



US00993172B2

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

(10) **Patent No.:** **US 9,993,172 B2**
(45) **Date of Patent:** **Jun. 12, 2018**

(54) **NONINVASIVE CARDIAC THERAPY
EVALUATION**

50/30 (2018.01); *G16H 50/70* (2018.01); *A61B 5/0472* (2013.01); *A61B 5/1107* (2013.01); *A61B 5/6823* (2013.01); *A61N 1/3627* (2013.01); *A61N 1/36507* (2013.01)

(71) Applicant: **Medtronic, Inc.**, Minneapolis, MN (US)

(58) **Field of Classification Search**
CPC *A61N 1/3627*; *A61N 1/3684*
See application file for complete search history.

(72) Inventors: **Jeffrey Gillberg**, Coon Rapids, MN (US); **Subham Ghosh**, Blaine, MN (US)

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(73) Assignee: **Medtronic, Inc.**, Minneapolis, MN (US)

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(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 186 days.

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(21) Appl. No.: **14/228,024**

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(22) Filed: **Mar. 27, 2014**

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(65) **Prior Publication Data**

US 2015/0157232 A1 Jun. 11, 2015

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Related U.S. Application Data

(60) Provisional application No. 61/913,743, filed on Dec. 9, 2013.

(Continued)

(51) **Int. Cl.**
A61B 5/044 (2006.01)
A61B 5/0408 (2006.01)

Primary Examiner — Michael D Abreu

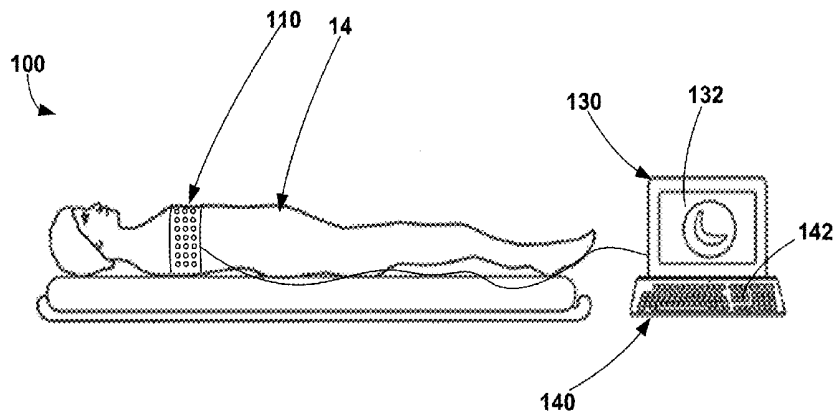
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(57) **ABSTRACT**

(52) **U.S. Cl.**
CPC *A61B 5/0408* (2013.01); *A61B 5/0402* (2013.01); *A61B 5/044* (2013.01); *A61B 5/0452* (2013.01); *A61B 5/04085* (2013.01); *A61B 5/743* (2013.01); *A61N 1/368* (2013.01); *A61N 1/3702* (2013.01); *G06F 19/3481* (2013.01); *G16H 40/63* (2018.01); *G16H*

Systems, methods, and interfaces are described herein for assisting a user in noninvasive evaluation of patients for cardiac therapy and noninvasive evaluation of cardiac therapy being delivered. The systems, methods, and interfaces may provide graphical representations of cardiac electrical activation times about one or more portions of human anatomy and one or more cardiac health metrics.

21 Claims, 14 Drawing Sheets



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FIG. 1

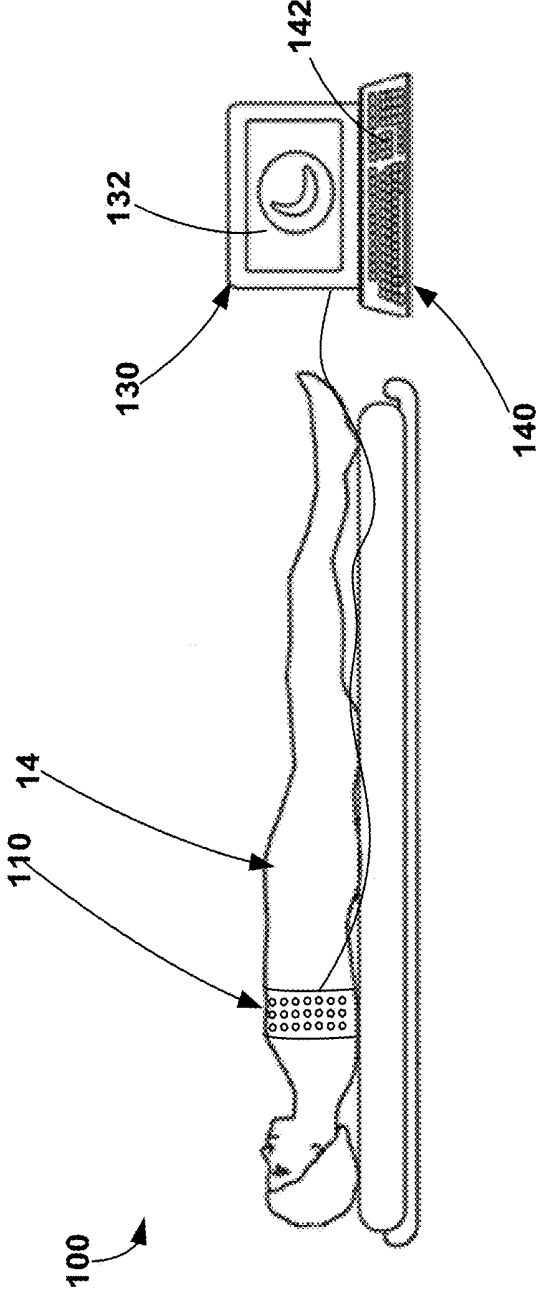


FIG. 2A

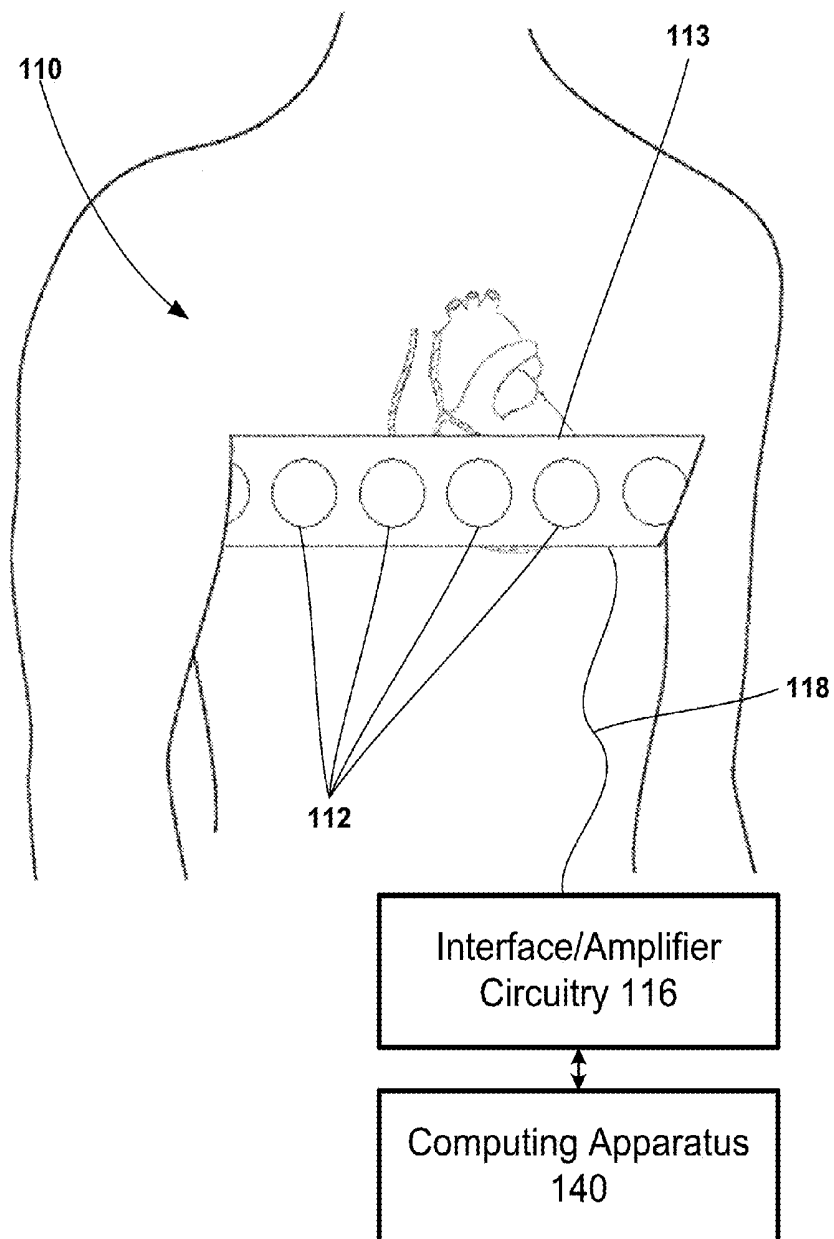


FIG. 2B

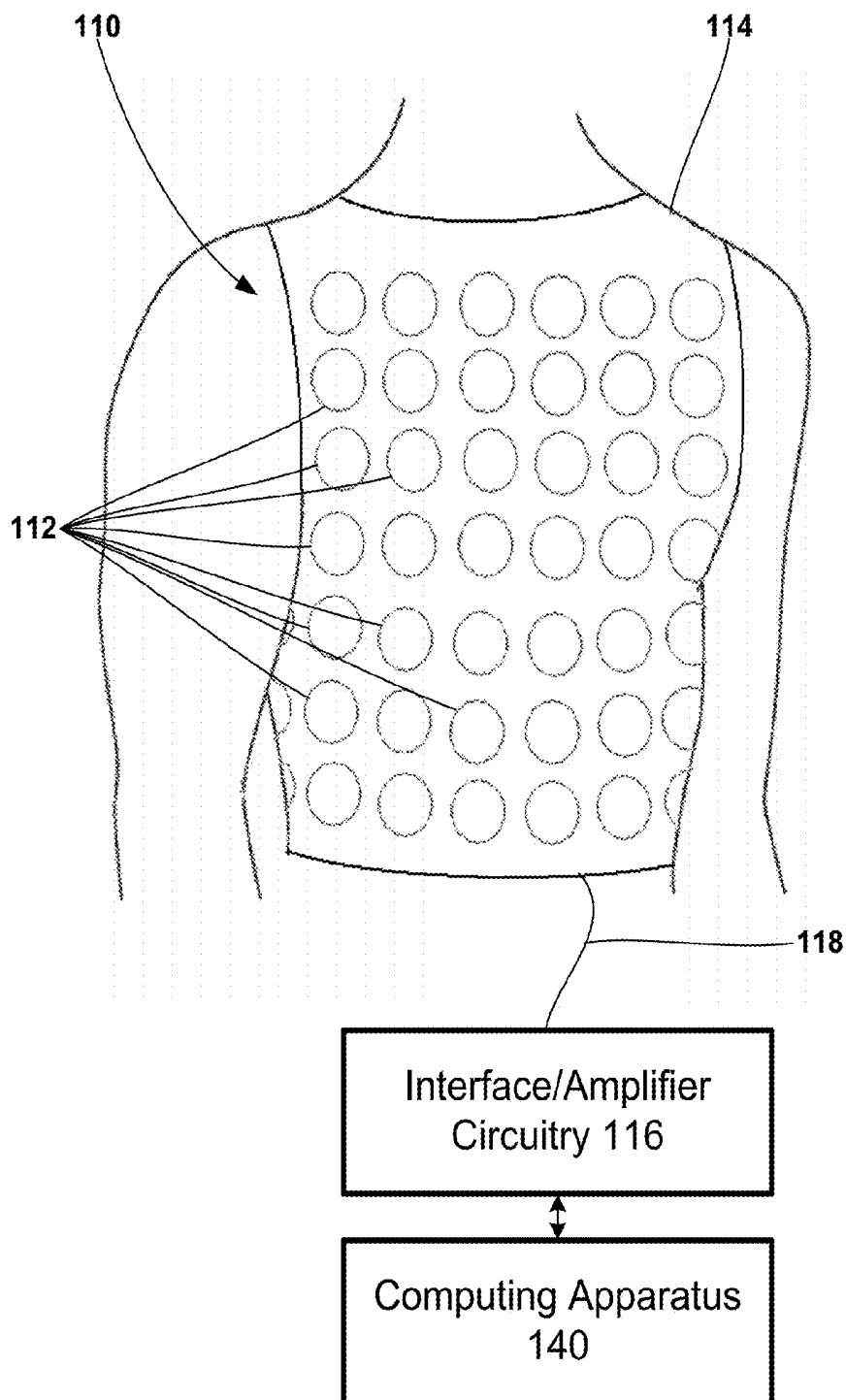


FIG. 3

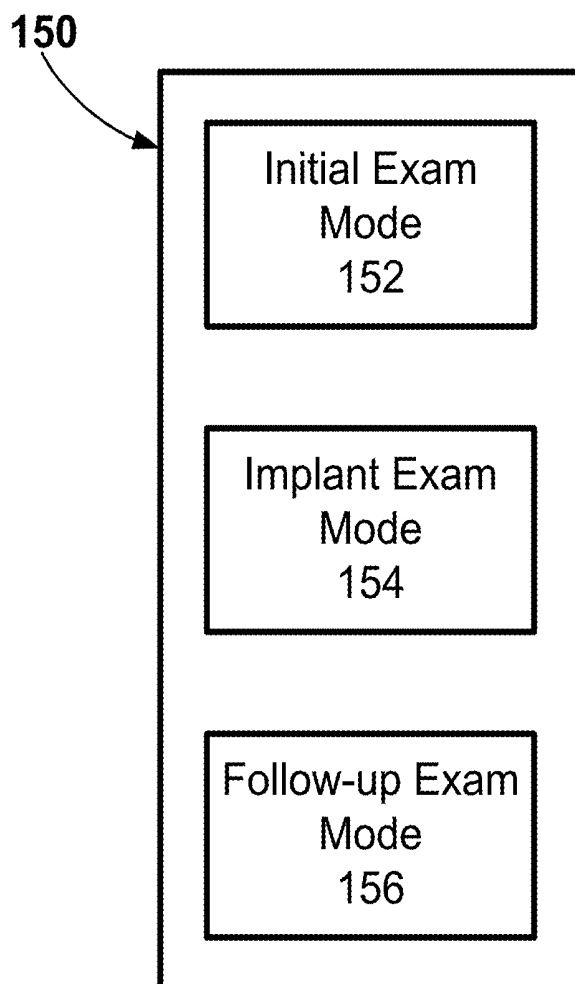


FIG. 4

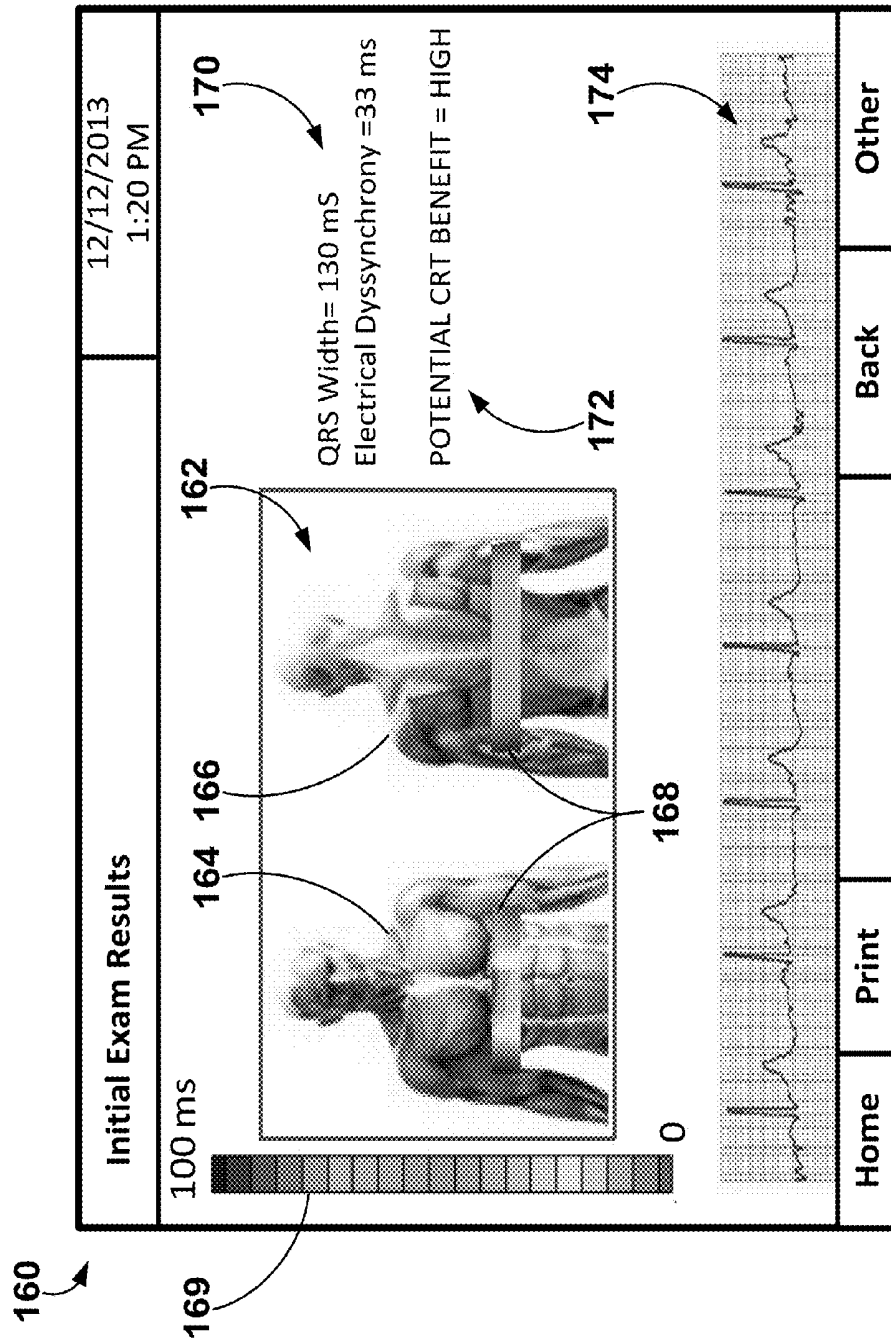


FIG. 5

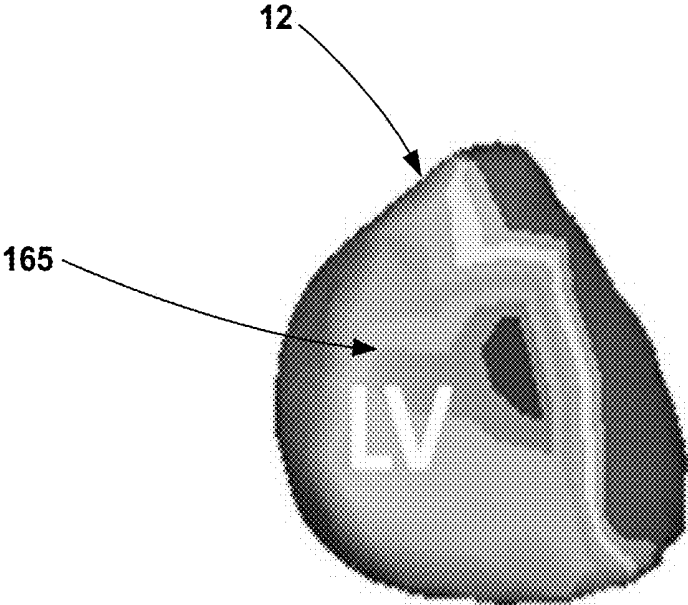


FIG. 6

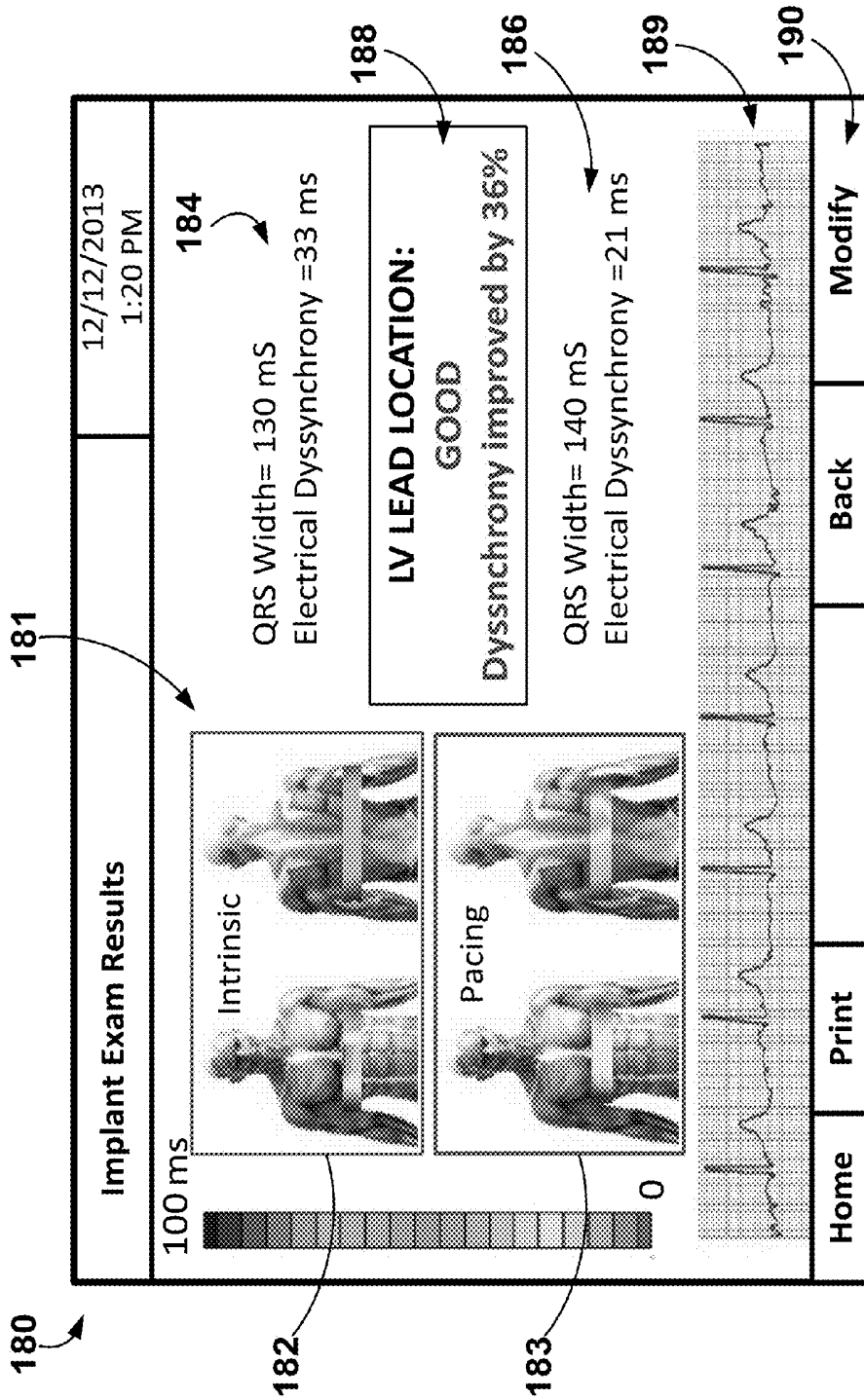


FIG. 7

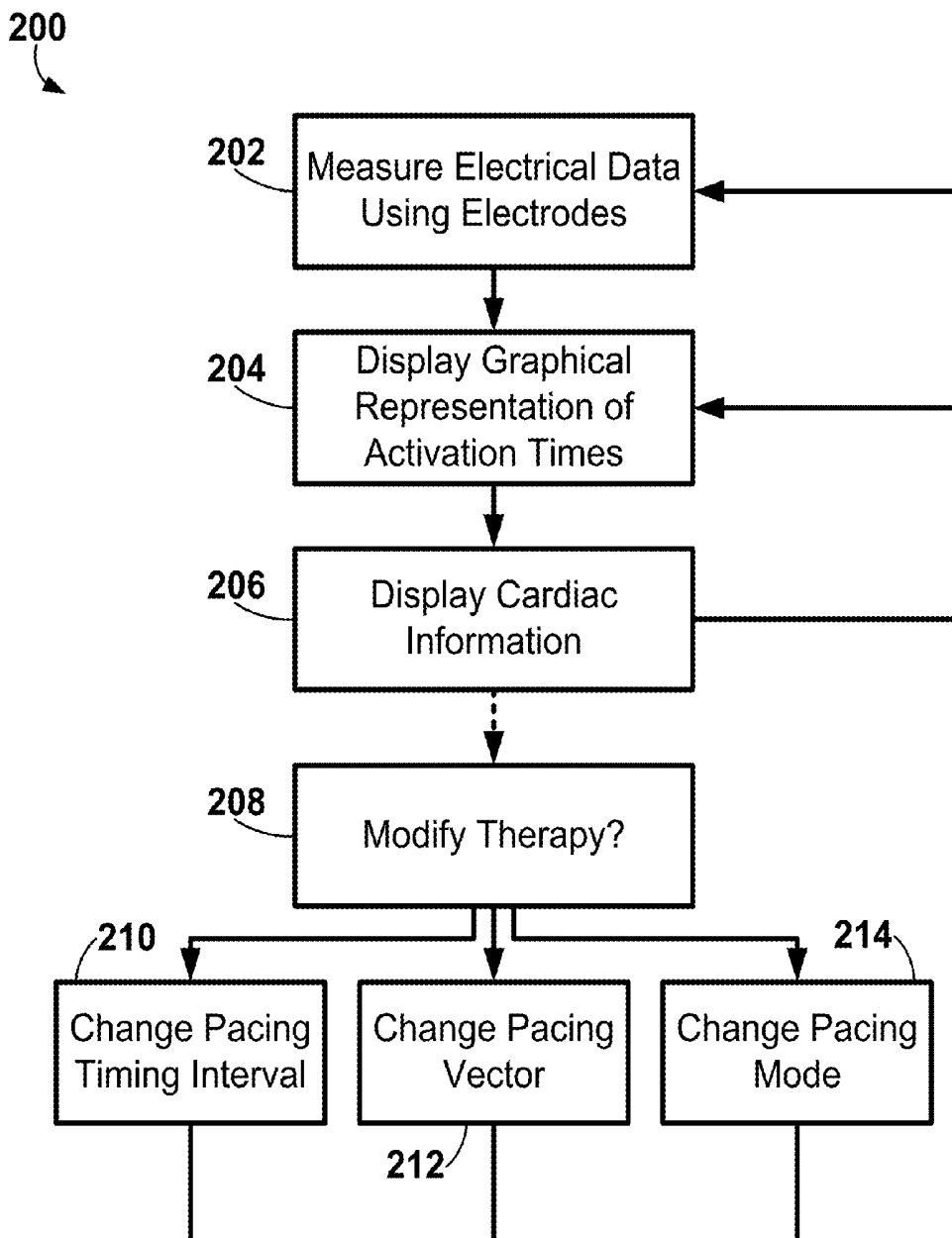


FIG. 8

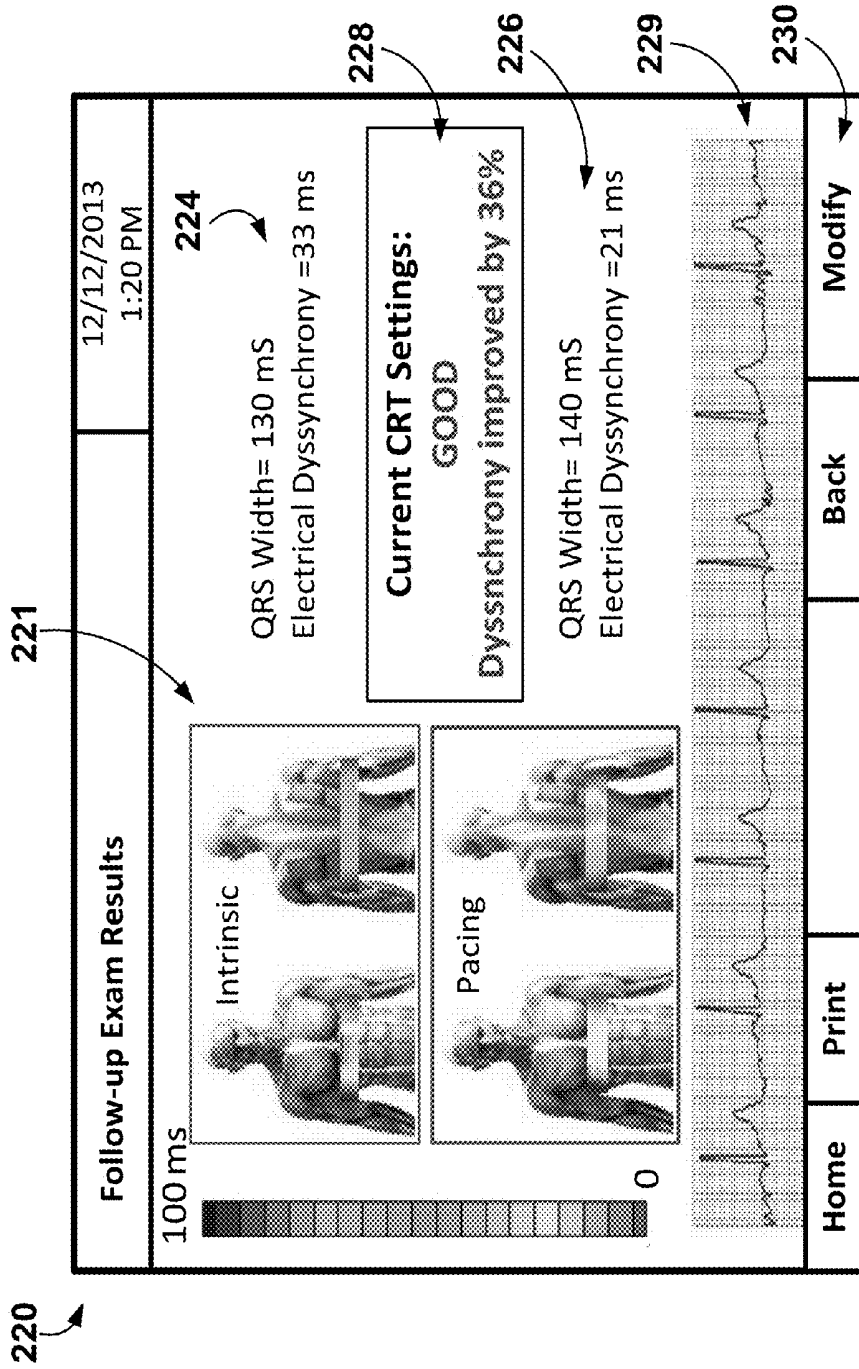


FIG. 9

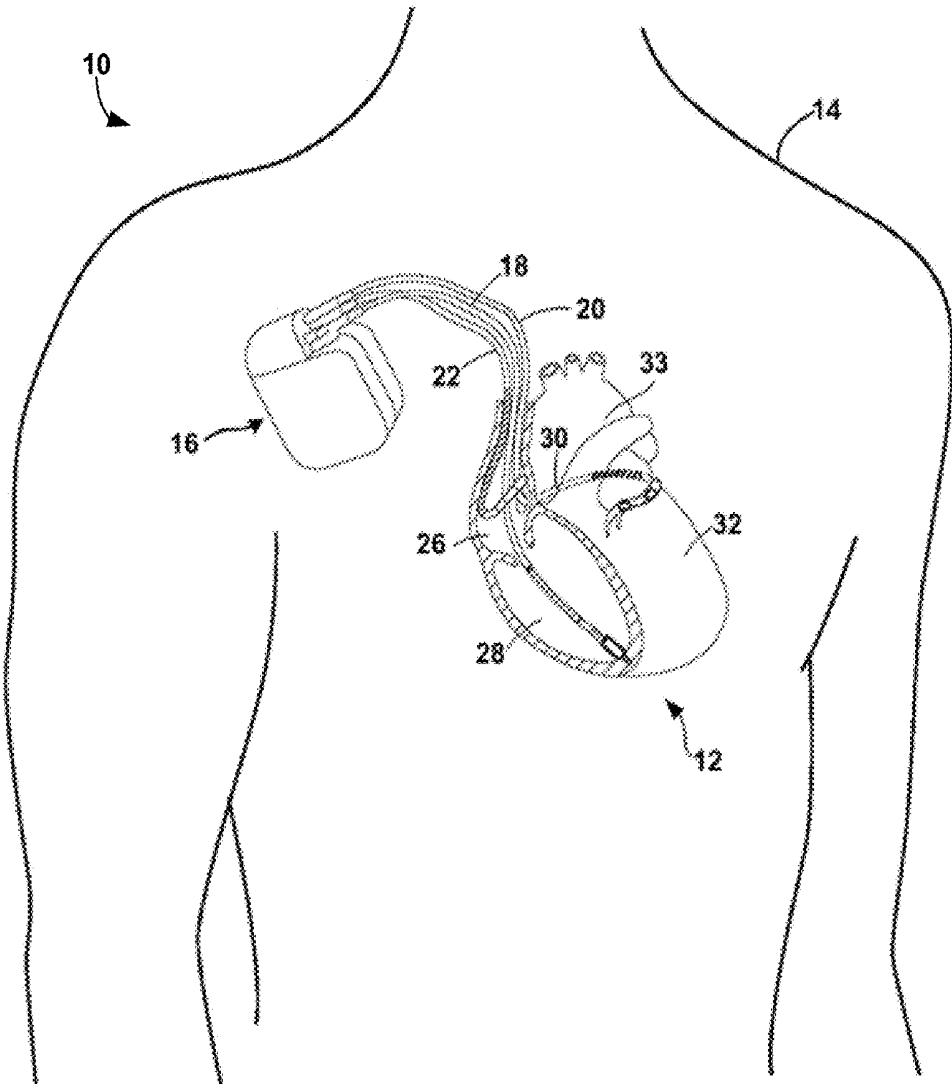


FIG. 10A

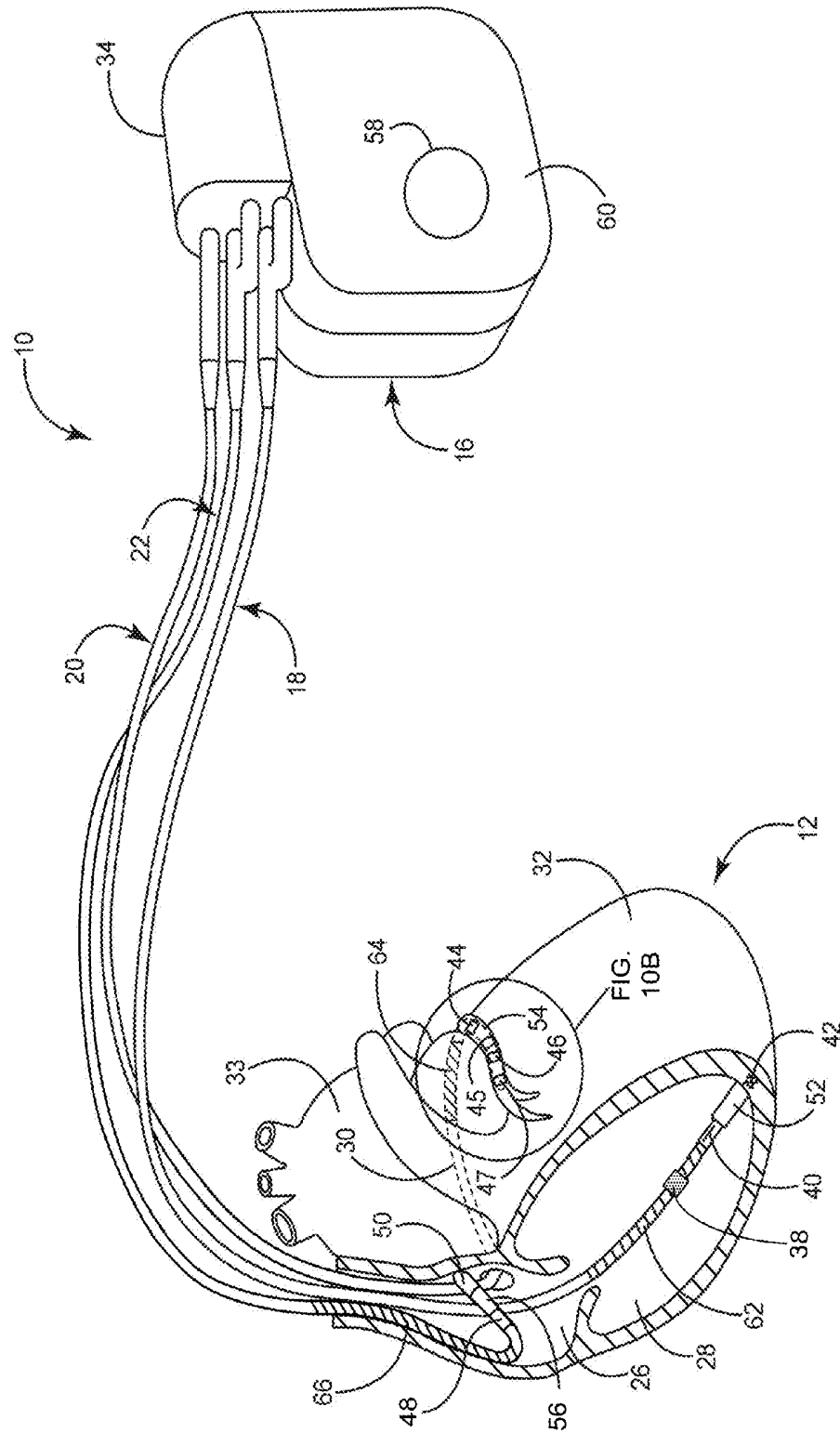


FIG. 10B

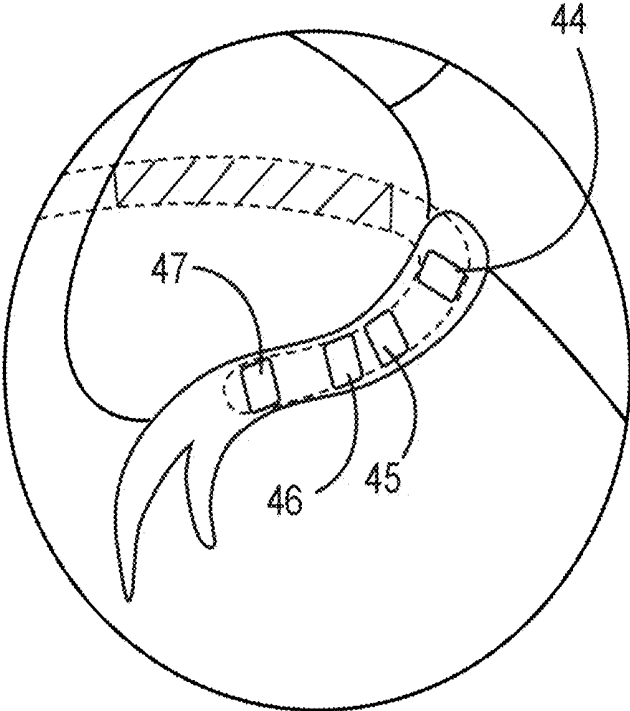
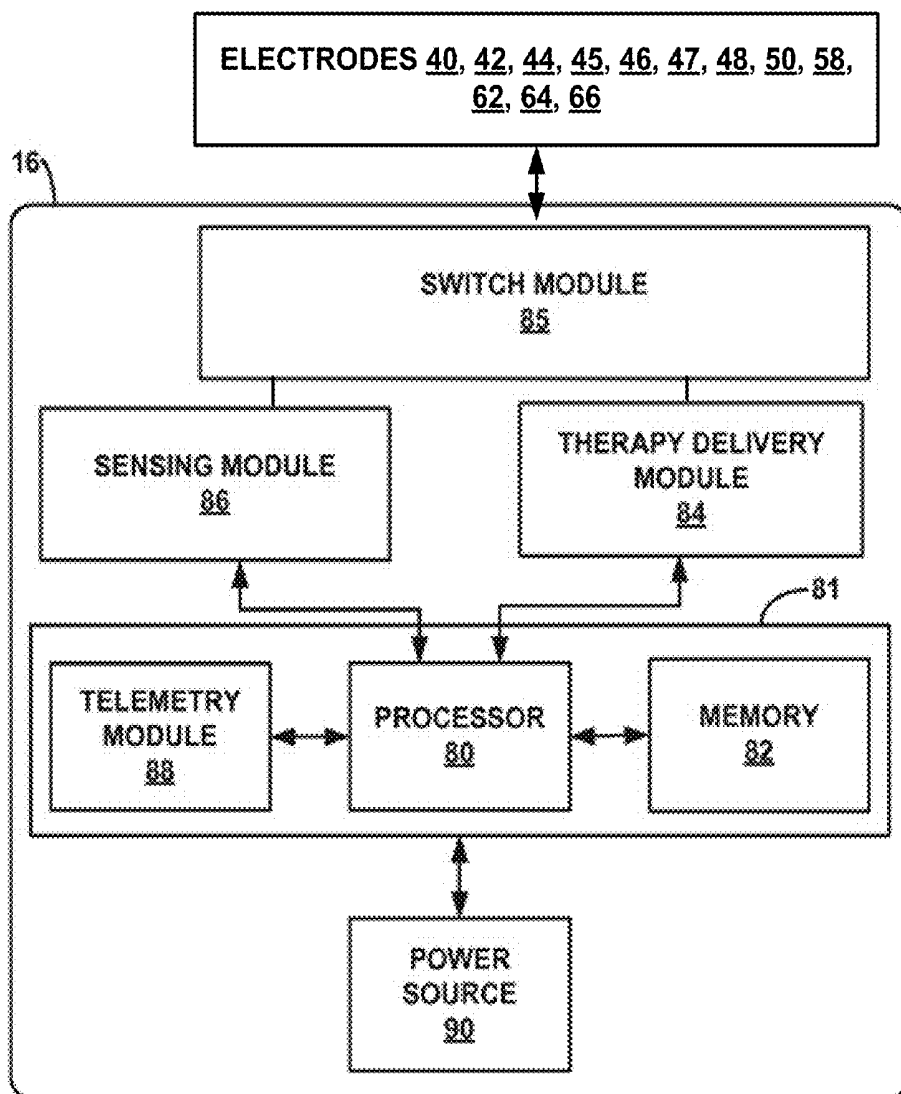


FIG. 11A



NONINVASIVE CARDIAC THERAPY EVALUATION

CROSS-REFERENCE TO RELATED APPLICATION

This application claims the benefit of U.S. Provisional Patent Application 61/913,743 entitled "Noninvasive Cardiac Therapy Evaluation" and filed on Dec. 9, 2013, which is incorporated herein by reference in its entirety.

BACKGROUND

The disclosure herein relates to systems, methods, and interfaces for use in the noninvasive evaluation of patients for cardiac therapy and noninvasive evaluation of cardiac therapy being performed on patients.

Cardiac therapy, such as cardiac resynchronization therapy (CRT), may correct symptoms of electrical dyssynchrony of a patient's heart by providing pacing therapy to one or both ventricles or atria, e.g., by providing pacing to encourage earlier activation of the left or right ventricles. By pacing the contraction of the ventricles, the ventricles may be controlled so that the ventricles contract in synchrony. Some patients undergoing cardiac therapy have experienced improved ejection fraction, increased exercise capacity, and an improved feeling of well-being.

Providing cardiac therapy to a patient may involve determining whether the patient will derive benefit from the cardiac therapy prior to implantation of a cardiac rhythm device, determining optimal site for placement of one or more ventricular pacing leads, and programming of device parameters, such as selection of electrodes on multi polar right or left ventricular leads, as well as selection of the timing of the pacing pulses delivered to the electrodes, such as atrioventricular (A-V) and intra-ventricular (V-V) delays.

SUMMARY

The exemplary systems, methods, and interfaces described herein may be configured to assist a user (e.g., a physician) in evaluating a patient for cardiac therapy and/or evaluating on-going cardiac therapy (e.g., cardiac therapy being performed on a patient). The systems, methods, and interfaces may be described as being noninvasive. For example, the systems, methods, and interfaces may not use implantable devices such as leads, probes, catheters, etc. to evaluate a patient for cardiac therapy and/or to evaluate on-going cardiac therapy. Instead, the systems, methods, and interfaces may use electrical measurements taken noninvasively using, e.g., a plurality of external electrodes attached to the skin of a patient about the patient's torso.

More specifically, the exemplary systems and methods may provide graphical user interfaces configured to assist a user in ascertaining, or assessing, whether a patient may benefit from cardiac therapy and/or whether cardiac therapy being delivered to the patient is beneficial. The exemplary graphical user interfaces may be configured to display electrical activation times about one or more portions of human anatomy, one or more metrics of the patient's cardiac health, and an indication of whether cardiac therapy may be beneficial for the patient and/or whether the on-going cardiac therapy appears to be beneficial (e.g., when compared to the patient's cardiac health prior to cardiac therapy, etc.).

An exemplary system for assisting in noninvasive evaluation of a patient for cardiac therapy may include electrode apparatus, a display apparatus, and computing apparatus

coupled to the electrode apparatus and the display apparatus. The electrode apparatus may include a plurality of external electrodes configured to be located proximate tissue of a patient (e.g., surface electrodes positioned in an array configured to be located proximate the skin of the torso of the patient). The display apparatus may include a graphical user interface configured to present cardiac electrical activation time information and cardiac health information. The computing apparatus may be configured to provide the graphical user interface displayed on the display apparatus to assist a user in noninvasively evaluating the patient for cardiac therapy. The computing apparatus may be further configured to: measure surrogate cardiac electrical activation times using one or more external electrodes of the plurality of external electrodes of the electrode apparatus proximate the patient's heart, display, on the graphical user interface, a graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy (e.g., color scaling the portion of human anatomy on the graphical user interface according to the measured surrogate cardiac electrical activation times), measure at least one metric of the patient's cardiac health (e.g., QRS width, electrical dyssynchrony, etc.) using one or more external electrodes of the plurality of external electrodes of the electrode apparatus proximate the patient's heart, display, on the graphical user interface, the at least one metric of the patient's cardiac health, and display, on the graphical user interface, an indication of whether cardiac therapy for the patient may be beneficial.

An exemplary computer-implemented method for assisting in noninvasive evaluation of a patient for cardiac therapy may include measuring surrogate cardiac electrical activation times using one or more external electrodes proximate the patient's heart (e.g., surface electrodes positioned in an array configured to be located proximate the skin of the torso of the patient), displaying a graphical user interface a graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy (e.g., color scaling the portion of human anatomy on the graphical user interface according to the measured surrogate cardiac electrical activation times), measuring at least one metric of the patient's cardiac health (e.g., QRS width, electrical dyssynchrony, etc.) using one or more external electrodes proximate the patient's heart, displaying, on the graphical user interface, the at least one metric of the patient's cardiac health, and displaying, on the graphical user interface, an indication of whether cardiac therapy for the patient may be beneficial.

Another exemplary system for assisting in noninvasive evaluation of a patient for cardiac therapy may include means for measuring surrogate cardiac electrical activation times representative of the electrical activation times of a patient's heart, means for measuring at least one metric of the patient's cardiac health (e.g., QRS width, electrical dyssynchrony, etc.), display means for displaying, on a graphical user interface, a graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy (e.g., color scaling the portion of human anatomy on the graphical user interface according to the measured surrogate cardiac electrical activation times), the at least one metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient may be beneficial.

In one or more exemplary embodiments, displaying the graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy may include displaying a graphical representation of the

measured surrogate cardiac electrical activation times about a graphical representation of at least one of an anterior side of a human torso and a posterior side of a human torso.

In one or more exemplary embodiments, displaying the graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy may include displaying a graphical representation of the measured surrogate cardiac electrical activation times about a graphical representation of at least one of an anterior side of a heart and a posterior side of a heart.

One or more exemplary embodiments may further include displaying, on the graphical user interface, at least one electrocardiogram of the patient, wherein each of the at least one electrocardiogram are captured using at least one different electrode, and wherein the at least one electrocardiogram is time-aligned on the graphical user interface and/or storing the surrogate cardiac electrical activation times, the at least one metric of the patient's cardiac health, and the at least one electrocardiogram for use in comparisons at a later time.

An exemplary system for assisting in noninvasive evaluation of cardiac therapy may include electrode apparatus, a display apparatus, and computing apparatus coupled to the electrode apparatus and the display apparatus. The electrode apparatus may include a plurality of external electrodes configured to be located proximate tissue of a patient (e.g., surface electrodes positioned in an array configured to be located proximate the skin of the torso of the patient). The display apparatus may include a graphical user interface configured to present cardiac electrical activation time information and cardiac health information. The computing apparatus may be configured to provide the graphical user interface displayed on the display apparatus to assist a user in noninvasively evaluating cardiac therapy. The computing apparatus may be further configured to measure surrogate cardiac electrical activation times using one or more external electrodes of the plurality of external electrodes of the electrode apparatus proximate the patient's heart, display, on the graphical user interface, a graphical representation of presently-measured surrogate cardiac electrical activation times about a first human anatomy portion and a graphical representation of previously-measured surrogate cardiac electrical activation times about a second human anatomy portion (e.g., the first human anatomy portion and the second human anatomy portion may depict the same human anatomy), measure at least one metric of the patient's cardiac health (e.g., QRS width, electrical dyssynchrony, etc.) using one or more external electrodes of the plurality of external electrodes of the electrode apparatus proximate the patient's heart, display, on the graphical user interface, at least one presently-measured metric of the patient's cardiac health and at least one previously-measured metric of the patient's cardiac health, and display, on the graphical user interface, an indication of whether cardiac therapy for the patient appears to be beneficial (e.g., a percentage improvement between the at least one previously-measured metric of the patient's cardiac health and the at least one presently-measured metric of the patient's cardiac health).

One exemplary method for assisting in noninvasive evaluation of cardiac therapy may include measuring surrogate cardiac electrical activation times using one or more external electrodes proximate the patient's heart and displaying a graphical user interface. The graphical user interface may include a graphical representation of presently-measured surrogate cardiac electrical activation times about a first human anatomy portion, and a graphical representation of previously-measured surrogate cardiac electrical activation

times about a second human anatomy portion (e.g., the first human anatomy portion and the second human anatomy portion may depict the same human anatomy). The exemplary method may further include measuring at least one metric of the patient's cardiac health (e.g., QRS width, electrical dyssynchrony, etc.) using one or more external electrodes proximate the patient's heart and displaying, on the graphical user interface, at least one presently-measured metric of the patient's cardiac health and at least one previously-measured metric of the patient's cardiac health, and displaying, on the graphical user interface, an indication of whether cardiac therapy for the patient appears to be beneficial (e.g., a percentage improvement between the at least one previously-measured metric of the patient's cardiac health and the at least one presently-measured metric of the patient's cardiac health).

One exemplary system for assisting in noninvasive evaluation of a patient for cardiac therapy may include means for measuring surrogate cardiac electrical activation times representative of the electrical activation times of a patient's heart, means for measuring at least one metric of the patient's cardiac health (e.g., QRS width, electrical dyssynchrony, etc.), and display means for displaying, on a graphical user interface, a graphical representation of presently-measured surrogate cardiac electrical activation times about a first human anatomy portion and a graphical representation of previously-measured surrogate cardiac electrical activation times about a second human anatomy portion (e.g., the first human anatomy portion and the second human anatomy portion may depict the same human anatomy), at least one presently-measured metric of the patient's cardiac health and at least one previously-measured metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient appears to be beneficial (e.g., a percentage improvement between the at least one previously-measured metric of the patient's cardiac health and the at least one presently-measured metric of the patient's cardiac health).

One or more exemplary embodiments may further include allowing the user to change at least one pacing parameter of an implantable medical device providing cardiac therapy to the patient (e.g., at least one of a pacing timing interval, a pacing vector, and a pacing mode) and/or displaying, on the graphical user interface, mechanical motion information of one or more regions of at least a portion of blood vessel anatomy of the patient's heart.

In one or more exemplary embodiments, displaying graphical representations of the measured surrogate cardiac electrical activation times about the first and second human anatomy portions may include displaying graphical representations of the measured surrogate cardiac electrical activation times about graphical representations of at least one of an anterior side of a human torso and a posterior side of a human torso.

In one or more exemplary embodiments, displaying graphical representations of the measured surrogate cardiac electrical activation times about the first and second human anatomy portions may include displaying graphical representations of the measured surrogate cardiac electrical activation times about graphical representations of at least one of an anterior side of a heart and a posterior side of a heart.

In one or more exemplary embodiments, displaying the graphical representations of the measured surrogate cardiac electrical activation times about a portion of human anatomy may include color scaling the portions of human anatomy on the graphical user interface according to the measured surrogate cardiac electrical activation times.

One or more exemplary embodiments may further include displaying, on a graphical user interface, at least one electrocardiogram of the patient. Each of the at least one electrocardiogram may be captured using at least one different electrode and may be time-aligned on the graphical user interface.

One exemplary system for assisting in noninvasive evaluation of cardiac therapy may include electrode apparatus (e.g., including a plurality of external electrodes configured to be located proximate tissue of a patient), a display apparatus (e.g., including a graphical user interface configured to present cardiac electrical activation time information and cardiac health information), and computing apparatus coupled to the electrode apparatus and the display apparatus. The computing apparatus may be configured to provide the graphical user interface displayed on the display apparatus. The computing apparatus may be further configured to measure surrogate cardiac electrical activation times using one or more external electrodes of the plurality of external electrodes of the electrode apparatus proximate the patient's heart, measure at least one metric of the patient's cardiac health using one or more external electrodes of the plurality of external electrodes of the electrode apparatus proximate the patient's heart, and allow a user to select one of initial examination mode for assisting a user in noninvasively evaluating the patient for cardiac therapy and follow-up examination mode for assisting a user in noninvasively evaluating cardiac therapy after implantation cardiac therapy. When in initial examination mode, the computing apparatus may be configured to display, on the graphical user interface, a graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy, at least one metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient may be beneficial. When in follow-up examination mode, the computing apparatus may be configured to display, on the graphical user interface, a graphical representation of presently-measured surrogate cardiac electrical activation times about a first human anatomy portion and a graphical representation of previously-measured surrogate cardiac electrical activation times about a second human anatomy portion and at least one presently-measured metric of the patient's cardiac health, at least one previously-measured metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient appears to be beneficial.

One exemplary a computer-implemented method for assisting in noninvasive evaluation of a patient for cardiac therapy may include measuring surrogate cardiac electrical activation times using one or more external electrodes proximate the patient's heart, measuring at least one metric of the patient's cardiac health using one or more external electrodes proximate the patient's heart, and allowing a user to select one of initial examination mode and follow-up examination mode. The initial examination mode may be configured for assisting a user in noninvasively evaluating the patient for cardiac therapy and the follow-up examination mode may be configured for assisting a user in noninvasively evaluating cardiac therapy after implantation cardiac therapy. When in initial examination mode, the exemplary method display, on a graphical user interface, a graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy, at least one metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient may be beneficial. When in follow-up examination mode, the exemplary method display, on a graphical user interface,

a graphical representation of presently-measured surrogate cardiac electrical activation times about a first human anatomy portion and a graphical representation of previously-measured surrogate cardiac electrical activation times about a second human anatomy portion and at least one presently-measured metric of the patient's cardiac health, at least one previously-measured metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient appears to be beneficial.

Another exemplary system for assisting in noninvasive evaluation of a patient for cardiac therapy may include means for measuring surrogate cardiac electrical activation times representative of the electrical activation times of a patient's heart, means for measuring at least one metric of the patient's cardiac health, and computing means for allowing a user to select one of initial examination mode and follow-up examination mode. The initial examination mode may be configured for assisting a user in noninvasively evaluating the patient for cardiac therapy and the follow-up examination mode may be configured for assisting a user in noninvasively evaluating cardiac therapy after implantation cardiac therapy. The exemplary system may further include display means for, when in initial examination mode, displaying, on a graphical user interface, a graphical representation of the measured surrogate cardiac electrical activation times about a portion of human anatomy, at least one metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient may be beneficial, and when in follow-up examination mode, displaying, on a graphical user interface, a graphical representation of presently-measured surrogate cardiac electrical activation times about a first human anatomy portion and a graphical representation of previously-measured surrogate cardiac electrical activation times about a second human anatomy portion and at least one presently-measured metric of the patient's cardiac health, at least one previously-measured metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient appears to be beneficial.

In one or more exemplary embodiments, a user may be allowed to select an implantation examination mode for assisting a user in noninvasively evaluating cardiac therapy during implantation and configuration of a cardiac therapy. When in implantation examination mode, a graphical representation of presently-measured surrogate cardiac electrical activation times about a first human anatomy portion and a graphical representation of previously-measured surrogate cardiac electrical activation times about a second human anatomy portion and at least one presently-measured metric of the patient's cardiac health, at least one previously-measured metric of the patient's cardiac health, and an indication of whether cardiac therapy for the patient appears to be beneficial may be displayed on a graphical user interface. In at least one embodiment, a user may be allowed to change at least one pacing parameter of an implantable medical device providing cardiac therapy to the patient.

The above summary is not intended to describe each embodiment or every implementation of the present disclosure. A more complete understanding will become apparent and appreciated by referring to the following detailed description and claims taken in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a diagram of an exemplary system including electrode apparatus, display apparatus, and computing apparatus.

FIGS. 2A-2B are diagrams of exemplary external electrode apparatus for measuring torso-surface potentials.

FIG. 3 is a block diagram of exemplary modes for noninvasive cardiac evaluation.

FIG. 4 is an exemplary initial examination graphical user interface depicting electrical activation information, cardiac health metrics, and an indication of the benefit of cardiac therapy.

FIG. 5 is a graphical representation of a human heart including activation times mapped thereon.

FIG. 6 is an exemplary implant examination graphical user interface depicting previously- and presently-measured electrical activation information and cardiac health metrics.

FIG. 7 is a block diagram of an exemplary method of noninvasive evaluation of cardiac therapy for a patient and configuration cardiac therapy.

FIG. 8 is an exemplary follow-up examination graphical user interface depicting previously- and presently-measured electrical activation information and cardiac health metrics.

FIG. 9 is a diagram of an exemplary system including an exemplary implantable medical device (IMD).

FIG. 10A is a diagram of the exemplary IMD of FIG. 9.

FIG. 10B is a diagram of an enlarged view of a distal end of the electrical lead disposed in the left ventricle of FIG. 10A.

FIG. 11A is a block diagram of an exemplary IMD, e.g., the IMD of FIGS. 9-10.

FIG. 11B is another block diagram of an exemplary IMD (e.g., an implantable pulse generator) circuitry and associated leads employed in the system of FIGS. 9-10 for providing three sensing channels and corresponding pacing channels.

DETAILED DESCRIPTION OF EXEMPLARY EMBODIMENTS

In the following detailed description of illustrative embodiments, reference is made to the accompanying figures of the drawing which form a part hereof, and in which are shown, by way of illustration, specific embodiments which may be practiced. It is to be understood that other embodiments may be utilized and structural changes may be made without departing from (e.g., still falling within) the scope of the disclosure presented hereby.

Exemplary systems, apparatus, and methods shall be described with reference to FIGS. 1-11. It will be apparent to one skilled in the art that elements or processes from one embodiment may be used in combination with elements or processes of the other embodiments, and that the possible embodiments of such methods, apparatus, and systems using combinations of features set forth herein is not limited to the specific embodiments shown in the Figures and/or described herein. Further, it will be recognized that the embodiments described herein may include many elements that are not necessarily shown to scale. Still further, it will be recognized that timing of the processes and the size and shape of various elements herein may be modified but still fall within the scope of the present disclosure, although certain timings, one or more shapes and/or sizes, or types of elements, may be advantageous over others.

From bipolar electrocardiogram (ECG) recordings, cardiac electrical activation times can be detected or estimated in proximity of a reference location (e.g., which can be a chosen location for the left ventricle lead during implant). Such electrical activation times may be measured and displayed, or conveyed, to an implanter by a system which acquires the ECG signals and generates the metric of elec-

trical activation times (e.g., depolarization) measured from various ECG locations. As described herein, at least in one or more embodiments, electrical activation times displayed on a graphical user interface may be used in noninvasive evaluation of a patient for cardiac therapy and/or evaluation of cardiac therapy.

Various exemplary systems, methods, and interfaces may be configured to use electrode apparatus including external electrodes, display apparatus, and computing apparatus to noninvasively assist a user (e.g., a physician) in the evaluation of a patient for cardiac therapy and/or evaluation of cardiac therapy. An exemplary system 100 including electrode apparatus 110, display apparatus 130, and computing apparatus 140 is depicted in FIG. 1.

The electrode apparatus 110 as shown includes a plurality of electrodes incorporated, or included within a band wrapped around the chest, or torso, of a patient 14. The electrode apparatus 110 is operatively coupled to the computing apparatus 140 (e.g., through one or wired electrical connections, wirelessly, etc.) to provide electrical signals from each of the electrodes to the computing apparatus 140 for analysis. Exemplary electrode apparatus may be described in U.S. Provisional Patent Application 61/913,759 entitled "Bioelectric Sensor Device and Methods" and filed on Dec. 9, 2013 and U.S. patent application Ser. No. entitled "Bioelectric Sensor Device and Methods" and filed on even date herewith, each of which is incorporated herein by reference in its entirety. Further, exemplary electrode apparatus 110 will be described in more detail in reference to FIGS. 2A-2B.

Although not described herein, the exemplary system 100 may further include imaging apparatus. The imaging apparatus may be any type of imaging apparatus configured to image, or provide images of, at least a portion of the patient in a noninvasive manner. For example, the imaging apparatus may not use any components or parts that may be located within the patient to provide images of at least a portion of the patient except noninvasive tools such as contrast solution. It is to be understood that the exemplary systems, methods, and interfaces described herein may noninvasively assist a user (e.g., a physician) in evaluation of a patient for cardiac therapy and/or on-going cardiac therapy delivered to a patient, and after the exemplary systems, methods, and interfaces have provided the noninvasive assistance, the exemplary systems, methods, and interfaces may then provide assistance to implant, or navigate, an implantable electrode into the patient, e.g., proximate the patient's heart, using imaging apparatus.

For example, after the exemplary systems, methods, and interfaces have provided noninvasive assistance, the exemplary systems, methods, and interfaces may then provide image guided navigation that may be used to navigate leads including electrodes, leadless electrodes, wireless electrodes, catheters, etc., within the patient's body. Further, although the exemplary systems, methods, and interfaces are described herein with reference to a patient's heart, it is to be understood that the exemplary systems, methods, and interfaces may be applicable to any other portion of the patient's body. Exemplary systems and methods that use imaging apparatus and/or electrode apparatus may be described in U.S. patent application Ser. No. 13/916,353 filed on Jun. 12, 2013 and entitled "Implantable Electrode Location Selection," U.S. patent application Ser. No. 13/916,377 filed on Jun. 12, 2013 and entitled "Implantable Electrode Location Selection," U.S. Provisional Patent Application No. 61/817,483 filed on Apr. 30, 2013 and entitled "Identifying Effective Electrodes," U.S. Provisional

Patent Application 61/817,480 filed on Apr. 30, 2013 and entitled "Identifying Optical Electrical Vectors," U.S. Provisional Patent Application 61/913,795 entitled "Systems, Methods, and Interfaces for Identifying Effective Electrodes" and filed on Dec. 9, 2013, U.S. patent application Ser. No. 14/227,955 entitled "Systems, Methods, and Interfaces for Identifying Effective Electrodes" and filed on Mar. 27, 2014, U.S. Provisional Patent Application 61/913,784 entitled "Systems, Methods, and Interfaces for Identifying Optimal Electrical Vectors" and filed on Dec. 9, 2013, and U.S. patent application Ser. No. 14/227,919 entitled "Systems, Methods, and Interfaces for Identifying Optimal Electrical Vectors" and filed on Mar. 27, 2014, each of which is incorporated herein by reference in its entirety.

Exemplary imaging apparatus may be configured to capture x-ray images and/or any other alternative imaging modality. For example, the imaging apparatus may be configured to capture images, or image data, using isocentric fluoroscopy, bi-plane fluoroscopy, ultrasound, computed tomography (CT), multi-slice computed tomography (MSCT), magnetic resonance imaging (MRI), high frequency ultrasound (HIFU), optical coherence tomography (OCT), intra-vascular ultrasound (IVUS), two dimensional (2D) ultrasound, three dimensional (3D) ultrasound, four dimensional (4D) ultrasound, intraoperative CT, intraoperative MRI, etc. Further, it is to be understood that the imaging apparatus may be configured to capture a plurality of consecutive images (e.g., continuously) to provide video frame data. In other words, a plurality of images taken over time using the imaging apparatus may provide motion picture data. Additionally, the images may also be obtained and displayed in two, three, or four dimensions. In more advanced forms, four-dimensional surface rendering of the heart or other regions of the body may also be achieved by incorporating heart data or other soft tissue data from an atlas map or from pre-operative image data captured by MRI, CT, or echocardiography modalities. Image datasets from hybrid modalities, such as positron emission tomography (PET) combined with CT, or single photon emission computer tomography (SPECT) combined with CT, could also provide functional image data superimposed onto anatomical data to be used to confidently reach target locations within the heart or other areas of interest.

Examples of systems and/or imaging apparatus may be described in U.S. Pat. App. Pub. No. 2005/0008210 to Evron et al. published on Jan. 13, 2005, U.S. Pat. App. Pub. No. 2006/0074285 to Zarkh et al. published on Apr. 6, 2006, U.S. Pat. App. Pub. No. 2011/0112398 to Zarkh et al. published on May 12, 2011, U.S. Pat. App. Pub. No. 2013/0116739 to Brada et al. published on May 9, 2013, U.S. Pat. No. 6,980,675 to Evron et al. issued on Dec. 27, 2005, U.S. Pat. No. 7,286,866 to Okerlund et al. issued on Oct. 23, 2007, U.S. Pat. No. 7,308,297 to Reddy et al. issued on Dec. 11, 2011, U.S. Pat. No. 7,308,299 to Burrell et al. issued on Dec. 11, 2011, U.S. Pat. No. 7,321,677 to Evron et al. issued on Jan. 22, 2008, U.S. Pat. No. 7,346,381 to Okerlund et al. issued on Mar. 18, 2008, U.S. Pat. No. 7,454,248 to Burrell et al. issued on Nov. 18, 2008, U.S. Pat. No. 7,499,743 to Vass et al. issued on Mar. 3, 2009, U.S. Pat. No. 7,565,190 to Okerlund et al. issued on Jul. 21, 2009, U.S. Pat. No. 7,587,074 to Zarkh et al. issued on Sep. 8, 2009, U.S. Pat. No. 7,599,730 to Hunter et al. issued on Oct. 6, 2009, U.S. Pat. No. 7,613,500 to Vass et al. issued on Nov. 3, 2009, U.S. Pat. No. 7,742,629 to Zarkh et al. issued on Jun. 22, 2010, U.S. Pat. No. 7,747,047 to Okerlund et al. issued on Jun. 29, 2010, U.S. Pat. No. 7,778,685 to Evron et al. issued on Aug. 17, 2010, U.S. Pat. No. 7,778,686 to Vass et al. issued on

Aug. 17, 2010, U.S. Pat. No. 7,813,785 to Okerlund et al. issued on Oct. 12, 2010, U.S. Pat. No. 7,996,063 to Vass et al. issued on Aug. 9, 2011, U.S. Pat. No. 8,060,185 to Hunter et al. issued on Nov. 15, 2011, and U.S. Pat. No. 8,401,616 to Verard et al. issued on Mar. 19, 2013, each of which are incorporated herein by reference in their entirety.

The display apparatus **130** and the computing apparatus **140** may be configured to display and analyze data such as, e.g., surrogate electrical activation information or data, electrocardiogram data, etc. gathered, or collected, using the electrode apparatus **110**. In at least one embodiment, the computing apparatus **140** may be a server, a personal computer, or a tablet computer. The computing apparatus **140** may be configured to receive input from input apparatus **142** and transmit output to the display apparatus **130**. Further, the computing apparatus **140** may include data storage that may allow for access to processing programs or routines and/or one or more other types of data, e.g., for driving a graphical user interface configured to noninvasively assist a user in location selection of an implantable electrode, etc.

The computing apparatus **140** may be operatively coupled to the input apparatus **142** and the display apparatus **130** to, e.g., transmit data to and from each of the input apparatus **142** and the display apparatus **130**. For example, the computing apparatus **140** may be electrically coupled to each of the input apparatus **142** and the display apparatus **130** using, e.g., analog electrical connections, digital electrical connections, wireless connections, bus-based connections, network-based connections, internet-based connections, etc. As described further herein, a user may provide input to the input apparatus **142** to manipulate, or modify, one or more graphical depictions displayed on the display apparatus **130** to view and/or select one or more pieces of information related to a patient's cardiac health as further described herein.

Although as depicted the input apparatus **142** is a keyboard, it is to be understood that the input apparatus **142** may include any apparatus capable of providing input to the computing apparatus **140** to perform the functionality, methods, and/or logic described herein. For example, the input apparatus **142** may include a mouse, a trackball, a touchscreen (e.g., capacitive touchscreen, a resistive touchscreen, a multi-touch touchscreen, etc.), etc. Likewise, the display apparatus **130** may include any apparatus capable of displaying information to a user, such as a graphical user interface **132** including graphical depictions of anatomy of a human heart, images or graphical depictions of the patient's heart, graphical depictions of locations of one or more electrodes, graphical depictions of a human torso, images or graphical depictions of the patient's torso, alphanumeric representations of one or more values, graphical depictions or actual images of implanted electrodes and/or leads, etc. For example, the display apparatus **130** may include a liquid crystal display, an organic light-emitting diode screen, a touchscreen, a cathode ray tube display, etc.

The graphical user interfaces **132** displayed by the display apparatus **130** may include, or display, one or more regions used to display graphical depictions, to display images, to allow selection of one or more regions or areas of such graphical depictions and images, etc. As used herein, a "region" of a graphical user interface **132** may be defined as a portion of the graphical user interface **132** within which information may be displayed or functionality may be performed. Regions may exist within other regions, which may be displayed separately or simultaneously. For example, smaller regions may be located within larger regions, regions may be located side-by-side, etc. Addition-

ally, as used herein, an "area" of a graphical user interface **132** may be defined as a portion of the graphical user interface **132** located with a region that is smaller than the region it is located within.

The processing programs or routines stored and/or executed by the computing apparatus **140** may include programs or routines for computational mathematics, matrix mathematics, decomposition algorithms, compression algorithms (e.g., data compression algorithms), calibration algorithms, image construction algorithms, signal processing algorithms (e.g., various filtering algorithms, Fourier transforms, fast Fourier transforms, etc.), standardization algorithms, comparison algorithms, vector mathematics, or any other processing required to implement one or more exemplary methods and/or processes described herein. Data stored and/or used by the computing apparatus **140** may include, for example, electrical signal/waveform data from the electrode apparatus **110**, electrical activation times from the electrode apparatus **110**, graphics (e.g., graphical elements, icons, buttons, windows, dialogs, pull-down menus, graphic areas, graphic regions, 3D graphics, etc.), graphical user interfaces, results from one or more processing programs or routines employed according to the disclosure herein, or any other data that may be necessary for carrying out the one and/or more processes or methods described herein.

In one or more embodiments, the exemplary systems, methods, and interfaces may be implemented using one or more computer programs executed on programmable computers, such as computers that include, for example, processing capabilities, data storage (e.g., volatile or non-volatile memory and/or storage elements), input devices, and output devices. Program code and/or logic described herein may be applied to input data to perform functionality described herein and generate desired output information. The output information may be applied as input to one or more other devices and/or methods as described herein or as would be applied in a known fashion.

The one or more programs used to implement the systems, methods, and/or interfaces described herein may be provided using any programmable language, e.g., a high level procedural and/or object orientated programming language that is suitable for communicating with a computer system. Any such programs may, for example, be stored on any suitable device, e.g., a storage media, that is readable by a general or special purpose program running on a computer system (e.g., including processing apparatus) for configuring and operating the computer system when the suitable device is read for performing the procedures described herein. In other words, at least in one embodiment, the exemplary systems, methods, and/or interfaces may be implemented using a computer readable storage medium, configured with a computer program, where the storage medium so configured causes the computer to operate in a specific and predefined manner to perform functions described herein. Further, in at least one embodiment, the exemplary systems, methods, and/or interfaces may be described as being implemented by logic (e.g., object code) encoded in one or more non-transitory media that includes code for execution and, when executed by a processor, is operable to perform operations such as the methods, processes, and/or functionality described herein.

The computing apparatus **140** may be, for example, any fixed or mobile computer system (e.g., a controller, a micro-controller, a personal computer, mini computer, tablet computer, etc.). The exact configuration of the computing apparatus **130** is not limiting, and essentially any device capable

of providing suitable computing capabilities and control capabilities (e.g., graphics processing, etc.) may be used. As described herein, a digital file may be any medium (e.g., volatile or non-volatile memory, a CD-ROM, a punch card, magnetic recordable tape, etc.) containing digital bits (e.g., encoded in binary, trinary, etc.) that may be readable and/or writeable by computing apparatus **140** described herein. Also, as described herein, a file in user-readable format may be any representation of data (e.g., ASCII text, binary numbers, hexadecimal numbers, decimal numbers, graphically, etc.) presentable on any medium (e.g., paper, a display, etc.) readable and/or understandable by a user.

In view of the above, it will be readily apparent that the functionality as described in one or more embodiments according to the present disclosure may be implemented in any manner as would be known to one skilled in the art. As such, the computer language, the computer system, or any other software/hardware which is to be used to implement the processes described herein shall not be limiting on the scope of the systems, processes or programs (e.g., the functionality provided by such systems, processes or programs) described herein.

The electrical activation times of the patient's heart may be useful to evaluate a patient's cardiac health and/or to evaluate cardiac therapy being delivered to a patient. Electrical activation information or data of one or more regions of a patient's heart may be determined using electrode apparatus **110** as shown in FIG. 1 and in FIGS. 2A-2B. The exemplary electrode apparatus **110** may be configured to measure body-surface potentials of a patient **14** and, more particularly, torso-surface potentials of a patient **14**. As shown in FIG. 2A, the exemplary electrode apparatus **110** may include a set, or array, of electrodes **112**, a strap **113**, and interface/amplifier circuitry **116**. The electrodes **112** may be attached, or coupled, to the strap **113** and the strap **113** may be configured to be wrapped around the torso of a patient **14** such that the electrodes **112** surround the patient's heart. As further illustrated, the electrodes **112** may be positioned around the circumference of a patient **14**, including the posterior, lateral, posterolateral, and anterior locations of the torso of a patient **14**.

Further, the electrodes **112** may be electrically connected to interface/amplifier circuitry **116** via wired connection **118**. The interface/amplifier circuitry **116** may be configured to amplify the signals from the electrodes **112** and provide the signals to the computing apparatus **140**. Other exemplary systems may use a wireless connection to transmit the signals sensed by electrodes **112** to the interface/amplifier circuitry **116** and, in turn, the computing apparatus **140**, e.g., as channels of data. For example, the interface/amplifier circuitry **116** may be electrically coupled to each of the computing apparatus **140** and the display apparatus **130** using, e.g., analog electrical connections, digital electrical connections, wireless connections, bus-based connections, network-based connections, internet-based connections, etc.

Although in the example of FIG. 2A the electrode apparatus **110** includes a strap **113**, in other examples any of a variety of mechanisms, e.g., tape or adhesives, may be employed to aid in the spacing and placement of electrodes **112**. In some examples, the strap **113** may include an elastic band, strip of tape, or cloth. In other examples, the electrodes **112** may be placed individually on the torso of a patient **14**. Further, in other examples, electrodes **112** (e.g., arranged in an array) may be part of, or located within, patches, vests, and/or other manners of securing the electrodes **112** to the torso of the patient **14**. Still further, in other examples, the electrodes **112** may be part of, or located within, two

sections of material or two “patches.” One of the two sections or patches may be located on the anterior side of the torso of the patient 14 (to, e.g., measure surrogate cardiac electrical activation times representative of the anterior side of the patient’s heart) and the other section or patch may be located on the posterior side of the torso of the patient 14 (to, e.g., measure surrogate cardiac electrical activation times representative of the posterior side of the patient’s heart).

The electrodes 112 may be configured to surround the heart of the patient 14 and record, or monitor, the electrical signals associated with the depolarization and repolarization of the heart after the signals have propagated through the torso of a patient 14. Each of the electrodes 112 may be used in a unipolar configuration to sense the torso-surface potentials that reflect the cardiac signals. The interface/amplifier circuitry 116 may also be coupled to a return or indifferent electrode (not shown) that may be used in combination with each electrode 112 for unipolar sensing. In some examples, there may be about 12 to about 50 electrodes 112 spatially distributed around the torso of patient. Other configurations may have more or fewer electrodes 112.

The computing apparatus 140 may record and analyze the torso-surface potential signals sensed by electrodes 112 and amplified/conditioned by the interface/amplifier circuitry 116. The computing apparatus 140 may be configured to analyze the signals from the electrodes 112 to provide surrogate electrical activation information or data such as surrogate cardiac electrical activation times, e.g., representative of actual, or local, electrical activation times of one or more regions of the patient’s heart as will be further described herein. For example, electrical signals measured at the left anterior surface location of a patient’s torso may be representative, or surrogates, of electrical signals of the left anterior left ventricle region of the patient’s heart, electrical signals measured at the left lateral surface location of a patient’s torso may be representative, or surrogates, of electrical signals of the left lateral left ventricle region of the patient’s heart, electrical signals measured at the left posterolateral surface location of a patient’s torso may be representative, or surrogates, of electrical signals of the posterolateral left ventricle region of the patient’s heart, and electrical signals measured at the posterior surface location of a patient’s torso may be representative, or surrogates, of electrical signals of the posterior left ventricle region of the patient’s heart. In one or more embodiments, measurement of activation times can be performed by measuring time between the onset of cardiac depolarization (e.g., onset of QRS complex) to the next the onset of cardiac depolarization. In one or more embodiments, measurement of activation times can be performed by picking an appropriate fiducial point (e.g., peak values, minimum values, minimum slopes, maximum slopes, zero crossings, threshold crossings, etc. of a near or far-field EGM) and measuring time between fiducial points (e.g., within the electrical activity).

Additionally, the computing apparatus 140 may be configured to provide graphical user interfaces depicting the surrogate electrical activation times obtained using the electrode apparatus 110. Exemplary systems, methods, and/or interfaces may noninvasively use the electrical information collected using the electrode apparatus 110 to evaluate a patient for cardiac therapy and/or evaluate cardiac therapy being delivered to the patient.

FIG. 2B illustrates another exemplary electrode apparatus 110 that includes a plurality of electrodes 112 configured to surround the heart of the patient 14 and record, or monitor, the electrical signals associated with the depolarization and repolarization of the heart after the signals have propagated

through the torso of the patient 14. The electrode apparatus 110 may include a vest 114 upon which the plurality of electrodes 112 may be attached, or to which the electrodes 112 may be coupled. In at least one embodiment, the plurality, or array, of electrodes 112 may be used to collect electrical information such as, e.g., surrogate electrical activation times. Similar to the electrode apparatus 110 of FIG. 2A, the electrode apparatus 110 of FIG. 2B may include interface/amplifier circuitry 116 electrically coupled to each of the electrodes 112 through a wired connection 118 and be configured to transmit signals from the electrodes 112 to computing apparatus 140. As illustrated, the electrodes 112 may be distributed over the torso of a patient 14, including, for example, the anterior, lateral, and posterior surfaces of the torso of the patient 14.

The vest 114 may be formed of fabric with the electrodes 112 attached to the fabric. The vest 114 may be configured to maintain the position and spacing of electrodes 112 on the torso of the patient 14. Further, the vest 114 may be marked to assist in determining the location of the electrodes 112 on the surface of the torso of the patient 14. In some examples, there may be about 25 to about 256 electrodes 112 distributed around the torso of the patient 14, though other configurations may have more or fewer electrodes 112.

As described herein, the electrode apparatus 110 may be configured to measure electrical information (e.g., electrical signals) representing different regions of a patient’s heart. More specifically, activation times of different regions of a patient’s heart can be approximated from surface electrocardiogram (ECG) activation times measured using surface electrodes in proximity to surface areas corresponding to the different regions of the patient’s heart.

The exemplary systems, methods, and interfaces may be used in one or more different settings to provide noninvasive assistance to a user in the evaluation of a patient for cardiac therapy and/or evaluation of on-going cardiac therapy (e.g., cardiac therapy being presently-delivered to a patient). Further, the exemplary systems, methods, and interfaces may be used to assist a user in the configuration and/or adjustment of one or more cardiac therapy settings for the cardiac therapy to be delivered to a patient or being presently-delivered to a patient.

To provide this functionality, the exemplary systems, methods, and interfaces may be described as being configurable in a plurality of modes of operation 150 as depicted in FIG. 3. For example, the exemplary systems, methods, and interfaces may be configured in an initial examination mode 152, an implantation examination mode 154, and a follow-up examination mode 156.

The initial examination mode 152 may be configured for the noninvasive evaluation of a patient for cardiac therapy. For example, the initial examination mode 152 may be designed for an initial consultation/evaluation of a patient considering cardiac therapy and will be described further herein in reference to the exemplary initial examination graphical user interface of FIG. 4.

The implantation examination mode 154 may be configured for the noninvasive evaluation of cardiac therapy being delivered to a patient and/or being currently configured for a patient (e.g., during configuration of cardiac resynchronization therapy (CRT) after implantation of the CRT device). For example, the implantation examination mode 154 may be designed for use by a physician in evaluating cardiac therapy during implantation of a cardiac therapy device, immediately after implantation of a cardiac therapy device and/or during configuration of the cardiac therapy immediately following implantation. Further, exemplary systems

configurable in the implantation examination mode **154** may include additional functionality such as imaging and navigation functionality for electrode and/or lead placement as described herein. The implantation examination mode **154** will be described further herein in reference to the exemplary implantation examination graphical user interface of FIG. **6**.

The follow-up examination mode **156** may be configured for the noninvasive evaluation of cardiac therapy being delivered to a patient after a cardiac therapy device has been delivering cardiac therapy to a patient for a period of time (e.g., after implantation, not immediately following implantation or initial configuration of the cardiac therapy, etc.). For example, the follow-up examination mode **156** may be designed for use by a physician in evaluating cardiac therapy in follow-up appointments days, weeks, months, or years after a cardiac therapy has been implanted in a patient and will be described further herein in reference to the exemplary follow-up examination graphical user interface of FIG. **8**.

The exemplary systems, methods, and interfaces may provide, or be configurable in, one or more of the plurality of modes **150**. For example, an exemplary system may provide, or be configurable in, all of the plurality of modes **150**. Further, for example, an exemplary system may provide, or be configurable in, two of the three modes **150** such as, e.g., the initial examination mode **152** and the follow-up mode **156**. In at least one embodiment, a system designed for use in an operating room of a medical care facility (e.g., for the surgical implantation and configuration of a cardiac therapy device) may include each of the three modes **150**. In at least one embodiment, a system designed for use in a consultation room of a medical care facility (e.g., for general evaluation and consultation of a patient) may include the initial examination mode **152** and the follow-up mode **156** but not the implantation mode **154**.

An exemplary graphical interface (GUI) **160** for use in an initial examination or consultation is depicted in FIG. **4**. As shown, the GUI **160** may include, among other things, a graphical representation **162** of measured surrogate electrical activation times, metrics **170** of cardiac health, and an indication **172** of whether cardiac therapy for the patient may be beneficial. The initial examination may include applying electrode apparatus such as described herein with reference to FIGS. **1-2** to a patient, measuring surrogate electrical activation times and ECG information using the electrode apparatus, and displaying the data for the evaluation of the patient for cardiac therapy on the GUI **160**.

The graphical representation **162** of measured surrogate cardiac electrical activation times may be depicted in a variety of fashions. As shown, the surrogate electrical activation times are shown as a color-coded, or color-scaled, segment **168** extending over, or wrapped around, a graphical representation of a human torso **164**, **166**. More specifically, an anterior side of a human torso **164** and a posterior side of a human torso **166** are depicted, each including a color-coded segment **168** graphically depicting surrogate electrical activation times measured, e.g., using the electrical apparatus described herein with reference to FIGS. **1-2**. Further, the graphical representation **162** of measured surrogate cardiac electrical activation times shown on the anterior side of a human torso **164** may be measured using electrodes located on, or proximate to, the anterior side of the patient's torso, and likewise, the graphical representation **162** of measured surrogate cardiac electrical activation times shown on the posterior side of the human torso **166** may be measured using electrodes located on, or proximate to, the

posterior side of the patient's torso. In other words, the graphical representation **162** of measured surrogate cardiac electrical activation times shown on the anterior side of the human torso **164** correlates to actual electrical signals measured using electrodes configured to measure electrical signals on the anterior side of the patient's torso, and the graphical representation **162** of measured surrogate cardiac electrical activation times shown on the posterior side of the human torso **166** correlates to actual electrical signals measured using electrodes configured to measure electrical signals on the posterior side of the patient's torso. The graphical representation **162** further includes a color-coded scale **169** corresponding to the color-coded segments **168**, to, e.g., provide basis for the coloring of the color-coded segments **168**.

Additional exemplary graphical representations of surrogate electrical activation times may be described in U.S. Patent Application Publication No. 2012/0284003 A1 published on Nov. 8, 2012 and entitled "Assessing Intra-Cardiac Activation Patterns" and U.S. Patent Application Publication No. 2012/0283587 A1 published on Nov. 8, 2012 and entitled "Assessing Intra-Cardiac Activation Patterns and Electrical Dyssynchrony," each of which are hereby incorporated by reference in their entireties.

In other embodiments, the surrogate electrical activation times may be color-coded across the entire graphical depiction of a human torso and/or any smaller or larger part of human anatomy. Further, in at least one embodiment, the graphical depictions of a human torso **164**, **166** may be actual images of the patient being evaluated. The surrogate cardiac electrical activation times may be further depicted alphanumerically over a graphical depiction of human anatomy. For example, a plurality of surrogate cardiac electrical activation times in milliseconds may be graphically overlaid over the torsos **164**, **166**.

In one or more embodiments, the graphical depiction of a portion of human anatomy displayed on the exemplary graphical user interfaces may include a graphical representation of a human heart. For example, a posterior side of a human heart **12** is depicted in FIG. **5** with surrogate electrical activation times color-coded across the surface of the heart **12**. As shown, the posterolateral left ventricle region **165** shows late activation (e.g., about 150 milliseconds). In other embodiments, both a posterior and anterior side of a human heart may be graphically depicted and overlaid with electrical activation information.

The exemplary GUI **160** of FIG. **4** further includes metrics **170** of the patient's cardiac health. As shown, the metrics **170** are QRS width and electrical dyssynchrony, which both may be useful in the evaluation of the patient for cardiac therapy. Although in this embodiment, the metrics **170** include two metrics, it is to be understood that exemplary graphical user interfaces may include one metric or more than two metrics related to the patient's cardiac health. For example, the metrics **170** may include one or more of a standard deviation, range, interquartile deviation, one or more statistical averages (e.g., mean, median, mode) of activation times of all electrodes or a subset of electrodes proximate a specific portion of anatomy (such as, e.g., electrodes located at the left side of the patient that are surrogate of the left ventricle), one or more timing intervals between certain fiducial points (e.g., onset of depolarization to first negative or positive peak on one or more electrocardiograms, etc.), etc. Further, the cardiac health metrics may include one or more metrics described in U.S. Provisional Patent Application No. 61/834,133 entitled "METRICS OF ELECTRICAL DYSSYNCHRONY AND ELECTRICAL

ACTIVATION PATTERNS FROM SURFACE ECG ELECTRODES” and filed on Jun. 12, 2013, which is hereby incorporated by reference in its entirety.

The exemplary GUI 160 of FIG. 4 further includes an indication 172 of whether cardiac therapy for the patient may be beneficial. The indication 172 of whether cardiac therapy for the patient may be beneficial may be defined as a suggestion or recommendation based on the measured electrical data from the electrode apparatus with respect to a potential cardiac therapy. In the exemplary GUI 160, the indication 172 of whether cardiac therapy for the patient may be beneficial notes the following: “POTENTIAL CRT BENEFIT=HIGH,” and thus, the indication 172 indicates that this patient may have a high likelihood of benefiting from cardiac resynchronization therapy.

Additionally, the exemplary GUI 160 of FIG. 4 includes an electrocardiogram area 174 that depicts one or more electrocardiograms of the patient measured using the electrode apparatus. Although only one electrocardiogram is depicted in the electrocardiogram area 174, a plurality of electrocardiograms may be depicted in the electrocardiogram area 174 and/or other areas of the GUI 160. Further, one or more (e.g., a set or subset) of available electrocardiograms (e.g., measured using different electrodes or electrode sets located in different locations about a patient) may be selected by a user (e.g., physician) for viewing on the GUI 160. When the electrocardiogram area 174 includes more than one electrocardiogram, the electrocardiograms may be time-aligned (e.g., the electrograms may be aligned along the same time period, the electrograms may be aligned by cardiac cycle, etc.). Additionally, each electrogram may cover either a single beat or multiple beats, which may be selected or configured by a user. Further, the one or more electrocardiograms may be stored from a previous examination such that a user may compare previously-recorded electrocardiograms to presently-monitored electrocardiograms.

The data such as electrical activation time data, metrics of cardiac health, electrocardiograms, etc. measured when using the initial examination mode may be stored within the computing apparatus for use at a later time, e.g., to compare the patient’s cardiac health before cardiac therapy to after cardiac therapy, to adjust or configure cardiac therapy for a patient, etc. For example, the data collected during the initial examination/consultation depicted in the initial examination graphical user interface 160 of FIG. 4 is used in the exemplary implantation graphical user interface 180 of FIG. 6 and the exemplary follow-up graphical user interface 220 of FIG. 8.

During implantation of a cardiac therapy device such, e.g., as a cardiac resynchronization therapy device, the exemplary systems, methods, and interfaces may be configured to provide noninvasive evaluation of the cardiac therapy being implanted. An exemplary graphical interface (GUI) 180 for use during implantation is depicted in FIG. 6. As shown, the GUI 180 may include, among other things, graphical representations 181 of previously-measured and presently measured surrogate electrical activation times, previously-measured and presently-measured metrics 184, 186 of cardiac health, and an indication 188 of whether the cardiac therapy appears to be beneficial. Although the surrogate electrical activation times and cardiac health metrics are described as being previously-measured or presently-measured, it is to be understood that “previously-measured” data is designed to encompass, or describe, any measurements prior to the “presently-measured” data, and, likewise, “presently-measured” data is designed to encompass, or

describe, any measurements after to the “previously-measured” data. In one or more embodiments, the “previously-measured” data may represent data measured during intrinsic rhythm before cardiac therapy (e.g., implantation of a CRT device). In other words, the “previously-measured” data may represent baseline values. In one or more embodiments, the “previously-measured” data may represent data measured with cardiac therapy being disabled or before any adjustments or modifications are made to cardiac therapy being presently delivered (e.g., by CRT device) and the “presently-measured” data may represent data measured after the adjustments or modifications are made.

Similar to the initial examination, electrode apparatus such as described herein with reference to FIGS. 1-2 may be applied to a patient, surrogate electrical activation times and ECG information using the electrode apparatus may be measured, and the surrogate electrical activation times and ECG information data may be displayed on the GUI 180. In at least one embodiment, the GUI 180 may be described as providing a before-and-after representation of the cardiac health of a patient before cardiac therapy and after cardiac therapy.

The graphical representations 181 of previously-measured and presently measured surrogate electrical activation times include a graphical representation 182 of previously-measured surrogate electrical activation times and a graphical representation 183 of presently-measured surrogate electrical activation times. As labeled on the GUI 180, the graphical representation 182 of previously-measured surrogate electrical activation times represents the patient’s intrinsic rhythm (e.g., without pacing, without cardiac therapy, etc.) and the graphical representation 183 of presently-measured surrogate electrical activation times represents the patient’s paced rhythm. The graphical representations 182, 183 may be substantially similar to the graphical representation 162 of the GUI 160 described herein with reference to FIG. 4.

The GUI 180 further includes previously-measured metrics 184 of cardiac health and presently-measured metrics 186 of cardiac health for the patient such that a user may compare the values presented thereby to, e.g., evaluate the cardiac therapy being delivered. As shown, similar to the initial examination GUI 160, the metrics 184, 186 include QRS width and electrical dyssynchrony. As shown, the QRS width has increased by 10 milliseconds and the electrical dyssynchrony has decreased 12 milliseconds from the intrinsic data to the paced data, which may, e.g., indicate that the cardiac therapy being delivered to the patient is beneficial. The GUI 180 further includes an indication 188 of whether the cardiac therapy appears to be beneficial. In this example, the indication 188 notes: “LV LEAD LOCATION: GOOD Dyssynchrony improved by 36%.” Although this indication 188 is based on an electrical dyssynchrony comparison between the previously-measured electrical dyssynchrony, other exemplary indications of whether cardiac therapy appears to be beneficial for a patient may include, or utilize, different data such as, e.g., comparisons of ECG waveform between previously measured ECG waveform shape (e.g., as may be described in “Analysis of Ventricular Activation Using Surface Electrocardiography to Predict Left Ventricular Reverse Volumetric Remodeling During Cardiac Resynchronization Therapy” by Sweeney, et al., *Circulation: Arrhythmia/Electrophysiology*, Feb. 9, 2010: pages 626-634, which is incorporated herein by reference in its entirety), changes in ECG vector loop areas (e.g., as may be described in “Vectorcardiography as a Tool for Easy Optimization of Cardiac Resynchronization Therapy in Canine Left Bundle

Branch Block Hearts” by van Deursen et al., *Circulation: Arrhythmia/Electrophysiology*, June 2012: pages 544-552, which is incorporated herein by reference in its entirety), changes in QRS duration, etc.

Additionally, the GUI **180** of FIG. **6** may include an electrocardiogram area **189** similar to the electrocardiogram area **174** described herein with reference to FIG. **4**. Further, the electrocardiogram area **189** may include one or more presently- and/or previously-measured electrocardiograms, which may be selected by a user. When the electrocardiogram area **189** includes more than one electrocardiogram, the electrocardiograms may be time-aligned as described herein.

A user (e.g., a physician) may use the GUI **180** during implantation of a cardiac therapy device and/or in the configuration of the cardiac therapy device. For example, the user may use the GUI **180** to confirm that the implanted cardiac therapy device (e.g., cardiac resynchronization therapy device) is providing beneficial therapy to the patient. Additionally, the user may want to change one or more settings of configurations of the cardiac therapy device while actively monitoring whether the cardiac therapy is beneficial to the patient. For example, the user may want to change a pacing timing interval, change a pacing vector, and/or change a pacing mode while actively monitoring the cardiac therapy using the GUI **180**.

An exemplary method of noninvasive evaluation of cardiac therapy for a patient and configuration of cardiac therapy **200** using, e.g., GUI **180** of FIG. **6**, is depicted in FIG. **7**. As shown, the method **200** may measure electrical data using electrode apparatus **202** such as, e.g., electrical activation times, electrocardiograms, etc. The electrical activation times may be displayed **204** on the GUI **180** such as the graphical depictions/representations of surrogate electrical activation times about a human anatomy portion. Further, the cardiac information/metrics based on the measured electrical data may also be displayed **206** on the GUI **180** such as electrical dyssynchrony.

Based on the information displayed by the GUI **180**, a user may decide to modify the cardiac therapy **208**. In at least one embodiment, the GUI **180** may provide the interface to modify/program the cardiac therapy device (e.g., modify or adjust one or more various configurations, parameters, etc. of the cardiac therapy device). For example, as shown in FIG. **6**, the GUI **180** may include a modify area, or button, **190** that, upon selection, may trigger or initiate another graphical user interface used to modify or program the cardiac therapy device.

As described herein, a user may want to change a pacing timing interval **210**, change a pacing vector **212**, and/or change a pacing mode **214**. The timing intervals may include A-V intervals, V-V intervals, etc. for two, three, and/or four chambers of the heart (e.g., two or more of the right atrium, the right ventricle, the left atrium, and the left ventricle) and/or timing intervals for multiple pulses delivered for a single cardiac cycle to the right and left ventricle from different electrodes (e.g. RV electrode to LV1 electrode, LV1 electrode to LV2 electrode, etc.). Changing the pacing vector **212** may include changing one or more electrodes that may be used for various pacing vectors during the programmed therapy. For example, one electrode of a pacing vector may be switched from a tip electrode to ring electrode. Further, any of one or more pacing electrodes (e.g., electrodes located on leads, leadless/wireless electrodes, etc.) used in cardiac therapy systems may be selected or changed for any pacing vector. Changing the pacing mode **214** may include changing the type of pacing being provided. For example, a

cardiac therapy device may be delivering LV-only pacing, and a user may switch the LV-only pacing to bi-ventricular pacing. Similarly, a pacing configuration may be changed to pace from a single location within one chamber of the heart to pacing more than two or more locations within that same chamber with or without programmed timing delays.

After the user has adjusted or modified the cardiac therapy, the graphical depiction **183** of the presently-measured activation times and the presently-measured metrics **186** may be updated to reflect the results from the new adjustments or modifications to the cardiac therapy. Thus, a user may immediately know whether any adjustments or modifications have appeared to benefit the patient.

Further, although the graphical depiction **182** of the previously-measured activation times and the previously-measured cardiac metrics **184** of the GUI **180** depicted in FIG. **6** show measurements from intrinsic rhythm, the GUI **180** could be configured (e.g., by user selection, etc.) to show previously-measured data that was measured or recorded prior to any adjustments or modifications being made to the cardiac therapy. Still further, the indication **188** of whether the cardiac therapy appears to be beneficial may also be updated to reflect the change.

After implantation of a cardiac therapy device such, e.g., as a cardiac resynchronization therapy device, the exemplary systems, methods, and interfaces may be configured to provide noninvasive evaluation of the cardiac therapy after implantation (e.g., days, weeks, months, years, etc. after implantation). An exemplary graphical interface (GUI) **220** for use during a follow-up examination is depicted in FIG. **8**. As shown, the GUI **220** may include graphical representations **221** of previously-measured and presently measured surrogate electrical activation times, previously-measured and presently-measured metrics **224**, **226** of cardiac health, an indication **228** of whether the cardiac therapy appears to be beneficial, electrocardiogram area **229**, and a modify area/button **230** that may be substantially similar to the graphical representations **181**, previously-measured and presently measured metrics **184**, **186**, an indication **188**, and electrocardiogram area **189**, and modify area/button **190** of the GUI **180** described herein with respect to FIG. **6**. The electrocardiogram area **189** may be configured to depict a plurality of different electrocardiograms (e.g., time-aligned, etc.). For example, the electrocardiogram area **189** may display electrocardiographic waveform data, such as several time-aligned single- or multiple beat electrocardiographic waveforms from the same cardiac cycle(s) measured from different electrode(s) around the patient’s torso, which may be selected by a user.

The implantable electrodes that may be implanted using the exemplary systems, methods, and graphical user interfaces described herein may be used with respect to the implantation and configuration an implantable medical device (IMD) and/or one or more leads configured to be located proximate one or more portions of a patient’s heart. For example, the exemplary systems, methods, and interfaces may be used in conjunction with an exemplary therapy system **10** described herein with reference to FIGS. **9-11**.

FIG. **9** is a conceptual diagram illustrating an exemplary therapy system **10** that may be used to deliver pacing therapy to a patient **14**. Patient **14** may, but not necessarily, be a human. The therapy system **10** may include an implantable medical device **16** (IMD), which may be coupled to leads **18**, **20**, **22**. The IMD **16** may be, e.g., an implantable pacemaker, cardioverter, and/or defibrillator, that provides electrical signals to the heart **12** of the patient **14** via electrodes coupled to one or more of the leads **18**, **20**, **22** (e.g.,

electrodes that may be implanted in accordance with the description herein, such as, with use of noninvasive selection of implantation site regions).

The leads **18**, **20**, **22** extend into the heart **12** of the patient **14** to sense electrical activity of the heart **12** and/or to deliver electrical stimulation to the heart **12**. In the example shown in FIG. **15**, the right ventricular (RV) lead **18** extends through one or more veins (not shown), the superior vena cava (not shown), and the right atrium **26**, and into the right ventricle **28**. The left ventricular (LV) coronary sinus lead **20** extends through one or more veins, the vena cava, the right atrium **26**, and into the coronary sinus **30** to a region adjacent to the free wall of the left ventricle **32** of the heart **12**. The right atrial (RA) lead **22** extends through one or more veins and the vena cava, and into the right atrium **26** of the heart **12**.

The IMD **16** may sense, among other things, electrical signals attendant to the depolarization and repolarization of the heart **12** via electrodes coupled to at least one of the leads **18**, **20**, **22**. The IMD **16** may be configured to determine or identify effective electrodes located on the leads **18**, **20**, **22** using the exemplary methods and processes described herein. In some examples, the IMD **16** provides pacing therapy (e.g., pacing pulses) to the heart **12** based on the electrical signals sensed within the heart **12**. The IMD **16** may be operable to adjust one or more parameters associated with the pacing therapy such as, e.g., AV delay and other various timings, pulse wide, amplitude, voltage, burst length, etc. Further, the IMD **16** may be operable to use various electrode configurations to deliver pacing therapy, which may be unipolar, bipolar, quadripolar, or further multipolar. For example, a multipolar lead may include several electrodes that can be used for delivering pacing therapy. Hence, a multipolar lead system may provide, or offer, multiple electrical vectors to pace from. A pacing vector may include at least one cathode, which may be at least one electrode located on at least one lead, and at least one anode, which may be at least one electrode located on at least one lead (e.g., the same lead, or a different lead) and/or on the casing, or can, of the IMD. While improvement in cardiac function as a result of the pacing therapy may primarily depend on the cathode, the electrical parameters like impedance, pacing threshold voltage, current drain, longevity, etc. may be more dependent on the pacing vector, which includes both the cathode and the anode. The IMD **16** may also provide defibrillation therapy and/or cardioversion therapy via electrodes located on at least one of the leads **18**, **20**, **22**. Further, the IMD **16** may detect arrhythmia of the heart **12**, such as fibrillation of the ventricles **28**, **32**, and deliver defibrillation therapy to the heart **12** in the form of electrical pulses. In some examples, IMD **16** may be programmed to deliver a progression of therapies, e.g., pulses with increasing energy levels, until a fibrillation of heart **12** is stopped.

FIGS. **10A-10B** are conceptual diagrams illustrating the IMD **16** and the leads **18**, **20**, **22** of therapy system **10** of FIG. **15** in more detail. The leads **18**, **20**, **22** may be electrically coupled to a therapy delivery module (e.g., for delivery of pacing therapy), a sensing module (e.g., for sensing one or more signals from one or more electrodes), and/or any other modules of the IMD **16** via a connector block **34**. In some examples, the proximal ends of the leads **18**, **20**, **22** may include electrical contacts that electrically couple to respective electrical contacts within the connector block **34** of the IMD **16**. In addition, in some examples, the leads **18**, **20**, **22** may be mechanically coupled to the

connector block **34** with the aid of set screws, connection pins, or another suitable mechanical coupling mechanism.

Each of the leads **18**, **20**, **22** includes an elongated insulative lead body, which may carry a number of conductors (e.g., concentric coiled conductors, straight conductors, etc.) separated from one another by insulation (e.g., tubular insulative sheaths). In the illustrated example, bipolar electrodes **40**, **42** are located proximate to a distal end of the lead **18**. In addition, the bipolar electrodes **44**, **45**, **46**, **47** are located proximate to a distal end of the lead **20** and the bipolar electrodes **48**, **50** are located proximate to a distal end of the lead **22**.

The electrodes **40**, **44**, **45**, **46**, **47**, **48** may take the form of ring electrodes, and the electrodes **42**, **50** may take the form of extendable helix tip electrodes mounted retractably within the insulative electrode heads **52**, **54**, **56**, respectively. Each of the electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50** may be electrically coupled to a respective one of the conductors (e.g., coiled and/or straight) within the lead body of its associated lead **18**, **20**, **22**, and thereby coupled to respective ones of the electrical contacts on the proximal end of the leads **18**, **20**, **22**.

Additionally, electrodes **44**, **45**, **46** and **47** may have an electrode surface area of about 5.3 mm² to about 5.8 mm². Electrodes **44**, **45**, **46**, and **47** may also be referred to as LV1, LV2, LV3, and LV4, respectively. The LV electrodes (i.e., left ventricle electrode 1 (LV1) **44**, left ventricle electrode 2 (LV2) **45**, left ventricle electrode 3 (LV3) **46**, and left ventricle 4 (LV4) **47** etc.) on the lead **20** can be spaced apart at variable distances. For example, electrode **44** may be a distance of, e.g., about 21 millimeters (mm), away from electrode **45**, electrodes **45** and **46** may be spaced a distance of, e.g. about 1.3 mm to about 1.5 mm, away from each other, and electrodes **46** and **47** may be spaced a distance of, e.g. 20 mm to about 21 mm, away from each other.

The electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50**, **58** may further be used to sense electrical signals (e.g., morphological waveforms within electrograms (EGM)) attendant to the depolarization and repolarization of the heart **12**. The sensed electrical signals may be used to determine which of the electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50** are the most effective in improving cardiac function. The electrical signals are conducted to the IMD **16** via the respective leads **18**, **20**, **22**. In some examples, the IMD **16** may also deliver pacing pulses via the electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50** to cause depolarization of cardiac tissue of the patient's heart **12**. In some examples, as illustrated in FIG. **10A**, the IMD **16** includes one or more housing electrodes, such as housing electrode **58**, which may be formed integrally with an outer surface of a housing **60** (e.g., hermetically-sealed housing) of the IMD **16** or otherwise coupled to the housing **60**. Any of the electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48** and **50** may be used for unipolar sensing or pacing in combination with housing electrode **58**. In other words, any of electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50**, **58** may be used in combination to form a sensing vector, e.g., a sensing vector that may be used to evaluate and/or analyze the effectiveness of pacing therapy. It is generally understood by those skilled in the art that other electrodes can also be selected to define, or be used for, pacing and sensing vectors. Further, any of electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50**, **58**, which are not being used to deliver pacing therapy, may be used to sense electrical activity during pacing therapy.

As described in further detail with reference to FIG. **10A**, the housing **60** may enclose a therapy delivery module that may include a stimulation generator for generating cardiac pacing pulses and defibrillation or cardioversion shocks, as

well as a sensing module for monitoring the patient's heart rhythm. The leads **18**, **20**, **22** may also include elongated electrodes **62**, **64**, **66**, respectively, which may take the form of a coil. The IMD **16** may deliver defibrillation shocks to the heart **12** via any combination of the elongated electrodes **62**, **64**, **66** and the housing electrode **58**. The electrodes **58**, **62**, **64**, **66** may also be used to deliver cardioversion pulses to the heart **12**. Further, the electrodes **62**, **64**, **66** may be fabricated from any suitable electrically conductive material, such as, but not limited to, platinum, platinum alloy, and/or other materials known to be usable in implantable defibrillation electrodes. Since electrodes **62**, **64**, **66** are not generally configured to deliver pacing therapy, any of electrodes **62**, **64**, **66** may be used to sense electrical activity (e.g., for use in determining electrode effectiveness, for use in analyzing pacing therapy effectiveness, etc.) and may be used in combination with any of electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50**, **58**. In at least one embodiment, the RV elongated electrode **62** may be used to sense electrical activity of a patient's heart during the delivery of pacing therapy (e.g., in combination with the housing electrode **58** forming a RV elongated coil, or defibrillation electrode-to-housing electrode vector).

The configuration of the exemplary therapy system **10** illustrated in FIGS. **9-11** is merely one example. In other examples, the therapy system may include epicardial leads and/or patch electrodes instead of or in addition to the transvenous leads **18**, **20**, **22** illustrated in FIG. **9**. Additionally, in other examples, the therapy system **10** may be implanted in/around the cardiac space without transvenous leads (e.g., leadless/wireless pacing systems) or with leads implanted (e.g., implanted transvenously or using approaches) into the left chambers of the heart (in addition to or replacing the transvenous leads placed into the right chambers of the heart as illustrated in FIG. **9**). Further, in one or more embodiments, the IMD **16** need not be implanted within the patient **14**. For example, the IMD **16** may deliver various cardiac therapies to the heart **12** via percutaneous leads that extend through the skin of the patient **14** to a variety of positions within or outside of the heart **12**. In one or more embodiments, the system **10** may utilize wireless pacing (e.g., using energy transmission to the intracardiac pacing component(s) via ultrasound, inductive coupling, RF, etc.) and sensing cardiac activation using electrodes on the can/housing and/or on subcutaneous leads.

In other examples of therapy systems that provide electrical stimulation therapy to the heart **12**, such therapy systems may include any suitable number of leads coupled to the IMD **16**, and each of the leads may extend to any location within or proximate to the heart **12**. For example, other examples of therapy systems may include three transvenous leads located as illustrated in FIGS. **9-11**. Still further, other therapy systems may include a single lead that extends from the IMD **16** into the right atrium **26** or the right ventricle **28**, or two leads that extend into a respective one of the right atrium **26** and the right ventricle **28**.

FIG. **11A** is a functional block diagram of one exemplary configuration of the IMD **16**. As shown, the IMD **16** may include a control module **81**, a therapy delivery module **84** (e.g., which may include a stimulation generator), a sensing module **86**, and a power source **90**.

The control module **81** may include a processor **80**, memory **82**, and a telemetry module **88**. The memory **82** may include computer-readable instructions that, when executed, e.g., by the processor **80**, cause the IMD **16** and/or the control module **81** to perform various functions attributed to the IMD **16** and/or the control module **81** described

herein. Further, the memory **82** may include any volatile, non-volatile, magnetic, optical, and/or electrical media, such as a random access memory (RAM), read-only memory (ROM), non-volatile RAM (NVRAM), electrically-erasable programmable ROM (EEPROM), flash memory, and/or any other digital media. An exemplary capture management module may be the left ventricular capture management (LVCM) module described in U.S. Pat. No. 7,684,863 entitled "LV THRESHOLD MEASUREMENT AND CAPTURE MANAGEMENT" and issued Mar. 23, 2010, which is incorporated herein by reference in its entirety.

The processor **80** of the control module **81** may include any one or more of a microprocessor, a controller, a digital signal processor (DSP), an application specific integrated circuit (ASIC), a field-programmable gate array (FPGA), and/or equivalent discrete or integrated logic circuitry. In some examples, the processor **80** may include multiple components, such as any combination of one or more microprocessors, one or more controllers, one or more DSPs, one or more ASICs, and/or one or more FPGAs, as well as other discrete or integrated logic circuitry. The functions attributed to the processor **80** herein may be embodied as software, firmware, hardware, or any combination thereof.

The control module **81** may be used to determine the effectiveness of the electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50**, **58**, **62**, **64**, **66** using the exemplary methods and/or processes described herein according to a selected one or more programs, which may be stored in the memory **82**. Further, the control module **81** may control the therapy delivery module **84** to deliver therapy (e.g., electrical stimulation therapy such as pacing) to the heart **12** according to a selected one or more therapy programs, which may be stored in the memory **82**. More, specifically, the control module **81** (e.g., the processor **80**) may control various parameters of the electrical stimulus delivered by the therapy delivery module **84** such as, e.g., AV delays, pacing pulses with the amplitudes, pulse widths, frequency, or electrode polarities, etc., which may be specified by one or more selected therapy programs (e.g., AV delay adjustment programs, pacing therapy programs, pacing recovery programs, capture management programs, etc.). As shown, the therapy delivery module **84** is electrically coupled to electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50**, **58**, **62**, **64**, **66**, e.g., via conductors of the respective lead **18**, **20**, **22**, or, in the case of housing electrode **58**, via an electrical conductor disposed within housing **60** of IMD **16**. Therapy delivery module **84** may be configured to generate and deliver electrical stimulation therapy such as pacing therapy to the heart **12** using one or more of the electrodes **40**, **42**, **44**, **45**, **46**, **47**, **48**, **50**, **58**, **62**, **64**, **66**.

For example, therapy delivery module **84** may deliver pacing stimulus (e.g., pacing pulses) via ring electrodes **40**, **44**, **45**, **46**, **47**, **48** coupled to leads **18**, **20**, and **22**, respectively, and/or helical tip electrodes **42** and **50** of leads **18** and **22**. Further, for example, therapy delivery module **84** may deliver defibrillation shocks to heart **12** via at least two of electrodes **58**, **62**, **64**, **66**. In some examples, therapy delivery module **84** may be configured to deliver pacing, cardioversion, or defibrillation stimulation in the form of electrical pulses. In other examples, therapy delivery module **84** may be configured deliver one or more of these types of stimulation in the form of other signals, such as sine waves, square waves, and/or other substantially continuous time signals.

The IMD **16** may further include a switch module **85** and the control module **81** (e.g., the processor **80**) may use the switch module **85** to select, e.g., via a data/address bus,

which of the available electrodes are used to deliver therapy such as pacing pulses for pacing therapy, or which of the available electrodes are used for sensing. The switch module **85** may include a switch array, switch matrix, multiplexer, or any other type of switching device suitable to selectively couple the sensing module **86** and/or the therapy delivery module **84** to one or more selected electrodes. More specifically, the therapy delivery module **84** may include a plurality of pacing output circuits. Each pacing output circuit of the plurality of pacing output circuits may be selectively coupled, e.g., using the switch module **85**, to one or more of the electrodes **40, 42, 44, 45, 46, 47, 48, 50, 58, 62, 64, 66** (e.g., a pair of electrodes for delivery of therapy to a pacing vector). In other words, each electrode can be selectively coupled to one of the pacing output circuits of the therapy delivery module using the switching module **85**.

The sensing module **86** is coupled (e.g., electrically coupled) to sensing apparatus, which may include, among additional sensing apparatus, the electrodes **40, 42, 44, 45, 46, 47, 48, 50, 58, 62, 64, 66** to monitor electrical activity of the heart **12**, e.g., electrocardiogram (ECG)/electrogram (EGM) signals, etc. The ECG/EGM signals may be used to measure or monitor activation times (e.g., ventricular activations times, etc.), heart rate (HR), heart rate variability (HRV), heart rate turbulence (HRT), deceleration/acceleration capacity, deceleration sequence incidence, T-wave alternans (TWA), P-wave to P-wave intervals (also referred to as the P-P intervals or A-A intervals), R-wave to R-wave intervals (also referred to as the R-R intervals or V-V intervals), P-wave to QRS complex intervals (also referred to as the P-R intervals, A-V intervals, or P-Q intervals), QRS-complex morphology, ST segment (i.e., the segment that connects the QRS complex and the T-wave), T-wave changes, QT intervals, electrical vectors, etc.

The switch module **85** may be also be used with the sensing module **86** to select which of the available electrodes are used, or enabled, to, e.g., sense electrical activity of the patient's heart (e.g., one or more electrical vectors of the patient's heart using any combination of the electrodes **40, 42, 44, 45, 46, 47, 48, 50, 58, 62, 64, 66**). Likewise, the switch module **85** may be also be used with the sensing module **86** to select which of the available electrodes are not to be used (e.g., disabled) to, e.g., sense electrical activity of the patient's heart (e.g., one or more electrical vectors of the patient's heart using any combination of the electrodes **40, 42, 44, 45, 46, 47, 48, 50, 58, 62, 64, 66**), etc. In some examples, the control module **81** may select the electrodes that function as sensing electrodes via the switch module within the sensing module **86**, e.g., by providing signals via a data/address bus.

In some examples, sensing module **86** includes a channel that includes an amplifier with a relatively wider pass band than the R-wave or P-wave amplifiers. Signals from the selected sensing electrodes may be provided to a multiplexer, and thereafter converted to multi-bit digital signals by an analog-to-digital converter for storage in memory **82**, e.g., as an electrogram (EGM). In some examples, the storage of such EGMs in memory **82** may be under the control of a direct memory access circuit.

In some examples, the control module **81** may operate as an interrupt driven device, and may be responsive to interrupts from pacer timing and control module, where the interrupts may correspond to the occurrences of sensed P-waves and R-waves and the generation of cardiac pacing pulses. Any necessary mathematical calculations may be performed by the processor **80** and any updating of the values or intervals controlled by the pacer timing and control

module may take place following such interrupts. A portion of memory **82** may be configured as a plurality of recirculating buffers, capable of holding one or more series of measured intervals, which may be analyzed by, e.g., the processor **80** in response to the occurrence of a pace or sense interrupt to determine whether the patient's heart **12** is presently exhibiting atrial or ventricular tachyarrhythmia.

The telemetry module **88** of the control module **81** may include any suitable hardware, firmware, software, or any combination thereof for communicating with another device, such as a programmer. For example, under the control of the processor **80**, the telemetry module **88** may receive downlink telemetry from and send uplink telemetry to a programmer with the aid of an antenna, which may be internal and/or external. The processor **80** may provide the data to be uplinked to a programmer and the control signals for the telemetry circuit within the telemetry module **88**, e.g., via an address/data bus. In some examples, the telemetry module **88** may provide received data to the processor **80** via a multiplexer.

The various components of the IMD **16** are further coupled to a power source **90**, which may include a rechargeable or non-rechargeable battery. A non-rechargeable battery may be selected to last for several years, while a rechargeable battery may be inductively charged from an external device, e.g., on a daily or weekly basis.

FIG. **11B** is another embodiment of a functional block diagram for IMD **16**. FIG. **11B** depicts bipolar RA lead **22**, bipolar RV lead **18**, and bipolar LV CS lead **20** without the LA CS pace/sense electrodes and coupled with an implantable pulse generator (IPG) circuit **31** having programmable modes and parameters of a bi-ventricular DDD/R type known in the pacing art. In turn, the sensor signal processing circuit **91** indirectly couples to the timing circuit **83** and via data and control bus to microcomputer circuitry **33**. The IPG circuit **31** is illustrated in a functional block diagram divided generally into a microcomputer circuit **33** and a pacing circuit **21**. The pacing circuit **21** includes the digital controller/timer circuit **83**, the output amplifiers circuit **51**, the sense amplifiers circuit **55**, the RF telemetry transceiver **41**, the activity sensor circuit **35** as well as a number of other circuits and components described below.

Crystal oscillator circuit **89** provides the basic timing clock for the pacing circuit **21**, while battery **29** provides power. Power-on-reset circuit **87** responds to initial connection of the circuit to the battery for defining an initial operating condition and similarly, resets the operative state of the device in response to detection of a low battery condition. Reference mode circuit **37** generates stable voltage reference and currents for the analog circuits within the pacing circuit **21**, while analog to digital converter ADC and multiplexer circuit **39** digitizes analog signals and voltage to provide real time telemetry if a cardiac signals from sense amplifiers **55**, for uplink transmission via RF transmitter and receiver circuit **41**. Voltage reference and bias circuit **37**, ADC and multiplexer **39**, power-on-reset circuit **87** and crystal oscillator circuit **89** may correspond to any of those presently used in current marketed implantable cardiac pacemakers.

If the IPG is programmed to a rate responsive mode, the signals output by one or more physiologic sensor are employed as a rate control parameter (RCP) to derive a physiologic escape interval. For example, the escape interval is adjusted proportionally to the patient's activity level developed in the patient activity sensor (PAS) circuit **35** in the depicted, exemplary IPG circuit **31**. The patient activity sensor **27** is coupled to the IPG housing and may take the

form of a piezoelectric crystal transducer as is well known in the art and its output signal is processed and used as the RCP. Sensor **27** generates electrical signals in response to sensed physical activity that are processed by activity circuit **35** and provided to digital controller/timer circuit **83**. Activity circuit **35** and associated sensor **27** may correspond to the circuitry disclosed in U.S. Pat. No. 5,052,388 entitled "METHOD AND APPARATUS FOR IMPLEMENTING ACTIVITY SENSING IN A PULSE GENERATOR" and issued on Oct. 1, 1991 and U.S. Pat. No. 4,428,378 entitled "RATE ADAPTIVE PACER" and issued on Jan. 31, 1984, each of which are incorporated herein by reference in their entirety. Similarly, the exemplary systems, apparatus, and methods described herein may be practiced in conjunction with alternate types of sensors such as oxygenation sensors, pressure sensors, pH sensors and respiration sensors, all well known for use in providing rate responsive pacing capabilities. Alternately, QT time may be used as the rate indicating parameter, in which case no extra sensor is required. Similarly, the exemplary embodiments described herein may also be practiced in non-rate responsive pacemakers.

Data transmission to and from the external programmer is accomplished by way of the telemetry antenna **57** and an associated RF transceiver **41**, which serves both to demodulate received downlink telemetry and to transmit uplink telemetry. Uplink telemetry capabilities will typically include the ability to transmit stored digital information, e.g. operating modes and parameters, EGM histograms, and other events, as well as real time EGMs of atrial and/or ventricular electrical activity and marker channel pulses indicating the occurrence of sensed and paced depolarizations in the atrium and ventricle, as are well known in the pacing art.

Microcomputer **33** contains a microprocessor **80** and associated system clock and on-processor RAM and ROM chips **82A** and **82B**, respectively. In addition, microcomputer circuit **33** includes a separate RAM/ROM chip **82C** to provide additional memory capacity. Microprocessor **80** normally operates in a reduced power consumption mode and is interrupt driven. Microprocessor **80** is awakened in response to defined interrupt events, which may include A-TRIG, RV-TRIG, LV-TRIG signals generated by timers in digital timer/controller circuit **83** and A-EVENT, RV-EVENT, and LV-EVENT signals generated by sense amplifiers circuit **55**, among others. The specific values of the intervals and delays timed out by digital controller/timer circuit **83** are controlled by the microcomputer circuit **33** by way of data and control bus from programmed-in parameter values and operating modes. In addition, if programmed to operate as a rate responsive pacemaker, a timed interrupt, e.g., every cycle or every two seconds, may be provided in order to allow the microprocessor to analyze the activity sensor data and update the basic A-A, V-A, or V-V escape interval, as applicable. In addition, the microprocessor **80** may also serve to define variable, operative AV delay intervals and the energy delivered to each ventricle.

In one embodiment, microprocessor **80** is a custom microprocessor adapted to fetch and execute instructions stored in RAM/ROM unit **82** in a conventional manner. It is contemplated, however, that other implementations may be suitable to practice the present invention. For example, an off-the-shelf, commercially available microprocessor or microcontroller, or custom application-specific, hardwired logic, or state-machine type circuit may perform the functions of microprocessor **80**.

Digital controller/timer circuit **83** operates under the general control of the microcomputer **33** to control timing

and other functions within the pacing circuit **320** and includes a set of timing and associated logic circuits of which certain ones pertinent to the present invention are depicted. The depicted timing circuits include URI/LRI timers **83A**, V-V delay timer **83B**, intrinsic interval timers **83C** for timing elapsed V-EVENT to V-EVENT intervals or V-EVENT to A-EVENT intervals or the V-V conduction interval, escape interval timers **83D** for timing A-A, V-A, and/or V-V pacing escape intervals, an AV delay interval timer **83E** for timing the A-LVp delay (or A-RVp delay) from a preceding A-EVENT or A-TRIG, a post-ventricular timer **83F** for timing post-ventricular time periods, and a date/time clock **83G**.

The AV delay interval timer **83E** is loaded with an appropriate delay interval for one ventricular chamber (e.g., either an A-RVp delay or an A-LVp delay as determined using known methods) to time-out starting from a preceding A-PACE or A-EVENT. The interval timer **83E** triggers pacing stimulus delivery, and can be based on one or more prior cardiac cycles (or from a data set empirically derived for a given patient).

The post-event timer **83F** time out the post-ventricular time period following an RV-EVENT or LV-EVENT or a RV-TRIG or LV-TRIG and post-atrial time periods following an A-EVENT or A-TRIG. The durations of the post-event time periods may also be selected as programmable parameters stored in the microcomputer **33**. The post-ventricular time periods include the PVARP, a post-atrial ventricular blanking period (PAVBP), a ventricular blanking period (VBP), a post-ventricular atrial blanking period (PVARP) and a ventricular refractory period (VRP) although other periods can be suitably defined depending, at least in part, on the operative circuitry employed in the pacing engine. The post-atrial time periods include an atrial refractory period (ARP) during which an A-EVENT is ignored for the purpose of resetting any AV delay, and an atrial blanking period (ABP) during which atrial sensing is disabled. It should be noted that the starting of the post-atrial time periods and the AV delays can be commenced substantially simultaneously with the start or end of each A-EVENT or A-TRIG or, in the latter case, upon the end of the A-PACE which may follow the A-TRIG. Similarly, the starting of the post-ventricular time periods and the V-A escape interval can be commenced substantially simultaneously with the start or end of the V-EVENT or V-TRIG or, in the latter case, upon the end of the V-PACE which may follow the V-TRIG. The microprocessor **80** also optionally calculates AV delays, post-ventricular time periods, and post-atrial time periods that vary with the sensor based escape interval established in response to the RCP(s) and/or with the intrinsic atrial rate.

The output amplifiers circuit **51** contains a RA pace pulse generator (and a LA pace pulse generator if LA pacing is provided), a RV pace pulse generator, and a LV pace pulse generator or corresponding to any of those presently employed in commercially marketed cardiac pacemakers providing atrial and ventricular pacing. In order to trigger generation of an RV-PACE or LV-PACE pulse, digital controller/timer circuit **83** generates the RV-TRIG signal at the time-out of the A-RVp delay (in the case of RV pre-excitation) or the LV-TRIG at the time-out of the A-LVp delay (in the case of LV pre-excitation) provided by AV delay interval timer **83E** (or the V-V delay timer **83B**). Similarly, digital controller/timer circuit **83** generates an RA-TRIG signal that triggers output of an RA-PACE pulse (or an LA-TRIG signal that triggers output of an LA-PACE pulse, if provided) at the end of the V-A escape interval timed by escape interval timers **83D**.

The output amplifiers circuit **51** includes switching circuits for coupling selected pace electrode pairs from among the lead conductors and the IND CAN electrode **20** to the RA pace pulse generator (and LA pace pulse generator if provided), RV pace pulse generator and LV pace pulse generator. Pace/sense electrode pair selection and control circuit **53** selects lead conductors and associated pace electrode pairs to be coupled with the atrial and ventricular output amplifiers within output amplifiers circuit **51** for accomplishing RA, LA, RV and LV pacing.

The sense amplifiers circuit **55** contains sense amplifiers corresponding to any of those presently employed in contemporary cardiac pacemakers for atrial and ventricular pacing and sensing. High impedance P-wave and R-wave sense amplifiers may be used to amplify a voltage difference signal that is generated across the sense electrode pairs by the passage of cardiac depolarization wavefronts. The high impedance sense amplifiers use high gain to amplify the low amplitude signals and rely on pass band filters, time domain filtering and amplitude threshold comparison to discriminate a P-wave or R-wave from background electrical noise. Digital controller/timer circuit **83** controls sensitivity settings of the atrial and ventricular sense amplifiers **55**.

The sense amplifiers are typically uncoupled from the sense electrodes during the blanking periods before, during, and after delivery of a pace pulse to any of the pace electrodes of the pacing system to avoid saturation of the sense amplifiers. The sense amplifiers circuit **55** includes blanking circuits for uncoupling the selected pairs of the lead conductors and the IND-CAN electrode **20** from the inputs of the RA sense amplifier (and LA sense amplifier if provided), RV sense amplifier and LV sense amplifier during the ABP, PVABP and VBP. The sense amplifiers circuit **55** also includes switching circuits for coupling selected sense electrode lead conductors and the IND-CAN electrode **20** to the RA sense amplifier (and LA sense amplifier if provided), RV sense amplifier and LV sense amplifier. Again, sense electrode selection and control circuit **53** selects conductors and associated sense electrode pairs to be coupled with the atrial and ventricular sense amplifiers within the output amplifiers circuit **51** and sense amplifiers circuit **55** for accomplishing RA, LA, RV and LV sensing along desired unipolar and bipolar sensing vectors.

Right atrial depolarizations or P-waves in the RA-SENSE signal that are sensed by the RA sense amplifier result in a RA-EVENT signal that is communicated to the digital controller/timer circuit **83**. Similarly, left atrial depolarizations or P-waves in the LA-SENSE signal that are sensed by the LA sense amplifier, if provided, result in a LA-EVENT signal that is communicated to the digital controller/timer circuit **83**. Ventricular depolarizations or R-waves in the RV-SENSE signal are sensed by a ventricular sense amplifier result in an RV-EVENT signal that is communicated to the digital controller/timer circuit **83**. Similarly, ventricular depolarizations or R-waves in the LV-SENSE signal are sensed by a ventricular sense amplifier result in an LV-EVENT signal that is communicated to the digital controller/timer circuit **83**. The RV-EVENT, LV-EVENT, and RA-EVENT, LA-SENSE signals may be refractory or non-refractory, and can inadvertently be triggered by electrical noise signals or aberrantly conducted depolarization waves rather than true R-waves or P-waves.

The techniques described in this disclosure, including those attributed to the IMD **16**, the computing apparatus **140**, and/or various constituent components, may be implemented, at least in part, in hardware, software, firmware, or any combination thereof. For example, various aspects of

the techniques may be implemented within one or more processors, including one or more microprocessors, DSPs, ASICs, FPGAs, or any other equivalent integrated or discrete logic circuitry, as well as any combinations of such components, embodied in programmers, such as physician or patient programmers, stimulators, image processing devices, or other devices. The term "module," "processor," or "processing circuitry" may generally refer to any of the foregoing logic circuitry, alone or in combination with other logic circuitry, or any other equivalent circuitry.

Such hardware, software, and/or firmware may be implemented within the same device or within separate devices to support the various operations and functions described in this disclosure. In addition, any of the described units, modules, or components may be implemented together or separately as discrete but interoperable logic devices. Depiction of different features as modules or units is intended to highlight different functional aspects and does not necessarily imply that such modules or units must be realized by separate hardware or software components. Rather, functionality associated with one or more modules or units may be performed by separate hardware or software components, or integrated within common or separate hardware or software components.

When implemented in software, the functionality ascribed to the systems, devices and techniques described in this disclosure may be embodied as instructions on a computer-readable medium such as RAM, ROM, NVRAM, EEPROM, FLASH memory, magnetic data storage media, optical data storage media, or the like. The instructions may be executed by one or more processors to support one or more aspects of the functionality described in this disclosure.

This disclosure has been provided with reference to illustrative embodiments and is not meant to be construed in a limiting sense. As described previously, one skilled in the art will recognize that other various illustrative applications may use the techniques as described herein to take advantage of the beneficial characteristics of the apparatus and methods described herein. Various modifications of the illustrative embodiments, as well as additional embodiments of the disclosure, will be apparent upon reference to this description.

What is claimed:

1. A system for assisting in noninvasive evaluation of cardiac therapy comprising:

electrode apparatus comprising a plurality of external surface electrodes positioned in an array, wherein the electrode apparatus is configured to locate the plurality of external surface electrodes proximate the skin of the patient;

a display apparatus comprising a graphical user interface configured to present cardiac electrical activation time information and cardiac health information; and

computing apparatus coupled to the electrode apparatus and the display apparatus and configured to provide the graphical user interface displayed on the display apparatus to assist a user in noninvasively evaluating cardiac therapy, wherein the computing apparatus is further configured to:

noninvasively measure surrogate cardiac electrical activation times using one or more external surface electrodes of the plurality of external surface electrodes of the electrode apparatus proximate the patient's heart, wherein the surrogate cardiac elec-

trical activation times are representative of depolarization of cardiac tissue that propagates through the torso of the patient,

noninvasively measure at least one metric of the patient's cardiac health using one or more external surface electrodes of the plurality of external surface electrodes of the electrode apparatus proximate the patient's heart,

determine whether cardiac therapy for the patient is beneficial based on a comparison of presently-measured noninvasive measurements made using the one or more external surface electrodes of the plurality of external surface electrodes and previously-measured noninvasive measurements made using the one or more external surface electrodes of the plurality of external surface electrodes of the electrode apparatus, and

simultaneously display, on the graphical user interface, an indication of whether cardiac therapy for the patient is beneficial based on the comparison of presently-measured noninvasive measurements and previously-measured noninvasive measurements,

a graphical depiction of a portion of a first human anatomy and presently-measured surrogate cardiac electrical activation times about the first human anatomy portion that were noninvasively measured using the one or more external surface electrodes of the plurality of external surface electrodes proximate the skin of the patient,

a graphical depiction of a portion of a second human anatomy and previously-measured surrogate cardiac electrical activation times about the second human anatomy portion that were noninvasively measured using the one or more external surface electrodes of the plurality of external surface electrodes proximate the skin of the patient,

the at least one presently-measured metric of the patient's cardiac health that was noninvasively measured using the one or more external surface electrodes of the plurality of external surface electrodes proximate the skin of the patient, and

the at least one previously-measured metric of the patient's cardiac health that was noninvasively measured using the one or more external surface electrodes of the plurality of external surface electrodes proximate the skin of the patient.

2. The system of claim 1, wherein the indication of whether cardiac therapy for the patient is beneficial comprises a percentage improvement between the at least one previously-measured metric of the patient's cardiac health and the at least one presently-measured metric of the patient's cardiac health.

3. The system of claim 1, wherein the computing apparatus is further configured to allow the user to change at least one pacing parameter of an implantable medical device providing cardiac therapy to the patient.

4. The system of claim 3, wherein the at least one pacing parameter comprises at least one of a pacing timing interval, a pacing vector, and a pacing mode.

5. The system of claim 1, wherein the computing apparatus is further configured to display, on the graphical user interface, mechanical motion information of one or more regions of at least a portion of blood vessel anatomy of the patient's heart.

6. The system of claim 1, wherein the first human anatomy portion and the second human anatomy portion depict the same human anatomy.

7. The system of claim 1, wherein displaying graphical depictions of the measured surrogate cardiac electrical activation times about the first and second human anatomy portions comprises displaying graphical depictions of the measured surrogate cardiac electrical activation times about graphical depictions of at least one of an anterior side of a human torso and a posterior side of a human torso.

8. The system of claim 1, wherein displaying graphical depictions of the measured surrogate cardiac electrical activation times about the first and second human anatomy portions comprises displaying graphical depictions of the measured surrogate cardiac electrical activation times about graphical depictions of at least one of an anterior side of a heart and a posterior side of a heart.

9. The system of claim 1, wherein the at least one metric comprises QRS width and electrical dyssynchrony.

10. The system of claim 1, wherein displaying the graphical depictions of the measured surrogate cardiac electrical activation times about a portion of human anatomy comprises color scaling the portions of human anatomy on the graphical user interface according to the measured surrogate cardiac electrical activation times.

11. The system of claim 1, wherein the computing apparatus is further configured to display, on the graphical user interface, at least one electrocardiogram of the patient, wherein each of the at least one electrocardiogram are captured using at least one different electrode, and wherein the at least one electrocardiogram is time-aligned on the graphical user interface.

12. A method for assisting in noninvasive evaluation of cardiac therapy comprising:

noninvasively measuring surrogate cardiac electrical activation times using one or more external surface electrodes proximate the skin of the patient and proximate the patient's heart, wherein the surrogate cardiac electrical activation times are representative of depolarization of cardiac tissue that propagates through the torso of the patient;

noninvasively measuring at least one metric of the patient's cardiac health using one or more external surface electrodes proximate the skin of the patient and proximate the patient's heart;

determining whether cardiac therapy for the patient is beneficial based on a comparison of presently-measured noninvasive measurements made using the one or more external surface electrodes proximate the skin of the patient and previously-measured noninvasive measurements made using the one or more external surface electrodes proximate the skin of the patient; and

simultaneously displaying, on a graphical user interface, an indication of whether cardiac therapy for the patient is beneficial based on the comparison,

a graphical depiction of a portion of a first human anatomy and the presently-measured surrogate cardiac electrical activation times about the first human anatomy portion that were noninvasively measured using the one or more external surface electrodes proximate the skin of the patient,

a graphical depiction of a portion of a second human anatomy and the previously-measured surrogate cardiac electrical activation times about the second human anatomy portion that were noninvasively measured using the one or more external surface electrodes proximate the skin of the patient,

the at least one presently-measured metric of the patient's cardiac health that was noninvasively measured using the one or more external surface electrodes proximate the skin of the patient, and

the at least one previously-measured metric of the patient's cardiac health that was noninvasively measured using the one or more external surface electrodes proximate the skin of the patient.

13. The method of claim 12, wherein the indication of whether cardiac therapy for the patient is beneficial comprises a percentage improvement between the at least one previously-measured metric of the patient's cardiac health and the at least one presently-measured metric of the patient's cardiac health.

14. The method of claim 12, wherein the method further comprises allowing the user to change at least one pacing parameter of an implantable medical device providing cardiac therapy to the patient.

15. The method of claim 12, wherein the method further comprises displaying, on the graphical user interface, mechanical motion information of one or more regions of at least a portion of blood vessel anatomy of the patient's heart.

16. The method of claim 12, wherein displaying graphical depictions of the measured surrogate cardiac electrical activation times about the first and second human anatomy portions comprises displaying graphical depictions of the measured surrogate cardiac electrical activation times about graphical depictions of at least one of an anterior side of a human torso, a posterior side of a human torso, an anterior side of a heart, and a posterior side of a heart.

17. The method of claim 12, wherein the at least one metric comprises QRS width and electrical dyssynchrony.

18. The method of claim 12, wherein displaying the graphical depictions of the measured surrogate cardiac electrical activation times about a portion of human anatomy comprises color scaling the portions of human anatomy on the graphical user interface according to the measured surrogate cardiac electrical activation times.

19. The method of claim 12, wherein the method further comprises displaying, on the graphical user interface, at least one electrocardiogram of the patient, wherein each of the at least one electrocardiogram are captured using at least one different electrode, and wherein the at least one electrocardiogram is time-aligned on the graphical user interface.

20. A system for assisting in noninvasive evaluation of a patient for cardiac therapy comprising:

means for noninvasively measuring surrogate cardiac electrical activation times representative of electrical activation times of a patient's heart from the patient's skin;

means for noninvasively measuring at least one metric of the patient's cardiac health from the patient's skin, wherein the surrogate cardiac electrical activation times are representative of depolarization of cardiac tissue that propagates through the torso of the patient;

means for determining whether cardiac therapy for the patient is beneficial based on a comparison of presently-measured noninvasive measurements made and previously-measured noninvasive measurements that were noninvasively measured from the patient's skin; and

display means for simultaneously displaying, on a graphical user interface,

a graphical depiction of a portion of a first human anatomy and presently-measured surrogate cardiac electrical activation times about the first human anatomy portion that were noninvasively measured from the patient's skin,

a graphical depiction of a portion of a second human anatomy and previously-measured surrogate cardiac electrical activation times about the second human anatomy portion that were noninvasively measured from the patient's skin,

the at least one presently-measured metric of the patient's cardiac health that was noninvasively measured from the patient's skin,

the at least one previously-measured metric of the patient's cardiac health that was noninvasively measured from the patient's skin, and

an indication of whether cardiac therapy for the patient is beneficial based on the comparison.

21. The system of claim 1, wherein the indication of whether cardiac therapy for the patient is beneficial comprises one or more of an indication that a location of an implantable lead delivering cardiac therapy is providing beneficial cardiac therapy to the patient, an indication that one or more settings of the cardiac therapy being delivered to the patient are beneficial, and an indication that one or more adjustments to the cardiac therapy are beneficial to the patient.

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专利名称(译)	无创心脏疗法评估		
公开(公告)号	US9993172	公开(公告)日	2018-06-12
申请号	US14/228024	申请日	2014-03-27
[标]申请(专利权)人(译)	美敦力公司		
申请(专利权)人(译)	美敦力公司, INC.		
当前申请(专利权)人(译)	美敦力公司, INC.		
[标]发明人	GILLBERG JEFFREY GHOSH SUBHAM		
发明人	GILLBERG, JEFFREY GHOSH, SUBHAM		
IPC分类号	A61B5/044 G06F19/00 A61B5/0402 A61B5/00 A61N1/37 A61B5/0452 A61N1/368 A61B5/0408 A61N1/365 A61B5/11 A61N1/362 A61B5/0472		
CPC分类号	A61B5/0408 A61B5/044 A61B5/0452 A61B5/04085 A61B5/743 A61N1/368 A61N1/3702 G16H40/63 G16H50/30 G16H50/70 A61B5/0402 A61N1/36507 A61B5/0472 A61B5/1107 A61B5/6823 A61N1/3627 G16H10/60 G06F19/00 G06F19/3481 G16H20/40		
优先权	61/913743 2013-12-09 US		
其他公开文献	US20150157232A1		
外部链接	Espacenet USPTO		

摘要(译)

本文描述了系统, 方法和界面, 用于帮助用户无创评估患者的心脏治疗以及正在递送的心脏治疗的非侵入性评估。系统, 方法和界面可以提供关于人体解剖结构的一个或多个部分以及一个或多个心脏健康度量的心脏电激活时间的图形表示。

