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Cao et al.

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(54) **AUTOMATIC THRESHOLDS FOR ATRIAL TACHYARRHYTHMIA DETECTION IN AN IMPLANTABLE MEDICAL DEVICE**

(58) **Field of Classification Search**
None
See application file for complete search history.

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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 264 days.

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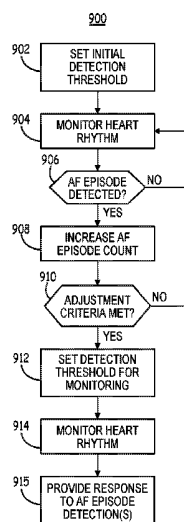
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CPC **A61B 5/0464** (2013.01); **A61B 5/002** (2013.01); **A61B 5/0006** (2013.01); **A61B 5/02405** (2013.01); **A61B 5/044** (2013.01); **A61B 5/04012** (2013.01); **A61B 5/0422** (2013.01); **A61B 5/0432** (2013.01); **A61B 5/0456** (2013.01); **A61B 5/0472** (2013.01);
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(57) **ABSTRACT**

A system for detecting an atrial tachyarrhythmia episode includes a medical device having sensing circuitry configured to receive a cardiac electrical signal from electrodes coupled to the medical device and a processor configured to detect an atrial tachyarrhythmia episode in response to a time duration of the cardiac electrical signal classified as an atrial tachyarrhythmia being greater than or equal to a first detection threshold. The processor is configured to determine if detection threshold adjustment criteria are met based on at least the detected first atrial tachyarrhythmia episode and adjust the first detection threshold to a second detection threshold different than the first detection threshold in response to the detection threshold adjustment criteria being met.

35 Claims, 17 Drawing Sheets



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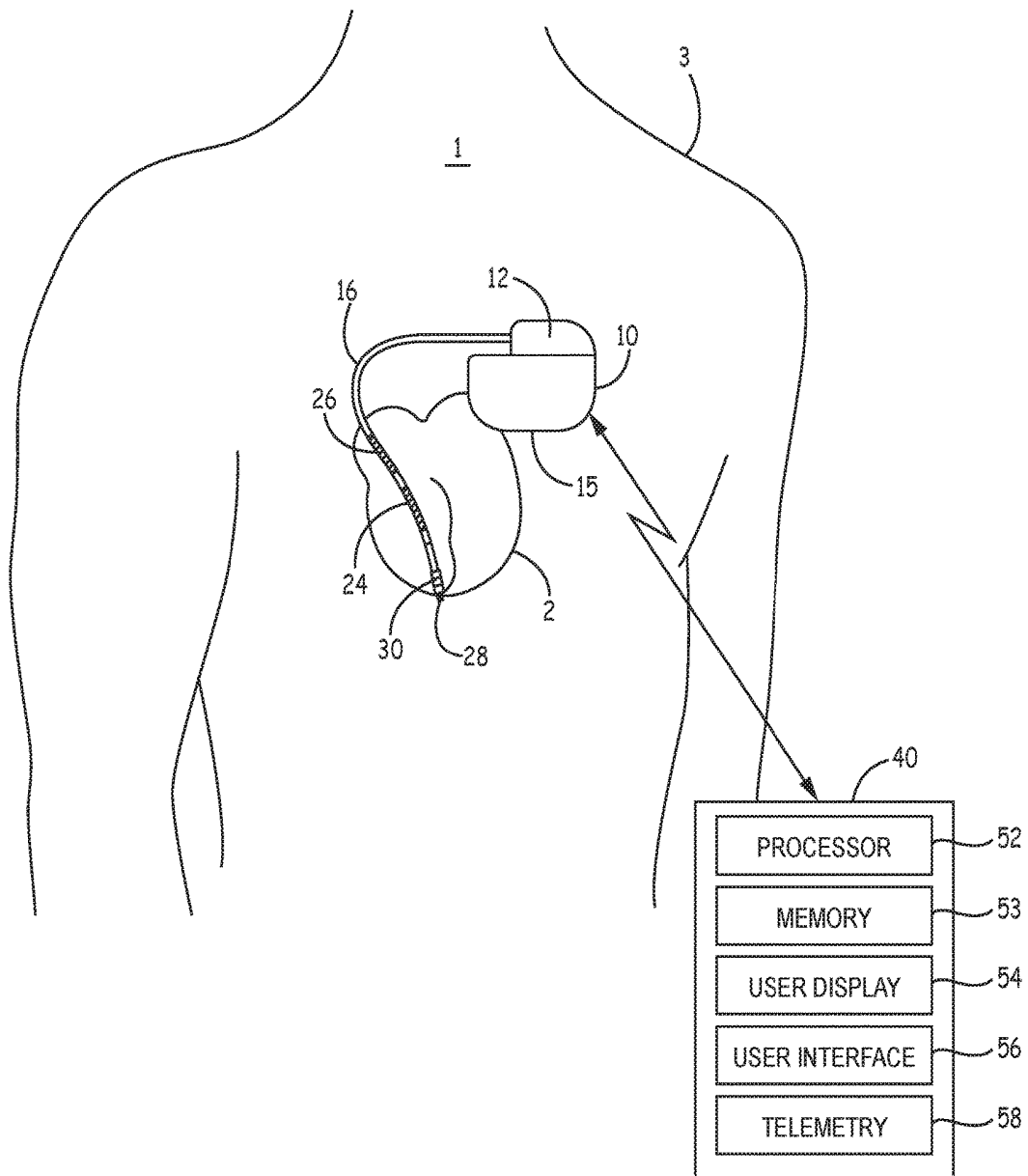


FIG. 1

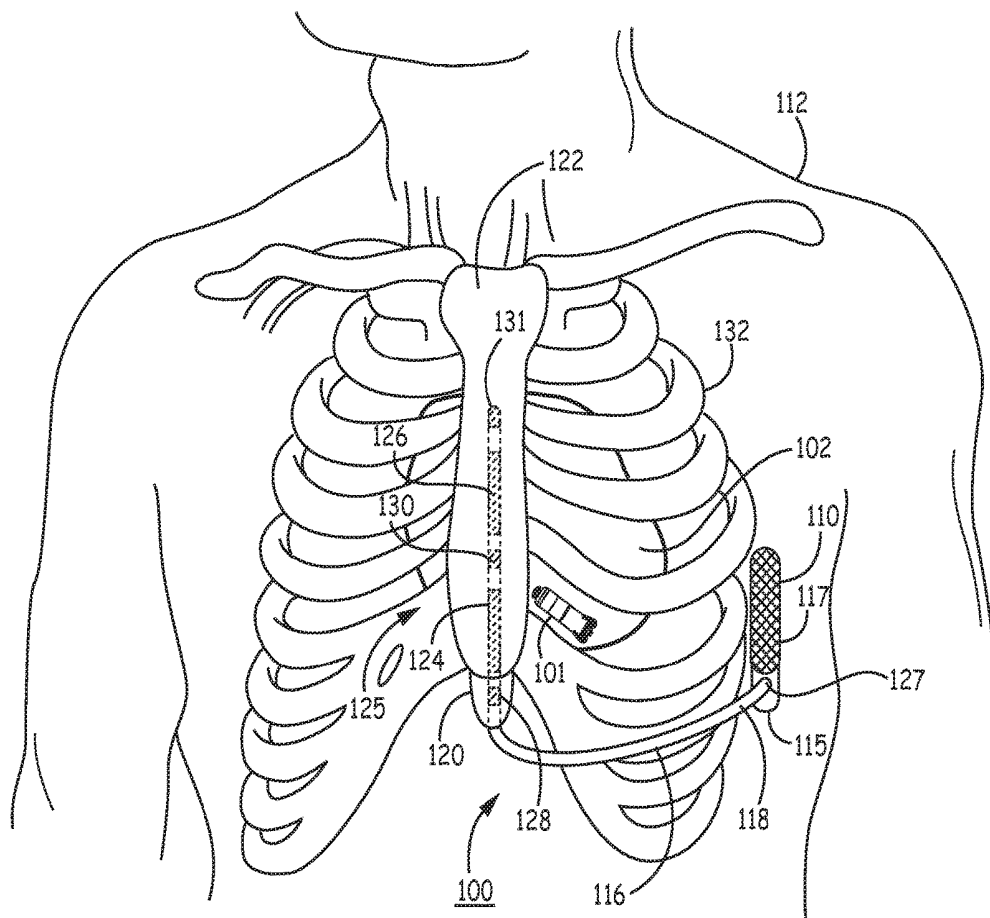


FIG. 2A

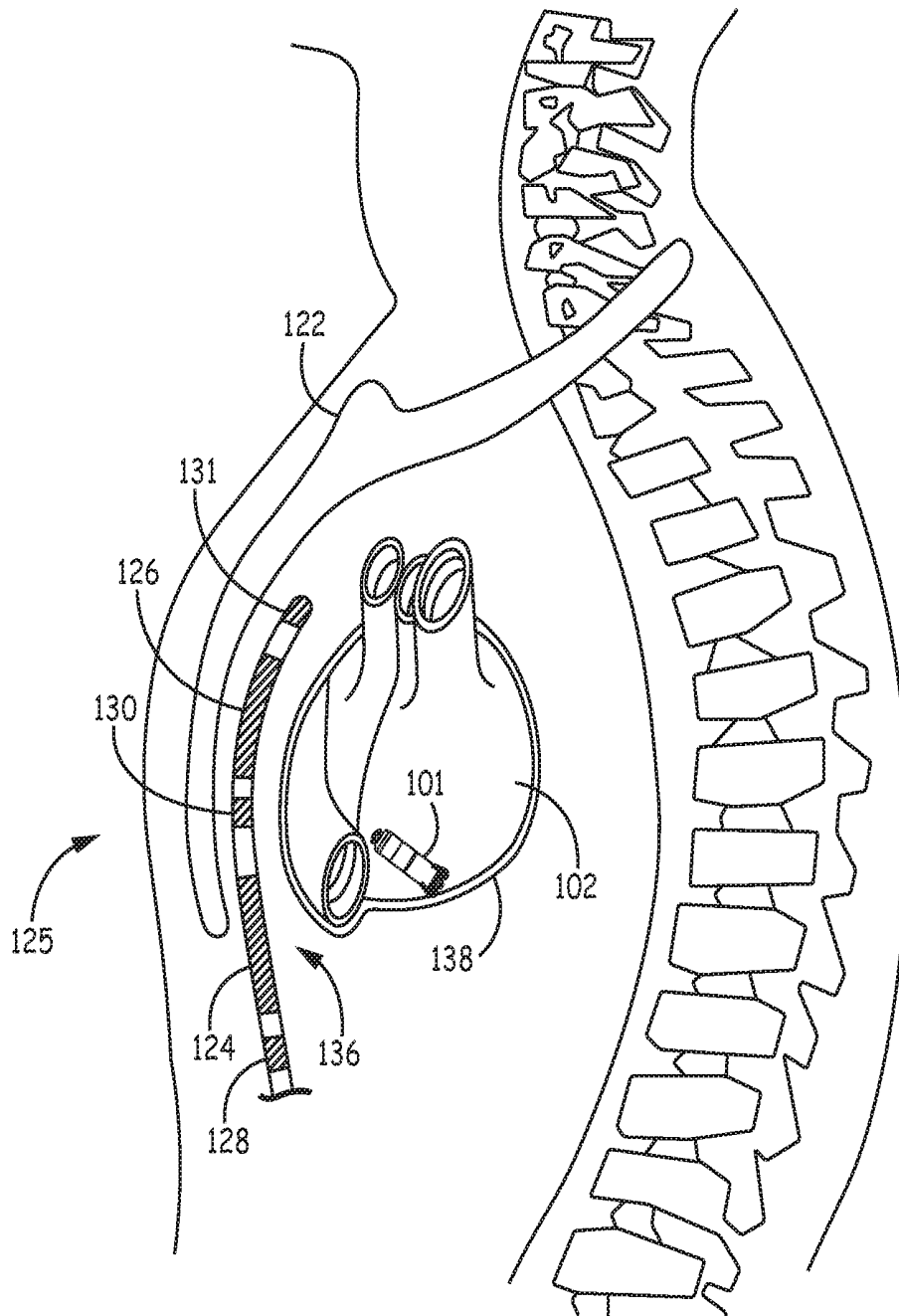


FIG. 2B

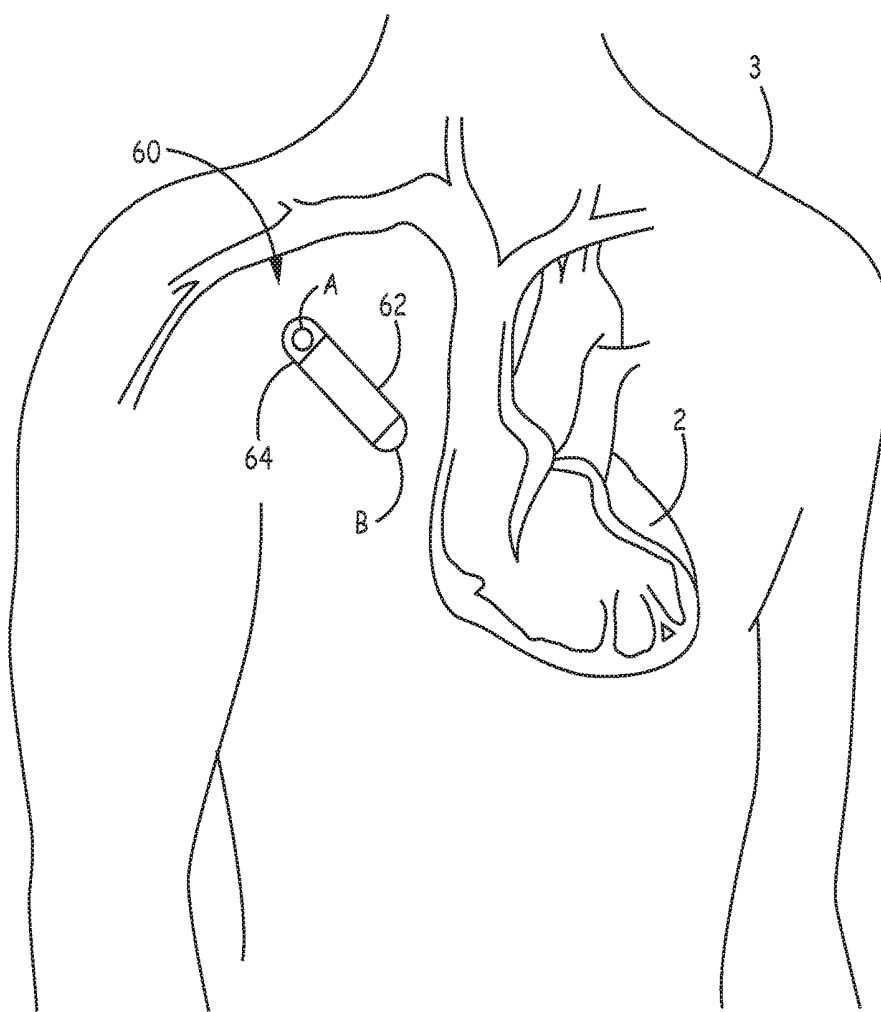


FIG. 3

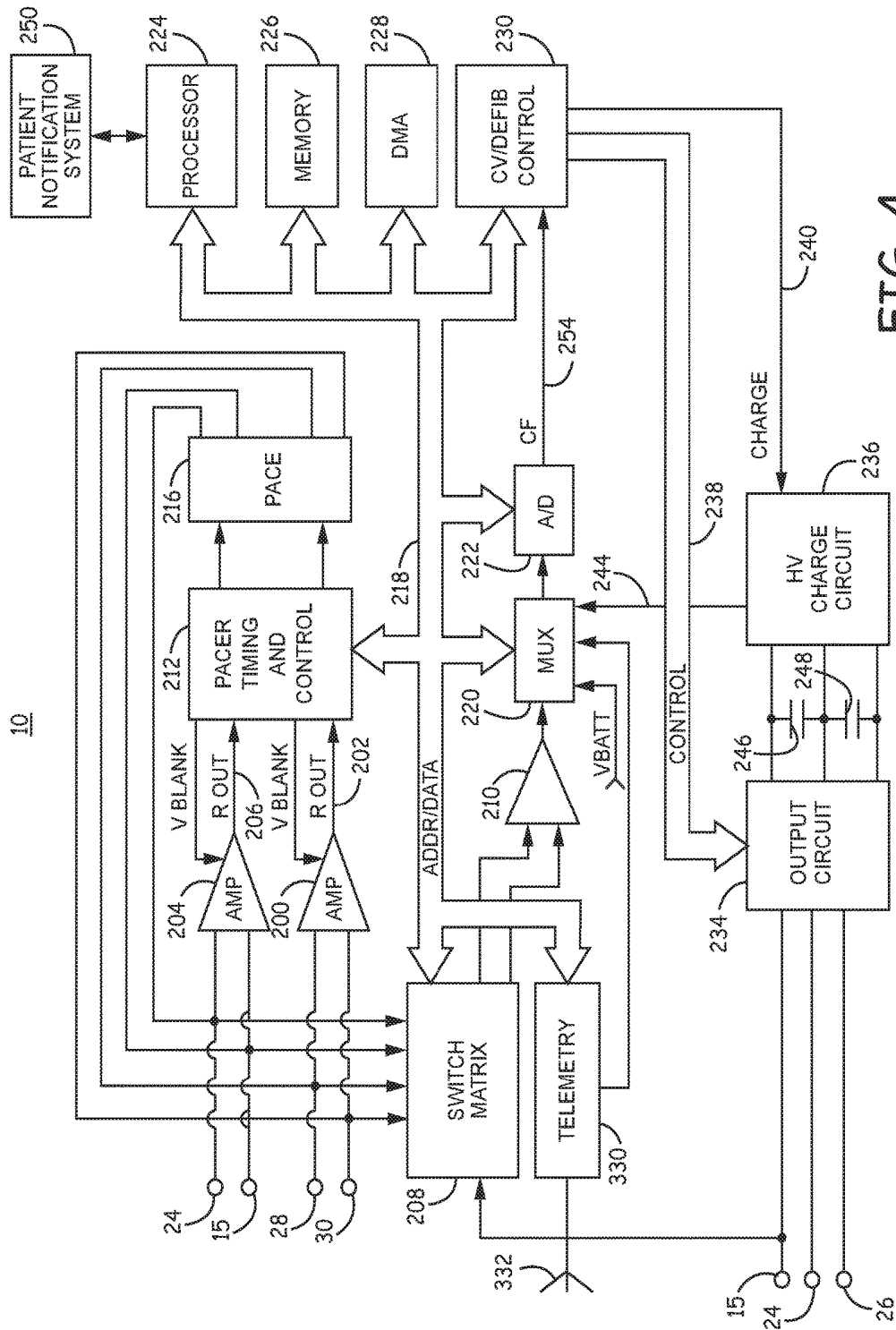


FIG. 4

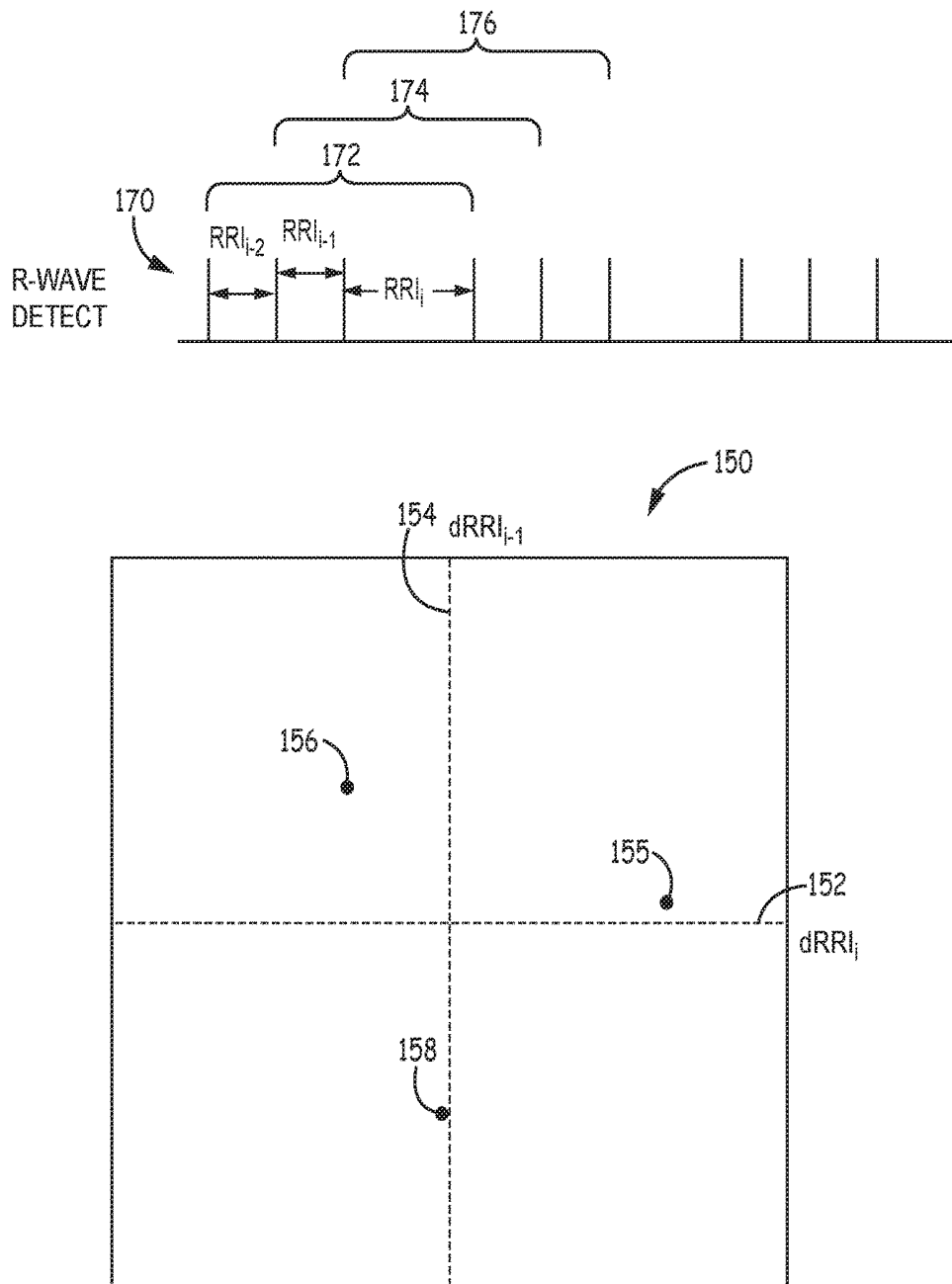


FIG. 5

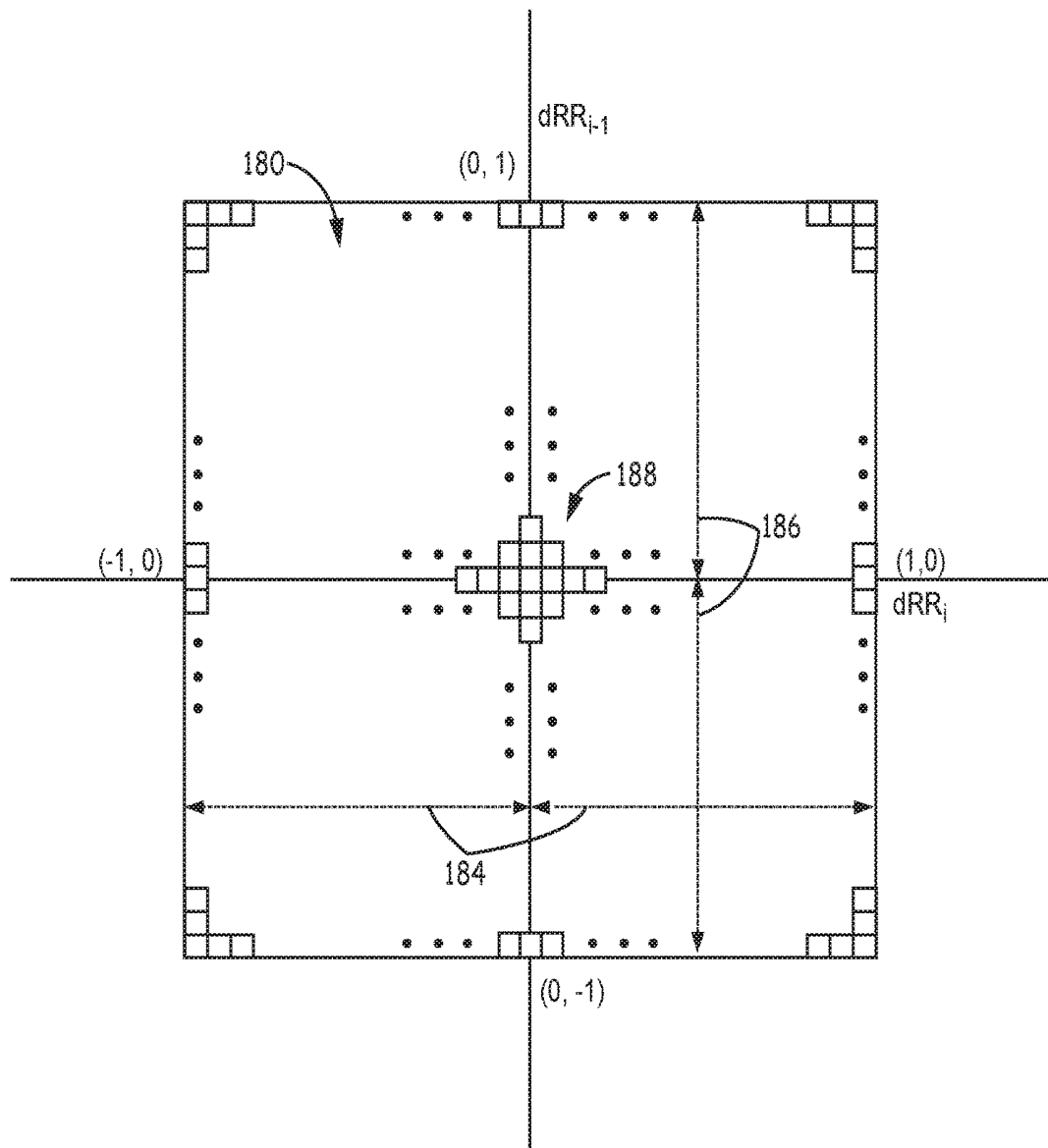


FIG. 6

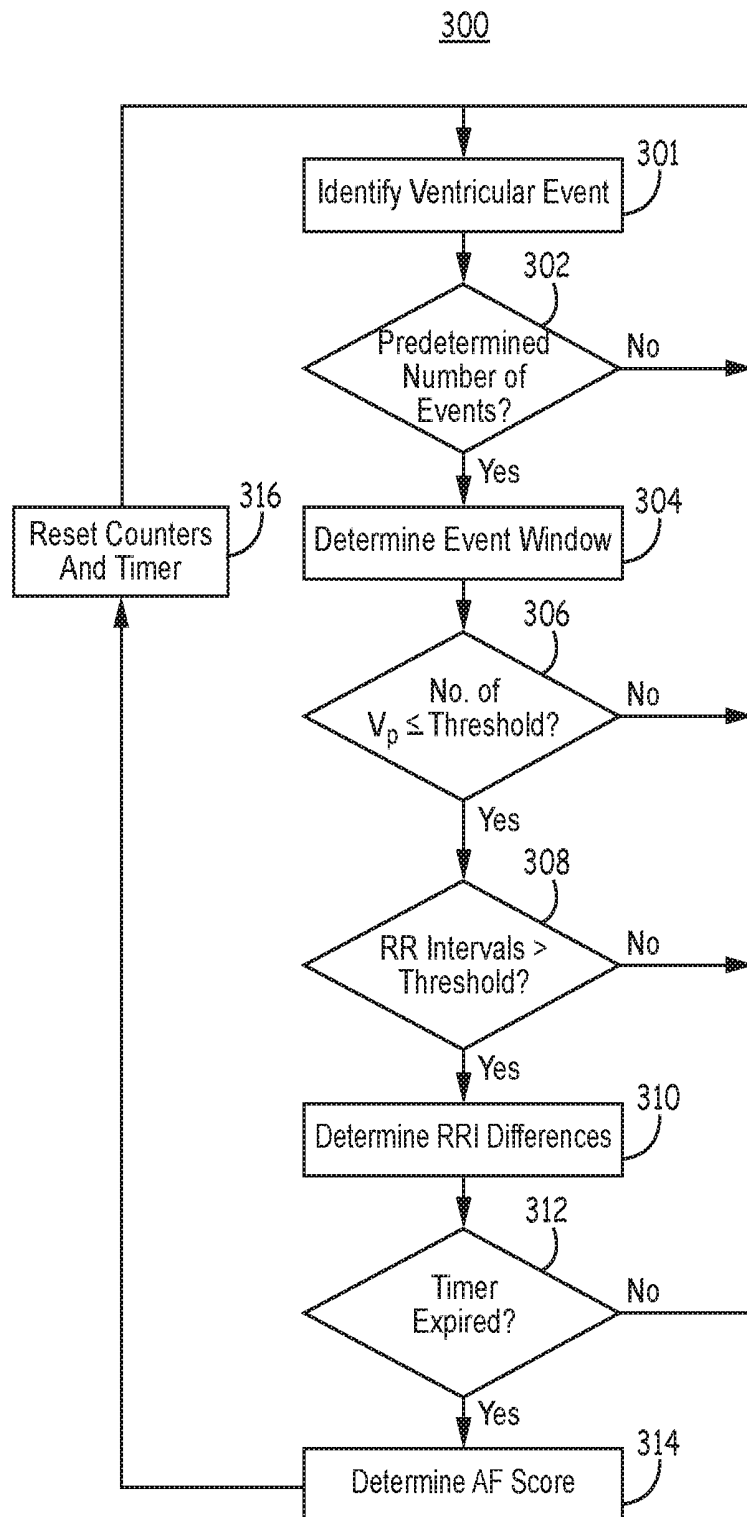


FIG. 7

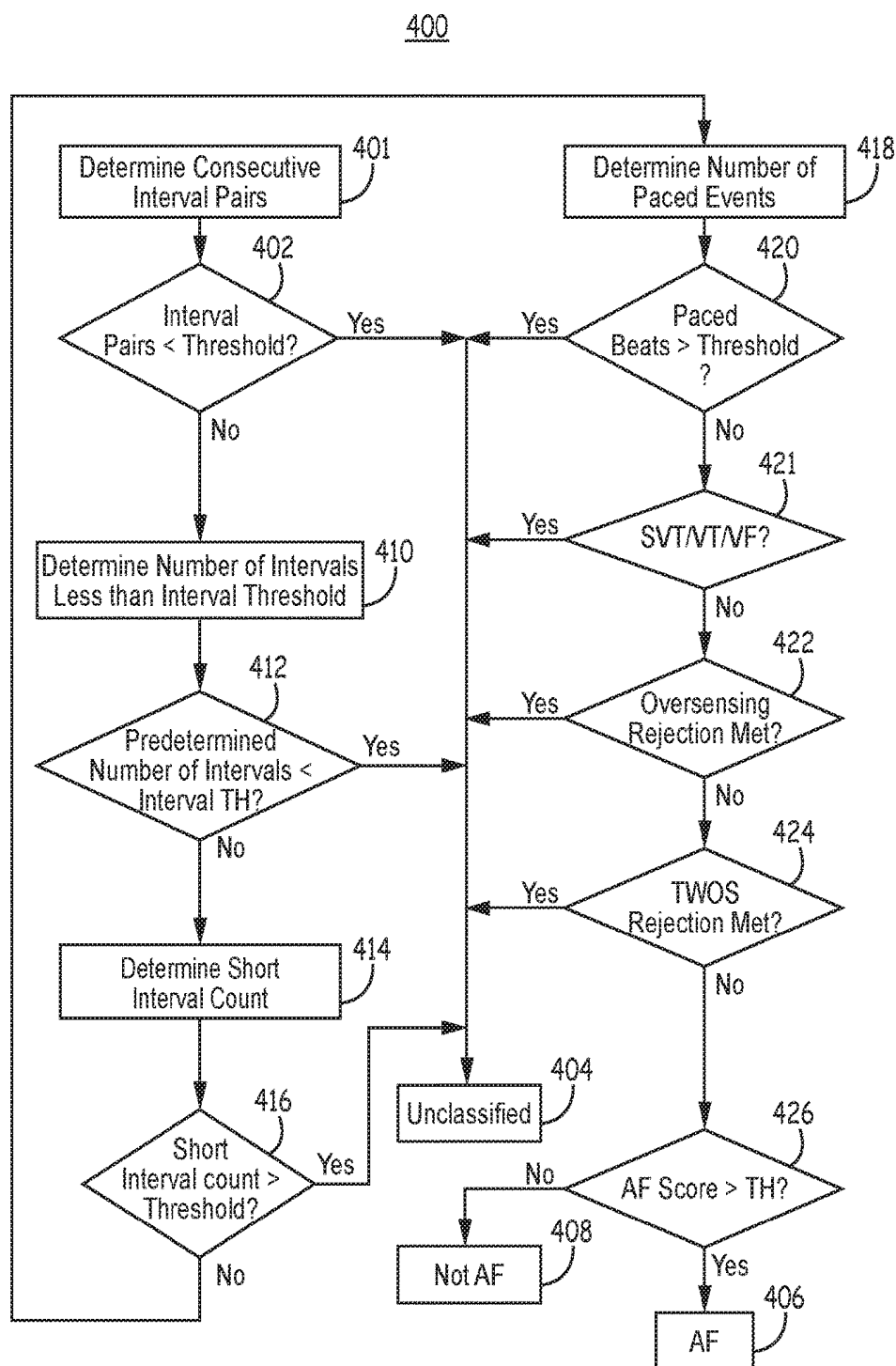


FIG. 8

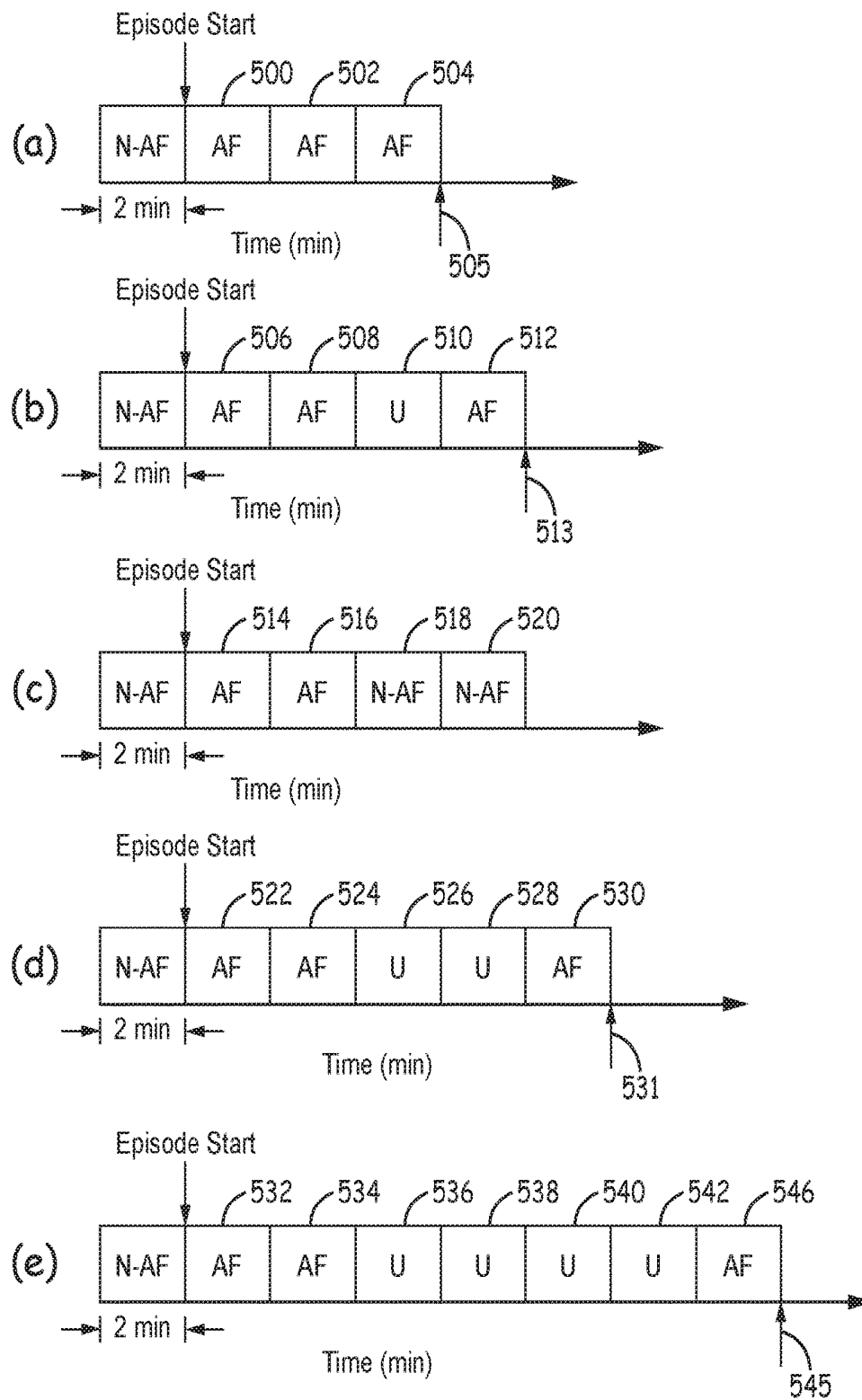


FIG. 9

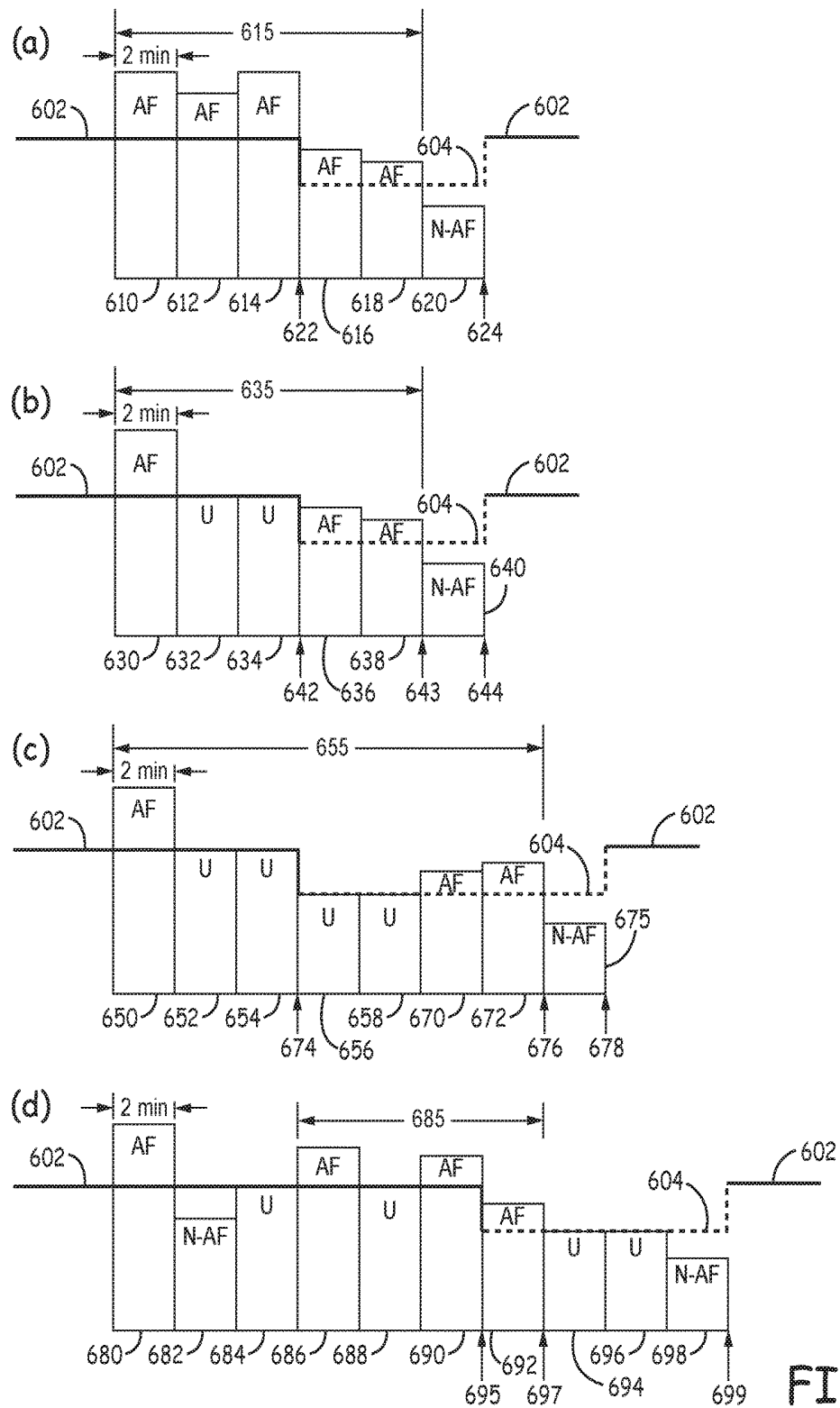


FIG. 10

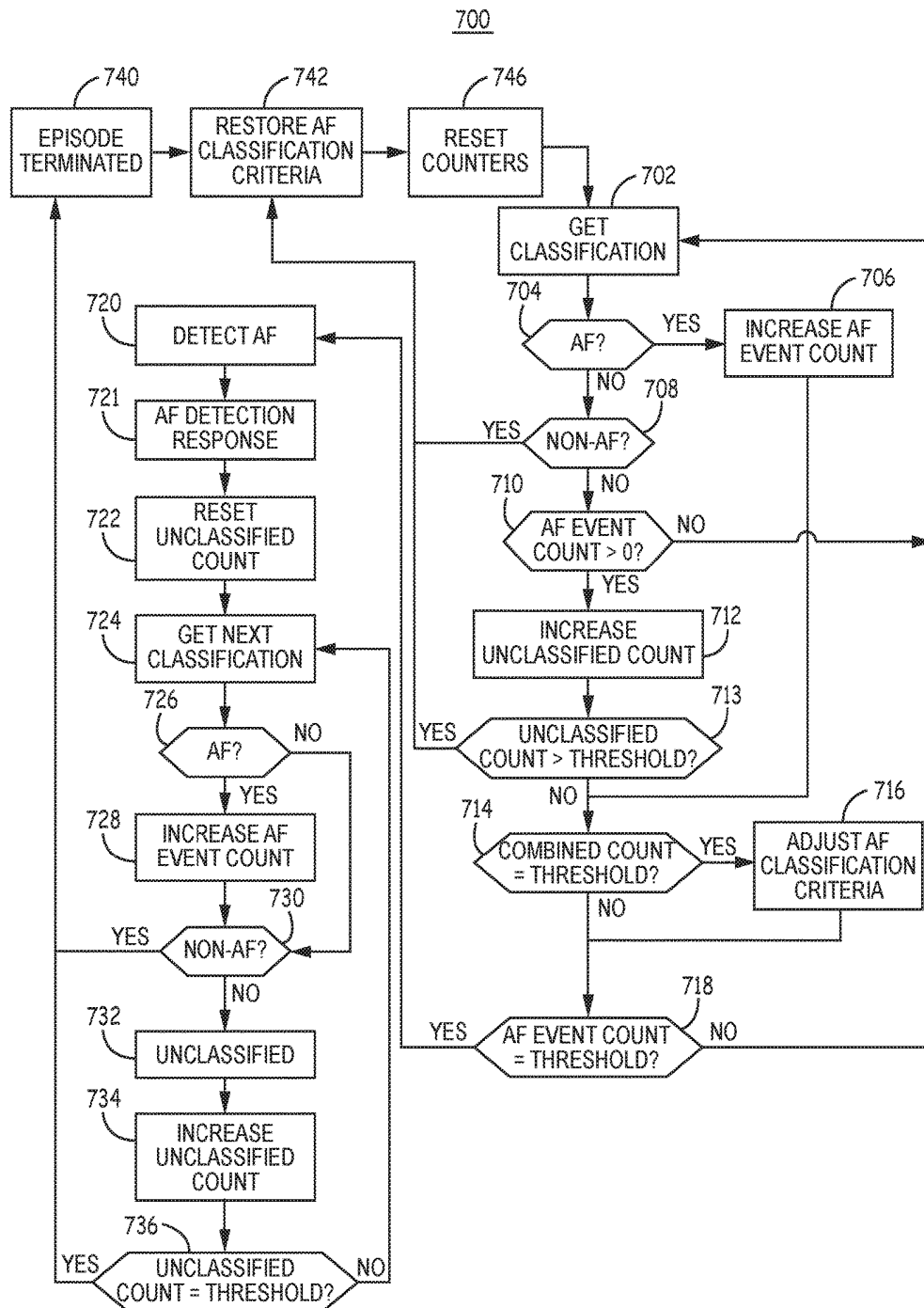


FIG. 11

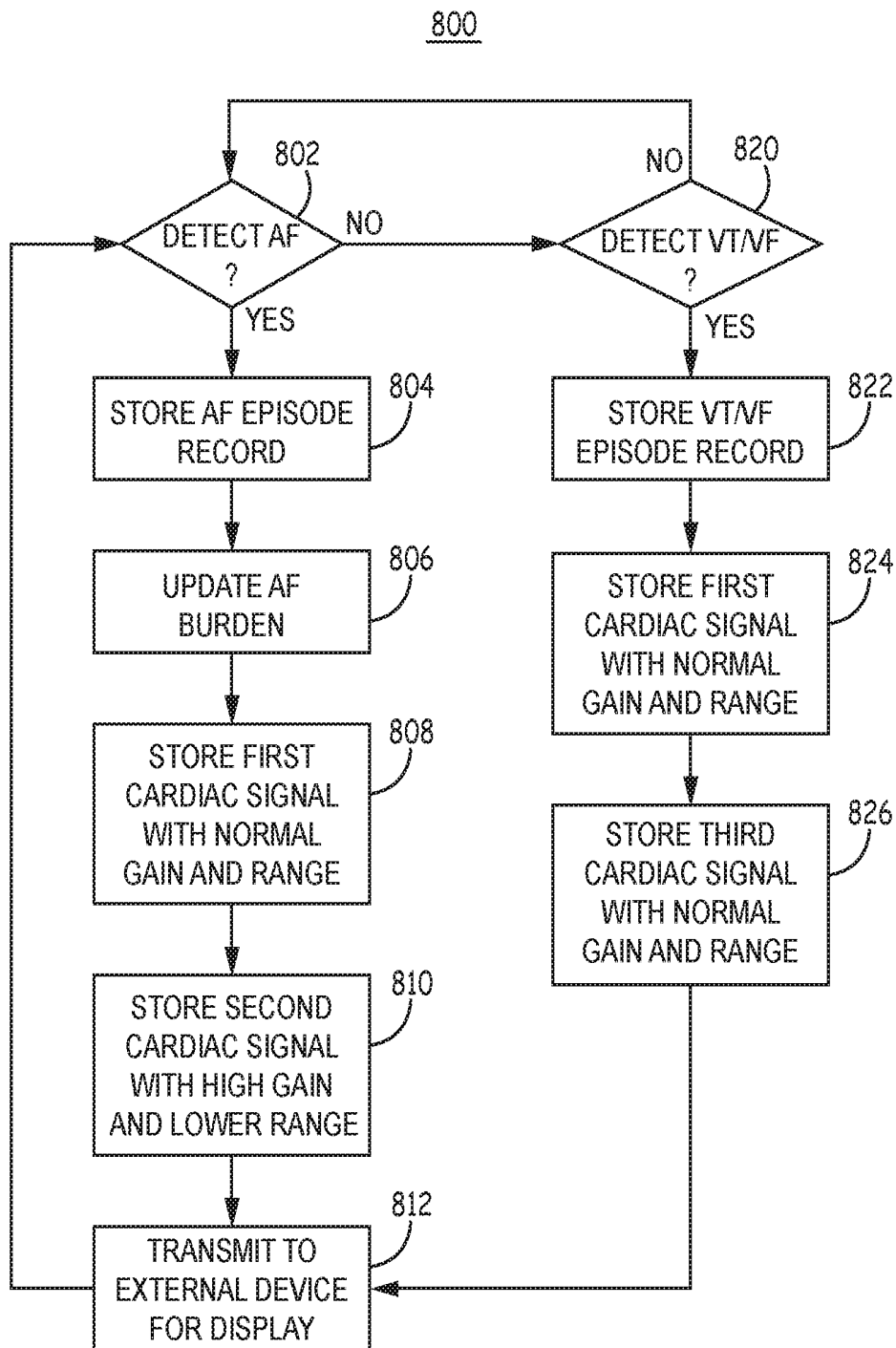


FIG. 12

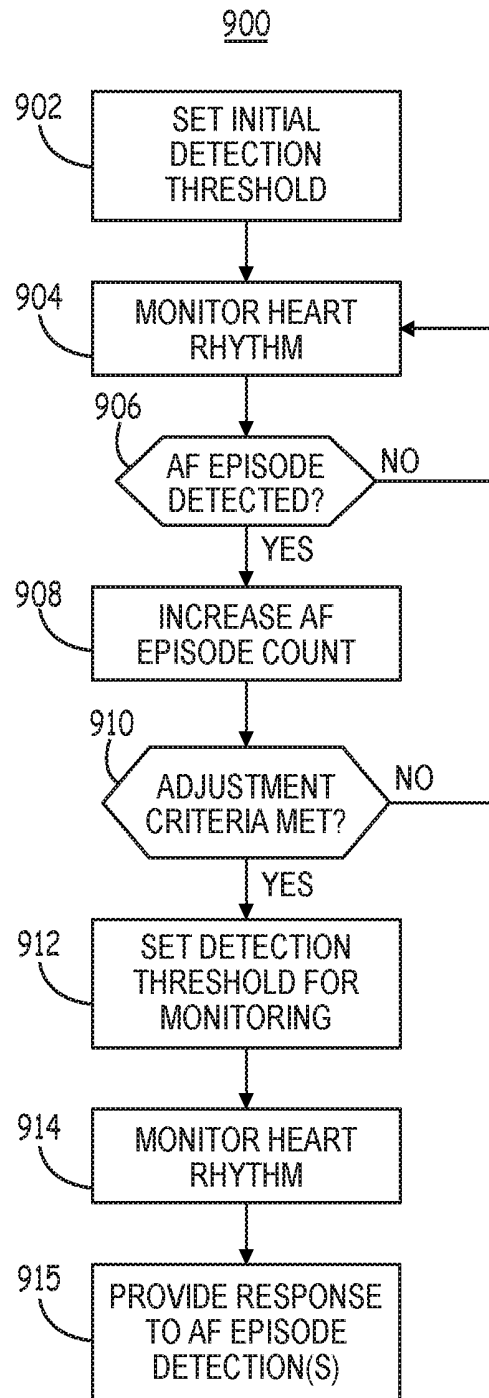


FIG. 13

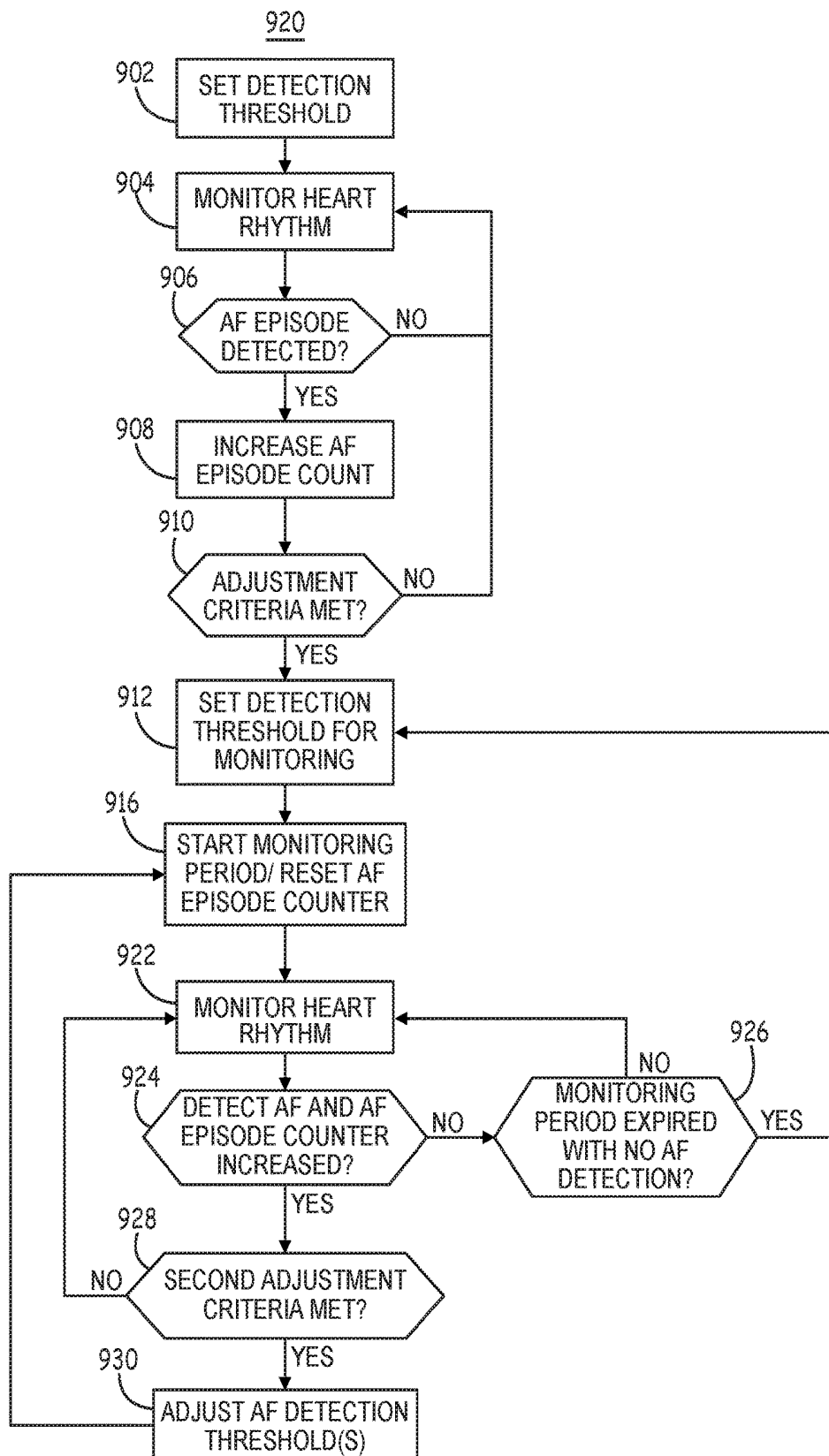


FIG. 14

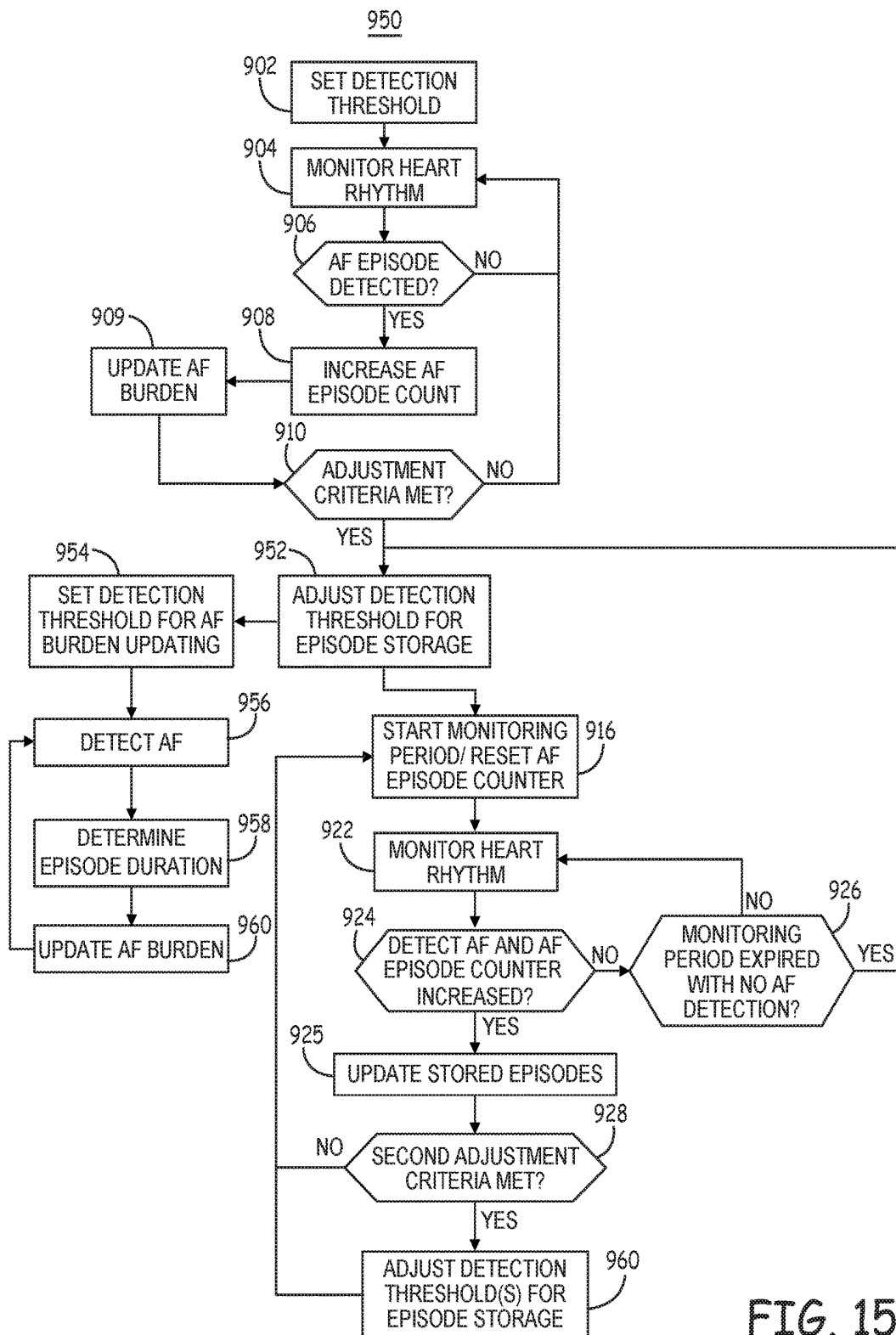


FIG. 15

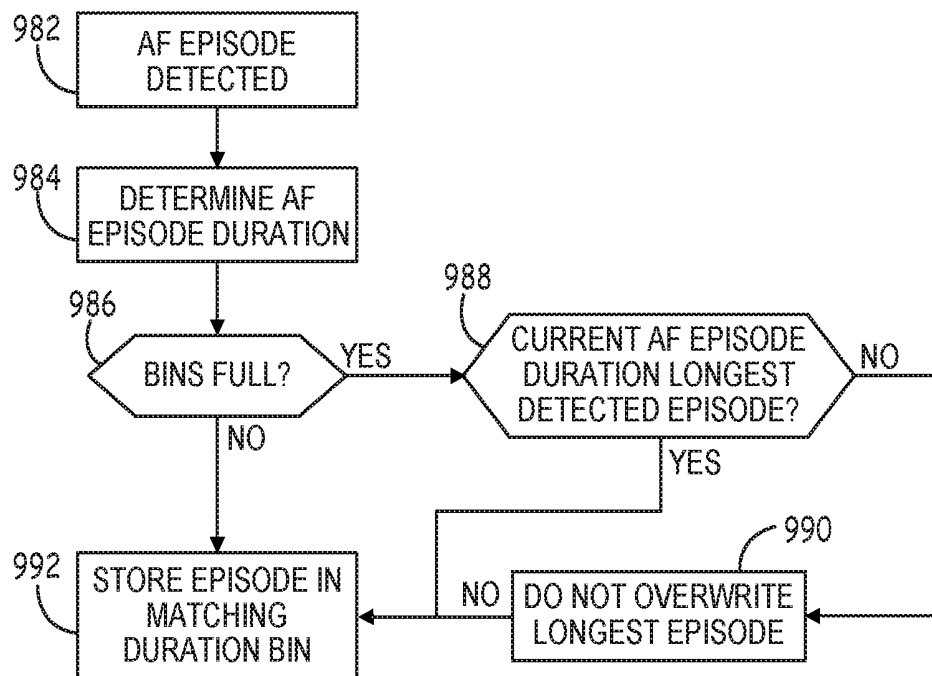
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FIG. 16

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AUTOMATIC THRESHOLDS FOR ATRIAL TACHYARRHYTHMIA DETECTION IN AN IMPLANTABLE MEDICAL DEVICE

RELATED APPLICATIONS

This application claims the benefit of the filing date of U.S. Provisional Patent Application Ser. No. 62/367,177, filed Jul. 27, 2016, the content of which is incorporated by reference herein its entirety.

TECHNICAL FIELD

The disclosure relates generally to cardiac medical devices and, in particular, to an implantable cardiac medical device and method for automatically adjusting a threshold for detecting atrial tachyarrhythmia episodes from sensed cardiac electrical signals.

BACKGROUND

During normal sinus rhythm (NSR), the heart beat is regulated by electrical signals produced by the sino-atrial (SA) node located in the right atrial wall. Each atrial depolarization signal produced by the SA node spreads across the atria, causing the depolarization and contraction of the atria, and arrives at the atrioventricular (A-V) node. The A-V node responds by propagating a ventricular depolarization signal through the bundle of His of the ventricular septum and thereafter to the bundle branches and the Purkinje muscle fibers of the right and left ventricles.

Atrial tachyarrhythmia includes the disorganized form of atrial fibrillation and varying degrees of organized atrial tachycardia, including atrial flutter. Atrial fibrillation (AF) occurs because of multiple focal triggers in the atrium or because of changes in the substrate of the atrium causing heterogeneities in conduction through different regions of the atria. The ectopic triggers can originate anywhere in the left or right atrium or pulmonary veins. The AV node will be bombarded by frequent and irregular atrial activations but will only conduct a depolarization signal when the AV node is not refractory. The ventricular cycle lengths will be irregular and will depend on the different states of refractoriness of the AV-node.

As more serious consequences of persistent atrial arrhythmias have come to be understood, such as an associated risk of relatively more serious ventricular arrhythmias and stroke, there is a growing interest in monitoring and treating atrial arrhythmias. Implantable cardiac monitors and implantable cardioverter defibrillators (ICDs) may be configured to acquire cardiac electrical signals that can be analyzed for detecting atrial arrhythmias.

SUMMARY

In general, the disclosure is directed to techniques for detecting atrial tachyarrhythmia episodes by an implantable medical device. A medical device operating according to the techniques disclosed herein analyzes a cardiac electrical signal over a plurality of time periods and classifies each of the time periods based on characteristics of the cardiac electrical signal, such as characteristics of the RR-intervals occurring during each of the plurality of time periods. The device may automatically adjust a threshold number of time periods that are classified as atrial tachyarrhythmia which are required in order to detect the atrial tachyarrhythmia.

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In one example, the disclosure provides a method for detecting an atrial tachyarrhythmia episode by a medical device, comprising receiving a cardiac electrical signal via electrodes coupled to sensing circuitry of the medical device, detecting an atrial tachyarrhythmia episode by a processor of the medical device in response to a time duration of the cardiac electrical signal classified as an atrial tachyarrhythmia being greater than or equal to a first detection threshold, determining if detection threshold adjustment criteria are met based on at least the detected atrial tachyarrhythmia episode; and adjusting the first detection threshold to a second detection threshold different than the first detection threshold in response to the detection threshold adjustment criteria being met.

In another example, the disclosure provides a medical device for detecting an atrial tachyarrhythmia. The medical device includes sensing circuitry configured to receive a cardiac electrical signal from electrodes coupled to the sensing circuitry, and a processor configured to detect an atrial tachyarrhythmia episode in response to a time duration of the cardiac electrical signal classified as an atrial tachyarrhythmia being greater than or equal to a first detection threshold, determine if detection threshold adjustment criteria are met based on at least the detected atrial tachyarrhythmia episode, and adjust the first detection threshold to a second detection threshold different than the first detection threshold in response to the detection threshold adjustment criteria being met.

In another example, the disclosure provides a non-transitory, computer-readable storage medium storing instructions for causing a processor included in a medical device to perform a method for detecting an atrial tachyarrhythmia episode. The method includes receiving a cardiac electrical signal via electrodes coupled to sensing circuitry of the medical device, detecting an atrial tachyarrhythmia episode in response to a time duration of the cardiac electrical signal classified as an atrial tachyarrhythmia being greater than or equal to a first detection threshold, determining if detection threshold adjustment criteria are met based on at least the detected atrial tachyarrhythmia episode, and adjusting the first detection threshold to a second detection threshold different than the first detection threshold in response to the detection threshold adjustment criteria being met.

This summary is intended to provide an overview of the subject matter described in this disclosure. It is not intended to provide an exclusive or exhaustive explanation of the apparatus and methods described in detail within the accompanying drawings and description below. Further details of one or more examples are set forth in the accompanying drawings and the description below.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a conceptual diagram of an implantable cardioverter defibrillator (ICD) system for detecting atrial arrhythmias according to one example.

FIGS. 2A and 2B are conceptual diagrams of an alternative ICD system that may be configured to detect atrial tachyarrhythmia according to the techniques disclosed herein.

FIG. 3 is a conceptual diagram of an implantable medical device (IMD) system for detecting atrial tachyarrhythmia according to another example.

FIG. 4 is a functional schematic diagram of an ICD, such as the ICD of FIG. 1 or the ICD of FIGS. 2A and 2B.

FIG. 5 is a schematic diagram of methods used for detecting cardiac events by any of the ICDs of FIGS. 1A, 1B and 2 or the monitoring device of FIG. 1C according to one example.

FIG. 6 is a diagram of a two-dimensional histogram representing a Lorenz plot area used in the techniques disclosed herein for detecting atrial tachyarrhythmia.

FIG. 7 is flowchart of a method for determining a factor for classifying time periods for detecting atrial tachyarrhythmia according to one example.

FIG. 8 is a flowchart of a method for classifying a predetermined time period for use in detecting atrial tachyarrhythmia according to one example.

FIG. 9 is a schematic diagram of atrial fibrillation detection that may be performed by the ICDs shown in FIGS. 1, 2A and 2B or the monitor of FIG. 3.

FIG. 10 is a schematic diagram of a method for detecting atrial fibrillation by an ICD or implantable monitoring device according to another example.

FIG. 11 is a flowchart of a method for detecting atrial fibrillation, according to one example.

FIG. 12 is a flow chart of a method performed by an ICD or implantable monitor for providing a response to detecting atrial tachyarrhythmia according to one example.

FIG. 13 is a flow chart of a method performed by an implantable medical device system, such as the ICD system of FIG. 1 or the ICD system of FIGS. 2A and 2B, for automatically adjusting a detection threshold required to detect an atrial tachyarrhythmia, such as AF.

FIG. 14 is a flow chart of a method for automatically adjusting a detection threshold for detecting AF according to another example.

FIG. 15 is a flow chart of a method for detecting AF and determining AF burden according to another example.

FIG. 16 is a flow chart of a method for storing AF episode data in the memory of an ICD according to one example.

DETAILED DESCRIPTION

In the following description, references are made to illustrative examples for carrying out the methods described herein. It is understood that other variations from these examples may be utilized without departing from the scope of the disclosure. In these various examples, a cardiac electrical signal is used for determining successive ventricular cycle lengths for use in detecting atrial arrhythmias. Ventricular cycle lengths may be determined as intervals between successive R-waves that are sensed from the cardiac electrical signal and attendant to the depolarization of the ventricles. The differences between the successive ventricular cycle lengths, or RR intervals (RRIs), are analyzed for determining evidence of atrial tachyarrhythmia, e.g., atrial fibrillation (AF). As described herein, a time period of the cardiac signal may be classified as AF, non-AF, or unclassified based on an analysis of the RRIs and other factors. When a predetermined number of time periods of the cardiac signal are classified as AF, a medical device operating according to the techniques disclosed herein may detect an AF episode. The device, however, may adjust a classification criterion applied for classifying a time period of the cardiac signal prior to detecting AF and detect AF based on the adjusted classification criterion used for classifying subsequent time periods. Furthermore, the device may automatically adjust the number of time periods required to be classified as AF in order to detect the AF episode.

Aspects of the methods described herein can be incorporated in a variety of implantable or external medical devices having cardiac signal monitoring capabilities, which may or may not include therapy delivery capabilities. Such devices include single chamber, dual chamber or bi-ventricular pacing systems or ICDs that sense the R-waves and deliver an electrical stimulation therapy to the ventricles. The atrial arrhythmia detection methods presently disclosed may also be incorporated in implantable cardiac monitors having implantable electrodes or external cardiac monitors having electrocardiogram (ECG) electrodes coupled to the patient's skin to detect R-waves, e.g., Holter monitors, or within computerized systems that analyze pre-recorded ECG or cardiac electrogram (EGM) data. Embodiments may further be implemented in a patient monitoring system, such as a centralized computer system which processes cardiac electrical signals and other data sent to it by implantable or wearable monitoring devices.

FIG. 1 is a conceptual diagram of an implantable medical device (IMD) system 1 for detecting atrial tachyarrhythmias according to one example. The IMD system 1 includes an implantable cardioverter defibrillator (ICD) 10 coupled to a patient's heart 2 via a transvenous electrical medical lead 16. ICD 10 includes a connector block 12 that may be configured to receive the proximal end of lead 16, which is advanced transvenously for positioning electrodes for sensing and stimulation in the right ventricular chamber of heart 2 in the example shown. The techniques disclosed herein may be implemented in a single chamber ICD system 1 that is coupled only to a ventricular lead such as right ventricular (RV) lead 16 for receiving cardiac electrical signals including at least R-waves attendant to the ventricular depolarizations of heart 2.

RV lead 16 is positioned such that its distal end is in the right ventricle for sensing RV cardiac signals and delivering pacing or shocking pulses in the right ventricle. For these purposes, RV lead 16 is equipped with pacing and sensing electrodes shown as a ring electrode 30 and a tip electrode 28. In some examples, tip electrode 28 is an extendable helix electrode mounted retractably within an electrode head. RV lead 16 is further shown to carry defibrillation electrodes 24 and 26, which may be elongated coil electrodes used to deliver high voltage cardioversion/defibrillation (CV/DF) shocks. Defibrillation electrode 24 is referred to herein as the "RV defibrillation electrode" or "RV coil electrode" because it may be carried along RV lead 16 such that it is positioned substantially within the right ventricle when distal pacing and sensing electrodes 28 and 30 are positioned for pacing and sensing in the right ventricle. Defibrillation electrode 26 is referred to herein as a "superior vena cava (SVC) defibrillation electrode" or "SVC coil electrode" because it may be carried along RV lead 16 such that it is positioned at least partially along the SVC when the distal end of RV lead 16 is advanced within the right ventricle.

Each of electrodes 24, 26, 28 and 30 are connected to a respective insulated conductor extending within the body of lead 16. The proximal end of the insulated conductors are coupled to corresponding connectors carried by a proximal lead connector assembly, e.g., an industry standard DF-4 connector, at the proximal end of lead 16, which may be inserted into a connector bore of connector block 12 for providing electrical connection to ICD 10.

The techniques disclosed herein for detecting atrial tachyarrhythmia may be successfully performed without requiring atrial signal sensing. As such, these techniques may be implemented in a single chamber ICD system, such as system 1, which includes a lead extending into the right

ventricle for positioning electrodes for sensing ventricular signals but does not include electrodes positioned in or along the atrial chambers for sensing atrial signals. In other examples, other transvenous leads may be present, e.g., a right atrial lead for sensing right atrial signals and delivering electrical stimulation pulses to the right atrium, and/or a left ventricular lead, which may be advanced transvenously into a cardiac vein via the coronary sinus, for sensing left ventricular signals and delivering electrical stimulation pulses to the left atrium. A multi-chamber ICD system in which aspects of the techniques described herein is generally disclosed in U.S. patent application Ser. No. 14/520,798, (Cao et al.), incorporated herein by reference in their entirety.

Electrodes **28** and **30** (and/or coil electrodes **24** and **26**) may be used for acquiring cardiac electrical signals needed for performing atrial tachyarrhythmia detection as described herein. R-waves sensed from cardiac electrical signals obtained by ICD **10** are used for determining RRs between consecutively sensed R-waves for detecting atrial tachyarrhythmia by a processor of ICD **10** based at least in part on an analysis of the RRs. ICD **10** may be configured to sense cardiac electrical signals from electrodes **24**, **26**, **28** and/or **30**, detect atrial tachyarrhythmia and provide an atrial tachyarrhythmia detection response such as storing atrial tachyarrhythmia episode data for transmission to an external device **40**. ICD **10** may additionally be configured to deliver ventricular bradycardia pacing, detect ventricular tachyarrhythmias, and deliver anti-tachycardia pacing therapy and cardioversion/defibrillation shock therapies to the RV via electrodes **24**, **26**, **28** and/or **30** carried by lead **16**.

The RV pacing and sensing electrodes **28** and **30** may be used as a bipolar pair, commonly referred to as a “tip-to-ring” configuration for sensing cardiac electrical signals. In some cases, RV tip electrode **28** may be selected with a coil electrode **24** or **26** to be used as an integrated bipolar pair, commonly referred to as a “tip-to-coil” configuration for sensing cardiac electrical signals. ICD **10** may, for example, select one or more sensing electrode vectors including a tip-to-ring sensing vector between electrodes **28** and **30** and a tip-to-coil or ring-to-coil sensing vector, e.g., between RV tip electrode **28** and SVC coil electrode **26**, between RV tip electrode **28** and RV coil electrode **24**, between RV ring electrode **30** and SVC coil electrode **26** or between RV ring electrode **30** and RV coil electrode **24**. In other examples, any of the electrodes **24**, **26**, **28** or **30** carried by RV lead **16** may be selected by ICD **10** in a unipolar sensing configuration with the ICD housing **15** serving as the indifferent electrode, commonly referred to as the “can” or “case” electrode. It is recognized that numerous sensing and electrical stimulation electrode vectors may be available using the various electrodes carried by lead **16** and coupled to ICD **10**. ICD **10** may be configured to selectively couple one or more sensing electrode vectors to sensing circuitry enclosed by housing **15**, e.g., sensing circuitry including one or more amplifiers, filters, rectifiers, comparators, sense amplifiers, analog-to-digital converters and/or other circuitry configured to acquire a cardiac electrical signal for use in detecting cardiac arrhythmias.

In other examples, the ICD housing **15** may serve as a subcutaneous defibrillation electrode in combination with one or both of the coil electrodes **24** and/or **26** for delivering CV/DF shocks to the atria or ventricles. It is recognized that alternate lead systems may be substituted for the single RV lead system illustrated in FIG. **1**. While a particular single-chamber ICD and transvenous lead system **1** is illustrated in FIG. **1**, methodologies included in the present disclosure

may be adapted for use with any single chamber, dual chamber, or multi-chamber ICD or pacemaker system, subcutaneous implantable device, or other internal or external cardiac monitoring device.

An external device **40** is shown in telemetric communication with ICD **10** by an RF communication link **42**. External device **40** is often referred to as a “programmer” because it is typically used by a physician, technician, nurse, clinician or other qualified user for programming operating parameters in ICD **10**. External device **40** may be located in a clinic, hospital or other medical facility. External device **40** may alternatively be embodied as a home monitor or a handheld device that may be used in a medical facility, in the patient’s home, or another location. Operating parameters, such as sensing and therapy delivery control parameters, may be programmed into ICD **10** using external device **40**.

External device **40** includes a processor **52**, memory **53**, user display **54**, user interface **56** and telemetry circuitry **58**. Processor **52** controls external device operations and processes data and signals received from ICD **10**. According to techniques disclosed herein, processor **52** may receive atrial tachyarrhythmia data obtained by ICD **10** and transmitted from ICD **10** to external telemetry circuitry **58**. As described below in conjunction with FIGS. **12** and **16**, ICD **10** may be configured to store cardiac signal data associated with detected atrial tachyarrhythmia episodes and transmit the cardiac signal data to external device **40**. Processor **52** provides user display **54** with at least a portion of the cardiac electrical signal data for generating a display of the cardiac electrical signal detected as atrial tachyarrhythmia for observation and review by a clinician.

The user display **54** provides a display of the cardiac signal data and may include a graphical user interface that facilitates programming of one or more sensing parameters and/or atrial arrhythmia detection parameters as well as other arrhythmia detection and therapy control parameters by a user interacting with external device **40**. External device **40** may display other data and information relating to ICD functions to a user for reviewing ICD operation and programmed parameters as well as cardiac electrical signals or other physiological data that is retrieved from ICD **10** during an interrogation session. User interface **56** may include a mouse, touch screen, or other pointing device, keyboard and/or keypad to enable a user to interact with external device **40** to initiate a telemetry session with ICD **10** for retrieving data from and/or transmitting data to ICD **10** and for selecting and programming desired sensing and therapy delivery control parameters into ICD **10**.

Telemetry circuitry **58** includes a transceiver and antenna configured for bidirectional communication with an implantable transceiver and antenna included in ICD **10**. Telemetry circuitry **58** is configured to operate in conjunction with processor **52** for encoding and decoding transmitted and received data relating to ICD functions via communication link **42**. Communication link **42** may be established between ICD **10** and external device **40** using a radio frequency (RF) link such as BLUETOOTH®, Wi-Fi, Medical Implant Communication Service (MICS) or other RF bandwidth. In some examples, external device **40** may include a programming head that is placed proximate ICD **10** to establish and maintain a communication link, and in other examples external device **40** and ICD **10** may be configured to communicate using a distance telemetry algorithm and circuitry that does not require the use of a programming head and does not require user intervention to maintain a communication link.

It is contemplated that external device **40** may be in wired or wireless connection to a communications network via telemetry circuitry **58** for transferring data to a remote database or computer to allow remote management of the patient **3**. Remote patient management systems may be configured to utilize the presently disclosed techniques to enable a clinician to review cardiac electrical signal data and atrial tachyarrhythmia episode data received from ICD **10** and to select and program control parameters transmitted to ICD **10**. Reference is made to commonly-assigned U.S. Pat. No. 6,599,250 (Webb et al.), U.S. Pat. No. 6,442,433 (Linberg et al.), U.S. Pat. No. 6,418,346 (Nelson et al.), and U.S. Pat. No. 6,480,745 (Nelson et al.) for general descriptions and examples of remote patient management systems that enable remote patient monitoring and device programming. Each of these patents is incorporated herein by reference in their entirety.

FIGS. **2A** and **2B** are conceptual diagrams of an alternative ICD system **100** that may be configured to detect atria arrhythmias according to the techniques disclosed herein. FIG. **2A** is a front view of an extra-cardiovascular ICD system **100** implanted within patient **112**. FIG. **2B** is a side view of ICD system **100** implanted within patient **112**. ICD system **100** includes an ICD **110** connected to an extra-cardiovascular electrical stimulation and sensing lead **116**. ICD system **100** may further include an intracardiac pacemaker **101** configured to deliver pacing pulses to a heart chamber, for example from within the right ventricle or within the left ventricle.

ICD **110** includes a housing **115** that forms a hermetic seal that protects internal components of ICD **110**. Internal device components may include circuitry shown and described in conjunction with FIG. **4** below, such as sense amplifier(s), A/D converter, pacing output circuitry, high voltage output circuitry and a microprocessor and memory and/or other control circuitry. The housing **115** of ICD **110** may be formed of a conductive material, such as titanium or titanium alloy. The housing **115** may function as a housing electrode (sometimes referred to as a can electrode). In examples described herein, housing **115** may be used as an active can electrode for use in delivering cardioversion/defibrillation (CV/DF) shocks or other high voltage pulses generated by high voltage charging circuitry of ICD **110**. In other examples, housing **115** may be available for use in sensing cardiac signals and/or for delivering unipolar, low voltage cardiac pacing pulses by a pacer output circuit in conjunction with lead-based cathode electrodes. In other instances, the housing **115** of ICD **110** may include multiple electrodes on an outer portion of the housing. The outer portion(s) of the housing **115** functioning as an electrode(s) may be coated with a material, such as titanium nitride.

ICD **110** includes a connector assembly **117** (also referred to as a connector block or header) that includes electrical feedthroughs crossing housing **115** to provide electrical connections between conductors extending within the lead body **118** of lead **116** and electronic components included within the housing **115** of ICD **110**. As described below in conjunction with FIG. **4**, housing **115** may house one or more processors, memories, telemetry transceivers, sensing circuitry such as sense amplifiers and analog-to digital converters, therapy delivery circuitry such as pacer timing and control, CV/DF control, pace output and HV output circuits and associated charging circuits, a switch matrix, a data bus, one or more batteries or other power sources and other components for sensing cardiac electrical signals,

detecting a heart rhythm, and controlling and delivering electrical stimulation pulses to treat an abnormal heart rhythm.

Lead **116** includes an elongated lead body **118** having a proximal end **127** that includes a lead connector (not shown) configured to be connected to ICD connector assembly **117** and a distal portion **125** that includes one or more electrodes. In the example illustrated in FIGS. **2A** and **2B**, the distal portion **125** of lead **116** includes defibrillation electrodes **124** and **126** and pace/sense electrodes **128**, **130** and **131**. Electrical conductors (not illustrated) extend through one or more lumens of the elongated lead body **118** of lead **116** from the lead connector at the proximal lead end **127** to electrodes **124**, **126**, **128**, **130** and **131** located along the distal portion **125** of the lead body **118**. The lead body **118** of lead **116** may be formed from a non-conductive material, including silicone, polyurethane, fluoropolymers, mixtures thereof, and other appropriate materials, and shaped to form one or more lumens within which the one or more conductors extend. However, the techniques disclosed herein are not limited to such constructions or to any particular lead body design.

The respective conductors electrically couple the electrodes **124**, **126**, **128**, **130** and **131** to circuitry, such as a switch matrix or other switching circuitry for selection and coupling to a sense amplifier or other cardiac event detection circuitry and/or to a therapy output circuit, e.g., a pacing output circuit or a HV output circuit for delivering CV/DF shock pulses. Connections between electrode conductors and ICD circuitry is made via connections in the connector assembly **117**, including associated electrical feedthroughs crossing housing **115**. The electrical conductors transmit therapy from an output circuit within ICD **110** to one or more of defibrillation electrodes **124** and **126** and/or pace/sense electrodes **128**, **130** and **131** and transmit sensed electrical signals from one or more of defibrillation electrodes **124** and **126** and/or pace/sense electrodes **128**, **130** and **131** to the sensing circuitry within ICD **110**.

ICD **110** may obtain electrical signals corresponding to electrical activity of heart **102** via a combination of sensing vectors that include combinations of electrodes **128**, **130**, and/or **131**. In some examples, housing **115** of ICD **110** is used in combination with one or more of electrodes **128**, **130** and/or **131** in a sensing electrode vector. ICD **110** may even obtain cardiac electrical signals using a sensing vector that includes one or both defibrillation electrodes **124** and/or **126**, e.g., between electrodes **124** and **126** or one of electrodes **124** or **126** in combination with one or more of electrodes **128**, **130**, **131**, and/or the housing **115**.

ICD **110** analyzes the cardiac electrical signals received from one or more of the sensing vectors to monitor for abnormal rhythms, such as AF, VT and VF. ICD **110** generates and delivers electrical stimulation therapy in response to detecting a ventricular tachyarrhythmia (e.g., VT or VF). ICD **110** may deliver ATP in response to VT detection, and in some cases may deliver ATP prior to a CV/DF shock or during high voltage capacitor charging in an attempt to avert the need for delivering a CV/DF shock. ICD **110** may deliver a CV/DF shock pulse when VF is detected or when VT is not terminated by ATP.

Electrodes **124** and **126** (and in some examples housing **115**) are referred to herein as defibrillation electrodes because they are utilized, individually or collectively, for delivering high voltage stimulation therapy (e.g., cardioversion or defibrillation shocks). Electrodes **124** and **126** may be elongated coil electrodes and generally have a relatively high surface area for delivering high voltage electrical

stimulation pulses compared to low voltage pacing and sensing electrodes **28**, **30** and **31**. However, electrodes **124** and **126** and housing **115** may also be utilized to provide pacing functionality, sensing functionality or both pacing and sensing functionality in addition to or instead of high voltage stimulation therapy. In this sense, the use of the term “defibrillation electrode” herein should not be considered as limiting the electrodes **124** and **126** for use in only high voltage cardioversion/defibrillation shock therapy applications. Electrodes **124** and **126** may be used in a pacing electrode vector for delivering extra-cardiovascular pacing pulses such as ATP pulses, post-shock pacing or other pacing therapies and/or in a sensing vector used to sense cardiac electrical signals for detecting atrial and ventricular arrhythmias, including AF, atrial flutter, VT and VF.

Electrodes **128**, **130** and **131** are relatively smaller surface area electrodes for delivering low voltage pacing pulses and for sensing cardiac electrical signals. Electrodes **128**, **130** and **131** are referred to as pace/sense electrodes because they are generally configured for use in low voltage applications, e.g., used as either a cathode or anode for delivery of pacing pulses and/or sensing of cardiac electrical signals.

The pace/sense electrodes **128**, **130** and/or **131** may be located at different locations along the length of lead **116** than shown. In other examples, lead **116** may include less than three pace/sense electrodes or more than three pace/sense electrodes and/or a single defibrillation electrode or more than two electrically isolated or electrically coupled defibrillation electrodes or electrode segments. Various example configurations of extra-cardiovascular leads and electrodes and dimensions that may be implemented in conjunction with the AF detection techniques disclosed herein are described in commonly-assigned U.S. patent application Ser. No. 14/519,436, U.S. patent application Ser. No. 14/695,255 and provisionally-filed U.S. Pat. Application No. 62/089,417, all of which are incorporated herein by reference in their entirety.

Lead **16** extends subcutaneously or submuscularly over the ribcage **132** medially from the connector assembly **127** of ICD **110** toward a center of the torso of patient **112**, e.g., toward xiphoid process **120** of patient **112**. At a location near xiphoid process **120**, lead **116** bends or turns and extends superiorly within anterior mediastinum **136** in a substernal position. Lead **116** of system **100** is implanted at least partially underneath sternum **122** of patient **112**.

Anterior mediastinum **136** may be viewed as being bounded laterally by pleurae, posteriorly by pericardium **138**, and anteriorly by sternum **122**. In some instances, the anterior wall of anterior mediastinum **136** may also be formed by the transversus thoracis muscle and one or more costal cartilages. Anterior mediastinum **136** includes a quantity of loose connective tissue (such as areolar tissue), adipose tissue, some lymph vessels, lymph glands, substernal musculature, small side branches of the internal thoracic artery or vein, and the thymus gland. In one example, the distal portion **125** of lead **116** extends along the posterior side of sternum **122** substantially within the loose connective tissue and/or substernal musculature of anterior mediastinum **136**.

A lead implanted such that the distal portion **125** is substantially within anterior mediastinum **136** may be referred to as a “substernal lead.” In the example illustrated in FIGS. **2A** and **2B**, lead **116** extends substantially centered under sternum **122**. In other instances, however, lead **116** may be implanted such that it extends in a position that is offset laterally from the center of sternum **122**. In some instances, lead **116** may extend laterally such that distal

portion **125** of lead **116** is underneath/below the ribcage **132** in addition to or instead of sternum **122**. In other examples, the distal portion **125** of lead **116** may be implanted in other extra-cardiovascular, intra-thoracic locations, including the pleural cavity or around the perimeter of and adjacent to but typically not within the pericardium **138** of heart **102**.

In other examples, lead **116** may remain outside the thoracic cavity and extend subcutaneously or submuscularly over the ribcage **132** and/or sternum **122**. The path of lead **116** may depend on the location of ICD **110**, the arrangement and position of electrodes carried by the lead distal portion **125**, and/or other factors.

ICD **110** is shown implanted subcutaneously on the left side of patient **112** along the ribcage **132**. ICD **110** may, in some instances, be implanted between the left posterior axillary line and the left anterior axillary line of patient **112**. ICD **110** may, however, be implanted at other subcutaneous or submuscular locations in patient **112**. For example, ICD **110** may be implanted in a subcutaneous pocket in the pectoral region. In this case, lead **116** may extend subcutaneously or submuscularly from ICD **110** toward the manubrium of sternum **122** and bend or turn and extend inferior from the manubrium to the desired location subcutaneously or submuscularly. In yet another example, ICD **110** may be placed abdominally.

In some patients, an intracardiac pacemaker **101** may be present in the right ventricle, right atrium or along the left ventricle. Pacemaker **101** may be configured to deliver pacing pulses in the absence of sensed intrinsic heart beats, in response to detecting VT, or according to other pacing therapy algorithms. For example, pacemaker **101** may be implanted in the right ventricle of the patient for providing single chamber ventricular pacing. The techniques disclosed herein for classifying time periods of a cardiac signal for detecting atrial tachyarrhythmia may be utilized in the presence of ventricular pacing delivered by ICD **110** and/or by an intracardiac pacemaker such as pacemaker **101**. Pacemaker **101** may generally correspond to the intra-cardiac pacemaker disclosed in U.S. Pat. No. 8,923,963 (Bonner, et al.), incorporated herein by reference in its entirety. ICD **110** may be configured to detect pacing pulses delivered by pacemaker **101**. The frequency of pacing pulses delivered by pacemaker **101** may be a factor determined in classifying a cardiac electrical signal time period for AF detection purposes.

Pacemaker **101** may have limited processing power and therapy delivery capacity compared to ICD **110** such that the advanced cardiac rhythm detection techniques disclosed herein may be implemented in ICD **110** rather than in pacemaker **101**. The methods disclosed herein as being performed by ICD **10** or ICD **110**, however, are not to be considered limited to being implemented in an ICD. Aspects of the atrial tachyarrhythmia detection techniques disclosed herein may be implemented in pacemaker **101**, all or in part.

FIG. **3** is a conceptual diagram of a cardiac monitoring device **60** which may employ aspects of the atrial tachyarrhythmia detection techniques disclosed herein. Monitoring device **60** is shown implanted subcutaneously in the upper thoracic region of a patient's body **3** and displaced from the patient's heart **2**. The housing **62** of cardiac monitor **60** (shown enlarged in scale compared to the patient's body **3**) includes a non-conductive header module **64** attached to a hermetically sealed housing **62**. The housing **62** contains the circuitry of the cardiac monitor **60** and is generally electrically conductive but may be covered in part by an electrically insulating coating. A first, subcutaneous, sense electrode, **A**, is formed on the surface of the header module **64**.

In some examples, one or more electrodes may be incorporated in the header module **64**. A second, subcutaneous, sense electrode, B, is formed by at least a portion of the housing **62**. For example, electrode B may be an exposed portion of housing **62** when housing **62** is coated by an electrically insulating coating. The conductive housing electrode B may be directly connected with the sensing circuitry.

An electrical feedthrough extends through the mating surfaces of the header module **64** and the housing **62** to electrically connect the first sense electrode A with sensing circuitry enclosed within the housing **62**. The electrical signals attendant to the depolarization and re-polarization of the heart **2** are referred to as the cardiac electrical signals and are sensed across the sense electrodes A and B and include at least R-waves attendant to the ventricular depolarizations of heart **2**. The cardiac monitoring device **60** may be sutured to subcutaneous tissue at a desired orientation of its electrodes A and B to the axis of the heart **2** to detect and record the cardiac electrical signals in a sensing vector A-B for subsequent processing and uplink telemetry transmission to an external device **40** (shown in FIG. 1A).

In one embodiment, the spacing between electrodes A and B may range from 60 mm to 25 mm. In other embodiments, the electrode spacing may range from 55 mm to 30 mm, or from 55 mm to 35 mm. The volume of the implantable cardiac monitoring device **60** may be three cubic centimeters or less, 1.5 cubic centimeters or less or any volume between three and 1.5 cubic centimeters. The length of cardiac monitoring device **60** may range from 30 to 70 mm, 40 to 60 mm or 45 to 60 mm and may be any length between 30 and 70 mm. The width of a major surface such a cardiac monitoring device **60** may range from 3 to 10 mm and may be any thickness between 3 and 10 mm. The thickness of cardiac monitoring device **60** may range from 2 to 9 mm or 2 to 5 mm and may be any thickness between 2 and 9 mm.

The sensing circuitry included in housing **62** is configured to detect R-waves for monitoring for atrial tachyarrhythmia according to the techniques disclosed herein. Such sensing circuitry may include a pre-filter and amplifier, an analog-to-digital filter, a rectifier, a sense amplifier, a comparator and/or other components configured to receive cardiac electrical signals and detect R-waves from the signals. Aspects of a cardiac monitoring device of the type that may employ atrial tachyarrhythmia detection techniques disclosed herein are generally disclosed in U.S. Publication No. 2015/0088216 (Gordon, et al.) and U.S. Pat. No. 7,027,858 (Cao, et al.), both incorporated herein by reference in its entirety.

In general, the hermetically sealed housing **62** includes a lithium battery or other power source, a processor and memory or other control circuitry that controls device operations and records arrhythmic cardiac electrical signal episode data in memory registers or bins, and a telemetry transceiver and antenna circuit that receives downlink telemetry commands from and transmits stored data in a telemetry uplink to an external device **40** (FIG. 1). The circuitry and memory may be implemented in discrete logic or a microcomputer based system with A/D conversion of sampled cardiac electrical signal amplitude values. One implantable cardiac monitor that can be modified in accordance with the presently disclosed techniques is described in U.S. Pat. No. 6,412,490 (Lee et al.), incorporated herein by reference in its entirety, as well as the cardiac monitors disclosed in any of the above-incorporated references.

FIG. 4 is a functional schematic diagram of an ICD, such as ICD **10** of FIG. 1 or ICD **110** of FIGS. 2A and 2B. This diagram should be taken as illustrative of the type of device with which the techniques disclosed herein may be embodied

and not as limiting. The example shown in FIG. 4 is a microprocessor-controlled device, but the disclosed methods may also be practiced with other types of devices such as those employing dedicated digital circuitry.

With regard to the electrode system illustrated in FIG. 1, ICD **10** is provided with a number of connection terminals for achieving electrical connection to the RV lead **6** and its respective electrodes. Housing **15** may be used as an indifferent electrode during unipolar stimulation or sensing or as an active can electrode during shock delivery. Electrodes **24**, **26** and housing **15** may be selectively coupled to the high voltage output circuit **234** to facilitate the delivery of high energy shocking pulses to the heart using one or more of the coil electrodes **24** and **26** and optionally the housing **15**.

RV tip electrode **28** and the RV ring electrode **30** may be coupled to a ventricular sense amplifier **200** for sensing ventricular signals. The ventricular sense amplifier **200** may take the form of automatic gain controlled amplifiers with adjustable sensitivity. ICD **10** and, more specifically, microprocessor **224** may automatically adjust the sensitivity of ventricular sense amplifier **200** in response to detection of oversensing in order to reduce the likelihood of oversensing of cardiac events and/or non-cardiac noise.

Ventricular sense amplifier **200** may receive timing information from pacer timing and control circuitry **212**. For example, ventricular sense amplifier **200** may receive blanking period input, e.g., V_BLANK, which indicates the amount of time the amplifier is "turned off" in order to prevent saturation due to an applied pacing pulse or defibrillation shock. The general operation of the ventricular sense amplifier **200** may correspond to that disclosed in U.S. Pat. No. 5,117,824 (Keimel, et al.), incorporated herein by reference in its entirety.

Whenever a signal received by ventricular sense amplifier **200** exceeds a ventricular sensitivity, a signal is generated on the R-out signal line **202**. As described below, a signal on the R-out signal line **202**, which may be referred to as a ventricular sense event (Vs event) signal, may be received by microprocessor **224** and used for determining RRI differences.

Switch matrix **208** is used to select which of the available electrodes **24**, **26**, **28** and **30** (or **124**, **126**, **128** and **130** of FIG. 2A) are coupled to a wide band amplifier **210** for use in digital signal analysis. Selection of the electrodes is controlled by the microprocessor **224** via data/address bus **218**. The selected electrode configuration may be varied as desired for the various sensing, pacing, cardioversion and defibrillation functions of the ICD **10**. For example, while RV electrodes **28** and **30** are shown coupled to sense amplifier **200** and pace output circuit **216** suggesting dedicated pace/sense electrodes and coil electrodes **24** and **26** are shown coupled to HV output circuit **234** suggesting dedicated CV/DV shock electrodes, it is recognized that switching circuitry included in switch matrix **208** may be used to select any of the available electrodes in a sensing electrode vector, a pacing electrode vector, or a CV/DF shock vector as described previously.

Signals from the electrodes selected for coupling to bandpass amplifier **210** are provided to multiplexer **220**, and thereafter converted to multi-bit digital signals by A/D converter **222**, for storage in random access memory **226** under control of direct memory access circuit **228** via data/address bus **218**. Microprocessor **224** may employ digital signal analysis techniques to characterize the digitized signals stored in random access memory **226** to recognize and classify the patient's heart rhythm employing any of numerous signal processing methodologies for ana-

lyzing cardiac signals and cardiac event waveforms, e.g., R-waves. One tachyarrhythmia detection system is described in U.S. Pat. No. 5,545,186 (Olson et al.), incorporated herein by reference in its entirety.

ICD 10 or ICD 110 may include a second sensing channel including sense amplifier 204 receiving a V BLANK signal from pace timing and control 212 and providing an R OUT signal line 206. In the example shown, a sensing electrode vector including electrodes 28 and 30 is coupled to amplifier 200 and a second sensing electrode vector including electrode 24 and housing 15 is coupled to the second amplifier 204. Switch matrix 208 may select which sensing electrode vector selected from any of electrodes 24, 26, 28 and 30 and housing 15 (or electrodes 124, 126, 128 130 and 131 and housing 115 in the case of ICD 110) is coupled to each of the two sensing channels represented by sense amplifiers 200 and 204 and which pacing electrode vector is coupled to pace output circuit 216.

It is to be understood that the circuitry shown in FIG. 4 may be modified according to the particular device requirements. For example, the single chamber ICD 10 of FIG. 1 or extravascular ICD 110 of FIG. 2A may include one ventricular sense amplifier 200 and terminals for electrically coupling to electrodes 24, 26, 28 and 30 and/or housing 15 (or electrodes 124, 126, 128, 130 or 131 and/or housing 115 in any desired sensing electrode vector combination.

Upon detection of an arrhythmia, an episode of cardiac signal data, along with sensed intervals and corresponding annotations of sensed events, may be stored in random access memory 226. The cardiac electrical signals received via sensing electrode pairs may be stored in RAM 226. In some cases, a near-field and a far-field signal are received by the two amplifiers 200 and 204. Typically, a near-field sensing electrode pair includes a tip electrode and a ring electrode located in the ventricle, electrodes 28 and 30. A far-field sensing electrode pair includes electrodes spaced further apart such as any of: the defibrillation coil electrodes 24 or 26 with housing 15; a tip electrode 28 with housing 15; a tip electrode 28 with a defibrillation coil electrode 24 or 26. The use of near-field and far-field signal detecting arrhythmia episodes is described in U.S. Pat. No. 5,193,535 (Bardy), incorporated herein by reference in its entirety. Annotation of sensed events, which may be displayed and stored with cardiac signal data, is described in U.S. Pat. No. 4,374,382 (Markowitz), incorporated herein by reference in its entirety.

The techniques disclosed herein may be applied to one or more cardiac electrical signals acquired using any combination of the available electrodes. In some examples, the sensing circuitry of ICD 10 (or ICD 110) includes more than two sensing channels for acquiring more than two cardiac electrical signals. For example, a first cardiac electrical signal is acquired between the ICD housing 15 and RV coil electrode 24, a second cardiac electrical signal is acquired between the RV coil electrode 24 and the SVC coil electrode 26, and a third cardiac electrical signal is acquired between the RV tip electrode 28 and the RV ring electrode 30. All three signals may be collected and used by microprocessor 224 for analyzing R-waves and RRIs and detecting atrial and/or ventricular arrhythmias. As discussed below in conjunction with FIG. 12, at least two cardiac signals may be stored in RAM 226 when a tachyarrhythmia episode is detected for transmission by telemetry circuit 330 to external device 40. When atrial tachyarrhythmia is detected, with or without simultaneous detection of ventricular tachyarrhythmia, the two signals may be stored having two different gain settings to provide two different signals for display on

external device 40. One signal displayed at a higher gain may result in R-wave clipping but enables relatively small amplitude P-waves to be more readily observed, which enables any relationship between the detected atrial and ventricular tachyarrhythmia (if present) to be observed by a clinician through comparison of the two different signals. When ventricular tachyarrhythmia is detected without atrial tachyarrhythmia detection, two signals may be stored both having a gain setting that avoids clipping of R-waves.

The telemetry circuit 330 includes a transceiver for receiving downlink telemetry from and sending uplink telemetry to external device 40 using antenna 332. Telemetry circuit 330 provides bi-directional telemetric communication with an external device 40 as described above.

ICD 10 may receive programmable operating parameters and algorithms via telemetry circuit 330 for storage in RAM 226 and accessed by microprocessor 224 for controlling ICD functions. For example, cardiac rhythm detection parameters and therapy control parameters used by ICD 10 may be programmed via telemetry circuit 330.

Data stored or acquired by ICD 10, including physiological signals or associated data derived therefrom, results of device diagnostics, and histories of detected arrhythmia episodes and delivered therapies, may be retrieved from ICD 10 by the external device 40 following an interrogation command received by telemetry circuit 330. Data to be uplinked to the external device and control signals for the telemetry circuit 330 are provided by microprocessor 224 via address/data bus 218. Received telemetry is provided to microprocessor 224 via multiplexer 220. Numerous types of telemetry systems known for use in implantable medical devices may be implemented in ICD 10.

Other circuitry shown in FIG. 2 is illustrative of therapy delivery circuitry that may be included in an ICD or other implantable medical device employing the atrial arrhythmia detection technique disclosed herein when the device is configured for providing cardiac pacing, cardioversion and defibrillation therapies. For example, the pacer timing and control circuitry 212 may include programmable digital counters which control the basic time intervals associated with various single, dual or multi-chamber pacing modes or anti-tachycardia pacing therapies delivered in the atria or ventricles. Pacer timing and control circuitry 212 also sets the amplitude, pulse width, polarity or other characteristics of the cardiac pacing pulses under the control of microprocessor 224.

During pacing, escape interval counters within pacer timing and control circuitry 212 are reset upon sensing of R-waves or P-waves as indicated by signals on lines 202 and 206, respectively. In accordance with the selected mode of pacing, pacing pulses are generated by atrial pace output circuit 214 and ventricular pace output circuit 216. The pace output circuits 214 and 216 are coupled to the desired electrodes for pacing via switch matrix 208. The escape interval counters are reset upon generation of pacing pulses, and thereby control the basic timing of cardiac pacing functions, including anti-tachycardia pacing.

The durations of the escape intervals are determined by microprocessor 224 via data/address bus 218. The value of the count present in the escape interval counters when reset by sensed R-waves or P-waves can be used to measure R-R intervals and P-P intervals for detecting the occurrence of a variety of arrhythmias. Microprocessor 224 may also track the number of pacing pulses delivered, particularly the number of ventricular pacing pulses delivered, during predetermined time periods as a factor used in classifying the cardiac electrical signal during the time period.

The microprocessor **224** includes associated read-only memory (ROM) in which stored programs controlling the operation of the microprocessor **224** reside. A portion of the random access memory (RAM) **226** may be configured as a number of recirculating buffers capable of holding a series of measured intervals for analysis by the microprocessor **224** for predicting or diagnosing an arrhythmia.

In response to the detection of tachycardia, anti-tachycardia pacing therapy can be delivered by loading a regimen from microprocessor **224** into the pacer timing and control circuitry **212** according to the type of tachycardia detected. In the event that higher voltage cardioversion or defibrillation pulses are required, microprocessor **224** activates the cardioversion and defibrillation control circuitry **230** to initiate charging of the high voltage capacitors **246** and **248** via charging circuit **236** under the control of high voltage charging control line **240**. The voltage on the high voltage capacitors is monitored via a voltage capacitor (VCAP) line **244**, which is passed through the multiplexer **220**. When the voltage reaches a predetermined value set by microprocessor **224**, a logic signal is generated on the capacitor full (CF) line **254**, terminating charging. The defibrillation or cardioversion pulse is delivered to the heart under the control of the pacer timing and control circuitry **212** by an output circuit **234** via a control bus **238**. The output circuit **234** determines the electrodes used for delivering the cardioversion or defibrillation pulse and the pulse wave shape.

In some examples, the ICD **10** may be equipped with a patient notification system **250**. Any patient notification method known for use in implantable medical devices may be used such as generating perceivable twitch stimulation or an audible sound. A patient notification system may include an audio transducer that emits audible sounds including voiced statements or musical tones stored in analog memory and correlated to a programming or interrogation operating algorithm or to a warning trigger event as generally described in U.S. Pat. No. 6,067,473 (Greeninger et al.), incorporated herein by reference in its entirety. In some examples, ICD **10** provides a response to an atrial tachyarrhythmia detection by generating a patient notification via system **250** and/or a clinician notification using telemetry circuit **330**. An atrial tachyarrhythmia response provided by ICD **10** may include determining an AF burden as the total combined duration of all detected AF episodes during a predetermined monitoring time interval, e.g., 24 hours, and generating a patient notification and/or clinician notification when the AF burden exceeds a threshold.

In the following description, AF detection techniques are described with reference to the circuitry of FIG. 4 of ICD **10** of FIG. 1 or ICD **110** of FIGS. 2A and 2B. It is to be understood, however, that the methods and techniques of the descriptions that follow may be implemented in a cardiac monitoring device such as the device of FIG. 3 or even an intracardiac pacemaker such as the intracardiac pacemaker **110** of FIGS. 2A and 2B, all of which devices may include a microprocessor, memory and sensing circuitry, as generally described in conjunction with FIG. 4, for performing these atrial tachyarrhythmia detection techniques, particularly for detecting AF.

FIG. 5 is a schematic diagram of methods used for detecting AF by a medical device, such as ICD **10**, ICD **110**. In order to determine whether AF is occurring, the microprocessor **224** (FIG. 4) may determine differences between RRIs based on sensed R-waves (e.g., R OUT signal line **202** in FIG. 4). Microprocessor **224** may make the decision as to whether an AF event is occurring based at least in part on the resulting pattern or signature of RRI differences. As

described below, when the resulting signature of RRI differences acquired over a predetermined time period indicates AF is occurring, the cardiac signal time period is classified as AF. AF is detected when a required number of time periods are classified as AF. Techniques disclosed herein may be utilized as part of an overall tachyarrhythmia detection and discrimination algorithm implemented in ICD **10** or ICD **110** or the other devices described above or in other implantable or external cardiac devices.

The concept of using a signature of RRI differences for detecting AF is illustrated by the generation of a Lorenz scatter plot as shown in FIG. 5. Microprocessor **224** determines the differences between consecutive pairs of RR-intervals (δ RRs) which can be plotted for a time series of RRIs. The Lorenz plot **150** is a Cartesian coordinate system defined by δ RR_{*i*} along the x-axis **152** and δ RR_{*i-1*} along the y-axis **154**. δ RR_{*i*} is the difference between the *i*th RRI and the previous RRI, RRI_{*i-1*}. δ RR_{*i-1*} is the difference between RRI_{*i-1*} and the previous RRI, RRI_{*i-2*}. As such, each plotted point in a Lorenz plot is defined by an x-coordinate equaling δ RR_{*i*} and a y-coordinate equaling δ RR_{*i-1*}.

Each data point plotted on the Lorenz plot **150** represents an RRI pattern relating to three consecutive RRIs: RRI_{*i*}, RRI_{*i-1*} and RRI_{*i-2*}, measured between four consecutively sensed R-waves **172**. RRI information is not limited to detection of R-waves and determination of RRIs. The terms RRI and δ RR_{*i*} as used herein refer generally to a measurement of ventricular cycle length (VCL) and the difference between two consecutive VCL measurements, respectively, whether the VCL measurements were derived from a series of sensed R-waves from a cardiac electrical signal or a series of ventricular cycle event detections made from another physiological signal (e.g., a peak pressure determined from a pressure signal). For the sake of illustration, the methods described herein refer to R-wave detections for performing VCL measurements and the determination of (δ RR_{*i*}, δ RR_{*i-1*}) points.

As illustrated in FIG. 5, a series of R-waves **170** (represented by vertical bars) are sensed and in order to plot a point on the Lorenz plot area **150**, a point is determined from the ordered pair (δ RR_{*i*}, δ RR_{*i-1*}) by determining successive RRIs determined from the sensed R-waves **170**. In the example shown, a first series **172** of three consecutive RRIs (RRI_{*i-2*}, RRI_{*i-1*} and RRI_{*i*}) provides the first data point **155** on the Lorenz plot area **150**. δ RR_{*i-1*}, which is the difference between RRI_{*i-2*} and is near 0. δ RR_{*i*}, the difference between the RRI_{*i-1*} and RRI_{*i*}, is a positive change. Accordingly, a (δ RR_{*i*}, δ RR_{*i-1*}) point **155** having a y-coordinate near 0 and a positive x-coordinate is plotted in the Lorenz plot **150**, representing the first series **172** of four sensed R-waves (three RRIs).

The next series **174** of three RRIs provides the next (δ RR_{*i*}, δ RR_{*i-1*}) point **156** having a negative x-coordinate (the last RRI of series **174** being less than the immediately preceding RRI) and a positive y-coordinate (the middle RRI of series **174** being longer than the first RRI of series). This process of plotting (δ RR_{*i*}, δ RR_{*i-1*}) points continues with the three cycle series **176** providing data point **158** and so on.

Illustrative methods and devices for detecting AF using RRI differences determined from a ventricular cardiac electrical signal are generally described in commonly assigned U.S. patent application Ser. Nos. 14/520,798, 14/520,938 and 14/520,847 (Cao et al.), all of which are incorporated herein by reference in their entirety. An analysis of a Lorenz plot of the successive RRI differences may reveal an RRI variability pattern that is typical of AF. Other methods for detecting atrial tachyarrhythmias based on the irregularity of

ventricular cycles determined from RRI differences that exhibit discriminatory signatures when plotted in a Lorenz scatter plot, such as the plot shown in FIG. 5, are generally disclosed by Ritscher et al. in U.S. Pat. No. 7,031,765; Sarkar, et al. in U.S. Pat. Nos. 7,623,911 and 7,537,569; and by Houben in U.S. Pat. No. 7,627,368, all of which patents are; incorporated herein by reference in their entirety.

FIG. 6 is a diagram of a two-dimensional histogram representing a Lorenz plot area 150 used in the techniques disclosed herein for detecting atrial tachyarrhythmia. Generally, the Lorenz plot area 150 shown in FIG. 5 is numerically represented by a two-dimensional histogram 180 having predefined ranges 184 and 186 in both positive and negative directions for the δRR_i coordinates (corresponding to x-axis) and δRR_{i-1} coordinates (corresponding to y-axis), respectively. The two-dimensional histogram 180 is divided into bins 188 each having a predefined range of δRR_i and δRR_{i-1} values. In one example, the histogram range might extend from -1200 ms to +1200 ms for both δRR_i and δRR_{i-1} values, and the histogram range may be divided into bins extending for a range of 7.5 ms in each of the two dimensions resulting in a 160 bin $\times$ 160 bin histogram 180. The successive RRI differences determined over a classification time period are used to populate the histogram 180. Each bin stores a count of the number of (δRR_i , δRR_{i-1}) data points falling into each respective bin range. The bin counts may then be used by microprocessor 224 in determining RRI variability metrics and patterns for detecting a cardiac rhythm type.

An RRI variability metric is determined from the histogram bin counts. Generally, the more histogram bins that are occupied, or the more sparse the distribution of (δRR_i , δRR_{i-1}) points, the more irregular the VCL is during the data acquisition time period. As such, one metric of the RRI variability that can be used for detecting AF, which is associated with highly irregular VCL, may take into account the number of histogram bins that have a count of at least one, which is referred to as an "occupied" bin. In one example, an RRI variability metric for detecting AF, referred to as an AF score, is determined by microprocessor 224 as generally described in the above-incorporated '911 patent. Briefly, the AF score may be defined by the equation:

$$\text{AF Score} = \frac{\text{Irregularity Evidence} - \text{Origin Count} - \text{PAC Evidence}}{\text{Evidence}}$$

wherein Irregularity Evidence is the number of occupied histogram bins outside a Zero Segment 188 defined around the origin of the Lorenz plot area. During normal sinus rhythm or highly organized atrial tachycardia, nearly all points will fall into the Zero Segment 188 because of relatively small, consistent differences between consecutive RRIs. A high number of occupied histogram bins outside the Zero segment 188 is therefore positive evidence for AF.

The Origin Count is the number of points in the Zero Segment 188 defined around the Lorenz plot origin. A high Origin Count indicates regular RRIs, a negative indicator of AF, and is therefore subtracted from the Irregularity Evidence term. In addition, a regular PAC evidence score may be computed as generally described in the above-incorporated '911 patent. The regular PAC evidence score is computed based on a cluster signature pattern of data points that is particularly associated with premature atrial contractions (PACs) that occur at regular coupling intervals and present regular patterns of RRIs, e.g., associated with bigeminy (short-short-long RRIs) or trigeminy (short-short-short-long RRIs). In other embodiments, the AF score and/or other RRI variability score for classifying an atrial rhythm may be

determined by microprocessor 224 as described in any of the above-incorporated '765, '316, '911, '569 and '368 patents. Methods for rejecting noise in determining Lorenz plot points and an AF score are generally disclosed in U.S. Pat. No. 8,639,316 (Sarkar, et al.), incorporated herein by reference in its entirety. Methods for adjusting the AF score based on the presence of ectopy may be used in the techniques disclosed herein and are generally disclosed in U.S. Pat. No. 8,977,350 (Sarkar, et al.), incorporated herein by reference in its entirety. Other techniques that may be used in computing an AF score are generally disclosed in U.S. patent application Ser. Nos. 14/695,135, 14/695,156, 14/695,171 and 14/695,111 (Sarkar, et al.), all filed on Apr. 24, 2015 and incorporated herein by reference in their entirety.

The AF score is compared to an AF score threshold for classifying a predetermined time period of a cardiac signal as AF or non-AF based on the RRI analysis. The AF score threshold may be selected and optimized based on historical clinical data of selected patient populations or historical individual patient data, and the optimal AF score threshold setting may vary from patient to patient. In an illustrative example, the AF score may have a possible range of 0 to 100. The AF score threshold may be set between 25 and 75. If the AF score meets or crosses an AF score threshold, the time period over which the RRIs were collected, and thus the cardiac signal occurring within the time period, is classified as an AF time period. The AF score threshold may be adjusted after classifying at least one time period of the cardiac signal as being AF and the adjusted AF score threshold may be used for classifying subsequent time periods, which may lead to an AF detection. The adjusted AF score threshold is less than the initial AF score threshold and may have a value ranging from 19 to 57 in the example given above where the maximum AF score is 100 and the initial AF score threshold is at least 26 and not more than 75. Thus, the adjusted AF score threshold may be between 65-85% of the initial AF score threshold and, in some instances between 70-75% of the initial AF score.

An AF detection is made when a threshold number of detection time periods are classified as AF. In one example, a single n-second or n-minute time period classified as AF based on the AF score meeting the AF score threshold may result in an AF detection. In other examples, a higher number of time periods, e.g., three two-minute time periods for a total of 6 minutes of the cardiac electrical signal, may be required to be classified as being AF before detecting the heart rhythm as AF. As described below in conjunction with FIGS. 13-15, the threshold number of AF classified time periods required to detect AF may be automatically adjusted by microprocessor 224 based on the history of AF episodes detected in a given patient.

The microprocessor 224 provides a response to the AF detection, which may include withholding, adjusting or delivering a therapy (e.g., withholding ATP or shock therapy for treating a ventricular tachyarrhythmia or delivering an atrial anti-tachyarrhythmia therapy if available), storing cardiac signal data that can be later retrieved by a clinician using external device 40, triggering patient notification system 250, transmitting data via telemetry circuit 330 to alert a clinician, patient or caregiver, and/or triggering other signal acquisition or analysis.

The RRI analysis may continue to be performed by microprocessor 224 after an AF detection is made to fill the histogram during the next detection time period. After each detection time period, the AF score may be re-determined and the histogram bins are re-initialized to zero for the next detection time period. The new AF score (or other RRI

variability metrics) determined at the end of each detection time period may be used to determine if the AF episode is sustained or terminated after the initial AF detection is made.

FIG. 7 is a flowchart 300 of a method for determining a factor for classifying time periods for detecting atrial arrhythmias according to one example. Flow chart 300 and other flow charts presented herein are intended to illustrate the functional operation of ICD 10 or another device performing the disclosed methods, and should not be construed as reflective of a specific form of software, firmware or hardware necessary to practice the methods. It is believed that the particular form of software, firmware and/or hardware will be determined primarily by the particular system architecture employed in the device and by the particular detection and therapy delivery methodologies employed by the device. Providing software, firmware and/or hardware to accomplish the techniques disclosed herein in the context of any modern medical device, given the disclosure herein, is within the abilities of one of skill in the art.

Methods described in conjunction with flow charts presented herein may be implemented in a non-transitory computer-readable medium that includes instructions for causing a programmable processor, such as microprocessor 224, to carry out the methods described. A "computer-readable medium" includes but is not limited to any volatile or non-volatile media, such as a RAM, ROM, CD-ROM, NVRAM, EEPROM, flash memory, and the like. The instructions may be implemented as one or more software modules, which may be executed by themselves or in combination with other software.

As illustrated in FIG. 7, the microprocessor 224 identifies ventricular events at block 301, such as R-waves based on Rout signal line 202, and identifies the ventricular event as being either an intrinsic sensed event Vs or a paced event Vp resulting from pacing being delivered by ICD 10 or 10' (or by ICD 110 or pacemaker 101). Depending upon the number of RR intervals chosen for determining RR interval differences, microprocessor 224 determines whether a predetermined number of events, either a ventricular pacing event Vp or intrinsic ventricular sensed event VS, have been identified at block 302. For example, according to one example, if the desired number of RR intervals for determining successive RR interval differences is three, the predetermined number of events utilized in block 302 would be four events, with the four events forming an event window. If the predetermined number of events has not been reached, "No" branch of block 302, microprocessor 224 determines the next ventricular event, at block 301, and the process is repeated.

Once the predetermined number of events are identified, "Yes" branch of block 302, an event window is identified based on the four events, at block 304, and a determination may be made as to whether the number of the events in the event window that are ventricular pace Vp events is less than or equal to a predetermined pacing event threshold at block 306. For example, according to one example, the pacing event threshold is set as one so that microprocessor 224 determines whether one or less of the identified events in the event window are ventricular pace events. If the number of identified events in the event window that are ventricular pace Vp events is not less than or equal to, i.e., is greater than, the predetermined pacing event threshold, "No" branch of block 306, microprocessor 224 identifies the next event at block 301, and the process is repeated.

If the number of events in the event window that are ventricular pace Vp events is less than or equal to the predetermined pacing event threshold, "Yes" branch of

block 306, microprocessor 224 determines whether each of the RR intervals associated with the events in the current event window are greater than a predetermined interval threshold at block 308. For example, according to one example, microprocessor 224 determines whether each of the RR intervals associated with the events in the event window is greater than 220 milliseconds. If each of the RR intervals associated with the events in the event window are not greater than the predetermined interval threshold, "No" branch of block 308, microprocessor 224 identifies the next event at block 301, and the process is repeated using the next identified event and the resulting next event window.

If each of the RRs associated with the events in the event window are greater than the predetermined interval threshold, "Yes" branch of block 308, microprocessor 224 determines differences between successive RRs associated with the identified events in the event window, block 310. Once the RRI differences for the current event window have been determined at block 308, to populate a Lorenz plot histogram as described above, microprocessor 224 determines whether a predetermined time period has expired at block 312. Microprocessor 224 may set a timer or counter to control acquisition of RRI differences over a predetermined time period at the onset of the method of flow chart 300. In one example, the predetermined time period may be set to two minutes. If the time period has not expired, "No" branch of block 312, microprocessor 224 returns to block 301 to identify the next ventricular event and the process is repeated using the next event and the resulting next event window.

Once the timer has expired, "Yes" branch of block 312, microprocessor 224 determines an AF score at block 314, based on the determined RRI differences during the predetermined time period, e.g., two minutes. The AF score may be determined as described above with respect to FIG. 6 and/or the incorporated patents. As described below in conjunction with FIG. 8, the determined AF score for the predetermined time period is used to classify the time period (and thus the cardiac signal during the time period) as an AF time period, a non-AF time period or an unclassified time period. The stored RRI differences are then cleared and all counters and timers reset at block 316. A timer set to the predetermined time period, e.g., two minutes, is reset. Microprocessor 224 identifies the next ventricular event at block 300, and the process is repeated for the next time period using the next identified events and the next event windows.

FIG. 8 is a flowchart 400 of a method for classifying a predetermined time period according to one example. The example described in flowchart 400 of FIG. 7 will be described in the context of having predetermined time periods that are two minutes in length. However, the techniques described in FIG. 7 or elsewhere throughout this description can be for predetermined time periods that are longer or shorter than two minutes. Once the predetermined time period, e.g., a two-minute time period, has expired and a Lorenz plot has been populated with a point associated with ordered pairs representing each determined RR interval difference determined based on the intervals in each event window occurring during the two-minute time period as described in conjunction with FIGS. 5, 6 and 7, microprocessor 224 determines whether to classify the time period as being either an AF time period, a non-AF time period, or an unclassified time period (i.e., the time period can neither be classified as an AF time period nor a non-AF time period).

For example, microprocessor 224 may analyze one or more of several factors, in any combination or particular order, to make the determination.

As described in conjunction with the example of FIG. 8, among the factors that may be analyzed for classifying the two-minute time period are the number of valid RRI difference pairs, RRI lengths, number of paced beats, number of short intervals, presence of oversensing of ventricular events, presence of T-wave oversensing, detection of a ventricular tachyarrhythmia (e.g., SVT, VT or VF or more generally referred to as "other episodes"), and the AF score. However, microprocessor 224 may analyze only a subset of these factors and/or include other factors.

Microprocessor 224 may determine the number of RRI difference pairs acquired during the two-minute time period at block 401, where each RRI difference pair represents one point of the Lorenz plot. A determination is then made as to whether the total number of RRI difference pairs formed during the two-minute time period is greater than an interval pair threshold at block 402. According to one example, a threshold number of RRI difference pairs is set at 30, though other thresholds may be used. If the total number of RRI difference pairs (representing three consecutive RRIs) during the two-minute time period is less than the threshold, "Yes" branch at block 402, the two-minute time period is determined to be unclassified at block 404. In the example described above in which the threshold is set at 30, the "Yes" branch of block 402 means that less than 30 RRI difference pairs were determined during the two-minute time period, resulting in the Lorenz plot histogram being populated with less than 30 points. An AF score determined from fewer than the threshold number of RRI difference pairs may not yield a reliable AF score for the predetermined time period and therefore is not used to classify the time period as either AF or non-AF.

If the number of RRI difference pairs that are formed during the two-minute time period is not less than the interval pair threshold (30 in the example above), "No" branch of block 402, the interval pairs factor for classifying the time period as either AF or non-AF based on an AF score determined from the RRI difference pairs is satisfied. In other words, using the example above, 30 or RRI difference pairs were determined during the two-minute time period, resulting in the Lorenz plot histogram being populated with 30 or more points. The number of RRI difference pairs obtained during the pre-determined time period is adequate to reliably classify the time period as AF or non-AF based on the AF score.

According to another example, microprocessor 224 may additionally or alternatively determine, at block 410, the total number of RRIs during the predetermined time period that were determined to be less than the interval threshold applied at block 308 of flow chart 300. If more than a threshold number of RRIs, e.g., more than a predetermined number of RRIs or a predetermined percentage of the total number of RRIs occurring during the two-minute time period, are less than the interval threshold, "Yes" branch at block 412, the two-minute time period is determined to be unclassified, at block 404. If the number of RRIs less than the interval threshold does not reach or exceed a predetermined number, e.g., if less than 10 RRIs are less than the interval threshold during the two-minute time period, this RRI length factor is determined not to be satisfied at block 412 ("No" branch of block 412) for classifying the predetermined time interval as unclassified. Based on at least this factor, a classification of either AF or non-AF based on the AF score is warranted.

In order to classify the two-minute time period as either AF or non-AF, microprocessor 224 may determine, at block 414, a short interval count of the total number of RRIs from all of the event windows obtained during the two-minute time period that were less than or equal to a predetermined short interval threshold, such as 120 milliseconds or 130 milliseconds, for example. Microprocessor 224 determines whether the short interval count is greater than a short interval threshold, at block 416, such as 5 short intervals for example. Too many short intervals during the two-minute time period indicates the possibility of ventricular oversensing of non-physiological signals such as EMI or lead noise due to lead fracture. In this situation, the RRIs may be unreliable for determining an AF score and classifying the time period as AF or non-AF based on the AF score.

If the determined short interval count is greater than the short interval count threshold, "Yes" branch at block 416, the two-minute time period is determined to be unclassified at block 404. On the other hand, if the short interval count is less than the short interval count threshold, the time period can be classified based on the AF score, "No" branch at block 416. This short interval count factor minimizes false AF detection due to lead noise oversensing.

Microprocessor 224 may additionally or alternatively determine the number of events identified during the total two-minute time period within all of the event windows that were determined to be ventricular pace Vp events at block 418. A determination is made as to whether the determined number of ventricular pace Vp events identified during all event windows of the two-minute time period is greater than a total ventricular pace Vp event threshold at block 420. According to one example, the total ventricular pace Vp threshold is set as 30 ventricular pace Vp events, though other thresholds may be used.

If the number of ventricular pace Vp events during the two-minute time period is greater than the total ventricular pace Vp event threshold, "Yes" branch of block 420, microprocessor 224 classifies the two-minute time period as unclassified at block 404. Ventricular pacing pulses may include bradycardia pacing pulses and/or ATP pacing pulses and may be delivered by ICD 10 or 110 or by another implanted device, e.g., pacemaker 110. On the other hand, if the determined number of ventricular pace Vp events is not greater than Vp event threshold, "No" branch at block 420, the two-minute time period is not classified as unclassified; a classification of AF or non-AF based on the AF score may be made as long as no other factors lead to a determination of the time period being unclassified.

Microprocessor 224 may be configured to simultaneously evaluate R-waves and RRIs for detecting supraventricular tachycardia (SVT), VT and VF while the AF detection algorithms described herein are operating. ICD 10 may be configured to deliver therapies such as ATP in response to detecting VT. As such, if a ventricular tachyarrhythmia detection, e.g., SVT, VT or VF detection, is being made during or at the expiration of the current time period, as determined at block 421, the current time period is determined to be unclassified at block 404. If no other episode detections are being made, the process may advance to block 422.

The microprocessor 224 may additionally or alternatively determine whether ventricular event oversensing caused by noise was detected during the two-minute time period, at block 422. Detection of oversensing may be performed by microprocessor 224 using an implemented oversensing detection scheme, such as the oversensing detection methods generally described in U.S. Pat. No. 7,333,855 to

Gunderson et. al., incorporated herein by reference in its entirety. If oversensing detection criteria were met or were in the process of being met during the two-minute time period, "Yes" branch of block 422, the two-minute time period is determined to be unclassified at block 404. Detection of oversensing indicates that the RRs may be unreliable for determining an AF score and classifying the time period based on the AF score. If a detection of oversensing was not made or not in the process of being made during the two-minute time period, "No" branch of block 422, an AF or non-AF classification may be made based on the AF score as long as other factors do not lead to the time period be classified as unclassified.

Microprocessor 224 may determine whether T-wave oversensing occurred during the two-minute time period at block 424. The determination of T-wave oversensing may be performed by ICD 10 using an implemented T-wave oversensing detection scheme, such as the T-wave oversensing determination described in U.S. Pat. No. 7,831,304 (Cao, et al.), incorporated herein by reference in its entirety. If a determination of T-wave oversensing was made or was in process during the two-minute time period, "Yes" branch of block 424, the T-wave oversensing factor is satisfied as an indication of the two-minute time period being unclassified. Microprocessor 224 classifies the two-minute time period as unclassified at block 404. If a determination of T-wave oversensing was not made or was not in the process of being made during the two-minute time period, "No" branch of block 424, the T-wave oversensing factor is not satisfied. Microprocessor 224 advances to block 426 to classify the time period based on the AF score.

In this way, ICD 10 or ICD 110 may analyze the cardiac electrical signal over the two-minute time period for one or more of the described factors, which if satisfied would cause the two-minute time period to be classified as "unclassified." In other words, ICD 10 or 110 may analyze all of the described factors or only a subset of the described factors in making this determination. In some examples, if at least one the described factors for identifying the two-minute time period as being unclassified is met, the two-minute time period is classified as unclassified at block 404. If none of the factors evaluated in blocks 401 through 424 are determined to be satisfied according to predetermined criteria, the time period is classified as either AF or non-AF based on the AF score. As such, the AF score is determined based on the populated Lorenz plot histogram as described above. If the AF score is greater than an AF threshold at block 426, the two-minute time period is classified as AF at block 406. On the other hand, if the AF score is not greater than the AF threshold, "No" branch of block 426, the two-minute time period is classified as a non-AF at block 408.

It is understood that the determination of whether the time period is classified as unclassified (block 404), classified as AF (block 406), or classified as non-AF (block 408), may be made in any order, or at the same time, so that the determination of the two-minute time period as being an unclassified time period may be used to override an initial determination of the two-minute time period as being classified as AF or non-AF, or be made prior to determining the AF score for making a classification based on the AF score.

FIG. 9 is a schematic diagram of atrial fibrillation detection that may be performed by a medical device according to one example. The examples described in FIG. 9 will be described in the context of classifying predetermined time periods that are two minutes in length. However, the tech-

niques described in FIG. 9 can be implemented by classifying predetermined time periods that are longer or shorter than two minutes.

As illustrated in FIG. 9, the microprocessor 224 classifies the cardiac signal of each two-minute time period as being either AF, non-AF or unclassified using the method described in conjunction with FIG. 7. The classifications of the time periods are used to detect an AF episode. For example, once a predetermined number of two-minute time periods, such as three time periods, have been classified as AF, the device detects the AF episode. Therefore, as illustrated in the scenario of timing diagram (a) of FIG. 8, once the predetermined number of two-minute time periods, 500, 502 and 504, are classified as AF, microprocessor 224 detects the AF episode at time 505. The microprocessor 224 may track the number of two-minute time periods classified as AF by updating an AF event counter each time a time period is classified as AF.

However, in the scenario illustrated in timing diagram (b), two consecutive two-minute time periods 506 and 508 are classified as being AF, but the next two-minute time period 510 is determined to be unclassified, followed by a subsequent time period 512 being classified as AF. According to one example, microprocessor 224 may ignore the unclassified two-minute time period 510 and detect an AF episode at time 513 once the third time period 512 is classified as AF, so that an AF episode may be detected despite one or more unclassified time periods occurring between time periods classified as AF.

In the timing diagram of scenario (b), at the identification of two-minute time period 506, an AF event counter may be incremented to one. At the identification of subsequent two minute period 508, the AF event counter is incremented to two. At the classification of subsequent two-minute time period 510, since the cardiac signal was determined to be unclassified, the AF event counter remains at a count of two. At the classification of subsequent two-minute time period 512, the AF event counter is incremented to three, and an AF episode is detected in response to the AF event counter reaching the AF detection threshold, which is 3 in this example.

As illustrated in the timing diagram of scenario (c), the classification of one or more time periods as non-AF result in no detection of an AF episode. During the determination of whether the predetermined number of two-minute time periods are classified as AF, the microprocessor 224 updates the AF event counter each time an AF classification is made as described above. For example, upon classification of two-minute time period 514, the AF event counter is incremented to one, and at the classification of subsequent two-minute time period 516, the AF event counter is incremented to two. If two-minute time period 518 were also classified as AF, microprocessor 224 would detect an AF episode, since three two-minute time periods classified as AF would have occurred, e.g., as described in the timing diagram of scenario (a) above. However, since two-minute time period 518 is classified as non-AF, an AF episode detection is not made. The non-AF classification of time period 518 may be evidence that an AF episode causing AF classifications of time periods 514 and 516 is terminated or a non-sustained AF episode. In response to classifying time period 518 as non-AF, the AF event counter is reset to zero. In other examples, the AF counter may be decreased when a time period is classified as non-AF rather than immediately reset to zero.

In the timing diagram of scenario (d), at the classification of two-minute time period 522 as being AF, the AF event

counter is incremented to one, and at the classification of subsequent two-minute time period **524** as AF, the AF event counter is incremented to two. At the classifications of subsequent two-minute time periods **526** and **528**, both determined to be unclassified, the AF event counter remains unchanged at a count of two. Upon classification of subsequent two-minute time period **530** as being AF, the AF event counter is increased to three, and an AF episode is detected at time **531**.

Had any of time periods **524**, **526**, **528** or **530** been classified as non-AF, the AF event counter would have been reset to zero, and the process repeated starting with the next classified two-minute time period. However, in addition to resetting the AF event counter in response to a two-minute time period being classified as a non-AF time period, microprocessor **224** may also be configured to reset the AF event counter to zero if a predetermined number of two-minute time periods are determined to be unclassified. For example, five consecutive two-minute time periods determined to be unclassified may cause the AF event counter to be reset to zero. In other examples, more than five or fewer than five unclassified time periods, which may not be required to be consecutive, may cause the AF counter to be decremented or reset to zero. Therefore, in the timing diagram of scenario (e), at the identification of two-minute time period **532** as AF, the AF event counter is incremented to one. At the identification of subsequent two-minute time period **534** as AF, the AF event counter is incremented to two. At the identification of the four subsequent two-minute time periods **536**, **538**, **540** and **542**, all determined to be unclassified, the AF event count remains unchanged at two. In the example shown, the next two-minute time period **546** is classified as AF. The AF event counter is incremented from two to three, and an AF episode is detected at **545** by microprocessor **224** in response to the AF event counter reaching the threshold count, which is three in this example.

On the other hand, if the subsequent two-minute time period **546** had been determined to be unclassified, the AF event counter would be reset to zero in response to a threshold number (five in this example) of consecutive unclassified time periods. If time period **546** had been classified as a non-AF time period, the AF event counter would also be reset to zero. In either of these two cases, if the time periods **532** and **534** represent a true AF episode, the AF episode has terminated or is non-sustained as evidenced by the unclassified and/or non-AF classified time periods. The process is repeated starting with the next classified two minute interval.

FIG. **10** is a schematic diagram of a method for detecting atrial fibrillation that may be performed by ICD **10** or ICD **110** (or cardiac monitoring device **60**) according to another example. The examples described in FIG. **10** will be described in the context of having predetermined time periods that are two minutes in length and an AF detection threshold set equal to 3 time periods classified as AF. However, the techniques described may utilize different time period durations and/or different thresholds. As described below in conjunction with FIGS. **13-15**, the threshold number of time periods, or total time duration of the cardiac signal required to be classified as AF may be automatically adjusted by microprocessor **224**.

The microprocessor **224** classifies the cardiac signal within each two-minute time period as being either AF, non-AF or unclassified using the method described in conjunction with FIG. **8**. If factors that cause the two-minute time period to be unclassified are not satisfied, each two-minute time period is classified as AF or non-AF based on

the AF score. In the method of FIG. **10**, the threshold that the AF score is compared to for classifying a time period is not a fixed value but is dynamically adjusted by microprocessor **224** in response to classifications of two-minute time periods.

For example, once a predetermined number of time periods, such as one time period, has been classified as AF based on a first AF score threshold value, if the next predetermined number of time periods are classified as any combination of AF and/or unclassified, the AF score threshold is adjusted to a second, lower value. In other words, following an initial AF classification using the first, higher AF score threshold, microprocessor **224** decreases the AF score threshold to the second lower value at the expiration of a predetermined number of next consecutive time periods, e.g., two consecutive time periods following the initial AF classification in the examples illustrated in FIG. **9**, as long as none of the predetermined number of next consecutive time periods are classified as non-AF. In other instances, the predetermined number of next consecutive time periods may be less than two, e.g., zero or one, or more than two. If any of the predetermined number of next consecutive time periods following an initial AF classification are classified as non-AF, the AF score threshold remains at the first higher threshold value.

Therefore, as illustrated in the scenario of timing diagram (a) of FIG. **10**, an initial time period **610** is classified as AF based on the AF score determined for time period **610** being greater than a first AF score threshold **602** and analysis of other classification factors does not lead to an unclassified time period (as described with FIG. **8**). The first AF score threshold **602** stays in effect for at least two more consecutive time periods **612** and **614** in this example. If both of these time periods are also classified as AF, in response to an AF score exceeding the first AF score threshold **602**, AF is detected at time **622**. Additionally, microprocessor **224** adjusts the AF score threshold to a second, lower AF score threshold **604** at time **622**. The AF score of subsequent time periods will be compared to this lower threshold **604** for classifying the respective time periods.

The lower AF score threshold **604** may be set to a percentage of the initial AF score threshold **602**, e.g., approximately 75% of the initial AF score. To illustrate, when the maximum possible value of the AF score is 100, the first AF score threshold may be set at 75 and adjusted to a second, lower AF score of 57. In another example, the first AF score threshold is 60 and the second is 45. In still other examples, the first AF score threshold is 60 and the second is 45, the first is 50 and the second is 38, the first is 40 and the second is 30, or the first is 25 and the second is 19. A user may program the AF score thresholds based on selection of a least sensitive, e.g., first threshold of 75 and second threshold of 57, to most sensitive, e.g., first threshold 25 and second threshold 19, with the other example given above corresponding to a less sensitive setting (first threshold 60 and second threshold 45), balanced sensitivity (first threshold 50 and second threshold 38), and more sensitivity (first threshold 40 and second threshold 30). In other examples, the actual values of the first and second thresholds may be programmable selected individually in any combination of a first range, e.g., from 25 to and including 75 for the first threshold, and a second range, e.g., from 19 to and including 57 for the second threshold, as long as the first threshold is greater than the second threshold value. In other examples, the second threshold may be set to be another percentage of the initial threshold, e.g., between 65-85%, 70-80%, or some other percentage.

By reducing the AF score for subsequent time periods, AF detection sensitivity is increased at appropriate times. AF detection specificity is maintained by using the first higher AF score threshold and applying the factors that lead to unclassified time periods. For example, the next two consecutive time periods **616** and **618** are both classified as AF based on an AF score exceeding the second threshold **604**, even though the first, higher threshold **602** is not met (and factors leading to an unclassified classification are not present). The AF episode is detected as still being in progress during time periods **616** and **618** even though the AF scores for these time periods **616** and **618** are each less than the first threshold **602**. The next time period **620** is classified as non-AF due to an AF score being less than the second threshold **604**. In response to the non-AF classification, microprocessor **224** adjusts the AF score threshold from the lower threshold **604** back to the higher threshold **602** at time **624**. Termination of the AF episode is detected in response to the non-AF classification. The AF episode duration **615** is the time interval from the start of the earliest time period **610** classified as AF that led to AF detection at time **622** to the end of last AF classification time period **618** that precedes termination detection at time **624**, i.e., that precedes the time period **620** classified as non-AF.

In the scenario illustrated in timing diagram (b), two consecutive two-minute time periods **632** and **634** are determined to be unclassified after an initial time period **630** that is classified as AF based on the first AF score threshold **602**. In response to neither of the two time periods **632** and **634** being classified as non-AF following the initial AF classification of time period **630**, microprocessor **224** adjusts the AF score threshold at time **642** to the second lower threshold **604**. Three consecutive classifications including at least an initial AF classification and no non-AF classifications cause an adjustment of the AF score threshold. As such, in one example, time periods **632** and **634** immediately and consecutively following the initial AF classified time period **630** may both be unclassified (as shown in this example), both be classified as AF, or one classified as AF and one unclassified to cause the AF score threshold to be adjusted at time **642**.

Since only one time period **630** has been classified as AF in the example shown, however, an AF detection is not made at time **642** when the AF score threshold is adjusted. The next two time periods **636** and **638** are classified as AF in response to an AF score being greater than the adjusted AF score threshold **604** (and factors that would cause an unclassified classification to be made not being identified). When the AF event counter reaches a count of three at time **643**, an AF detection is made. The next time period **640** is classified as non-AF in this example. Termination of the AF episode is detected, and the AF score threshold is adjusted from the lower value **604** back up to the higher value **602** at time **644** in response to the non-AF classification and resulting episode termination detection.

The episode duration **635** starts with the earliest time period **630** that was classified as AF and led to AF detection at time **643** and extends through the latest AF-classified time period **638** prior to termination detection at time **644**. The episode duration **635** includes unclassified time periods **632** and **634** that do not lead to detection of termination at time **644**. Unclassified time periods **632** and **634** occur between AF-classified time periods **630** and **636** and are therefore included in AF episode duration **635**. The time periods **632** and **634** may be classified as unclassified due to any of the other factors described in FIG. 8. In one particular example, one or both of the time periods **632** and **634** may be unclassified due to ventricular tachyarrhythmia detection

(block **421** of FIG. 8). By allowing time periods **632** and **634** to be classified as unclassified when ventricular tachyarrhythmia is being detected, the detection of AF and determination of the AF episode duration **635** are uninterrupted. The detection of an AF episode that is concurrent with a ventricular tachyarrhythmia episode provides important diagnostic information for the clinician to use in properly determining the patient's heart rhythm status and subsequent treatment.

In scenario (b) and other scenarios that follow, the time periods determined to be unclassified, e.g., time periods **632** and **634** are represented as having AF scores being equal to the currently set AF score threshold. It is to be understood, however, that an actual AF score, if determined, may be greater than, equal to, or less than the current value of the AF score threshold but is not used to classify the time period when the analysis of other factors cause the time period to be determined as unclassified as described in conjunction with FIG. 8. In some cases, if the time period is determined to be unclassified due to analysis of one or more factors as described in conjunction with FIG. 8, determination of an AF score for the current time period may not be made; classification of the time period as unclassified may preclude the need to determine the AF score in some examples.

In scenario (c), the AF score threshold is adjusted from a first threshold **602** to a second threshold **604** at time **674** after an initial AF classified time period **650** based on the first, higher threshold **602** followed by two consecutive time periods **652** and **654** that do not include a non-AF classification. Microprocessor **224** may increment an AF event counter in response to each AF classification and increment an unclassified event counter in response to each unclassified time period. Accordingly, in the example of scenario (c), at time **674** the AF event counter is at a count of one, and the unclassified event counter is at a count of two. After time period **656**, the unclassified event counter is at a count of three, and after time period **658** the unclassified event counter is at a count of four. The next two time periods **670** and **672** are classified as AF based on the respective AF scores exceeding the second, lower AF score threshold **604**. If the next time period **670** is also an unclassified time period, such that five unclassified time periods occur consecutively, the AF event counter and the unclassified event counter may be reset to zero, and the AF score threshold may be increased to the first, higher AF score threshold **602**. Microprocessor **224** may therefore adjust the AF score threshold and reset counters in response to detecting a predetermined number of consecutive unclassified time periods.

In the example shown, the next time period **670** is classified as AF so the unclassified event counter remains at a count of four. The AF event counter is increased to two after time period **670** and to three after time period **672**. AF is detected at time **676** in response to the AF event count reaching the detection threshold, which is three in this example. The AF score threshold remains at the second, lower threshold **604** until termination of the AF episode is detected in response to a non-AF classification, e.g., time period **675**, or a predetermined number of consecutive unclassified time periods, e.g., five consecutive unclassified time periods. At time **678**, the AF score threshold is adjusted to the first, higher threshold **602** in response to the non-AF classification of time period **675**. As shown by the example of scenarios (b) and (c), an AF episode may be detected after the AF threshold is adjusted to the second, lower threshold.

The episode duration **655** in scenario (c) begins with AF time period **650** and extends through AF time period **672**

which led to AF detection at time 676. This episode duration 655 includes the consecutive unclassified time periods 652, 654, 656, and 658 which do not lead to detection of AF termination at 678.

Scenario (d) shows another example of a series of two-minute time period classifications and the corresponding adjustment to the AF score threshold. An initial time period 680 is classified as AF based on the first, higher AF score threshold 602. The AF event counter is increased to a count of one. The next time period 682 is classified as non-AF based on the first AF score threshold 604. The AF event counter may be reset to zero in response to the non-AF classification. The unclassified time period 684 may not be counted by microprocessor 224 since the AF event counter is currently zero.

A subsequent sequence of AF-U-AF (time periods 686, 688 and 690, respectively) result in an AF event count of two and an unclassified event count of one. The two consecutive time periods 688 and 690, following the AF time period 686 and classified as unclassified and AF, respectively, result in a combined event count of the AF and unclassified time periods being equal to three. In response to this combined event count of three, microprocessor 224 adjusts the AF score threshold from the first, higher AF score threshold 602 to the second lower AF score threshold 604 at time 695. AF is not yet detected because the AF event count is two. The next time period 692 is classified as AF based on a comparison of the AF score to the second, lower AF score threshold 604. Microprocessor 224 increases the AF event count to three and detects AF at time 697 in response to the AF event count reaching the detection threshold. Upon detecting AF at time 697, the unclassified event counter is reset to zero. The unclassified event counter will count unclassified time segments beginning from zero after the AF detection in order to count consecutive unclassified time periods for detecting termination of the AF episode. The unclassified event count reaches two after time periods 694 and 696. The next time period 698 is classified as non-AF resulting in detecting termination of the AF episode at time 699. All event counters are reset to zero, and the AF score threshold is adjusted back to the first, higher threshold 602 at time 699.

The episode duration 685 starts with AF-classified time period 686 which is the earliest AF classified episode that led to AF detection at time 697. The episode duration includes unclassified time period 688 which did not contribute to detection of termination at block 699. Episode duration 685 ends with the last AF-classified time period 692 prior to detecting termination at time 699. Unclassified time periods 694 and 696 may, in some instances, not be included in the episode duration 685 because they immediately precede the non-AF classified time period 698 that results in episode termination detection with no intervening AF-classified episode. In other instances, however, those unclassified time periods may also be included in the episode duration 685.

The first and second AF score thresholds 602 and 604 may be fixed values or may be programmable by a user. In one example, a user may program the first and second AF score thresholds 602 and 604 to be increased or set to relatively higher respective values than currently programmed values or decreased or set to relatively lower values than the currently programmed values. Both the first and second AF score thresholds are adjusted together up or down by the same increment or decrement respectively, in response to the user-entered programming command. In other examples, a user may programmably select each of the first and second AF score thresholds tailored to individual patient need.

While only two different AF score thresholds 602 and 604 are illustrated in the example of FIG. 10, it is understood that the AF score threshold may be adjusted between three or more AF score threshold values in other examples. For instance, after AF detection is made at time 697, the AF score threshold set to the second, lower threshold 604 could be reduced to a third lowest AF score threshold to allow continuing detection of the AF episode using less stringent criteria than the initial AF detection criteria. In other examples, once the AF detection is made at time 697, the AF score threshold could be increased from the second, lower threshold 604 back up to the first, higher threshold 602 or to a third, intermediate threshold value between the second, lower threshold 604 and the first, higher threshold 602. The third intermediate threshold value may be applied for classifying subsequent time periods until termination of the AF episode is detected based on a predetermined number of time periods being classified as non-AF based on an AF score falling below the third intermediate threshold value.

In the example scenarios of FIG. 10, the AF score threshold is adjusted after at least two consecutive time period classifications of any combination of AF and unclassified immediately follow a preceding or initial AF classified time period. In other examples, the AF score threshold may be adjusted after fewer or more time periods. For example, a single time period classified as AF may cause the AF score threshold to be reduced to a second lower value. In other examples, at least one unclassified or AF time period following an immediately preceding AF time period may cause the AF score threshold to be adjusted. In still other examples, more than two time periods that are not classified as non-AF and consecutively follow a first time period classified as AF may be required before adjusting the AF score threshold.

FIG. 11 is a flowchart 700 of a method for detecting atrial fibrillation, according to one example. At block 702, the classification of the current time period is determined. If the classification is AF, as determined at block 704, microprocessor 224 increases the AF event counter at block 706 and advances to block 714 to compare a combined count of the AF event counter and the unclassified event counter to a threshold. If the combined counts do not meet the threshold at block 714, the AF event count is compared to the AF detection threshold at block 718. If the AF detection threshold is not reached, the process returns to block 702 to get the next time period classification.

If the time period is not classified as AF, "No" branch of block 704, and is classified as non-AF, "Yes" branch of block 708, the microprocessor 224 advances to block 742. The AF classification criteria, if previously adjusted, are restored to initial values. For example, if an AF score threshold has been previously adjusted to a second lower threshold, the AF score threshold is returned to a higher first threshold as described in conjunction with FIG. 10. At block 746, the AF event counter and the unclassified event counter are reset to zero if they have been previously incremented to a non-zero value.

If the current time period classification obtained at block 702 is neither AF nor non-AF, i.e., if the current time period is determined to be unclassified, "No" branch of block 708, and the AF counter is currently inactive with a value of zero, "No" branch of block 710, microprocessor 224 determines the classification of the next time period at block 702. If the AF event count is greater than zero as determined at block 710, indicating that an initial AF classification has been made, and the current time period is unclassified, microprocessor 224 increases the unclassified event count by one at block 712. The unclassified event count may be used for

controlling adjustment of AF classification criteria prior to an AF detection being made as described in conjunction with FIG. 10. If the unclassified event count has reached a predetermined threshold, "Yes" branch of block 713, microprocessor may restore initial AF classification criteria (if previously adjusted) at block 742 and reset the unclassified event counter and the AF event counter to zero at block 746. The process begins again at block 702 with the classification of the next time period.

After increasing the AF event count at block 706 or increasing the unclassified event count at block 712, if the unclassified event count has not reached the predetermined threshold, "No" of block 713, the combined event count may be compared to a threshold at block 714. When the combined count of the AF event counter and the unclassified event counter has reached a threshold at block 714, e.g., a combined count of three, the AF classification criteria may be adjusted at block 716. In one example, microprocessor 224 adjusts the AF classification criteria by decreasing the AF score threshold to a second lower threshold after classifying a first time period as AF and classifying the next two consecutive time periods as any combination of AF or unclassified based on the first higher AF score threshold as described above. The AF classification criteria may therefore be adjusted in response to three consecutive time periods being classified as AF, a sequence of AF-U-AF or a sequence of AF-U-U.

It is to be understood that in some examples once the combined count reaches a predetermined threshold at block 714, and the AF classification criteria have been adjusted at block 716 prior to an AF detection being made, the AF classification criteria are not adjusted again until AF episode termination is detected, e.g., based on a time period classified as being non-AF (block 708) or based on a predetermined number of unclassified time periods (block 713), e.g., five consecutive unclassified time periods. In other examples, additional adjustments to the AF score may be made before AF episode termination is detected, e.g., to a third AF score threshold or back to the first, highest AF score threshold, as described above.

At block 718, microprocessor 224 compares the AF event count to the AF detection threshold. When the AF detection threshold has not been reached, microprocessor 224 returns to block 702 to obtain the next time period classification. As described above, after adjusting the AF classification criteria at block 716, if a non-AF classification is made ("Yes" branch of block 708) before detecting AF, the AF classification criteria are restored to the initial classification criteria at block 742 and all AF event and unclassified event counters are reset to zero at block 746. If subsequent time periods are classified as AF, "Yes" branch of block 704, the AF event count is increased accordingly at block 706.

If the AF event count reaches a detection threshold, "Yes" branch of block 718, microprocessor 224 detects AF at block 720. An AF detection response is provided at block 721. The response to AF detection may include controlling pace timing and control 212 to deliver an atrial anti-arrhythmia therapy or withhold a ventricular therapy. The response to AF detection may additionally or alternatively include storing data relating to the AF episode, such as the time of onset, the total duration (as determined from the AF event counter upon detection of termination of the AF episode as discussed below or computed using the techniques described in conjunction with FIG. 10), storing an episode of the cardiac electrical signal in RAM 226 and/or other data relating to the AF event. The data may be transmitted to external device 40

(FIG. 1) for displaying or communicating the data to a clinician for use in managing the patient.

When an AF detection is made at block 720, the unclassified event counter is reset to a count of zero at block 722. Microprocessor 224 may begin counting subsequent time periods determined to be unclassified after AF detection is made for detecting termination of the AF episode. The next time period classification is obtained at block 724. If the next time period classification is AF, as determined at block 726, the AF event count is increased at block 728. AF classifications made after AF detection are based on the adjusted AF classification criteria. The AF event counter may continue to be increased with each AF classification made after detecting AF at block 720 for use in determining the duration of the AF episode and determining AF burden (e.g., the combined duration of all detected AF episodes over a given monitoring interval such as 24 hours). Such AF episode data may be transmitted to an external medical device for display or communication to a clinician thereby providing useful information to the clinician in making diagnostic and therapy management decisions.

If the classification of the next time period is not AF, "No" branch of block 726, but is non-AF, "Yes" branch of block 730, termination of the AF episode is detected at block 740. If the classification of the next time period is neither AF nor non-AF, "No" branch of block 730, i.e., if the time period is determined to be unclassified as indicated at block 732, the unclassified event counter is increased by one at block 734. The unclassified event counter is compared to a threshold at block 736. If the threshold is not reached, microprocessor 224 returns to block 724 to fetch the next time period classification. If the unclassified count reaches a threshold at block 736, e.g., five consecutive unclassified time periods, termination of the AF episode is detected at block 740.

If episode termination is detected, the initial AF classification criteria are restored at block 742, and the AF event and unclassified event counters are reset at block 746. The process begins again at block 702.

FIG. 12 is a flow chart 800 of a method performed by ICD 10 or ICD 110 for providing a response to detecting AF according to one example. If an AF detection is made at block 802, e.g., as described in conjunction with any of FIGS. 9-11, an AF episode record is stored at block 804. The AF episode record may include a Lorenz plot or histogram of RRI data that led to AF classifications and AF detection. The episode record stored at block 804 may further include the start time, termination time, and total duration of the AF episode. Referring again to FIG. 10, examples of episode durations 615, 635, 655, and 685 are shown extending from the start of the first respective AF classified time period that led to AF detection and ending with the last AF classified time period that leads to detection of termination of the AF episode. Unclassified episodes that lead to termination detection, e.g., unclassified episodes 694 and 696 in scenario (d) may not be included in the episode duration, whereas unclassified episodes that do not immediately precede termination detection, e.g., unclassified episodes 652, 654, 656 and 658 in scenario (c), are included in the episode duration, e.g., duration 655.

When unclassified time periods are included in the AF episode or lead to termination detection of the AF episode, the factor(s) leading to determining the time period as being unclassified may be stored with the AF episode record. For example, if the time period is determined to be unclassified due to other episodes being detected such as VT or VF, due to oversensing, due to too many short RRIs, or due to too many ventricular pacing pulses during the time period, this

factor may be stored to provide the clinician with useful information in diagnosing the patient's heart rhythm status for guiding therapy decisions for treating the patient's AF.

At block **806**, microprocessor **224** may determine AF burden of the patient by computing the total time AF was identified over a 24-hour time interval (or other predetermined monitoring interval). Computation of the AF burden may include counting or summing all time periods classified as AF or counting or summing only AF classified time periods that were included in a detected AF episode. AF burden may also include all unclassified time periods that occur during a detected AF episode. In some examples, unclassified time periods that lead to detecting termination of the AF episode that are not included in the AF episode duration are not included in the AF burden computation. For example, referring to FIG. **10**, scenario (d), unclassified time period **684** is not included in AF burden computation because it occurs before the AF episode indicated by episode duration **685**. Unclassified time period **688** is included in AF burden determination because it occurs during the AF episode. Unclassified time periods **694** and **696** are not included in determining AF burden because they lead to termination detection at time **699** and are not included in the episode duration **685**.

When AF is detected, microprocessor **224** may store a cardiac signal segment that is acquired during the detected AF episode. The cardiac signal segment is stored in memory at block **808** with the normal gain of sense amplifiers **200** and A/D converter **222** used during cardiac signal analysis and processing performed to identify RRLs, analyze the signal for oversensing, etc. For example, the cardiac signal stored at block **808** may be an EGM signal acquired using RV coil electrode **28** and ICD housing **15** in FIG. **1**. In system **100** of FIG. **2A**, the cardiac signal stored at block **808** may be an ECG signal acquired using defibrillation electrode **24** or defibrillation electrode **126** and housing **115**. The signal stored at normal range (e.g., with 8-bit resolution sampled at 128 Hz with A/D converter input range of ± 12 mV) may be used to provide an unclipped EGM or ECG signal for morphology analysis (e.g., wavelet template matching) and for storing unclipped cardiac signal episodes in response to detecting a tachyarrhythmia. The normal range signal stored at block **808** may be selected as a far-field or relatively global cardiac electrical signal that is used to produce a display of the electrical rhythm of the patient's heart clearly showing R-wave morphology and regularity of RRLs for the clinician to see a high level view of the signal and the patient's corresponding rhythm. However, depending on the sensing vector, the normal range cardiac signal stored at block **808** may not include observable or easily observed P-waves.

As such, when AF is detected, the microprocessor **224** stores a second cardiac signal in memory at block **810** with a lower range, higher gain setting, e.g., a range of ± 2 mV which may be controlled by adjusting the A/D converter input range. The high gain, lower range setting provides a clearer view of P-waves in the stored cardiac signal segment when displayed by external device **40**. The high gain, lower range setting may result in clipping of R-waves in the stored second cardiac signal. However, the first cardiac signal stored at normal range provides a reliable, unclipped display of the R-wave morphology.

The second cardiac signal stored with a high gain, lower range setting may be selected as a second far-field or relatively global signal. For example, in system **1** of FIG. **1**, the second cardiac signal may be acquired between the RV coil electrode **24** and the SVC coil electrode **26**, or the SVC

coil electrode **26** to the ICD housing **15**. In the system **100** of FIG. **2A**, the second cardiac signal may be acquired using the sensing electrode **128** and sensing electrode **130**, one of sensing electrodes **128**, **130** or **131** paired with one of defibrillation electrodes **124** or **126**, or one of electrodes **124**, **126**, **128**, **130** or **131** paired with housing **115**.

When ICD **10** or ICD **110** receives an interrogation command from an external device **40**, microprocessor **224** controls telemetry circuit **330** to transmit the AF episode record, AF burden information, and the stored first, normal gain cardiac signal and the second higher gain cardiac signal. The external device is configured to generate a display of the AF data for the patient to provide the clinician with valuable diagnostic information to support therapy decision-making processes.

If AF is not being detected, "No" branch of block **802**, microprocessor **224** may be detecting VT or VF at block **820**. If not detecting VT or VF, "No" branch of block **820**, microprocessor **224** continues monitoring for cardiac tachyarrhythmias at blocks **802** and **820**. If AF is not being detected at block **802** but microprocessor **224** is detecting VT or VF, "Yes" branch of block **820**, the VT or VF episode record is stored at block **822**, which may include rate, duration, start time, end time, delivered therapies and results, etc. At block **824**, a segment of the first cardiac signal with normal gain acquired during the detected episode is stored. The first cardiac signal at normal gain may be a far-field or relatively global signal as described above that provides a high level view of R-wave morphology and RRLs and may be the same signal with the same gain that is stored at block **808** in response to AF detection. At block **826**, a third cardiac electrical signal is stored with normal gain.

The third cardiac electrical signal may be a near-field or relatively local signal acquired with a different sensing vector than either of the first or second cardiac electrical signals stored at blocks **808** and **810**. For example, the third cardiac signal stored at block **826** may be an EGM signal acquired using RV tip electrode **28** and RV ring electrode **30** of system **1** shown in FIG. **1**. In system **100** of FIG. **2A**, the cardiac signal stored at block **826** may be an ECG signal acquired using sensing electrodes **128** and **130**. The third cardiac electrical signal is stored with normal gain but is acquired using a near-field or localized ventricular sensing vector that can be used to generate a display of a high quality ventricular signal when VT or VF is detected and no atrial tachyarrhythmia is detected.

In this way, ICD **10** or ICD **110** provides a unique response for storing data depending on whether an atrial tachyarrhythmia is being detected (with or without concurrent SVT, VT or VF) or a ventricular tachyarrhythmia is being detected (without concurrent AF detection). Storage of a high gain cardiac electrical signal and transmission to an external device **40** for display to a clinician provides the clinician with valuable diagnostic information relating to the detected AF episode. When the AF episode is being detected simultaneously with a ventricular tachyarrhythmia detection, the relationship between events leading to the two detections can be ascertained. However, when only a ventricular tachyarrhythmia is detected, storage and transmission of a near-field or localized ventricular signal may provide the clinician with important information regarding the ventricular rhythm.

In some examples, microprocessor **224** responds to an AF detection by selecting which cardiac electrical signals are stored as described above and transmits the signals with normal gain and range to external device **40**. Processor **52** of external device **40** may automatically generate a display on

user display **54** that includes the first, high gain, low range signal for observation of P-waves and the second, normal gain, normal range signal for unclipped observation of R-waves.

FIG. **13** is a flow chart **900** of a method performed by an implantable medical device system, such as a system including ICD **10** and lead **16** or ICD **100** and lead **116**, for automatically adjusting a detection threshold required to detect an atrial tachyarrhythmia, such as AF. The detection threshold may be a minimum time duration that the cardiac electrical signal received by sensing circuitry of the ICD must correspond to, or be classified as, the atrial tachyarrhythmia in order to detect the atrial tachyarrhythmia episode. This detection threshold may be defined as a time interval in some atrial tachyarrhythmia detection algorithms. In other examples, the detection threshold may be defined as a threshold number of time predetermined periods required to be classified as AF in order to detect an AF episode, e.g., as generally described in conjunction with FIGS. **9**, **10** and **11**. In the illustrative examples provided herein, the time periods are two minutes long and the threshold number of time periods that are required to be classified as AF in order to detect an AF episode is three time periods. In other words, a total time duration of at least six minutes of the cardiac electrical signal needs correspond to AF, e.g., be classified as AF based on an AF score and other factors, in order to detect AF. The six minutes are not necessarily consecutive since one or more two-minute time periods that are unclassified may occur between the three time periods classified as AF. In the flow chart of FIG. **13**, the microprocessor **224** of ICD **10** or **110** may automatically adjust the detection threshold by adjusting the number of time periods classified as AF that are required in order to detect an AF episode. For example, microprocessor **224** may be configured to automatically adjust the detection threshold by adjusting a count threshold applied to the AF event counter that is used to count the number of time periods that are classified as AF (see blocks **706** and **718** of FIG. **11**).

At block **902**, microprocessor **224** may set an initial detection threshold. For example, the initial detection threshold may be three two-minute time periods, for a total time duration of six minutes of the cardiac electrical signal being classified as AF. The detection threshold may generally represent the minimum duration of time that the cardiac signal is required to present an AF rhythm and, depending on the particular AF detection algorithm implemented in the ICD **10** or **110**, may be expressed as a time interval, e.g., 6 minutes, rather than the value of an AF event counter that counts the number of time periods classified as AF. The initial detection threshold may be set to a relatively low value in order to achieve a relatively high sensitivity for detecting AF episodes. In this way, a patient that is unknown to experience AF, or the characteristics of the AF or AF burden are unknown for a given patient, AF episodes will be detected with relatively high sensitivity initially, e.g., beginning after ICD implantation.

The processor **224** ICD **10** or **110** monitors the heart rhythm at block **904** according to the implemented AF detection algorithm using the initial detection threshold. For example, the heart rhythm may be monitored according to the techniques described above in conjunction with FIGS. **5-11** for detecting an AF episode based on the initial detection threshold of *n* time periods (e.g., three time periods) being classified as AF. If AF is detected at block **906** based on the initial detection threshold, "yes" branch of block **906**, an AF episode counter is increased at block **908**.

Microprocessor **224** determines if detection threshold adjustment criteria are met at block **910**. This determination may be made by comparing the AF episode counter to an AF monitoring threshold count at block **910**. If the count of AF episodes detected based on the initial detection threshold has not reached the AF monitoring threshold count at block **910**, the ICD continues monitoring the heart rhythm at block **904** using the initial detection threshold.

In one example, the AF monitoring threshold count applied at block **910** is a count of three AF episodes detected based on the initial detection threshold. If the AF episode count has not reached a value of three detected AF episodes, the heart rhythm continues to be monitored at block **904** using the initial detection threshold. A relatively sensitive AF detection threshold, e.g., a relatively short total time duration or number of time periods classified as AF continues to be used for detecting AF episodes for identifying how likely the patient is to experience AF and how often.

In other examples, the detection threshold adjustment criteria applied to detected AF episodes at block **910** may include a frequency and/or duration of detected AF episodes. For instance, a threshold count may be required to be reached within a predetermined time period. In one example, at least five AF episodes may be required to be detected within 30 days in order for the detection threshold adjustment criteria to be satisfied at block **910**. In another example, at least one detected AF episode may be required to have an episode duration that is greater than a predetermined threshold duration, e.g., at least 10 minutes. In yet another example, the detection threshold adjustment criteria applied at block **910** may be based on an AF burden threshold. For example, the criteria at block **910** may require that the summed episode durations of AF episodes detected based on the initial detection threshold within a predetermined time period meets an AF burden threshold. For instance, the summed episode durations of detected AF episodes may be required to reach a cumulative duration of at least 15 minutes within a 24-hour period.

The detection threshold adjustment criteria may include multiple criteria, and when any one of the criterion are met, the detection threshold may be adjusted. In one example, the adjustment criteria may include a first threshold number of detected AF episodes, a second threshold number of episodes detected within a predetermined time interval, a threshold episode duration, and/or a threshold AF burden. If any one of these thresholds is reached, the detection threshold may be adjusted. To illustrate, the detection threshold adjustment criteria may be met if the AF episode counter reaches a total count of 5 detected AF episodes; if the AF episode counter reaches a count of 3 detected AF episodes within 30 days; if a detected AF episode has an episode duration of at least 10 minutes; or if the AF burden has reached at least 15 minutes of cumulative AF episode durations within 24 hours. The number of detected AF episodes, the frequency of detected AF episodes, the maximum episode duration of detected AF episodes, and/or the AF burden may be used alone or in any combination for determining if detection threshold adjustment are met at block **910**. The example values of the thresholds given above are intended to be illustrative in nature with no limitation intended. Numerous other examples of combinations of multiple criteria and corresponding threshold values may be conceived based on the examples given herein.

If the detected AF episodes satisfy the detection threshold adjustment criteria at block **910**, microprocessor **224** adjusts the detection threshold at block **912**. The detection threshold may be adjusted to a value that is greater than or less than

the initial detection threshold. In one example, the detection threshold is adjusted to reduce the sensitivity to detecting AF by increasing the minimum time duration or number of AF-classified time periods that are required to detect an AF episode. For instance, if the initial detection threshold is three two-minute time periods, the adjusted detection threshold may be five two-minute time periods required to be classified as AF in order to detect an AF episode. The reduced sensitivity to AF detection may avoid false AF detection and reduce the number of AF episodes for which data is stored in memory of the ICD that requires review or verification by a clinician, easing clinician burden while still capturing important AF episode data.

At block 914, ICD 10 or 110 monitors the heart rhythm for detecting AF according to the implemented AF detection scheme, e.g., as described in conjunction with FIGS. 5 through 11, using the adjusted detection threshold. The longer time duration of the AF detection threshold (e.g., five two-minute time periods instead of three two-minute time periods) may reduce the sensitivity of AF detection but may increase the specificity of AF detection for long-term monitoring of the patient.

In other examples, the AF detection threshold may be adjusted to a lower value at block 912 that increases AF detection sensitivity. In some cases, a clinician may prefer AF detection episodes be detected with greater sensitivity once AF is identified in a given patient based on a less sensitive AF detection threshold. In this case, the AF detection threshold may be reduced to a shorter total time duration, or fewer time periods classified as AF, in order to detect an AF episode.

Microprocessor 224 may execute a response to detecting an AF episode at block 915. The response to an AF episode detection may include one or more of storing AF episode data which may include a segment of the cardiac electrical signal received during the detected AF episode, updating an AF burden, withholding, delaying or adjusting a therapy, and/or withholding delaying or adjusting a ventricular tachyarrhythmia detection.

FIG. 14 is a flow chart 920 of a method for automatically adjusting a detection threshold for detecting AF according to another example. Blocks 902 through 912 correspond to identically-numbered blocks of FIG. 13. After adjusting the detection threshold at block 912 in response to one or more AF episodes detected based on the initial detection threshold satisfying the detection threshold adjustment criteria at block 910, a monitoring period is started at block 916. The monitoring period may be a predetermined time period used to control how long the detection threshold remains at the adjusted value. The monitoring period may be a predetermined number of hours, days or weeks. In one example, the monitoring period is 90 days.

At block 922, the heart rhythm is monitored according to the implemented AF detection techniques, e.g., as described above in conjunction with FIGS. 5 through 11. If an AF episode is detected causing the AF episode counter to be increased, "yes" branch of block 924, microprocessor 224 may determine if second detection threshold adjustment criteria are met at block 928. If the second detection threshold adjustment criteria are not satisfied at block 928, the microprocessor returns to block 922 to continue monitoring the heart rhythm using the detection threshold set at block 912.

During the monitoring period, however, the AF detection threshold may be adjusted from the detection threshold set at block 912 when the number and/or frequency of AF episode detections made based on the adjusted detection

threshold satisfy second threshold adjustment criteria at block 928. For instance, if the AF episode counter reaches a count of two AF episodes detected using the adjusted AF detection threshold, the AF detection threshold may be adjusted again at block 930. In another example, the second detection threshold adjustment criteria applied at block 928 may require that at least two AF episodes are detected within a predetermined time period that is shorter than the monitoring period, e.g., two AF episode detections within thirty days. If the required frequency of AF episode detections occurs during the monitoring period, the AF detection threshold is adjusted at block 930. The second detection threshold adjustment criteria may require a threshold count of AF episode detections based on the adjusted detection threshold, a threshold frequency of AF episode detections based on the adjusted detection threshold, an AF episode duration equal to or greater than a duration threshold, an AF burden equal to or greater than an AF burden threshold, or any combination thereof.

The AF detection threshold may be adjusted to be less sensitive to increase the specificity of AF detection. For instance, the total time duration required to detect AF may be increased, e.g., from 10 minutes or 5 two-minute time periods classified as AF to 20 minutes or 10 two-minute time periods classified as AF. After adjusting the AF detection threshold at block 930, the microprocessor 224 returns to block 916 to start a new monitoring period. The AF episode counter used to track the number of AF episodes detected during a monitoring period may be reset to zero. The microprocessor 224 continues to monitor the heart rhythm at block 922 using the most recently adjusted AF detection threshold.

In some examples, in addition to or alternatively to adjusting the total time duration or number of time periods classified as AF at block 930, the AF score threshold may be adjusted at block 930. The AF score threshold may be set to first and second values as described in conjunction with FIG. 10 as two-minute time periods are being classified. One or both of the first AF score threshold 602 and the second AF score threshold 604 (shown in FIG. 10) may be adjusted at block 930 for use in classifying a two-minute time period. For example, the AF score threshold may initially be set to a first threshold 602 of 50 for classifying an initial time period as AF and reduced to a second threshold 604 of 38. At block 930, the AF score threshold may be increased to less sensitive settings where the first threshold 602 is set to 60 and the second threshold 604 is set to 45.

If the monitoring period expires ("yes" branch of block 926) before any AF episodes are detected based on the adjusted AF detection threshold ("no" branch of block 924), the microprocessor 224 returns to block 912 and resets the detection threshold (if previously adjusted at block 930) to the previous detection threshold used for chronic monitoring at block 912. For example, if a second adjustment of the AF detection threshold is made at block 930 and a new monitoring period is started at block 916, but the monitoring period expires at block 926 with no AF episodes being detected during the monitoring period, the AF detection threshold is returned to the first adjusted value set at block 912. In the illustrative example given above, an initial detection threshold may be 6 minutes set at block 902, which is adjusted to 10 minutes at block 912, and may be adjusted again to 20 minutes at block 930. If no AF episodes are detected during a monitoring period when the AF detection threshold corresponds to a total time duration of 20 minutes of the cardiac electrical signal being classified as AF, the AF

detection threshold may be adjusted back to the 10 minute duration (e.g., 5 two-minute time periods classified as AF).

In some cases, the AF detection threshold is adjusted back to the preceding AF detection threshold value if the monitoring period expires at block 926. Continuing with the illustrative example given above, if the AF detection threshold is currently set at 10 minutes or 5 two-minute time periods classified as AF, the AF detection threshold is adjusted back to 6 minutes or 3 two-minute time periods. In other examples, instead of returning to block 912 when the monitoring period expires with no AF detection, the process may return to block 902 to set the detection threshold to the initial value, e.g., 6 minutes or 3 two-minute time periods in the example given above.

While not explicitly shown in flow chart 920 of FIG. 14, it is to be understood that upon detecting an AF episode at block 906 and/or block 924, microprocessor 224 may control the ICD to provide a response to AF episode detection as described above in conjunction with block 915 of FIG. 13 or any of the AF episode detection responses described in conjunction with block 721 of FIG. 11.

FIG. 15 is a flow chart 950 of a method for detecting AF and determining AF burden by a cardiac medical device according to another example. The techniques described above for adjusting the AF detection threshold may be used to control detection of AF episodes for providing a response of storing AF episode data such as a cardiac signal segment, e.g., as described in conjunction with FIG. 12. Other AF episode detection responses, however, may be made by ICD 10 or ICD 110 based on a different AF detection threshold, which may not be automatically adjustable. Other responses to detecting an AF episode that may be made by ICD 10 or ICD 110 may include determining AF burden. AF burden may be determined by summing the AF episode durations for all AF episodes detected over a 24 hour period or other predetermined time period. In some cases, a different AF detection threshold is used for detecting AF episodes for determining AF burden than the AF detection threshold that is used for detecting AF episodes for which a cardiac electrical signal segment is stored.

In flow chart 950, the blocks 902, 904, 906, 908 and 910 correspond to identically numbered blocks described above in conjunction with flow chart 900 of FIG. 13. However, in addition to determining whether detection threshold adjustment criteria are met at block 910 after increasing the AF episode count at block 908, the AF burden may be updated by microprocessor 224 at block 909 by summing the duration of the AF episode detected at block 906 with the episode durations of any AF episodes detected previously within a predetermined AF burden time interval. For example, the AF burden may be determined and stored for each consecutive 24 hour time interval or for a running 24 hour time interval. In other examples, the AF burden may be determined over another moving predetermined time interval.

If the detection threshold adjustment criteria are met at block 910, as described above in conjunction with FIG. 13, the detection threshold for AF episode storage is adjusted at block 952. The adjustment to the detection threshold at block 952 may be an increase in the total time duration, e.g., defined as the number of two-minute time periods classified as AF as described above in conjunction with FIGS. 13-14. For example, the detection threshold used for triggering AF episode storage may be adjusted from 6 minutes (or 3 two-minute time periods) to 10 minutes (or 5 two-minute time periods) of the cardiac electrical signal being classified as AF.

The detection threshold used for updating the AF burden, however, may be kept the same as the initial detection threshold set at block 902 or adjusted to a detection threshold different than the detection threshold set at block 952 for AF episode storage. The detection threshold for updating the AF burden is set at block 954. In some cases, the detection threshold for updating AF burden is set or maintained at a more sensitive setting than the detection threshold set at block 952 for episode storage. For example, the detection threshold for updating AF burden may be kept at 6 minutes or a count of 3 two-minute time periods classified as AF, and the detection threshold set at block 952 for episode storage is set to a relatively less sensitive setting of 10 minutes or a count of 5 two-minute time periods classified as AF. In this way, the AF burden is determined with relatively high sensitivity and relatively low specificity while AF episodes that may be stored for review by a clinician are detected with relatively higher specificity and relatively lower sensitivity than the AF episodes detected for updating the AF burden.

When an AF episode is detected at block 956 based on the detection threshold set at block 954 (or kept the same as the initial detection threshold), the duration of the detected episode is determined at block 958. The duration of the detected episode may be determined using the methods described above in conjunction with FIGS. 9 and 10. Microprocessor 224 updates the AF burden at block 960 using the determined episode duration. The episode duration may be summed with the duration(s) of previous AF episode(s) that have been detected since the start of the AF burden time period, e.g., since the start of a 24-hour time period.

The process of updating the AF burden at block 960 continues by returning to block 956 to wait for the next AF episode detected based on the detection threshold set at block 954. The detection threshold used for detecting AF episodes used to update the AF burden may not be automatically adjusted by microprocessor 224 in some examples and may remain fixed at the initial detection threshold set at block 902 or set to a user-programmed value. In other examples, the detection threshold used for updating AF burden may be adjusted once at block 954 from the initial detection threshold set at block 902 after the detection threshold adjustment criteria are met at block 910 and not adjusted again. In still other examples, the detection threshold used for updating the AF burden, storing AF episode data such as a cardiac electrical signal segment and/or providing other responses to AF detection by ICD may be adjusted to desired values based on adjustment criteria being met in the manner described above in conjunction with FIGS. 13 and 14.

In the example of FIG. 15, however, the detection threshold is adjusted at block 952 to a relatively less sensitive setting than the AF detection threshold set at block 954 for updating AF burden. The adjusted detection threshold value set at block 652 is used for detecting AF episodes for which a cardiac signal segment is to be stored in RAM 226 (or other computer-readable storage media such as the examples given previously herein). Blocks 916, 922, 924, 926 and 928 of flow chart 950 may correspond to identically-numbered blocks described above in conjunction with FIG. 14. In the process of flow chart 950, microprocessor 224 may be configured to provide a response to an AF episode detected at block 924 by updating stored AF episode data at block 925. Methods for updating AF episode data stored in RAM 224 of ICD 10 or 110 are described below in conjunction with FIG. 16.

At block 960, the detection threshold may be adjusted when second AF detection threshold adjustment criteria are

met at block 928. The adjustment performed at block 960 may correspond to the adjustment described in conjunction with block 930 of FIG. 14, e.g., to a less sensitive setting for detecting AF, but in the example of FIG. 15, the detection threshold that is adjusted at block 960 is applied only for detecting AF episodes that trigger AF episode data storage at block 925 and is not used for detecting AF episodes that trigger other AF detection responses, such as causing the AF burden to be updated.

FIG. 16 is a flow chart 980 of a method for responding to AF episode detection by storing AF episode data according to one example. At block 982, an AF episode is detected. The detection of the AF episode at block 982 may be based on an initial AF detection threshold, e.g., a detection made at block 906 based on the initial detection threshold set at block 902 of any of FIG. 13, 14 or 15. In other instances, the detection of the AF episode at block 982 may be an AF detection made based on an adjusted AF detection threshold during heart rhythm monitoring at block 914 of FIG. 13. The process of flow chart 980 for updating detected AF episode storage may be performed at block 915 of FIG. 13.

In other examples, the detection of the AF episode at block 982 is an AF detection made at block 924 in response to an adjusted AF detection threshold as described above in conjunction with FIGS. 14 and 15. In other words, the process for updating AF episodes stored in the memory of the ICD 10 or 110 according to the flow chart 980 may be performed any time AF is detected based on an initial AF detection threshold and/or based on an AF detection threshold that has been adjusted for AF monitoring and AF episode storage. It is to be understood that an AF episode detected at block 956 based on an AF detection threshold set specifically for updating AF burden as described in conjunction with FIG. 15, may be ignored at block 982 of FIG. 16 when a different AF detection threshold has been set for triggering AF episode data storage.

At block 984, the duration of the AF episode detected at block 982 is determined. For example, the AF episode duration may be determined at block 984 using any of the methods described above in conjunction with FIG. 10. At block 986, microprocessor 224 determines if memory bins allocated for AF episode storage are full. If not, a cardiac signal segment acquired during the detected AF episode is stored in ICD memory, e.g., RAM 226, at block 992. In some examples, a predetermined number of bins may be allocated for multiple AF episode durations or episode duration ranges. Each episode duration or duration range may have multiple bins allocated for storing episodes of that specific duration or duration range. For example, three bins may be allocated for storing AF episodes having an episode duration that is equal to or greater than an initial detection threshold but less than an adjusted detection threshold and three bins may be allocated for storing AF episodes having a duration that is equal to or greater than the adjusted detection threshold.

In the examples given above, three bins may be allocated for storing cardiac electrical signal segments acquired during AF episodes having an episode duration that is at least the initial detection threshold of 6 minutes or 3 two-minute time periods but less than the adjusted detection threshold of 10 minutes or 5 two-minute time periods. Three other bins may be allocated for storing cardiac electrical signal segments that are acquired during an AF episode having an episode duration that is at least the adjusted detection threshold of 10 minutes or 5 two-minute time periods. In this way, a cardiac electrical signal segment may be stored for each of up to three AF episodes detected using the 6 minute

detection threshold and having an episode duration that is less than 10 minutes and cardiac electrical signal segments may be stored for each of up to three AF episodes having an episode duration that is equal to or greater than 10 minutes, regardless of the detection threshold used to detect the AF episode.

In other examples, memory bins may be allocated according to the detection threshold used to detect the AF episode. For example, three bins may be allocated for storing a cardiac electrical signal segment acquired during an AF episode detected based on the initial detection threshold, and three bins may be allocated for storing a cardiac electrical signal segment acquired during an AF episode detected based on an adjusted detection threshold. Additional bins may be allocated for each detection threshold used when two or more adjustments to the detection threshold are made.

The episode duration determined at block 984 may be used to determine which bin the cardiac electrical signal segment is written to at block 992. The cardiac electrical signal segment is written to a bin allocated to an episode duration or range matching or including the determined episode duration. In other examples, the detection threshold used to detect the AF episode is used to determine which bin the cardiac electrical signal segment is written to at block 992. The cardiac electrical signal segment is written to a bin allocated to the detection threshold used to detect the AF episode.

If the bins allocated to the corresponding AF episode duration or detection threshold are not full, "no" branch of block 986, the cardiac electrical signal segment is written to any of the unoccupied bins allocated for that AF episode duration or detection threshold. If the bins allocated for the corresponding episode duration or detection threshold are full, "yes" branch of block 986, microprocessor 224 determines at block 988 if the currently detected AF episode has an episode duration that is longer than any currently stored AF episodes. If a cardiac electrical signal segment is stored for an AF episode having a longer episode duration than the most recently detected AF episode, the data stored for the longest duration AF episode is not overwritten as indicated at block 990. Another occupied bin allocated to the corresponding duration or detection threshold of the most recently detected AF episode is overwritten at block 992 to store the cardiac electrical signal segment corresponding to the most recently detected AF episode. The bin that is overwritten may be the bin storing the oldest AF episode data such that the cardiac electrical signal segments (and any other AF episode data) are stored in a first-in-first-out basis with the exception of retaining AF episode cardiac signal segment stored for the longest duration AF episode, regardless of age by not overwriting the data stored for the longest duration AF episode when a newer AF episode is detected. In this way, data for both the most recently occurring AF episodes and data for the AF episode having the longest episode duration, regardless of how long ago the longest episode was detected, are accumulated in ICD memory.

If the most recently detected AF episode has a longer episode duration than any of the stored episodes, "yes" branch of block 988, a cardiac electrical signal segment is stored at block 992 by overwriting the oldest data in a bin allocated for the corresponding episode duration or detection threshold. It is recognized that when ICD 10 or 110 is interrogated by an external device, e.g., external device 40 (shown in FIG. 1), data stored in all bins may be transmitted to the external device and all bins allocated for storing AF episode data may be cleared.

When the method of FIG. 16 is used to control storing of cardiac signal segments in response to AF episode detection, the AF episodes detected using the adjusted AF detection threshold that has been increased to achieve relatively higher specificity and relatively lower sensitivity promotes storage of true AF episodes. The likelihood of storing data corresponding to false AF episodes is reduced, thereby reducing the burden placed on a clinician in reviewing AF episode data that does not correspond to true AF episodes.

While the examples disclosed herein refer primarily to the detection of AF, it is contemplated that aspects of the techniques of the present disclosure may be utilized for detecting and responding to other atrial tachyarrhythmia episodes, such as atrial flutter and atrial tachycardia.

Thus, an apparatus and methods have been presented in the foregoing description for detecting and responding to atrial tachyarrhythmia with reference to specific examples. It is appreciated that various modifications to the referenced examples may be made, including modifying the order of steps performed and/or modifying the combinations of operations shown in the flow charts presented herein, without departing from the scope of the following claims.

The invention claimed is:

1. A method of detecting an atrial tachyarrhythmia episode by a medical device comprising:
 - obtaining a cardiac electrical signal sensed via a plurality of electrodes coupled to sensing circuitry of the medical device;
 - determining a time duration of the cardiac electrical signal classified as an atrial tachyarrhythmia;
 - determining that the time duration is greater than or equal to a first detection threshold;
 - detecting a first atrial tachyarrhythmia episode by a processor of the medical device in response to the time duration being greater than or equal to the first detection threshold;
 - determining that detection threshold adjustment criteria are met based on at least the detected first atrial tachyarrhythmia episode; and
 - adjusting the first detection threshold to a second detection threshold different than the first detection threshold in response to the detection threshold adjustment criteria being met.
2. The method of claim 1, wherein detecting the first atrial tachyarrhythmia episode comprises:
 - identifying R-waves in the cardiac electrical signal attendant ventricular depolarizations;
 - determining classification factors from the R-waves identified over a plurality of predetermined time periods;
 - classifying each of the plurality of predetermined time periods as one of atrial tachyarrhythmia, non-atrial tachyarrhythmia or unclassified based on comparing the determined classification factors to classification criteria; and
 - determining a count of the number of the plurality of predetermined time periods that are classified as atrial tachyarrhythmia;
 - detecting the atrial tachyarrhythmia episode in response to the count reaching a threshold number of time periods.
3. The method of claim 1, the method of claim 1 wherein adjusting the first detection threshold comprises increasing a time duration threshold that the time duration of the cardiac electrical signal classified as the atrial tachyarrhythmia must be equal to or greater than in order to detect the atrial tachyarrhythmia.
4. The method of claim 1, wherein the detection threshold adjustment criteria comprise a predetermined time duration

threshold that is longer than the first detection threshold and further wherein determining that the detection threshold adjustment criteria are met comprises:

- determining an episode duration of the first atrial tachyarrhythmia episode,
- determining that the episode duration of the first atrial tachyarrhythmia episode is greater than or equal to the predetermined time duration threshold; and
- determining that the detection threshold adjustment criteria are met when the episode duration is greater than or equal to the predetermined time duration threshold.

5. The method of claim 1, wherein the detection threshold adjustment criteria comprise a predetermined number of atrial tachyarrhythmia episodes being detected based on the first detection threshold and further wherein determining that the detection threshold adjustment criteria are met comprises:

- determining a count of atrial tachyarrhythmia episodes that are detected based on the first detection threshold; and
- determining that the count reaches the predetermined number;
- determining that the detection threshold adjustment criteria are met when the count reaches the predetermined number.

6. The method of claim 5, further comprising:

- determining that the count reaches the predetermined number within a predetermined time period; and
- determining that the detection threshold adjustment criteria are met comprises determining that the detection threshold adjustment criteria are met when the count reaches the predetermined number within the predetermined time period.

7. The method of claim 1, further comprising determining that the detection threshold adjustment criteria are met in response to at least one of:

- the first atrial tachyarrhythmia episode detected based on the first detection threshold having an episode duration that is greater than a predetermined episode duration;
- a first predetermined number of atrial tachyarrhythmia episodes detected based on the first detection threshold;
- a second predetermined number of atrial tachyarrhythmia episodes detected based on the first detection threshold within a predetermined time period; or
- an atrial tachyarrhythmia burden exceeding an atrial tachyarrhythmia burden threshold, the method comprising determining the atrial tachyarrhythmia burden by summing a plurality of episode durations determined from a plurality of atrial tachyarrhythmia episodes detected based on the first detection threshold.

8. The method of claim 1, further comprising:

- detecting expiration of a predetermined time period without detecting an atrial tachyarrhythmia episode based on the second detection threshold;
- adjusting the second detection threshold back to the first detection threshold when the predetermined time period expiring without detecting an atrial tachyarrhythmia episode based on the second detection threshold.

9. The method of claim 1, further comprising:

- detecting at least one atrial tachyarrhythmia episode based on the second detection threshold;
- determining if second detection threshold adjustment criteria are met in response to the at least one atrial tachyarrhythmia episode detected based on the second detection threshold; and

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adjusting the second detection threshold to a third detection threshold greater than the second detection threshold in response to the second detection threshold adjustment criteria being met.

10. The method of claim 1, further comprising:

storing in a memory of the medical device a segment of the cardiac electrical signal received during the first atrial tachyarrhythmia episode; and
transmitting the segment to an external device for generating a display of the cardiac electrical signal segment.

11. The method of claim 1, further comprising:

determining a second time duration of the cardiac electrical signal classified as the atrial tachyarrhythmia;
determining that the second time duration is greater than or equal to the second detection threshold
detecting a second atrial tachyarrhythmia episode when the second time duration is greater than or equal to the second detection threshold;

storing in a memory of the medical device a segment of the cardiac electrical signal in response to detecting the second atrial tachyarrhythmia episode.

12. The method of claim 1, further comprising:

setting a third detection threshold;

monitoring for atrial tachyarrhythmia episodes based on the third detection threshold concurrently with monitoring for atrial tachyarrhythmia episodes based on the second detection threshold;

determining an atrial tachyarrhythmia burden in response to detecting an atrial tachyarrhythmia episode based on the third detection threshold; and

storing a segment of the cardiac electrical signal received during an atrial tachyarrhythmia episode detected based on the second detection threshold.

13. The method of claim 1, further comprising:

allocating a plurality of memory bins to each one of a plurality of episode duration ranges;

determining a first episode duration of the first atrial tachyarrhythmia episode;

storing a segment of the cardiac electrical signal received during the first atrial tachyarrhythmia episode in one of the plurality of memory bins allocated to one of the episode duration ranges that includes one of the first episode duration or the first detection threshold;

detecting a second atrial tachyarrhythmia episode based on a second time duration of the cardiac electrical signal classified as the atrial tachyarrhythmia being equal to or greater than the second detection threshold;

determining a second episode duration of the second atrial tachyarrhythmia episode; and

storing a second segment of the cardiac electrical signal received during the second atrial tachyarrhythmia episode in one of the plurality of memory bins allocated to one of the episode duration ranges that includes one of the second episode duration or the second detection threshold.

14. The method of claim 13, further comprising:

comparing the second episode duration to a longest episode duration of atrial tachyarrhythmia episodes previously detected based on one of the first detection threshold or the second detection threshold;

storing the second segment in one of the plurality of memory bins not currently occupied by a third segment of the cardiac electrical signal stored during the longest episode duration.

15. The method of claim 1, further comprising receiving the cardiac electrical signal via electrodes carried by a transvenous lead coupled to the medical device.

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16. The method of claim 1, further comprising receiving the cardiac electrical signal via implantable extra-cardiovascular electrodes coupled to the medical device.

17. The method of claim 1, further comprising providing a response to the detection of the atrial tachyarrhythmia episode, the response comprising at least one of:

withholding a therapy;

adjusting a therapy;

delivering a therapy;

triggering a patient notification system;

transmitting data via telemetry circuit to alert a clinician;

transmitting data via telemetry circuit to alert a patient;

and/or

triggering a second signal acquisition or analysis.

18. A medical device system for detecting an atrial tachyarrhythmia episode, comprising:

sensing circuitry configured to receive a cardiac electrical signal from a plurality of electrodes coupled to the sensing circuitry; and

a processor configured to:

detect a first atrial tachyarrhythmia episode in response to a time duration of the cardiac electrical signal classified as an atrial tachyarrhythmia being greater than or equal to a first detection threshold;

determine if detection threshold adjustment criteria are met based on at least the detected first atrial tachyarrhythmia episode; and

adjust the first detection threshold to a second detection threshold different than the first detection threshold in response to the detection threshold adjustment criteria being met.

19. The system of claim 18, wherein the processor is configured to detect the first atrial tachyarrhythmia episode by:

identifying R-waves in the cardiac electrical signal attendant ventricular depolarizations;

determining classification factors from the R-waves identified over a plurality of predetermined time periods;

classifying each of the plurality of predetermined time periods as one of atrial tachyarrhythmia, non-atrial tachyarrhythmia or unclassified based on comparing the determined classification factors to classification criteria; and

detecting the atrial tachyarrhythmia episode in response to the plurality of predetermined time periods that are classified as atrial tachyarrhythmia reaching a threshold number of time periods.

20. The system of claim 18, wherein the processor is further configured to adjust the detection threshold by increasing a time duration threshold that the duration of the cardiac electrical signal classified as the atrial tachyarrhythmia must be equal to or greater than in order to detect the atrial tachyarrhythmia.

21. The system of claim 18, wherein:

the detection threshold adjustment criteria comprise a predetermined time duration threshold that is longer than the first detection threshold; and

the processor is configured to determine that the detection threshold adjustment criteria are met by:

determining an episode duration of the first atrial tachyarrhythmia episode,

comparing the episode duration to the predetermined time duration threshold; and

determining that the detection threshold adjustment criteria are met based on the episode duration being equal to or greater than the predetermined time duration threshold.

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22. The system of claim 18, wherein:

the detection threshold adjustment criteria comprise a predetermined number of atrial tachyarrhythmia episodes being detected based on the first detection threshold; and

the processor is further configured to determine that the detection threshold adjustment criteria are met by:

determining a count of atrial tachyarrhythmia episodes that are detected based on the first detection threshold; and

determining that the detection threshold adjustment criteria are met in response to the count reaching the predetermined number.

23. The system of claim 22, wherein the processor is further configured to determine that the detection threshold adjustment criteria are met in response to the count reaching the predetermined number within a predetermined time period.

24. The system of claim 18, wherein the processor is further configured to determine that the detection threshold adjustment criteria are met in response to at least one of:

the first atrial tachyarrhythmia episode detected based on the first detection threshold having an episode duration that is greater than a predetermined episode duration; a first predetermined number of atrial tachyarrhythmia episodes being detected based on the first detection threshold;

a second predetermined number of atrial tachyarrhythmia episodes being detected based on the first detection threshold within a predetermined time period; or an atrial tachyarrhythmia burden exceeding an atrial tachyarrhythmia burden threshold, the processor configured to determine the atrial tachyarrhythmia burden by summing a plurality of episode durations determined from a plurality of atrial tachyarrhythmia episodes detected based on the first detection threshold.

25. The system of claim 18, wherein the processor is further configured to:

adjust the second detection threshold back to the first detection threshold in response to a predetermined time period expiring without detecting an atrial tachyarrhythmia episode based on the second detection threshold.

26. The system of claim 18, wherein the processor is further configured to:

detect at least one atrial tachyarrhythmia episode based on the second detection threshold;

determine if second detection threshold adjustment criteria are met in response to the at least one atrial tachyarrhythmia episode detected based on the second detection threshold; and

adjust the second detection threshold to a third detection threshold greater than the second detection threshold in response to the second detection threshold adjustment criteria being met.

27. The system of claim 18, further comprising:

a memory coupled to the processor; and

a telemetry circuit coupled to the processor and memory and configured to transmit cardiac electrical signal data stored in the memory to an external device;

wherein the processor is configured to:

store in the memory a segment of the cardiac electrical signal received during the first atrial tachyarrhythmia episode; and

control the telemetry circuit to transmit the segment to the external device for generating a display of the cardiac electrical signal segment.

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28. The system of claim 18, further comprising a memory coupled to the processor, wherein the processor is further configured to:

detect a second atrial tachyarrhythmia episode based on a second time duration of the cardiac electrical signal classified as the atrial tachyarrhythmia being equal to or greater than the second detection threshold;

store in the memory a segment of the cardiac electrical signal in response to detecting the second atrial tachyarrhythmia episode.

29. The system of claim 18, wherein the processor is further configured to:

set a third detection threshold;

monitor for atrial tachyarrhythmia episodes based on the third detection threshold concurrently with monitoring for atrial tachyarrhythmia episodes based on the second detection threshold;

determine an atrial tachyarrhythmia burden in response to detecting an atrial tachyarrhythmia episode based on the third detection threshold; and

store a segment of the cardiac electrical signal received during an atrial tachyarrhythmia episode detected based on the second detection threshold.

30. The system of claim 18, further comprising a memory coupled to the processor, wherein the processor is further configured to:

allocate a plurality of memory bins of the memory to each one of a plurality of episode duration ranges;

determine a first episode duration of the first atrial tachyarrhythmia episode;

store a segment of the cardiac electrical signal received during the first atrial tachyarrhythmia episode in one of the plurality of memory bins allocated to one of the episode duration ranges that includes one of the first episode duration or the first detection threshold;

detect a second atrial tachyarrhythmia episode based on a second time duration of the cardiac electrical signal classified as the atrial tachyarrhythmia being equal to or greater than the second detection threshold;

determine a second episode duration of the second atrial tachyarrhythmia episode; and

store a second segment of the cardiac electrical signal received during the second atrial tachyarrhythmia episode in one of the plurality of memory bins allocated to one of the episode duration ranges that includes one of the second episode duration or the second detection threshold.

31. The system of claim 30, wherein the processor is further configured to:

compare the second episode duration to a longest episode duration of atrial tachyarrhythmia episodes previously detected based on one of the first detection threshold or the second detection threshold;

store the second segment in one of the plurality of memory bins not currently occupied by a third segment of the cardiac electrical signal stored during the longest episode duration.

32. The system of claim 18, comprising an implantable cardioverter defibrillator and a transvenous lead coupled to the implantable cardioverter defibrillator and carrying at least one transvenous electrode, the sensing circuitry receiving the cardiac electrical signal via the at least one transvenous electrode, the implantable cardioverter defibrillator comprising a housing enclosing the sensing circuitry and the processor.

33. The system of claim 18, comprising an implantable cardioverter defibrillator and an extra-cardiovascular lead

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coupled to the implantable cardioverter defibrillator and carrying at least one extra-cardiovascular electrode, the sensing circuitry receiving the cardiac electrical signal via the at least one extra-cardiovascular electrode, the implantable cardioverter defibrillator comprising a housing enclosing the sensing circuitry and the processor.

34. The system of claim 18, wherein the processor is further configured to provide a response to the detection of the atrial tachyarrhythmia episode, the response comprising at least one of:

controlling therapy circuitry within the system to withholding a therapy, adjusting a therapy, or deliver a therapy;

controlling a patient notification system to generate a patient alert;

transmitting data via telemetry circuit to alert a clinician or a patient; and/or

triggering a second signal acquisition or analysis.

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35. A non-transitory, computer-readable storage medium storing instructions for causing a processor included in a medical device to perform a method for detecting an atrial tachyarrhythmia episode, the method comprising:

receiving a cardiac electrical signal via a plurality of electrodes coupled to sensing circuitry of the medical device;

detecting a first atrial tachyarrhythmia episode in response to a time duration of the cardiac electrical signal classified as an atrial tachyarrhythmia being greater than or equal to a first detection threshold;

determining if detection threshold adjustment criteria are met based on at least the detected first atrial tachyarrhythmia episode; and

adjusting the first detection threshold to a second detection threshold different than the first detection threshold in response to the detection threshold adjustment criteria being met.

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专利名称(译)	在植入式医疗设备中检测房性快速性心律失常的自动阈值		
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申请(专利权)人(译)	美敦力公司, INC.		
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摘要(译)

一种用于检测房性快速性心律失常发作的系统，包括：医疗设备，具有被配置为从耦合到医疗设备的电极接收心电信号的处理器；以及处理器，被配置为响应于心电信号的持续时间来检测房性快速性心律失常发作。归类为房性快速性心律失常大于或等于第一检测阈值。处理器被配置为至少基于检测到的第一房性快速性心律失常发作确定是否满足检测阈值调整标准，并且响应于满足检测阈值调整标准，将第一检测阈值调整为不同于第一检测阈值的第二检测阈值。。

