Significance values (δ) are determined for the points of changes in the signals' first or second derivative, as described herein with reference to FIGS. 1-9. As illustrated in signal pair comparison 1000, point of change 1012 in SIG1 that is above threshold 1010 is assigned as the activation onset 1004 for the time interval 1002 in SIG1 based on the first derivative. Alternatively, point of change 1014 in SIG1 that is above threshold 1010 is assigned as the activation onset 1004 for the time interval 1002 in SIG1 based on the second derivative. Subsequent time intervals are selected and activation onsets are assigned as described herein with reference to FIGS. 1-9 until the analysis signal (SIG1) is processed.

FIG. 11 is an illustration of an example mapping 1100 of processed signals in accordance with FIGS. 1-10. Raw signal 1101 represents a signal that is processed to assign activation onsets (vertical lines) as described herein. For reference purposes, a composite signal 1102 is shown, which results from the raw (analysis) signal 1101 and another (reference) signal (not shown). A monophasic action potential (MAP) voltage representation is generated for each processed signal 1101. Multiple signals are processed as described herein and MAPs generated based on the processed signals. The electrical activity of all MAPs is mapped in a sequence of example activation mappings 1106 to show activation onsets 1108, 1110, 1112 and 1114 at each time interval, respectively. These mappings can be displayed by computing device 116. Although only four mapping are shown for illustrative purposes, there can be fewer or a greater number of mappings 1106 based on the time intervals represented in the signals.

As shown by the arrows in the example mappings 1106 (e.g., activation onsets 1108-1114), the electrical activity indicates a rotational activation pattern of activation onsets (rotor) in the heart rhythm disorder. At least a portion of the area of the heart 120 indicated by the rotational activation pattern indicated by the arrows in FIG. 11 can be treated to eliminate the cause of the heart rhythm disorder, and therefore the heart rhythm disorder itself. Such treatment may be delivered by ablation using various energy sources (including but not limited to radiofrequency, cryoenergy, microwave, and ultrasound), gene therapy, stem cell therapy, pacing stimulation, drug or other therapy. It is noted that the MAP representation and activation map are examples to illustrate a rotational activation pattern. Other activation patterns can result from different example signals collected by the sensors from the heart 120.

FIG. 12 is a block diagram of an illustrative embodiment of a general computer system 1200. The computer system 1200 can be the signal processing device 114 and the computing device 116 of FIG. 1. The computer system 1200 can include a set of instructions that can be executed to cause the computer system 1200 to perform any one or more of the methods or computer based functions disclosed herein. The computer system 1200, or any portion thereof, may operate as a standalone device or may be connected, e.g., using a network or

other connection, to other computer systems or peripheral devices. For example, the computer system 1200 may be operatively connected to signal processing device 114 and analysis database 118.

The computer system 1200 may also be implemented as or incorporated into various devices, such as a personal computer (PC), a tablet PC, a personal digital assistant (PDA), a mobile device, a palmtop computer, a laptop computer, a desktop computer, a communications device, a control system, a web appliance, or any other machine capable of executing a set of instructions (sequentially or otherwise) that specify actions to be taken by that machine. Further, while a single computer system 1200 is illustrated, the term "system" shall also be taken to include any collection of systems or sub-systems that individually or jointly execute a set, or multiple sets, of instructions to perform one or more computer functions.

As illustrated in FIG. 12, the computer system 1200 may include a processor 1202, e.g., a central processing unit (CPU), a graphics-processing unit (GPU), or both. Moreover, the computer system 1200 may include a main memory 1204 and a static memory 1206 that can communicate with each other via a bus 1226. As shown, the computer system 1200 may further include a video display unit 1210, such as a liquid crystal display (LCD), an organic light emitting diode (OLED), a flat panel display, a solid state display, or a cathode ray tube (CRT). Additionally, the computer system 1200 may include an input device 1212, such as a keyboard, and a cursor control device 1214, such as a mouse. The computer system 1200 can also include a disk drive unit 1216, a signal generation device 1222, such as a speaker or remote control, and a network interface device 1208.

In a particular embodiment or aspect, as depicted in FIG. 12, the disk drive unit 1216 may include a computer-readable medium 1218 in which one or more sets of instructions 1220, e.g., software, can be embedded. Further, the instructions 1220 may embody one or more of the methods or logic as described herein. In a particular embodiment or aspect, the instructions 1220 may reside completely, or at least partially, within the main memory 1204, the static memory 1206, and/or within the processor 1202 during execution by the computer system 1200. The main memory 1204 and the processor 1202 also may include computer-readable media.

In an alternative embodiment or aspect, dedicated hardware implementations, such as application specific integrated circuits, programmable logic arrays and other hardware devices, can be constructed to implement one or more of the methods described herein. Applications that may include the apparatus and systems of various embodiments or aspects can broadly include a variety of electronic and computer systems. One or more embodiments or aspects described herein may implement functions using two or more specific interconnected hardware modules or devices with related control and data signals that can be communicated between and through the modules, or as portions of an application-specific integrated circuit. Accordingly, the present system encompasses software, firmware, and hardware implementations.

In accordance with various embodiments or aspects, the methods described herein may be implemented by software programs tangibly embodied in a processor-readable medium and may be executed by a processor. Further, in an exemplary, non-limited embodiment or aspect, implementations can include distributed processing, component/object distributed processing, and parallel processing. Alternatively, virtual computer system processing can be constructed to implement one or more of the methods or functionality as described herein.

It is also contemplated that a computer-readable medium includes instructions 1220 or receives and executes instructions 1220 responsive to a propagated signal, so that a device connected to a network 1224 can communicate voice, video or data over the network 1224. Further, the instructions 1220 may be transmitted or received over the network 1224 via the network interface device 1208.

While the computer-readable medium is shown to be a single medium, the term "computer-readable medium" includes a single medium or multiple media, such as a centralized or distributed database, and/or associated caches and servers that store one or more sets of instructions. The term "computer-readable medium" shall also include any medium that is capable of storing, encoding or carrying a set of instructions for execution by a processor or that cause a computer system to perform any one or more of the methods or operations disclosed herein.

In a particular non-limiting, example embodiment or aspect, the computer-readable medium can include a solid-state memory, such as a memory card or other package, which houses one or more non-volatile read-only memories. Further, the computer-readable medium can be a random access memory or other volatile re-writable memory. Additionally, the computer-readable medium can include a magneto-optical or optical medium, such as a disk or tapes or other storage device to capture carrier wave signals, such as a signal communicated over a transmission medium. A digital file attachment to an e-mail or other self-contained information archive or set of archives may be considered a distribution medium that is equivalent to a tangible storage medium. Accordingly, any one or more of a computer-readable medium or a distribution medium and other equivalents and successor media, in which data or instructions may be stored, are included herein.

In accordance with various embodiments or aspects, the methods described herein may be implemented as one or more software programs running on a computer processor. Dedicated hardware implementations including, but not limited to, application specific integrated circuits, programmable logic arrays, and other hardware devices can likewise be constructed to implement the methods described herein. Furthermore, alternative software implementations including, but not limited to, distributed processing or component/object distributed processing, parallel processing, or virtual machine processing can also be constructed to implement the methods described herein.

It should also be noted that software that implements the disclosed methods may optionally be stored on a tangible storage medium, such as: a magnetic medium, such as a disk or tape; a magneto-optical or optical medium, such as a disk; or a solid state medium, such as a memory card or other package that houses one or more read-only (non-volatile) memories, random access memories, or other re-writable (volatile) memories. The software may also utilize a signal containing computer instructions. A digital file attachment to e-mail or other self-contained information archive or set of archives is considered a distribution medium equivalent to a tangible storage medium. Accordingly, a tangible storage medium or distribution medium as listed herein, and other equivalents and successor media, in which the software implementations herein may be stored, are included herein.

Thus, system and method to reconstruct cardiac activation information have been described. Although specific example embodiments or aspects have been described, it will be evident that various modifications and changes may be made to these embodiments or aspects without departing from the broader scope of the invention. Accordingly, the specification and drawings are to be regarded in an illustrative rather than

a restrictive sense. The accompanying drawings that form a part hereof, show by way of illustration, and not of limitation, specific embodiments or aspects in which the subject matter may be practiced. The embodiments or aspects illustrated are described in sufficient detail to enable those skilled in the art to practice the teachings disclosed herein. Other embodiments or aspects may be utilized and derived therefrom, such that structural and logical substitutions and changes may be made without departing from the scope of this disclosure. This Detailed Description, therefore, is not to be taken in a limiting sense, and the scope of various embodiments or aspects is defined only by the appended claims, along with the full range of equivalents to which such claims are entitled.

Such embodiments or aspects of the inventive subject matter may be referred to herein, individually and/or collectively, by the term "invention" merely for convenience and without intending to voluntarily limit the scope of this application to any single invention or inventive concept if more than one is in fact disclosed. Thus, although specific embodiments or aspects have been illustrated and described herein, it should be appreciated that any arrangement calculated to achieve the same purpose may be substituted for the specific embodiments or aspects shown. This disclosure is intended to cover any and all adaptations or variations of various embodiments or aspects. Combinations of the above embodiments or aspects, and other embodiments or aspects not specifically described herein, will be apparent to those of skill in the art upon reviewing the above description.

The Abstract is provided to comply with 37 C.F.R. §1.72(b) and will allow the reader to quickly ascertain the nature and gist of the technical disclosure. It is submitted with the understanding that it will not be used to interpret or limit the scope or meaning of the claims.

In the foregoing description of the embodiments or aspects, various features are grouped together in a single embodiment for the purpose of streamlining the disclosure. This method of disclosure is not to be interpreted as reflecting that the claimed embodiments or aspects have more features than are expressly recited in each claim. Rather, as the following claims reflect, inventive subject matter lies in less than all features of a single disclosed embodiment or aspect. Thus the following claims are hereby incorporated into the Detailed Description, with each claim standing on its own as a separate example embodiment or aspect. It is contemplated that various embodiments or aspects described herein can be combined or grouped in different combinations that are not expressly noted in the Detailed Description. Moreover, it is further contemplated that claims covering such different combinations can similarly stand on their own as separate example embodiments or aspects, which can be incorporated into the Detailed Description.

The invention claimed is:

1. A method of reconstructing cardiac activation information, the method comprising:
 - processing, by a computing device, an analysis cardiac signal and a reference cardiac signal to determine a first point of change in a derivative of the analysis cardiac signal with respect to a derivative of the reference cardiac signal, the derivative of the analysis cardiac signal and the derivative of the reference cardiac signal being selected from zero, first and second order derivative;
 - processing, by the computing device, the analysis cardiac signal and the reference cardiac signal to determine a second point of change in a different derivative of the analysis cardiac signal with respect to a different derivative of the reference cardiac signal, the different derivative of the analysis cardiac signal and the different

21

derivative of the reference cardiac signal being selected from zero, first and second order derivative; and assigning an activation onset time in the analysis cardiac signal at a point based on a mathematical association of the first point of change and the second point of change to define cardiac activation indicating a beat in the analysis cardiac signal.

2. The method of claim 1, wherein the mathematical association is an average of the first point of change and the second point of change representing a majority from reference cardiac signals processed in relation to the analysis cardiac signal.

3. The method of claim 2, wherein the first point of change and the second point of change are within a predetermined time interval of each other.

4. The method of claim 2, wherein the predetermined time interval is ± 5 ms.

5. The method of claim 1, wherein the mathematical association is an average of the first point of change and the second point of change representing a plurality from reference cardiac signals processed in relation to the analysis cardiac signal.

6. The method of claim 5, wherein the first point of change and the second point of change are within a predetermined time interval of each other.

7. The method of claim 6, wherein the predetermined time interval is ± 5 ms.

8. The method of claim 1, wherein the mathematical association is a selection of one of the first point of change and the second point of change.

9. The method of claim 8, wherein the selection comprises: computing a first difference value of the first point of change minus a first threshold that is associated with the first point of change;

computing a second difference value of the second point of change minus a second threshold that is associated with the second point of change; and

selecting one of the first point of change and the second point of change based on a highest difference value among the first difference value and the second difference value.

10. The method of claim 8, wherein the selection is based on a highest order derivative that is above an associated threshold.

11. The method of claim 1, wherein the first point of change and the second point of change are determined at about the same time point for the analysis cardiac signal and the reference cardiac signal.

12. The method of claim 1, wherein the first point of change and the second point of change are determined from one or more of slope, amplitude, timing and shape for the analysis cardiac signal and the reference cardiac signal.

13. The method of claim 1, wherein determination of the first point of change comprises:

forming a composite cardiac signal from the analysis cardiac signal and the reference cardiac signal;

determining ratio values at a plurality of points in the analysis cardiac signal, each ratio value representing a difference between the derivative of the reference cardiac signal and a derivative of the composite cardiac signal to a difference between the derivative of the analysis cardiac signal and the derivative of the composite cardiac signal, the derivative of the composite cardiac signal being one of zero, first and second order derivative; and

22

selecting as the first point of change in the analysis cardiac signal a point having a largest ratio value from the determined ratio values.

14. The method of claim 1, wherein determination of the second point of change comprises:

forming a composite cardiac signal from the analysis cardiac signal and the reference cardiac signal;

determining ratio values at a plurality of points in the analysis cardiac signal, each ratio value representing a difference between the different derivative of the reference cardiac signal and a derivative of the composite cardiac signal to a difference between the different derivative of the analysis cardiac signal and the derivative of the composite cardiac signal, the derivative of the composite cardiac signal being one of zero, first and second order derivative; and

selecting as the second point of change in the analysis cardiac signal a point having a largest ratio value from the determined ratio values.

15. The method of claim 1, wherein the first point of change and the second point of change at or below a noise level is associated with one or more signals from a heart, respiratory system, gastrointestinal tract, neurological system and electronic interference.

16. The method of claim 1, further comprising performing processing and assigning to define multiple cardiac activations indicating beats in the analysis cardiac signal.

17. The method of claim 1, further comprising iteratively selecting the analysis cardiac signal and the reference cardiac signal from a plurality of cardiac signals.

18. The method of claim 1, further comprising: iteratively selecting pairs of cardiac signals from a plurality of cardiac signals, each pair having an analysis cardiac signal and different reference cardiac signal;

performing processing and assigning for each of the pairs to define multiple cardiac activations associated indicating beats for the analysis cardiac signal in each of the pairs; and

reconstructing a cardiac activation pattern based on assigned activation onset times of cardiac activations from the plurality of cardiac signals to indicate a source of a cardiac rhythm disorder.

19. A system to reconstruct cardiac activation information comprising:

at least one computing device configured to:

process an analysis cardiac signal and a reference cardiac signal to determine a first point of change in a derivative of the analysis cardiac signal with respect to a derivative of the reference cardiac signal, the derivative of the analysis cardiac signal and the derivative of the reference cardiac signal being selected from zero, first and second order derivative; process the analysis cardiac signal and the reference cardiac signal to determine a second point of change in a different derivative of the analysis cardiac signal with respect to a different derivative of the reference cardiac signal, the different derivative of the analysis cardiac signal and the different derivative of the reference cardiac signal being selected from zero, first and second order derivative; and

assign an activation onset time in the analysis cardiac signal at a point based on a mathematical association of the first point of change and the second point of change to define cardiac activation indicating a beat in the analysis cardiac signal.

20. The system of claim 19, wherein the mathematical association is an average of the first point of change and the

23

second point of change representing a majority from reference cardiac signals processed in relation to the analysis cardiac signal.

21. The system of claim 20, wherein the first point of change and the second point of change are within a predetermined time interval of each other.

22. The system of claim 21, wherein the predetermined time interval is ± 5 ms.

23. The system of claim 19, wherein the mathematical association is an average of the first point of change and the second point of change representing a plurality from reference cardiac signals processed in relation to the analysis cardiac signal.

24. The system of claim 23, wherein the first point of change and the second point of change are within a predetermined time interval of each other.

25. The system of claim 24, wherein the predetermined time interval is ± 5 ms.

26. The system of claim 19, wherein the mathematical association is a selection of one of the first point of change and the second point of change.

27. The system of claim 26, wherein at least one computing device is further configured to:

compute a first difference value of the first point of change minus a first threshold that is associated with the first point of change;

compute a second difference value of the second point of change minus a second threshold that is associated with the second point of change; and

select one of the first point of change and the second point of change based on a highest difference value among the first difference value and the second difference value.

28. The system of claim 26, wherein the selection is based on a highest order derivative that is above an associated threshold.

29. The system of claim 19, wherein the first point of change and the second point of change are determined at about the same time point for the analysis cardiac signal and the reference cardiac signal.

30. The system of claim 19, wherein the first point of change and the second point of change are determined from one or more of slope, amplitude, timing and shape for the analysis cardiac signal and the reference cardiac signal.

31. The system of claim 19, wherein at least one computing device is further configured to:

form a composite cardiac signal from the analysis cardiac signal and the reference cardiac signal;

determine ratio values at a plurality of points in the analysis cardiac signal, each ratio value representing a difference between the derivative of the reference cardiac signal and a derivative of the composite cardiac signal to a difference between the derivative of the analysis cardiac signal and the derivative of the composite cardiac signal, the derivative of the composite cardiac signal being one of zero, first and second order derivative; and

select as the first point of change in the analysis cardiac signal a point having a largest ratio value from the determined ratio values.

32. The system of claim 19, wherein at least one computing device is further configured to:

form a composite cardiac signal from the analysis cardiac signal and the reference cardiac signal;

determine ratio values at a plurality of points in the analysis cardiac signal, each ratio value representing a difference between the different derivative of the reference cardiac signal and a derivative of the composite cardiac signal to

24

a difference between the different derivative of the analysis cardiac signal and the derivative of the composite cardiac signal, the derivative of the composite cardiac signal being one of zero, first and second order derivative; and

select as the second point of change in the analysis cardiac signal a point having a largest ratio value from the determined ratio values.

33. The system of claim 19, wherein the first point of change and the second point of change at or below a noise level is associated with one or more signals from a heart, respiratory system, gastrointestinal tract, neurological system and electronic interference.

34. The system of claim 19, wherein at least one computing device is further configured to perform to perform processing and assigning to define multiple cardiac activations indicating beats in the analysis cardiac signal.

35. The system of claim 19, wherein at least one computing device is further configured to iteratively select the analysis cardiac signal and the reference cardiac signal from a plurality of cardiac signals.

36. The system of claim 19, wherein at least one computing device is further configured to:

iteratively select pairs of cardiac signals from a plurality of cardiac signals, each pair having an analysis cardiac signal and different reference cardiac signal;

perform processing and assigning for each of the pairs to define multiple cardiac activations associated indicating beats for the analysis cardiac signal in each of the pairs; and

reconstruct a cardiac activation pattern based on assigned activation onset times of cardiac activations from the plurality of cardiac signals to indicate a source of a cardiac rhythm disorder.

37. A method of treating a cardiac rhythm disorder, the method comprising:

iteratively accessing an analysis cardiac signal and a reference cardiac signal from a plurality of cardiac signals; processing, by a computing device, the analysis cardiac signal and the reference cardiac signal to determine a first point of change in a derivative of the analysis cardiac signal with respect to a derivative of the reference cardiac signal, the derivative of the analysis cardiac signal and the derivative of the reference cardiac signal being selected from zero, first and second order derivative;

processing, by the computing device, the analysis cardiac signal and the reference cardiac signal to determine a second point of change in a different derivative of the analysis cardiac signal with respect to a different derivative of the reference cardiac signal above a second threshold, the different derivative of the analysis cardiac signal and the different derivative of the reference cardiac signal being selected from zero, first and second order derivative;

assigning activation onset times in the analysis cardiac signal at points based on a mathematical association of first points of change and second points of change to define cardiac activations indicating beats in the analysis cardiac signal;

reconstructing a cardiac activation pattern based on the assigned activation onset times to indicate a source of the cardiac rhythm disorder; and

treating cardiac tissue at the source to suppress or eliminate the cardiac rhythm disorder.

* * * * *

专利名称(译)	用于重建心脏激动信息的系统和方法		
公开(公告)号	US9220427	公开(公告)日	2015-12-29
申请号	US14/670300	申请日	2015-03-26
[标]申请(专利权)人(译)	加利福尼亚大学董事会 托佩拉公司		
申请(专利权)人(译)	加利福尼亚大学董事会 TOPERA INC.		
当前申请(专利权)人(译)	加利福尼亚大学董事会 TOPERA INC. AMERICA美国为代表由退伍军人事务部，办公室总法律顾问部		
[标]发明人	NARAYAN SANJIV BRIGGS CAREY ROBERT SEHRA RUCHIR		
发明人	NARAYAN, SANJIV BRIGGS, CAREY ROBERT SEHRA, RUCHIR		
IPC分类号	A61B5/04 A61B5/042 A61B5/0452 A61B5/00 A61B5/0205 A61B5/046 A61B5/0464		
CPC分类号	A61B5/04012 A61B5/0205 A61B5/046 A61B5/04011 A61B5/0452 A61B5/0464 A61B5/7203 A61B5/7239 A61B5/0422 A61B2562/046 A61B5/7246 A61B5/7278 A61B5/7282 A61B5/0036 A61B5/024 A61B5/02405 A61B5/0245 A61B5/0255 A61B5/0402 A61B5/0468 A61B5/0472 A61B5/4836 A61B5/6852 A61B5/7207 A61B5/7217 A61B5/742		
优先权	13/438534 2013-11-26 US 13/217123 2012-04-24 US 61/481607 2011-05-02 US		
其他公开文献	US20150196212A1		
外部链接	Espacenet USPTO		

摘要(译)

在用于重建心脏激动信息的系统和方法中，处理分析心脏信号和参考心脏信号以确定分析心脏信号的导数相对于参考心脏信号的导数的第一变化点。处理分析心脏信号和参考心脏信号以确定分析心脏信号的不同导数相对于参考心脏信号的不同导数的第二变化点。基于第一变化点和第二变化点的数学关联在分析心脏信号中分配激活开始时间，以定义指示分析心脏信号中的搏动的心脏激动。

information, an analysis cardiac signal and a reference cardiac signal are processed to determine a first point of change in a derivative of the analysis cardiac signal with respect to a