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Bardy(10) **Pub. No.: US 2007/0203415 A1**(43) **Pub. Date: Aug. 30, 2007**(54) **SYSTEM AND METHOD FOR
DETERMINING EDEMA THROUGH
REMOTE PATIENT MONITORING****Publication Classification**(76) Inventor: **Gust H. Bardy, Seattle, WA (US)**

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STE.1005
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600/301**(21) Appl. No.: **11/789,388**(22) Filed: **Apr. 23, 2007****Related U.S. Application Data**

(63) Continuation of application No. 10/152,207, filed on May 20, 2002, now Pat. No. 7,207,945, which is a continuation of application No. 09/442,125, filed on Nov. 16, 1999, now Pat. No. 6,398,728.

(57)

ABSTRACT

A system and method for determining edema through remote patient monitoring is presented. Device measures regularly recorded by a medical device are accumulated as raw physiometry for a patient. Derived measures are determined at least in part from the device measures. A clinical indication of one of an absence, onset, progression, regression, and status quo of edema is evaluated based on the device and derived measures.

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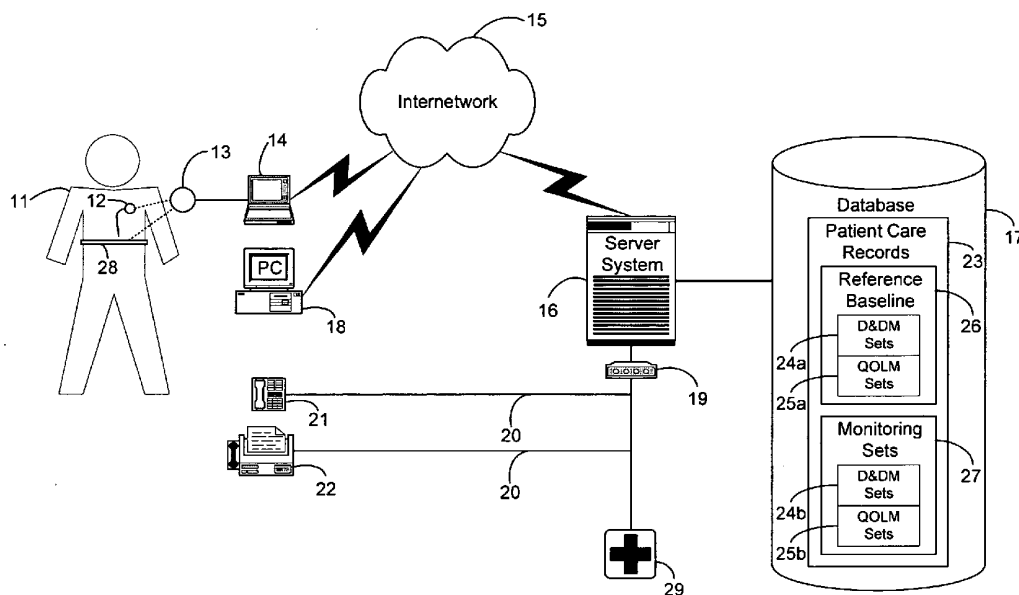


Fig. 1.

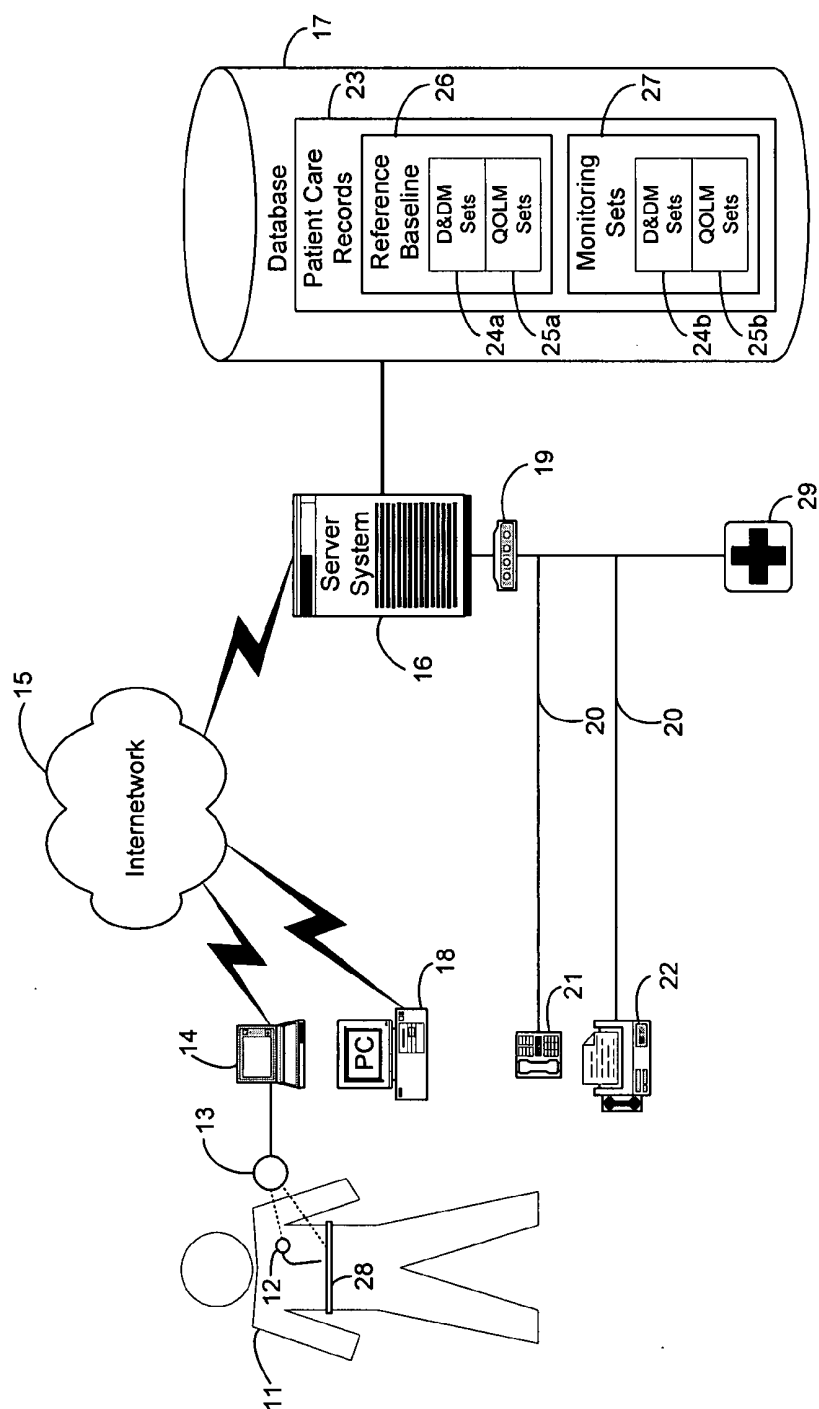


Fig. 2.

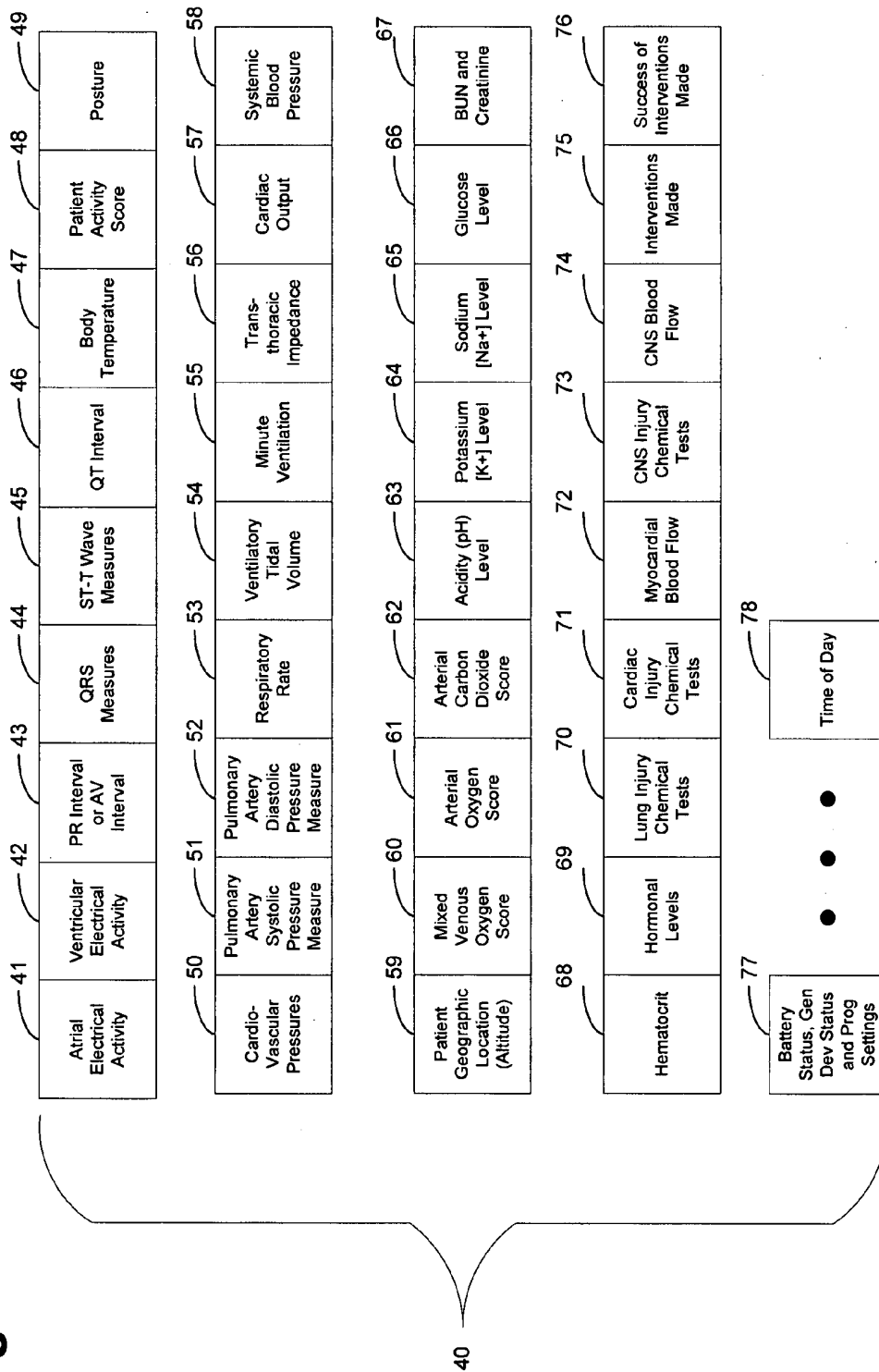


Fig. 3.

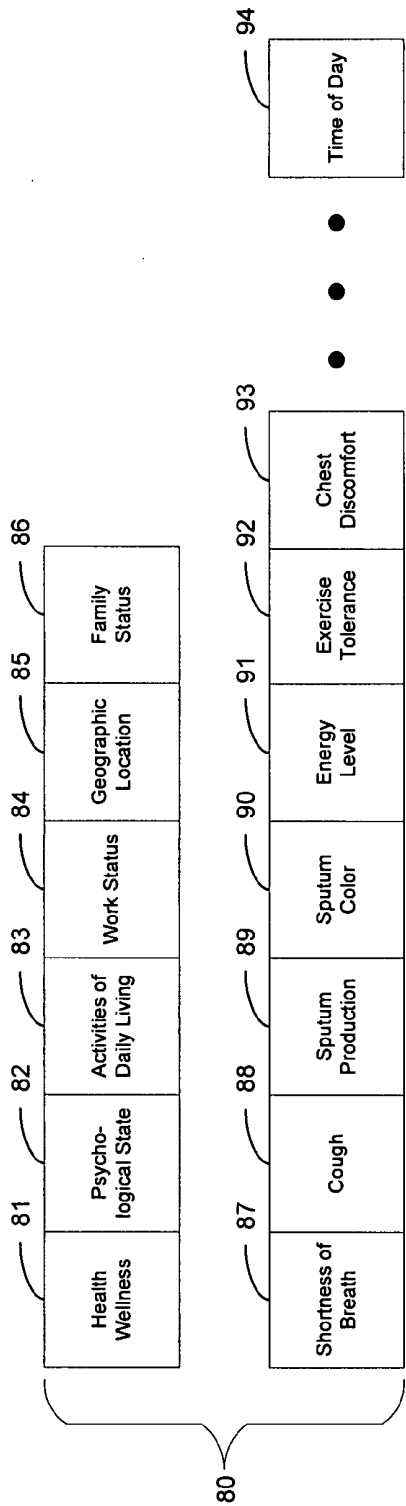


Fig. 4.

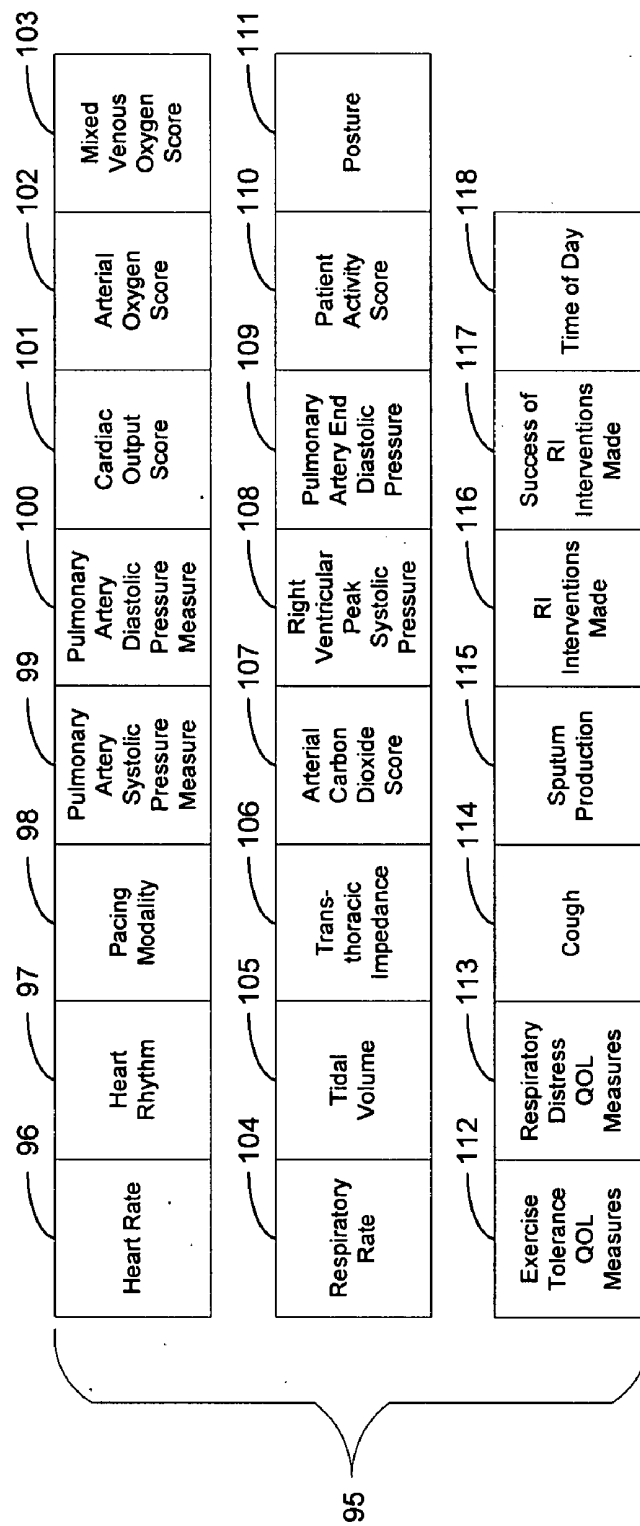


Fig. 5.

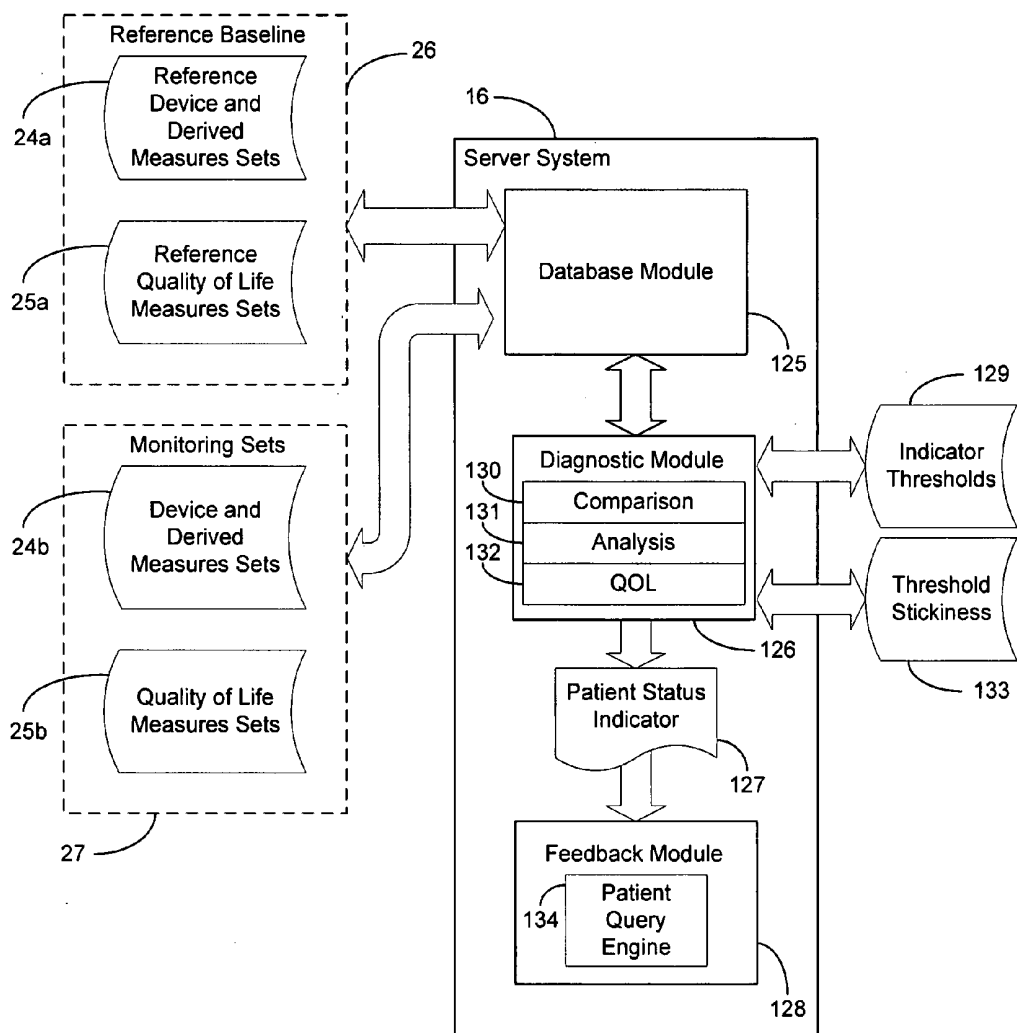


Fig. 6.

Patient 1						
Set 0				Set n-2	Set n-1	Set n
X_0	•	•	•	X_{n-2}	X_{n-1}	X_n
Y_0	•	•	•	Y_{n-2}	Y_{n-1}	Y_n
Z_0	•	•	•	Z_{n-2}	Z_{n-1}	Z_n

time →

Patient 2						
Set 0				Set n-2	Set n-1	Set n
$X_{0'}$	•	•	•	$X_{n-2'}$	$X_{n-1'}$	$X_{n'}$
$Y_{0'}$	•	•	•	$Y_{n-2'}$	$Y_{n-1'}$	$Y_{n'}$
$Z_{0'}$	•	•	•	$Z_{n-2'}$	$Z_{n-1'}$	$Z_{n'}$

time →

Patient 3						
Set 0				Set n-2	Set n-1	Set n
X_0''	•	•	•	X_{n-2}''	X_{n-1}''	X_n''
Y_0''	•	•	•	Y_{n-2}''	Y_{n-1}''	Y_n''
Z_0''	•	•	•	Z_{n-2}''	Z_{n-1}''	Z_n''

time →

Fig. 7.

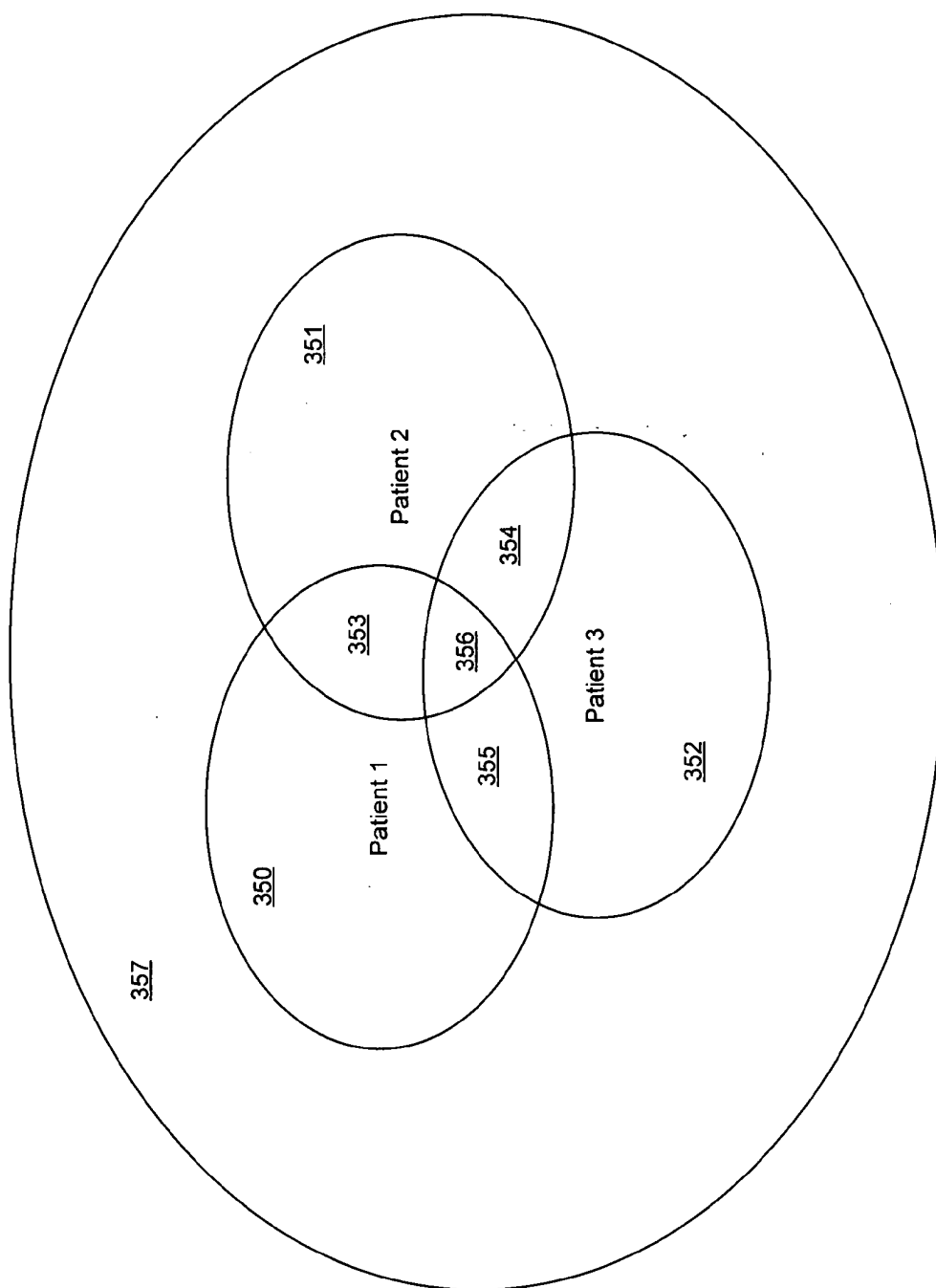


Fig. 8A.

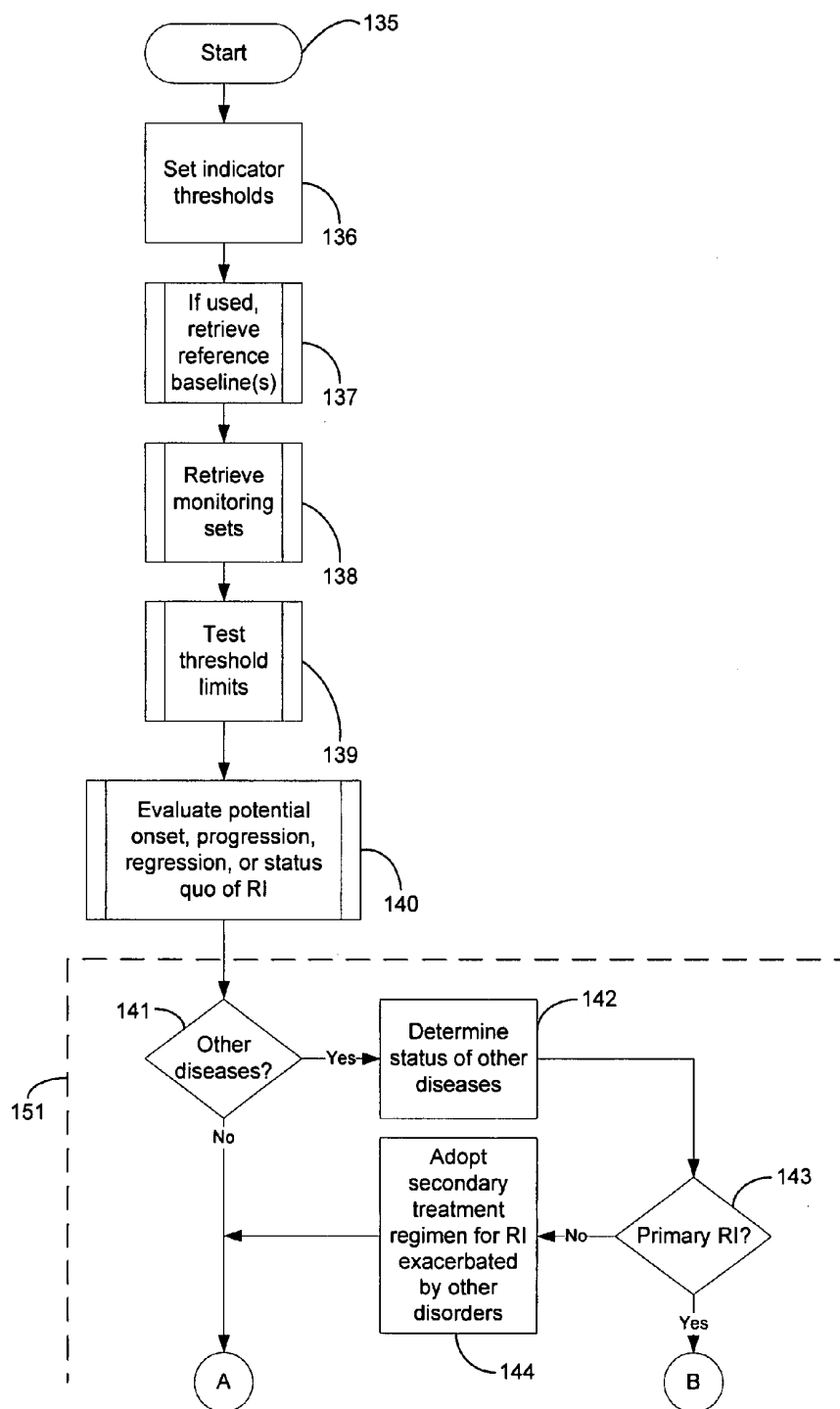


Fig. 8B.

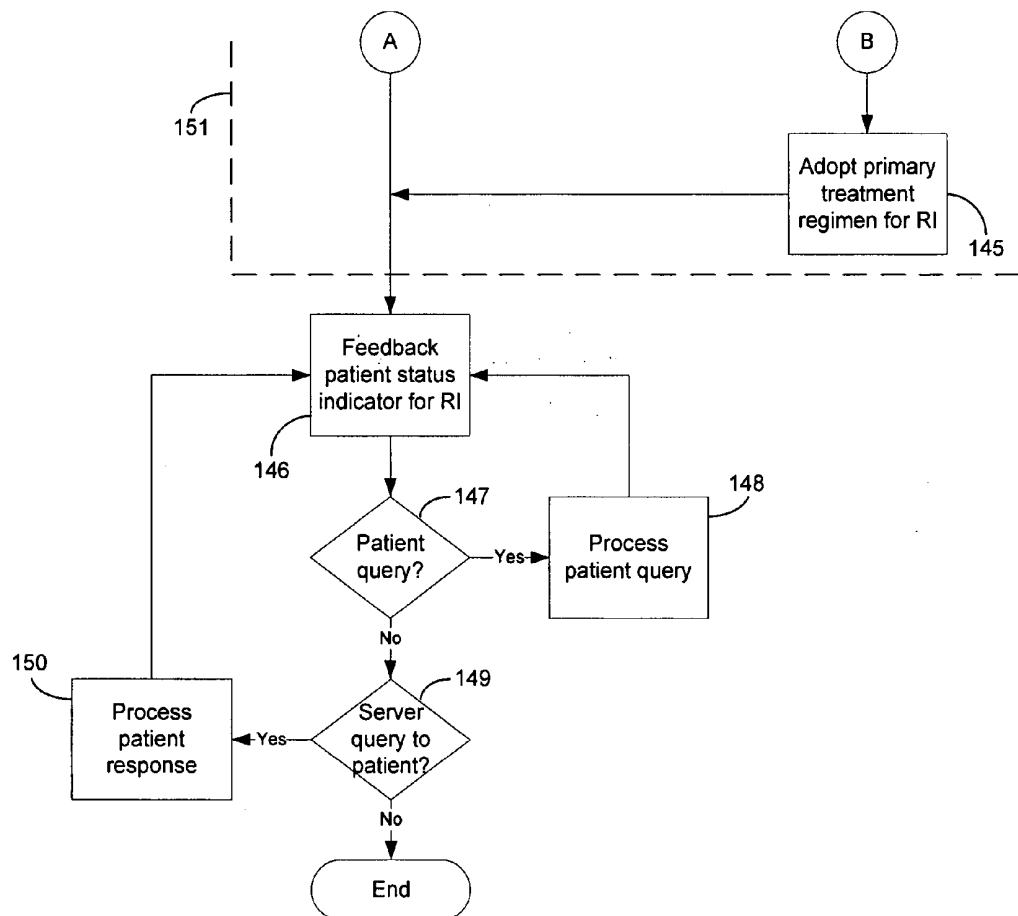


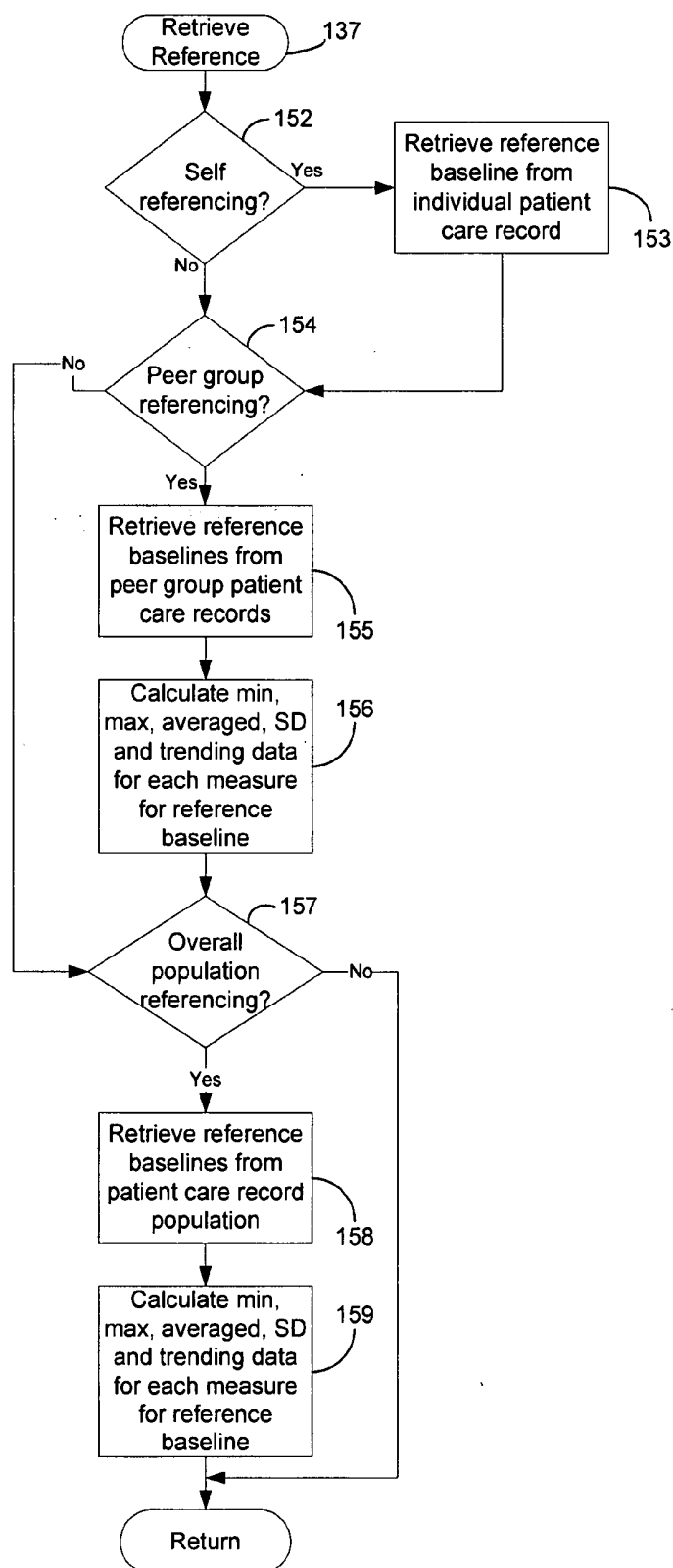
Fig. 9.

Fig. 10.

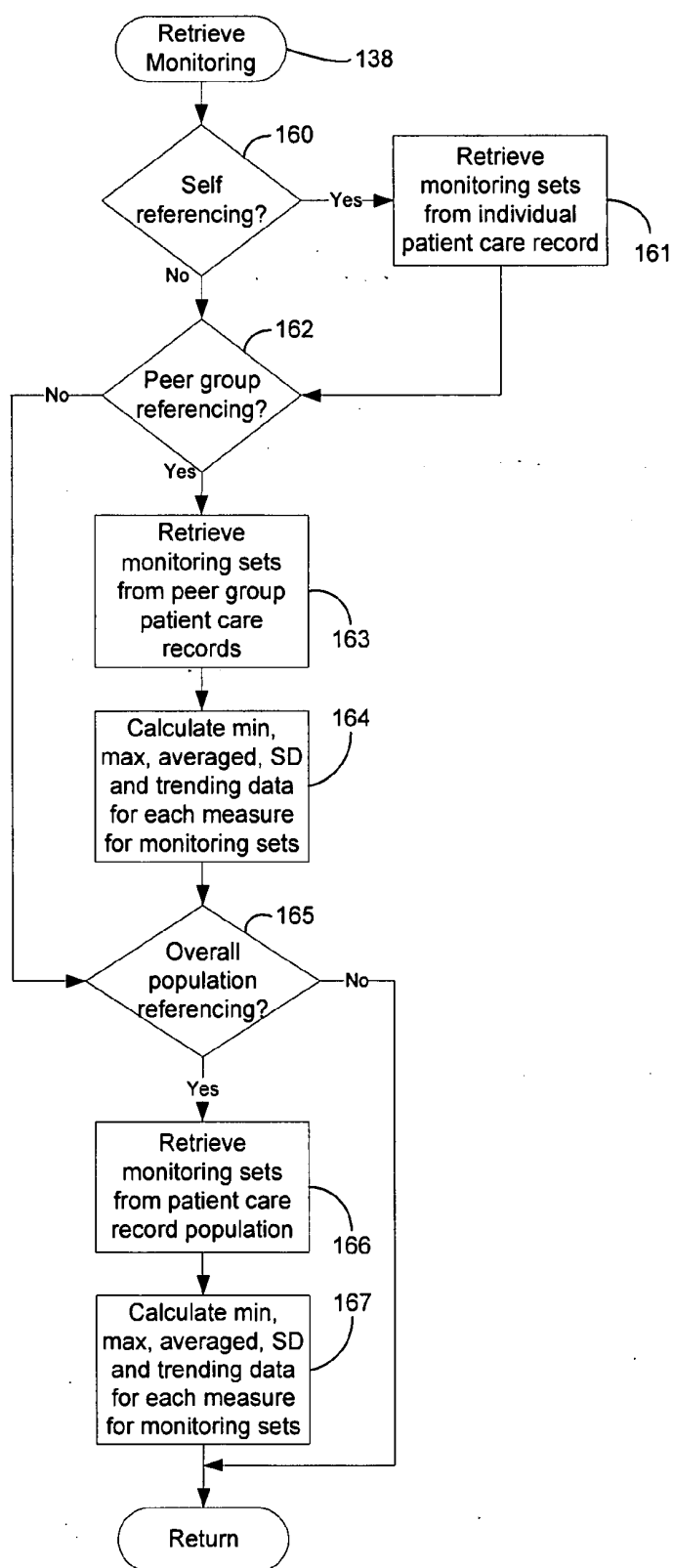


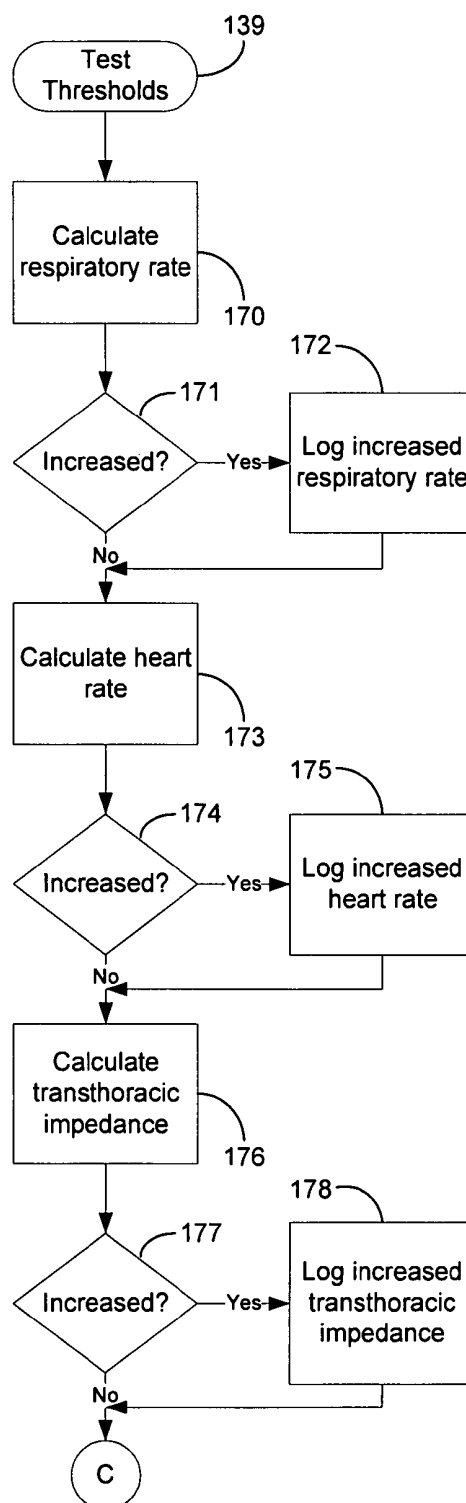
Fig. 11A.

Fig. 11B.

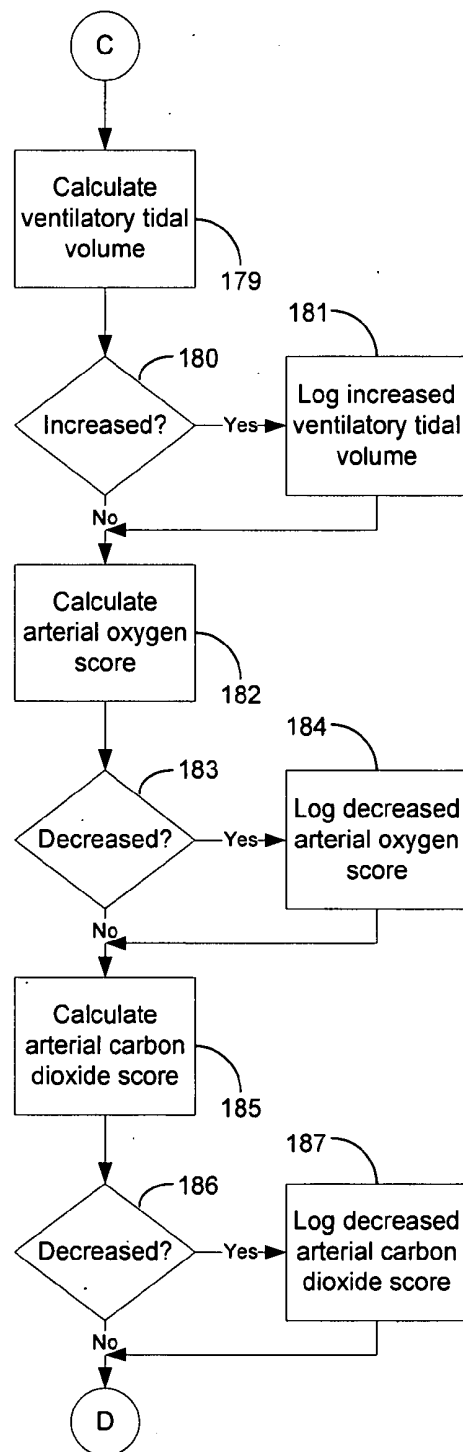


Fig. 11C.

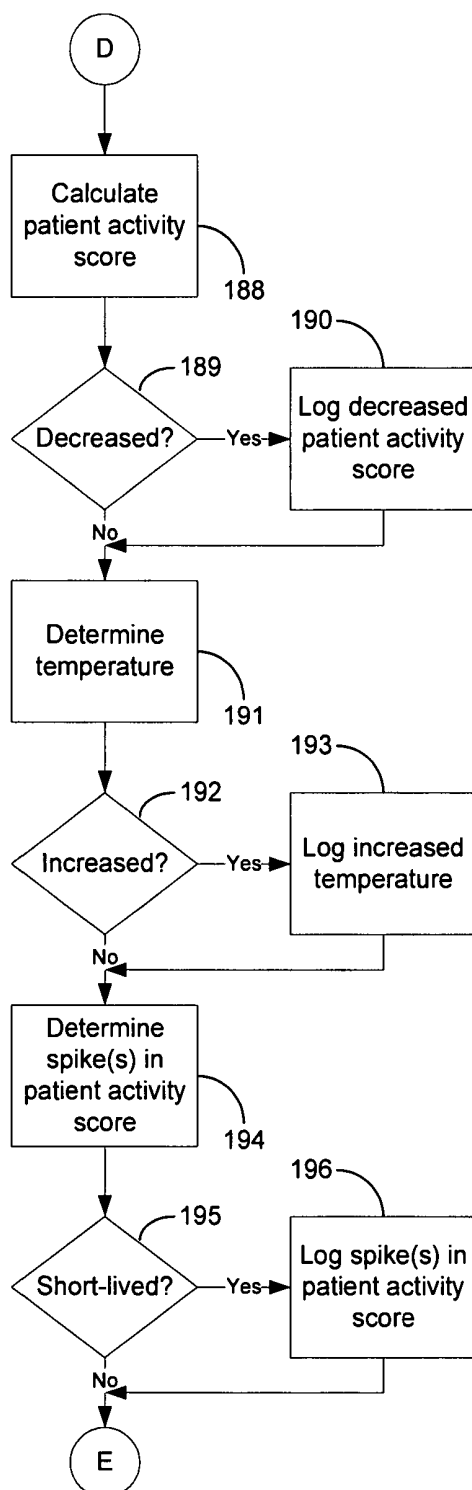


Fig. 11D.

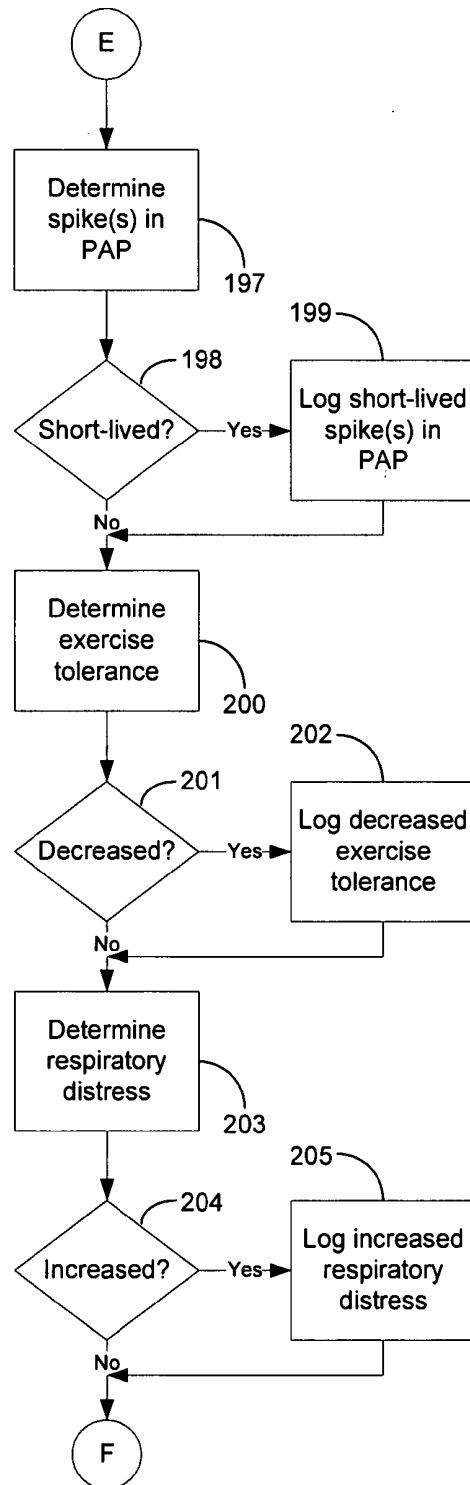


Fig. 11E.

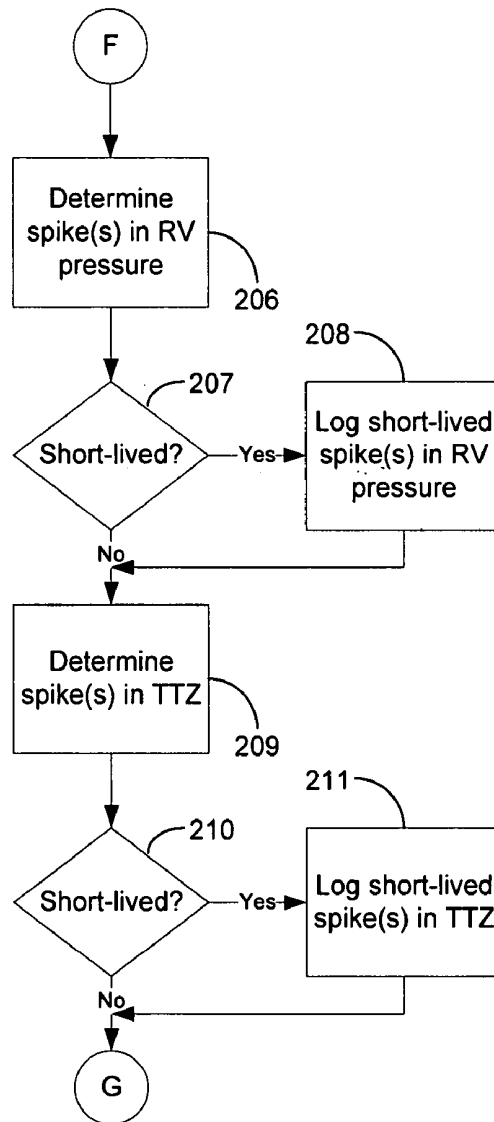


Fig. 11F.

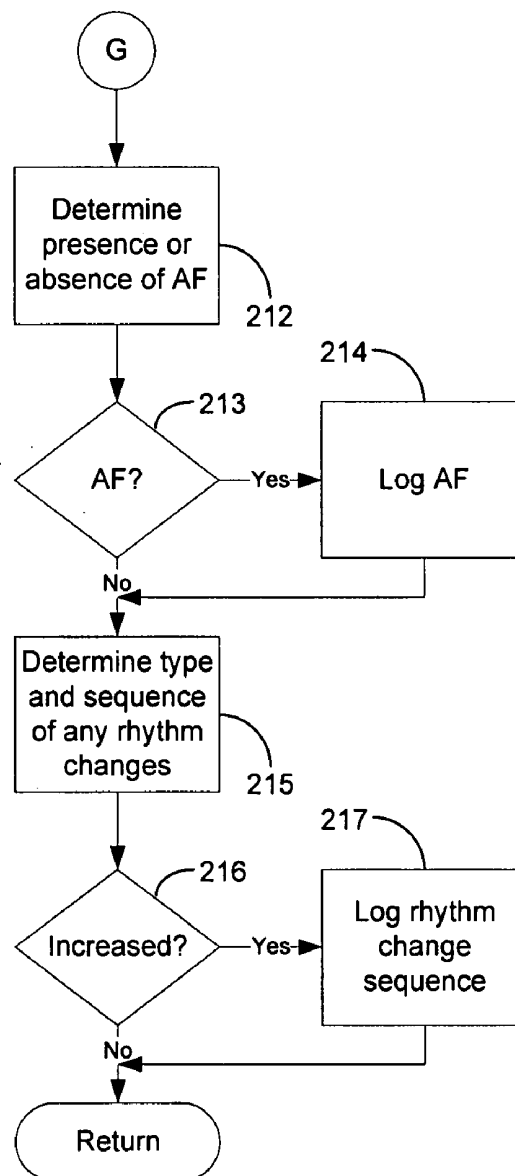


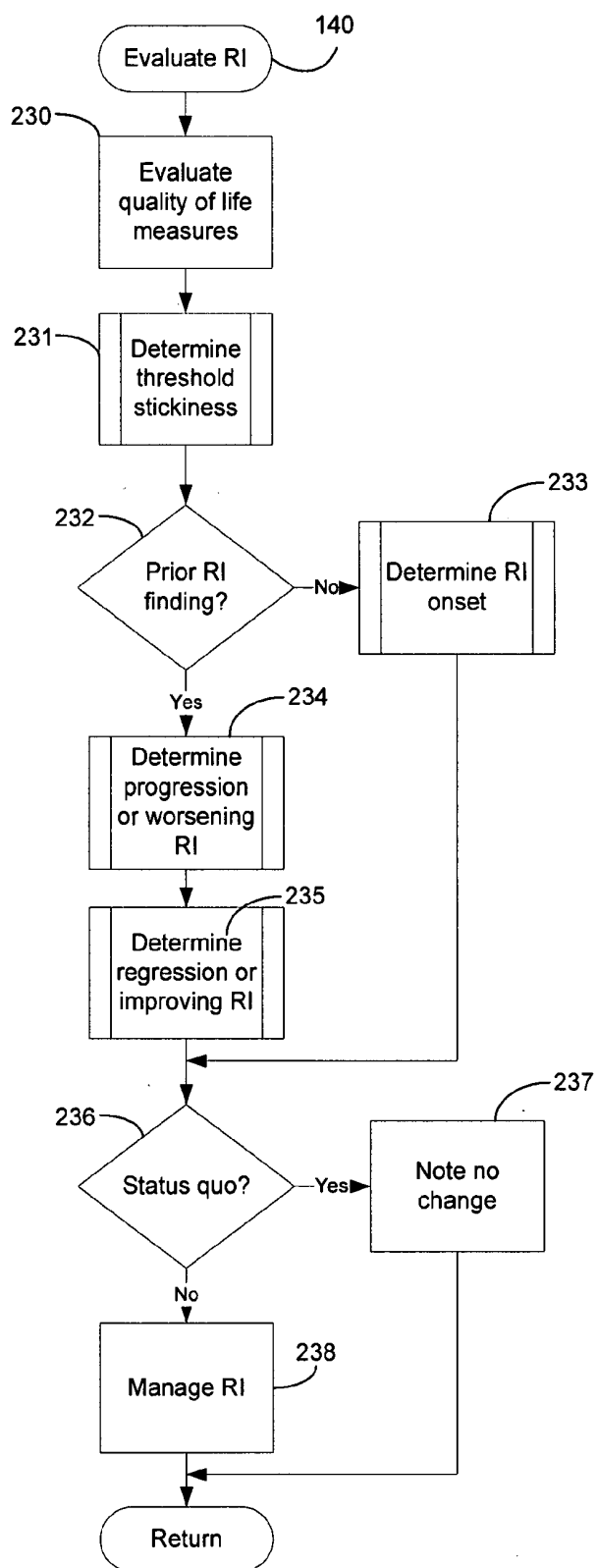
Fig. 12.

Fig. 13A.

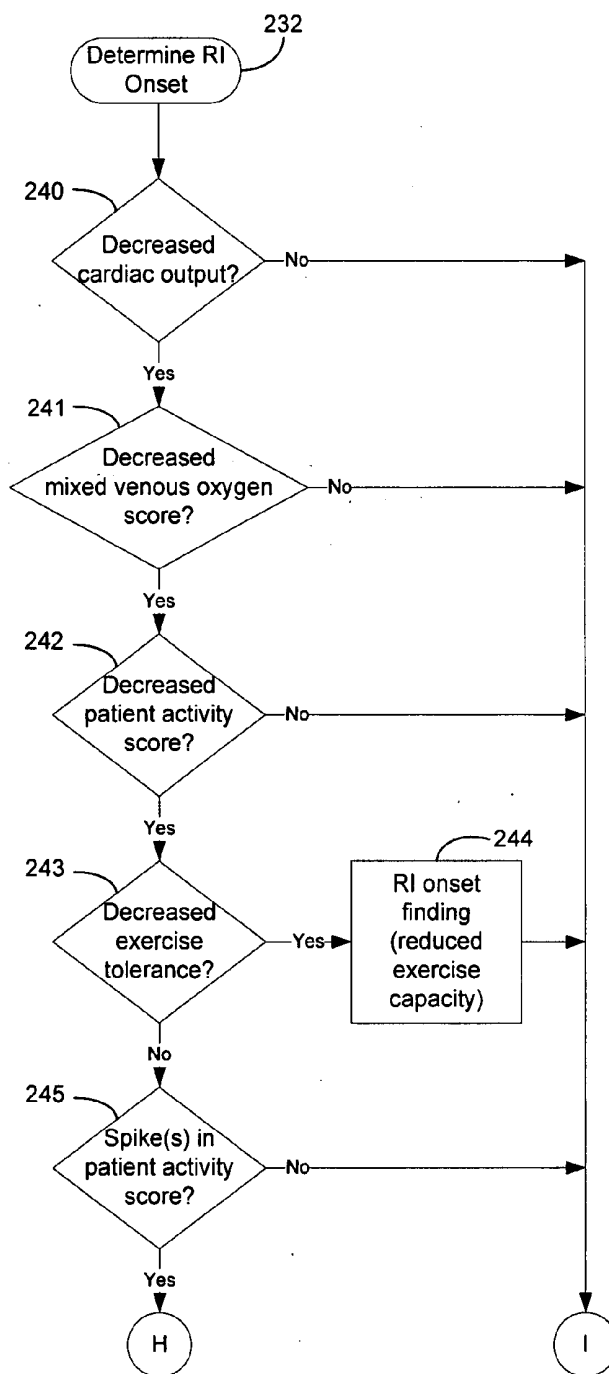


Fig. 13B.

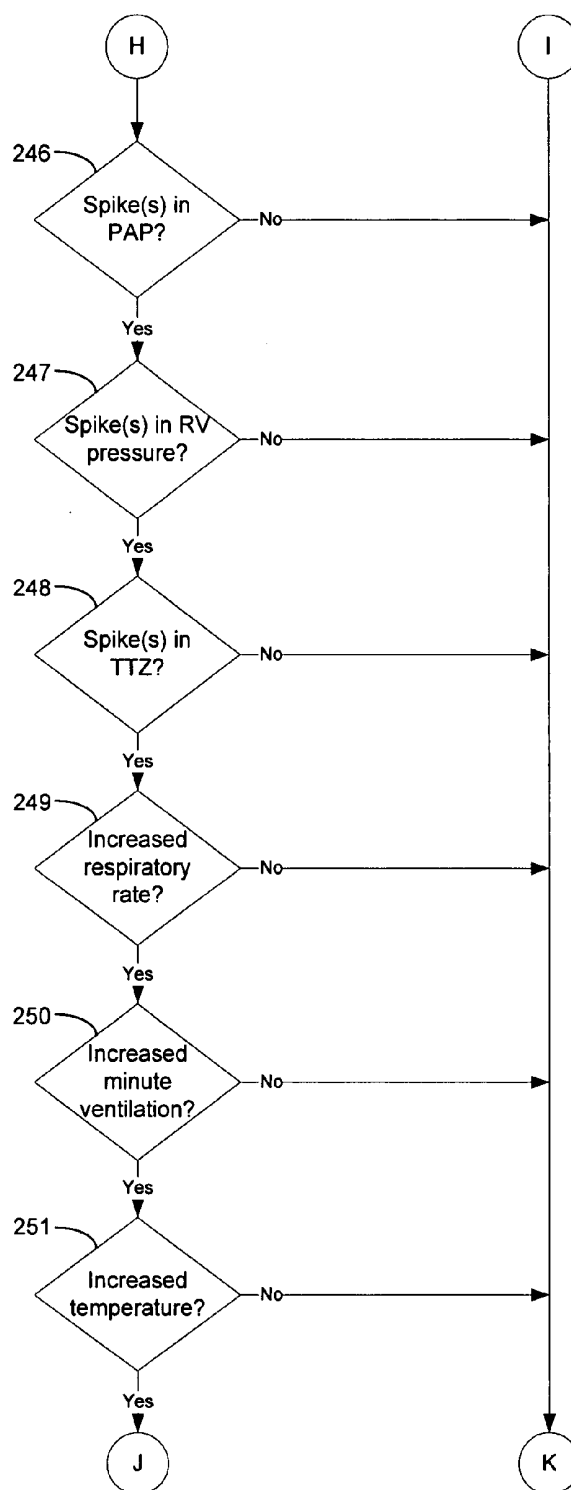


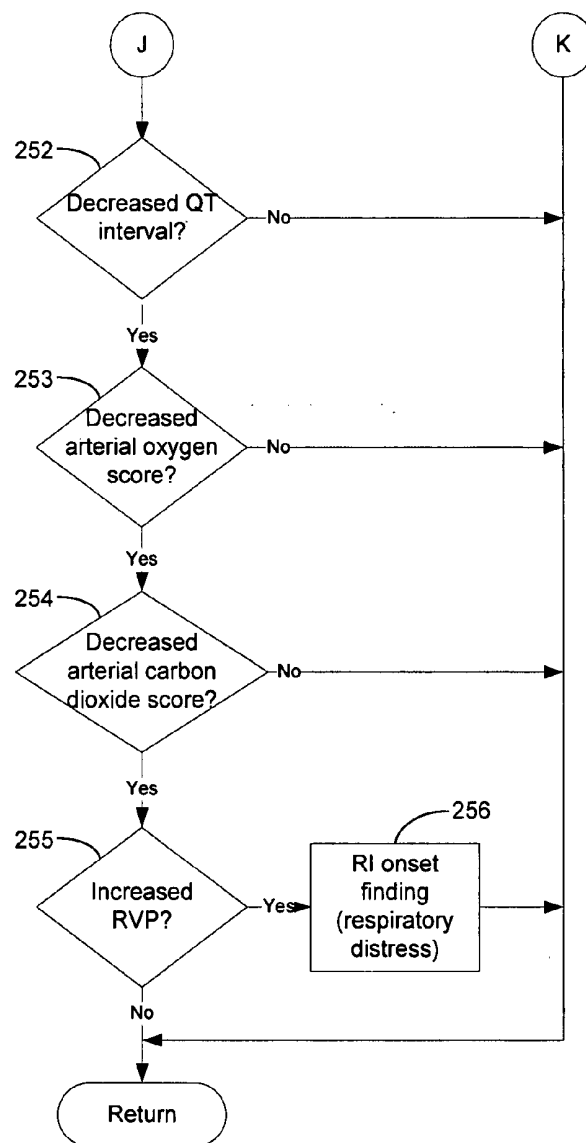
Fig. 13C.

Fig. 14A.

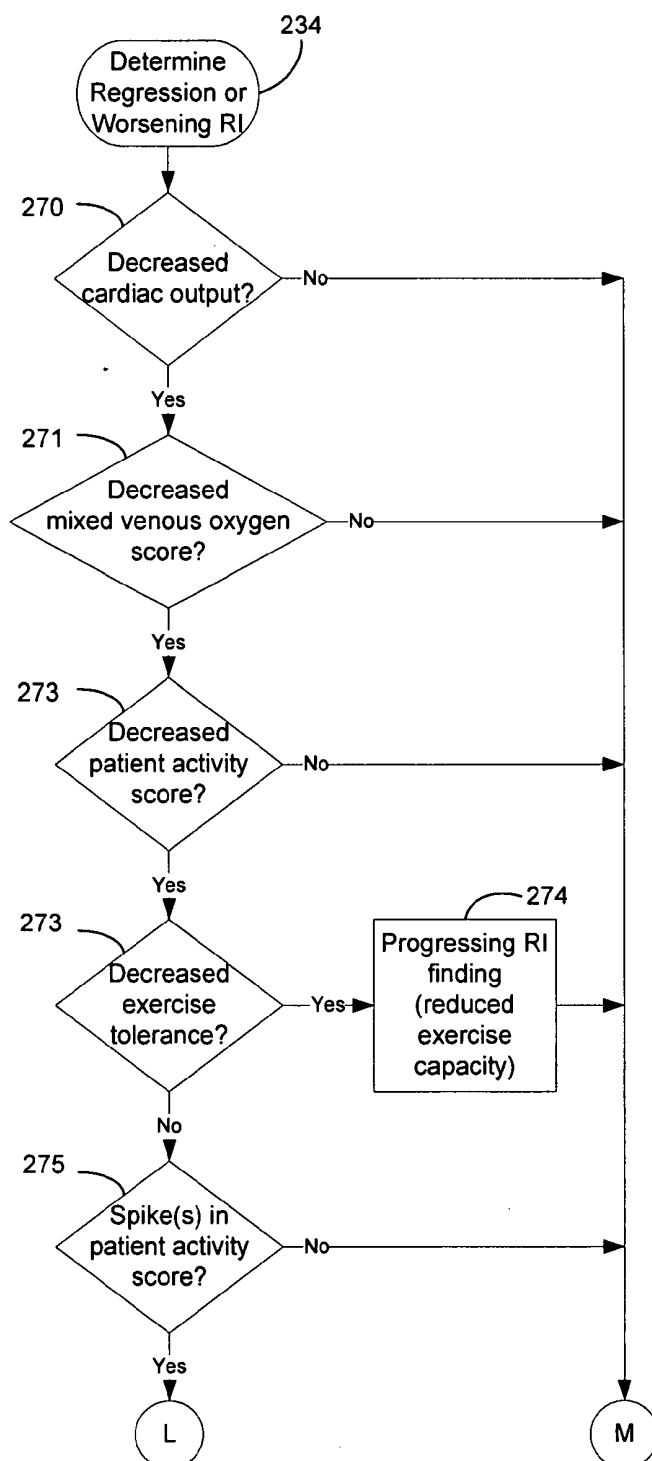


Fig. 14B.

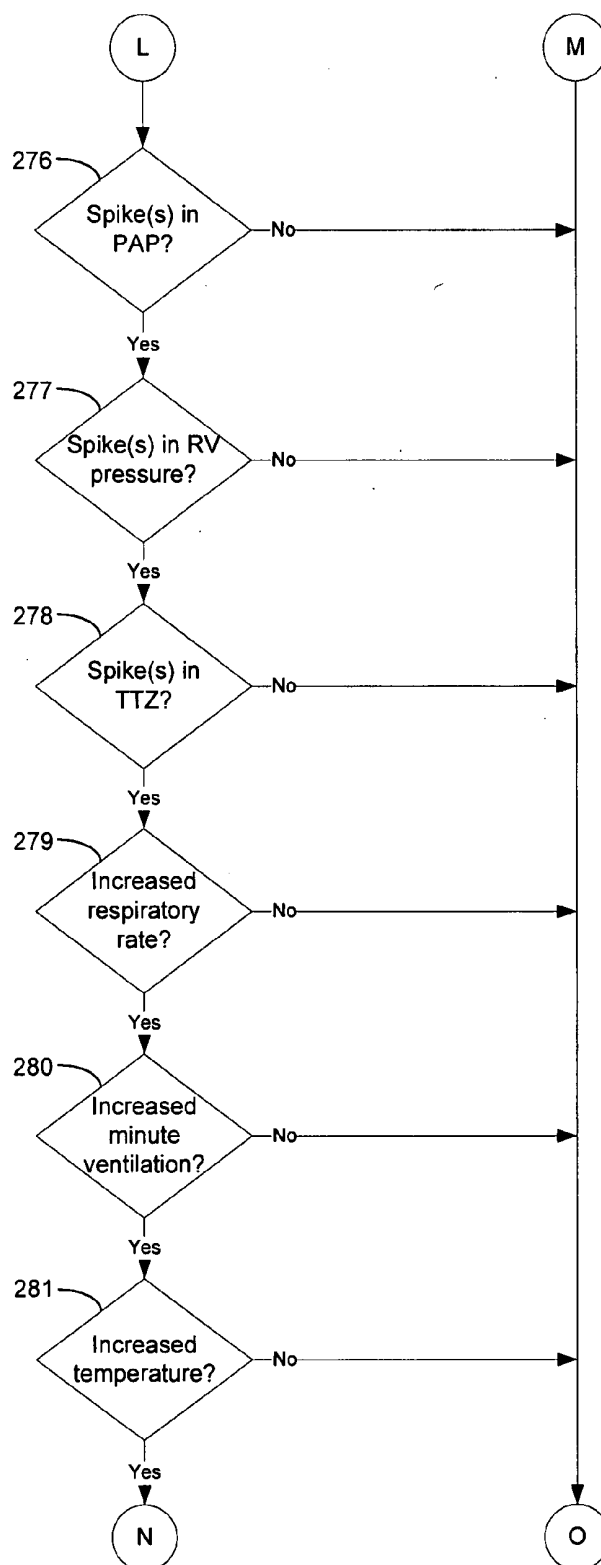


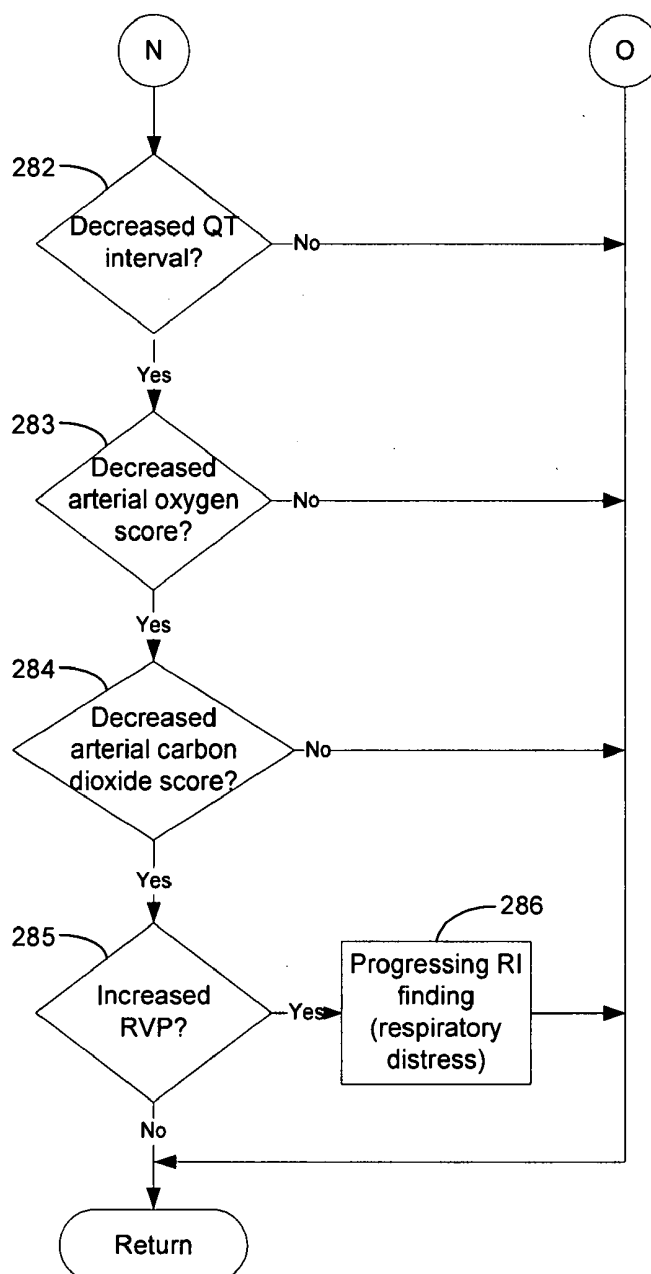
Fig. 14C.

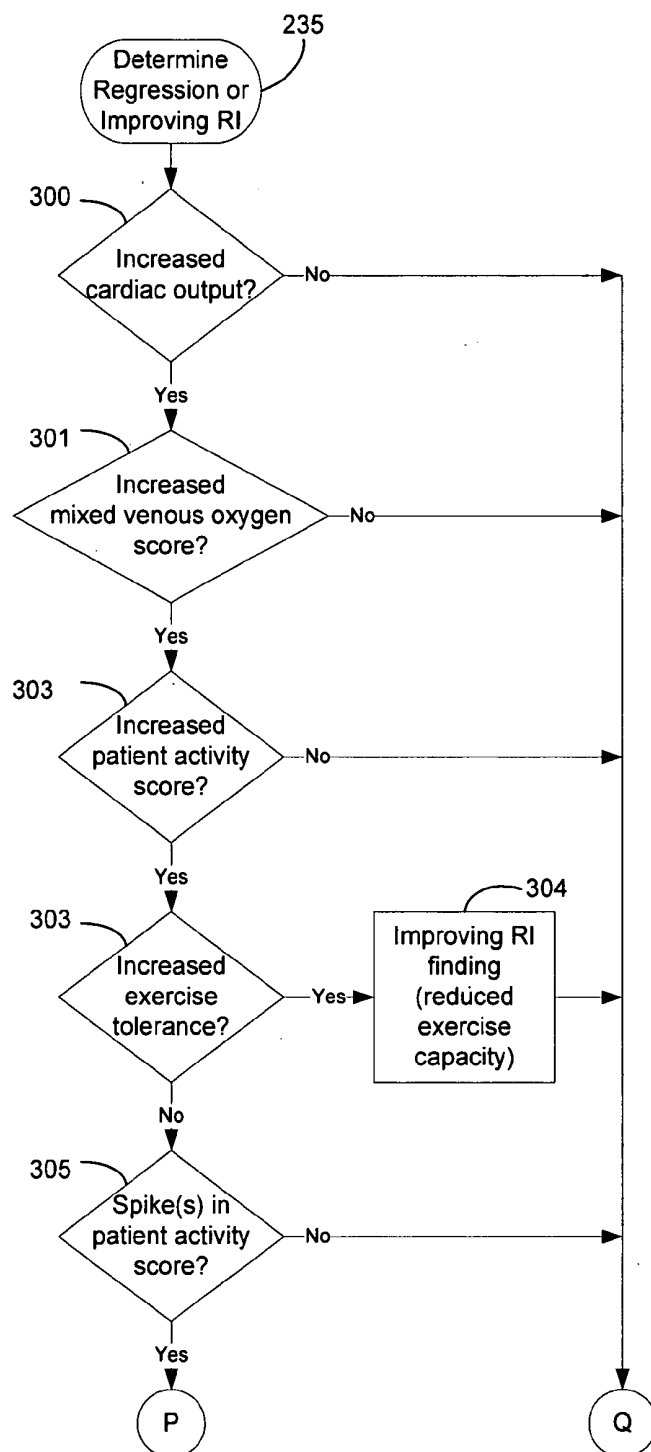
Fig. 15A.

Fig. 15B.

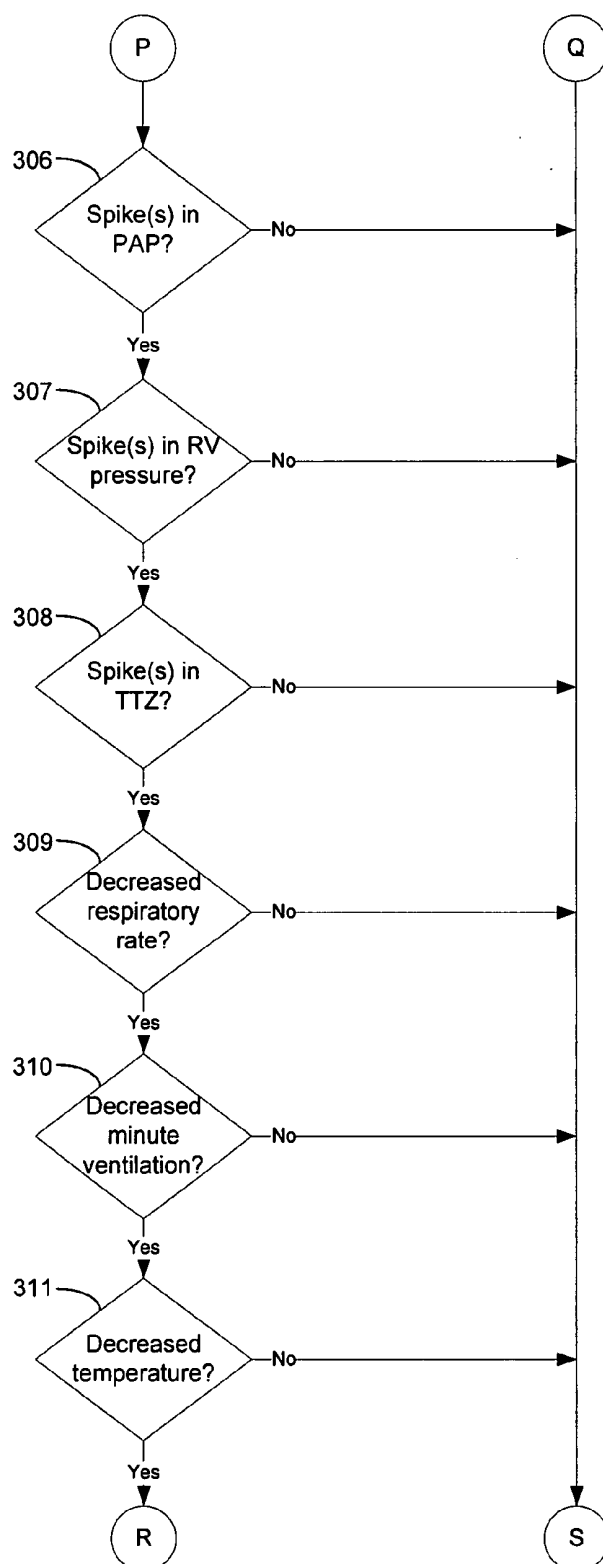


Fig. 15C.

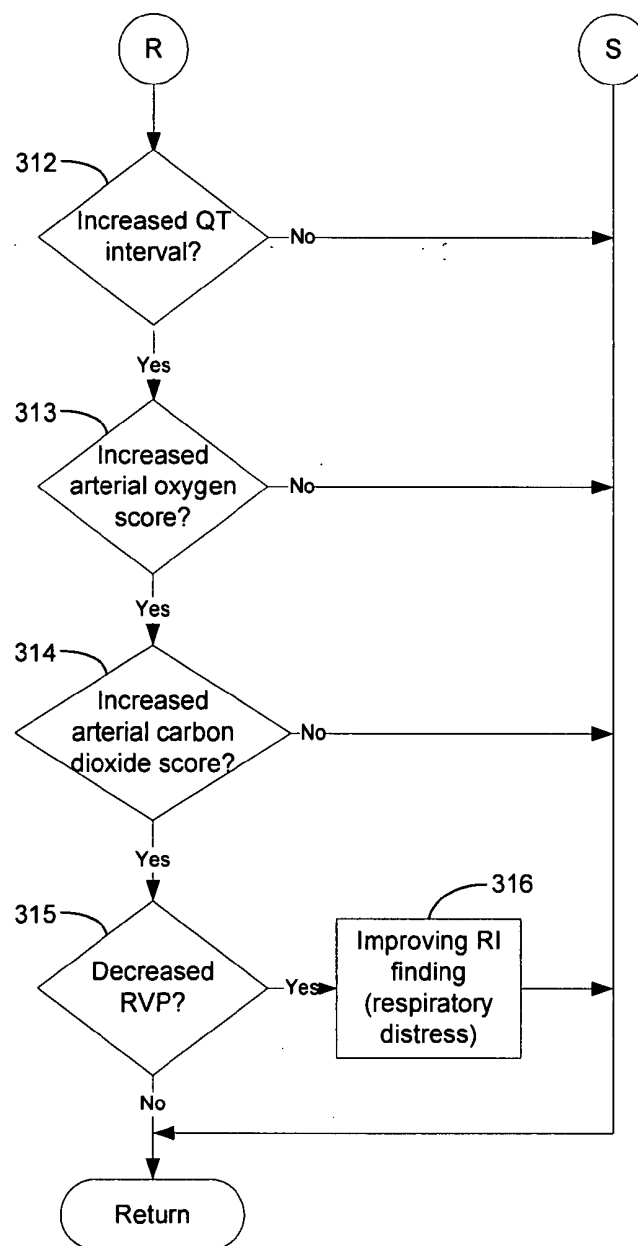
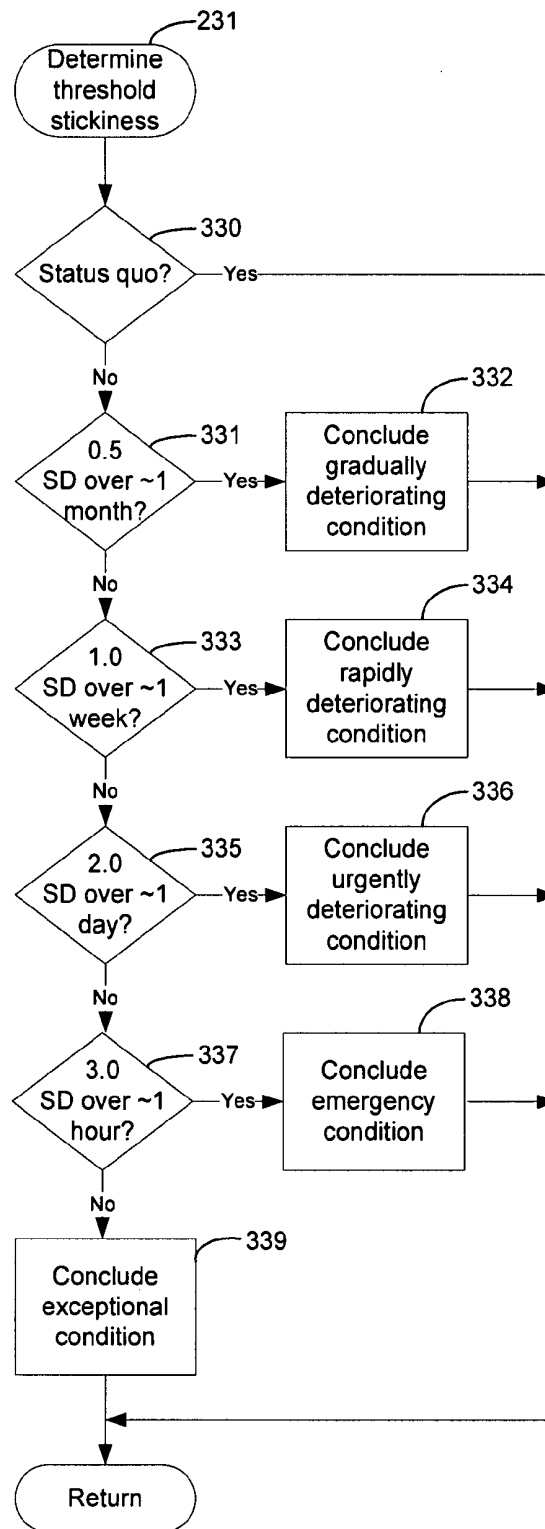


Fig. 16.



SYSTEM AND METHOD FOR DETERMINING EDEMA THROUGH REMOTE PATIENT MONITORING

CROSS-REFERENCE TO RELATED APPLICATION

[0001] This patent application is a continuation of U.S. patent application Ser. No. 10/152,207, filed May 20, 2002, pending; which is a continuation of U.S. Pat. No. 6,398,728, issued Jun. 4, 2002, the disclosures of which are incorporated by reference, and the priority filing dates of which are claimed.

FIELD OF THE INVENTION

[0002] The present invention relates in general to respiratory insufficiency diagnosis and analysis, and, in particular, to an automated collection and analysis patient care system and method for diagnosing and monitoring respiratory insufficiency and outcomes thereof throughout disease onset, progression, regression, and status quo.

BACKGROUND OF THE INVENTION

[0003] Presently, respiratory insufficiency due to primary diseases of the lungs is one of the leading causes of acute and chronic illness in the world. Clinically, respiratory insufficiency involves either difficulty in ventilation or in oxygenation. The former is manifest by increases in the arterial partial pressure of carbon dioxide and the latter is manifest by decreases in arterial partial pressure of oxygen. For purposes of this invention, the term "respiratory insufficiency" will refer to ventilatory insufficiency and/or to problems in oxygenation due to diseases of the lung. Common causes of respiratory insufficiency include bronchitis, emphysema, pneumonia, pulmonary emboli, congestive heart failure, tumor infiltration of the lung and abnormalities of the interstitium of the lungs that may be infectious in origin, due to immunological abnormalities, or as a result of exposure to environmental pathogens. The effects of respiratory insufficiency range from cough to impairment during physical exertion to a complete failure of lung function and respiratory arrest at any level of activity. Clinical manifestations of respiratory insufficiency include respiratory distress, such as shortness of breath and fatigue, cough, and reduced exercise capacity or tolerance.

[0004] Several factors make the early diagnosis and prevention of respiratory insufficiency, as well as the monitoring of the progression of respiratory insufficiency, relatively difficult. First, the onset of respiratory insufficiency is generally subtle and erratic. Often, the symptoms are ignored and the patient compensates by changing his or her daily activities. This situation is especially true in chronic lung disorders where the onset of symptoms can be very gradual. As a result, many respiratory insufficiency conditions or deteriorations in respiratory insufficiency remain undiagnosed until more serious problems arise seriously limiting the activities of daily living.

[0005] The susceptibility to suffer from respiratory insufficiency depends upon the patient's age, sex, physical condition, and other factors, such as smoking history, occupation, diabetes, co-existing heart disease, immunodepression, the presence or absence of cancer, surgical history, kidney function, and extent of pre-existing lung disease. No one

factor is dispositive. Finally, annual or even monthly lung checkups, including chest X-rays or other lung tests, provide, at best, a "snapshot" of patient wellness and the incremental and subtle clinicophysiological changes which portend the onset or progression of respiratory insufficiency often go unnoticed, even with regular health care. Documentation of subtle improvements following therapy that can guide and refine further evaluation and therapy can be equally elusive.

[0006] Nevertheless, taking advantage of frequently and regularly measured physiological measures, such as recorded manually by a patient, via an external monitoring or therapeutic device, or via implantable device technologies, can provide a degree of detection and prevention heretofore unknown. For instance, patients already suffering from some form of treatable heart disease often receive an implantable pulse generator (IPG), cardiovascular or heart failure monitor, therapeutic device, or similar external wearable device, with which rhythm and structural problems of the heart can be monitored and treated. These types of devices, although usually originally intended for use in treating some type of cardiac problem, can contain sufficient physiological data to allow accurate assessment of lung disorders. Such devices are useful for detecting physiological changes in patient conditions through the retrieval and analysis of telemetered signals stored in an on-board, volatile memory. Typically, these devices can store more than thirty minutes of per heartbeat and respiratory cycle data recorded on a per heartbeat, per respiration, binned average basis, or on a derived basis from, for example, atrial or ventricular electrical activity, minute ventilation, patient activity score, cardiac output score, arterial or mixed venous oxygen score, cardiopulmonary pressure measures, and the like. However, the proper analysis of retrieved telemetered signals requires detailed medical subspecialty knowledge, particularly by pulmonologists and cardiologists.

[0007] Alternatively, these telemetered signals can be remotely collected and analyzed using an automated patient care system. One such system is described in a related, commonly owned U.S. Pat. No. 6,312,378, issued Nov. 6, 2001, the disclosure of which is incorporated herein by reference. A medical device adapted to be implanted in an individual patient records telemetered signals that are then retrieved on a regular, periodic basis using an interrogator or similar interfacing device. The telemetered signals are downloaded via an internetwork onto a network server on a regular, e.g., daily, basis and stored as sets of collected measures in a database along with other patient care records. The information is then analyzed in an automated fashion and feedback, which includes a patient status indicator, is provided to the patient.

[0008] While such an automated system can serve as a valuable tool in providing remote patient care, an approach to systematically correlating and analyzing the raw collected telemetered signals, as well as manually collected physiological measures, through applied pulmonary and cardiovascular medical knowledge to accurately diagnose the onset of a particular medical condition, such as respiratory insufficiency, is needed, especially in patients with co-existing heart disease. One automated patient care system directed to a patient-specific monitoring function is described in U.S. Pat. No. 5,113,869 ('869) to Nappholz et al. The '869 patent discloses an implantable, programmable

electrocardiography (ECG) patient monitoring device that senses and analyzes ECG signals to detect ECG and physiological signal characteristics predictive of malignant cardiac arrhythmias. The monitoring device can communicate a warning signal to an external device when arrhythmias are predicted. However, the Nappholz device is limited to detecting tachycardias. Unlike requirements for automated respiratory insufficiency monitoring, the Nappholz device focuses on rudimentary ECG signals indicative of malignant cardiac tachycardias, an already well established technique that can be readily used with on-board signal detection techniques. Also, the Nappholz device is patient specific only and is unable to automatically take into consideration a broader patient or peer group history for reference to detect and consider the progression or improvement of lung disease. Moreover, the Nappholz device has a limited capability to automatically self-reference multiple data points in time and cannot detect disease regression even in the individual patient. Also, the Nappholz device must be implanted and cannot function as an external monitor. Finally, the Nappholz device is incapable of tracking the cardiovascular and cardiopulmonary consequences of any rhythm disorder.

[0009] Consequently, there is a need for a systematic approach to detecting trends in regularly collected physiological data indicative of the onset, progression, regression, or status quo of respiratory insufficiency diagnosed and monitored using an automated, remote patient care system. The physiological data could be telemetered signals data recorded either by an external or an implantable medical device or, alternatively, individual measures collected through manual means. Preferably, such an approach would be capable of diagnosing both acute and chronic respiratory insufficiency conditions, as well as the symptoms of other lung disorders. In addition, findings from individual, peer group, and general population patient care records could be integrated into continuous, on-going monitoring and analysis.

SUMMARY OF THE INVENTION

[0010] The present invention provides a system and method for diagnosing and monitoring the onset, progression, regression, and status quo of respiratory insufficiency using an automated collection and analysis patient care system. Measures of patient cardiopulmonary information are either recorded by an external or implantable medical device, such as an IPG, cardiovascular or heart failure monitor, or respiratory diagnostic or therapeutic device, or manually through conventional patient-operable means. The measures are collected on a regular, periodic basis for storage in a database along with other patient care records. Derived measures are developed from the stored measures. Select stored and derived measures are analyzed and changes in patient condition are logged. The logged changes are compared to quantified indicator thresholds to detect findings of respiratory distress or reduced exercise capacity indicative of the principal pathophysiological manifestations of respiratory insufficiency: elevated partial pressure of arterial carbon dioxide and reduced partial pressure of arterial oxygen.

[0011] One embodiment provides a system and method for determining edema through remote patient monitoring. Device measures regularly recorded by a medical device are accumulated as raw physiometry for a patient. Derived

measures are determined at least in part from the device measures. A clinical indication of one of an absence, onset, progression, regression, and status quo of edema is evaluated based on the device and derived measures.

[0012] The present invention provides a capability to detect and track subtle trends and incremental changes in recorded patient cardiopulmonary information for diagnosing and monitoring respiratory insufficiency. When coupled with an enrollment in a remote patient monitoring service having the capability to remotely and continuously collect and analyze external or implantable medical device measures, respiratory insufficiency detection, prevention and tracking regression from therapeutic maneuvers become feasible.

[0013] Still other embodiments of the present invention will become readily apparent to those skilled in the art from the following detailed description, wherein is described embodiments of the invention by way of illustrating the best mode contemplated for carrying out the invention. As will be realized, the invention is capable of other and different embodiments and its several details are capable of modifications in various obvious respects, all without departing from the spirit and the scope of the present invention. Accordingly, the drawings and detailed description are to be regarded as illustrative in nature and not as restrictive.

BRIEF DESCRIPTION OF THE DRAWINGS

[0014] FIG. 1 is a block diagram showing an automated collection and analysis patient care system for diagnosing and monitoring respiratory insufficiency and outcomes thereof in accordance with the present invention;

[0015] FIG. 2 is a database schema showing, by way of example, the organization of a device and derived measures set record for care of patients with respiratory insufficiency stored as part of a patient care record in the database of the system of FIG. 1;

[0016] FIG. 3 is a database schema showing, by way of example, the organization of a quality of life and symptom measures set record for care of patients with respiratory insufficiency stored as part of a patient care record in the database of the system of FIG. 1;

[0017] FIG. 4 is a database schema showing, by way of example, the organization of a combined measures set record for care of patients with respiratory insufficiency stored as part of a patient care record in the database of the system of FIG. 1;

[0018] FIG. 5 is a block diagram showing the software modules of the server system of the system of FIG. 1;

[0019] FIG. 6 is a record view showing, by way of example, a set of partial patient care records for care of patients with respiratory insufficiency stored in the database of the system of FIG. 1;

[0020] FIG. 7 is a Venn diagram showing, by way of example, peer group overlap between the partial patient care records of FIG. 6;

[0021] FIGS. 8A-8B are flow diagrams showing a method for diagnosing and monitoring respiratory insufficiency and outcomes thereof using an automated collection and analysis patient care system in accordance with the present invention;

[0022] FIG. 9 is a flow diagram showing the routine for retrieving reference baseline sets for use in the method of FIGS. 8A-8B;

[0023] FIG. 10 is a flow diagram showing the routine for retrieving monitoring sets for use in the method of FIGS. 8A-8B;

[0024] FIGS. 11A-11F are flow diagrams showing the routine for testing threshold limits for use in the method of FIGS. 8A-8B;

[0025] FIG. 12 is a flow diagram showing the routine for evaluating the onset, progression, regression, and status quo of respiratory insufficiency for use in the method of FIGS. 8A-8B;

[0026] FIGS. 13A-13C are flow diagrams showing the routine for determining an onset of respiratory insufficiency for use in the routine of FIG. 12;

[0027] FIGS. 14A-14C are flow diagrams showing the routine for determining progression or worsening of respiratory insufficiency for use in the routine of FIG. 12;

[0028] FIGS. 15A-15C are flow diagrams showing the routine for determining regression or improving of respiratory insufficiency for use in the routine of FIG. 12; and

[0029] FIG. 16 is a flow diagram showing the routine for determining threshold stickiness ("hysteresis") for use in the method of FIG. 12.

DETAILED DESCRIPTION

[0030] FIG. 1 is a block diagram showing an automated collection and analysis patient care system 10 for diagnosing and monitoring respiratory insufficiency in accordance with the present invention. An exemplary automated collection and analysis patient care system suitable for use with the present invention is disclosed in the related, commonly-owned U.S. Pat. No. 6,312,378, issued Nov. 6, 2001, the disclosure of which is incorporated herein by reference. Preferably, an individual patient 11 is a recipient of an implantable medical device 12, such as, by way of example, an IPG, cardiovascular, heart failure monitor, pulmonary monitor, or therapeutic device, with a set of leads extending into his or her heart and electrodes implanted throughout the cardiopulmonary system. Alternatively, an external monitoring or therapeutic medical device 28, a subcutaneous monitor or device inserted into other organs, a cutaneous monitor, or even a manual physiological measurement device, such as an respiratory monitor, electrocardiogram or heart rate monitor, could be used. The implantable medical device 12 and external medical device 28 include circuitry for recording into a short-term, volatile memory telemetered signals stored for later retrieval, which become part of a set of device and derived measures, such as described below, by way of example, with reference to FIG. 2. Exemplary implantable medical devices suitable for use in the present invention include the Discovery line of pacemakers, manufactured by Guidant Corporation, Indianapolis, Ind., and the Gem line of ICDs, manufactured by Medtronic Corporation, Minneapolis, Minn.

[0031] The telemetered signals stored in the implantable medical device 12 are preferably retrieved upon the completion of an initial observation period and subsequently thereafter on a continuous, periodic (daily) basis, such as

described in the related, commonly-owned U.S. Pat. No. 6,221,011, issued Apr. 24, 2001, the disclosure of which is incorporated herein by reference. A programmer 14, personal computer 18, or similar device for communicating with an implantable medical device 12 can be used to retrieve the telemetered signals. A magnetized reed switch (not shown) within the implantable medical device 12 closes in response to the placement of a wand 13 over the site of the implantable medical device 12. The programmer 14 sends programming or interrogating instructions to and retrieves stored telemetered signals from the implantable medical device 12 via RF signals exchanged through the wand 13. Similar communication means are used for accessing the external medical device 28. Once downloaded, the telemetered signals are sent via an internetwork 15, such as the Internet, to a server system 16 which periodically receives and stores the telemetered signals as device measures in patient care records 23 in a database 17, as further described below, by way of example, with reference to FIGS. 2 and 3. An exemplary programmer 14 suitable for use in the present invention is the Model 2901 Programmer Recorder Monitor, manufactured by Guidant Corporation, Indianapolis, Ind.

[0032] The patient 11 is remotely monitored by the server system 16 via the internetwork 15 through the periodic receipt of the retrieved device measures from the implantable medical device 12 or external medical device 28. The patient care records 23 in the database 17 are organized into two identified sets of device measures: an optional reference baseline 26 recorded during an initial observation period and monitoring sets 27 recorded subsequently thereafter. The device measures sets are periodically analyzed and compared by the server system 16 to indicator thresholds corresponding to quantifiable physiological measures of a pathophysiology indicative of respiratory insufficiency, as further described below with reference to FIG. 5. As necessary, feedback is provided to the patient 11. By way of example, the feedback includes an electronic mail message automatically sent by the server system 16 over the internetwork 15 to a personal computer 18 (PC) situated for local access by the patient 11. Alternatively, the feedback can be sent through a telephone interface device 19 as an automated voice mail message to a telephone 21 or as an automated facsimile message to a facsimile machine 22, both also situated for local access by the patient 11. Moreover, simultaneous notifications can also be delivered to the patient's physician, hospital, or emergency medical services provider 29 using similar feedback means to deliver the information.

[0033] The server system 10 can consist of either a single computer system or a cooperatively networked or clustered set of computer systems. Each computer system is a general purpose, programmed digital computing device consisting of a central processing unit (CPU), random access memory (RAM), non-volatile secondary storage, such as a hard drive or CD ROM drive, network interfaces, and peripheral devices, including user interfacing means, such as a keyboard and display. Program code, including software programs, and data are loaded into the RAM for execution and processing by the CPU and results are generated for display, output, transmittal, or storage, as is known in the art.

[0034] The database 17 stores patient care records 23 for each individual patient to whom remote patient care is being provided. Each patient care record 23 contains normal

patient identification and treatment profile information, as well as medical history, medications taken, height and weight, and other pertinent data (not shown). The patient care records **23** consist primarily of two sets of data: device and derived measures (D&DM) sets **24a**, **24b** and quality of life (QOL) sets **25a**, **25b**, the organization of which are further described below with respect to FIGS. 2 and 3, respectively. The device and derived measures sets **24a**, **24b** and quality of life and symptom measures sets **25a**, **25b** can be further logically categorized into two potentially overlapping sets. The reference baseline **26** is a special set of device and derived reference measures sets **24a** and quality of life and symptom measures sets **25a** recorded and determined during an initial observation period. Monitoring sets **27** are device and derived measures sets **24b** and quality of life and symptom measures sets **25b** recorded and determined thereafter on a regular, continuous basis. Other forms of database organization are feasible.

[0035] The implantable medical device **12** and, in a more limited fashion, the external medical device **28**, record patient information for care of patients with respiratory insufficiency on a regular basis. The recorded patient information is downloaded and stored in the database **17** as part of a patient care record **23**. Further patient information can be derived from recorded data, as is known in the art. FIG. 2 is a database schema showing, by way of example, the organization of a device and derived measures set record **40** for patient care stored as part of a patient care record in the database **17** of the system of FIG. 1. Each record **40** stores patient information which includes a snapshot of telemetered signals data which were recorded by the implantable medical device **12** or the external medical device **28**, for instance, on per heartbeat, binned average or derived bases; measures derived from the recorded device measures; and manually collected information, such as obtained through a patient medical history interview or questionnaire. The following non-exclusive information can be recorded for a patient: atrial electrical activity **41**, ventricular electrical activity **42**, PR interval or AV interval **43**, QRS measures **44**, ST-T wave measures **45**, QT interval **46**, body temperature **47**, patient activity score **48**, posture **49**, cardiovascular pressures **50**, pulmonary artery systolic pressure measure **51**, pulmonary artery diastolic pressure measure **52**, respiratory rate **53**, ventilatory tidal volume **54**, minute ventilation **55**, transthoracic impedance **56**, cardiac output **57**, systemic blood pressure **58**, patient geographic location (altitude) **59**, mixed venous oxygen score **60**, arterial oxygen score **61**, arterial carbon dioxide score **62**, acidity (pH) level **63**, potassium [K⁺] level **64**, sodium [Na⁺] level **65**, glucose level **66**, blood urea nitrogen (BUN) and creatinine **67**, hematocrit **68**, hormonal levels **69**, lung injury chemical tests **70**, cardiac injury chemical tests **71**, myocardial blood flow **72**, central nervous system (CNS) injury chemical tests **73**, central nervous system blood flow **74**, interventions made by the implantable medical device or external medical device **75**, and the relative success of any interventions made **76**. In addition, the implantable medical device or external medical device communicates device-specific information, including battery status, general device status and program settings **77** and the time of day **78** for the various recorded measures. Other types of collected, recorded, combined, or derived measures are possible, as is known in the art.

[0036] The device and derived measures sets **24a**, **24b** (shown in FIG. 1), along with quality of life and symptom

measures sets **25a**, **25b**, as further described below with reference to FIG. 3, are continuously and periodically received by the server system **16** as part of the on-going patient care monitoring and analysis function. These regularly collected data sets are collectively categorized as the monitoring sets **27** (shown in FIG. 1). In addition, select device and derived measures sets **24a** and quality of life and symptom measures sets **25a** can be designated as a reference baseline **26** at the outset of patient care to improve the accuracy and meaningfulness of the serial monitoring sets **27**. Select patient information is collected, recorded, and derived during an initial period of observation or patient care, such as described in the related, commonly-owned U.S. Pat. No. 6,221,011, issued Apr. 24, 2001, the disclosure of which is incorporated herein by reference.

[0037] As an adjunct to remote patient care through the monitoring of measured physiological data via the implantable medical device **12** or external medical device **28**, quality of life and symptom measures sets **25a** can also be stored in the database **17** as part of the reference baseline **26**, if used, and the monitoring sets **27**. A quality of life measure is a semi-quantitative self-assessment of an individual patient's physical and emotional well being and a record of symptoms, such as provided by the Duke Activities Status Indicator. These scoring systems can be provided for use by the patient **11** on the personal computer **18** (shown in FIG. 1) to record his or her quality of life scores for both initial and periodic download to the server system **16**. FIG. 3 is a database schema showing, by way of example, the organization of a quality of life record **80** for use in the database **17**. The following information is recorded for a patient: overall health wellness **81**, psychological state **82**, activities of daily living **83**, work status **84**, geographic location **85**, family status **86**, shortness of breath **87**, cough **88**, sputum production **89**, sputum color **90**, energy level **91**, exercise tolerance **92**, chest discomfort **93**, and time of day **94**, and other quality of life and symptom measures as would be known to one skilled in the art.

[0038] The patient may also add non-device quantitative measures, such as the six-minute walk distance, as complementary data to the device and derived measures sets **24a**, **24b** and the symptoms during the six-minute walk to quality of life and symptom measures sets **25a**, **25b**.

[0039] Other types of quality of life and symptom measures are possible, such as those indicated by responses to the Minnesota Living with Heart Failure Questionnaire described in E. Braunwald, ed., "Heart Disease—A Textbook of Cardiovascular Medicine," pp. 452-454, W.B. Saunders Co. (1997), the disclosure of which is incorporated herein by reference. Similarly, functional classifications based on the relationship between symptoms and the amount of effort required to provoke them can serve as quality of life and symptom measures, such as the New York Heart Association (NYHA) classifications I, II, III and IV, adapted for use for lung disease rather than heart disease, also described in *Ibid*.

[0040] On a periodic basis, the patient information stored in the database **17** is analyzed and compared to pre-determined cutoff levels, which, when exceeded, can provide etiological indications of respiratory insufficiency symptoms. FIG. 4 is a database schema showing, by way of example, the organization of a combined measures set

record **95** for use in the database **17**. Each record **95** stores patient information obtained or derived from the device and derived measures sets **24a**, **24b** and quality of life and symptom measures sets **25a**, **25b** as maintained in the reference baseline **26**, if used, and the monitoring sets **27**. The combined measures set **95** represents those measures most (but not exhaustively or exclusively) relevant to a pathophysiology indicative of respiratory insufficiency and are determined as further described below with reference to FIGS. **8A-8B**. The following information is stored for a patient: heart rate **96**, heart rhythm (e.g., normal sinus vs. atrial fibrillation) **97**, pacing modality **98**, pulmonary artery systolic pressure measure **99**, pulmonary artery diastolic pressure measure **100**, cardiac output score **101**, arterial oxygen score **102**, mixed venous oxygen score **103**, respiratory rate **104**, tidal volume **105**, transthoracic impedance **106**, arterial carbon dioxide score **107**, right ventricular peak systolic pressure **108**, pulmonary artery end diastolic pressure **109**, patient activity score **110**, posture **111**, exercise tolerance quality of life and symptom measures **112**, respiratory distress quality of life and symptom measures **113**, cough **114**, sputum production **115**, any interventions made to treat respiratory insufficiency **116**, including treatment by medical device, via drug infusion administered by the patient or by a medical device, surgery, and any other form of medical intervention as is known in the art, the relative success of any such interventions made **117**, and time of day **118**. Other types of comparison measures regarding respiratory insufficiency are possible as is known in the art. In the described embodiment, each combined measures set **95** is sequentially retrieved from the database **17** and processed. Alternatively, each combined measures set **95** could be stored within a dynamic data structure maintained transitively in the random access memory of the server system **16** during the analysis and comparison operations.

[**0041**] FIG. **5** is a block diagram showing the software modules of the server system **16** of the system **10** of FIG. **1**. Each module is a computer program written as source code in a conventional programming language, such as the C or Java programming languages, and is presented for execution by the CPU of the server system **16** as object or byte code, as is known in the art. The various implementations of the source code and object and byte codes can be held on a computer-readable storage medium or embodied on a transmission medium in a carrier wave. The server system **16** includes three primary software modules, database module **125**, diagnostic module **126**, and feedback module **128**, which perform integrated functions as follows.

[**0042**] First, the database module **125** organizes the individual patient care records **23** stored in the database **17** (shown in FIG. **1**) and efficiently stores and accesses the reference baseline **26**, monitoring sets **27**, and patient care data maintained in those records. Any type of database organization could be utilized, including a flat file system, hierarchical database, relational database, or distributed database, such as provided by database vendors, such as Oracle Corporation, Redwood Shores, Calif.

[**0043**] Next, the diagnostic module **126** makes findings of respiratory insufficiency based on the comparison and analysis of the data measures from the reference baseline **26** and monitoring sets **27**. The diagnostic module includes three modules: comparison module **130**, analysis module **131**, and quality of life module **132**. The comparison module **130**

compares recorded and derived measures retrieved from the reference baseline **26**, if used, and monitoring sets **27** to indicator thresholds **129**. The database **17** stores individual patient care records **23** for patients suffering from various health disorders and diseases for which they are receiving remote patient care. For purposes of comparison and analysis by the comparison module **130**, these records can be categorized into peer groups containing the records for those patients suffering from similar disorders, as well as being viewed in reference to the overall patient population. The definition of the peer group can be progressively refined as the overall patient population grows. To illustrate, FIG. **6** is a record view showing, by way of example, a set of partial patient care records for care of patients with respiratory insufficiency stored in the database **17** for three patients, Patient 1, Patient 2, and Patient 3. For each patient, three sets of peer measures, X, Y, and Z are shown. Each of the measures, X, Y, and Z could be either collected or derived measures from the reference baseline **26**, if used, and monitoring sets **27**.

[**0044**] The same measures are organized into time-based sets with Set 0 representing sibling measures made at a reference time $t=0$. Similarly, Set $n-2$, Set $n-1$ and Set n each represent sibling measures made at later reference times $t=n-2$, $t=n-1$ and $t=n$, respectively. Thus, for a given patient, such as Patient 1, serial peer measures, such as peer measure X_0 through X_n , represent the same type of patient information monitored over time. The combined peer measures for all patients can be categorized into a health disorder- or disease-matched peer group. The definition of disease-matched peer group is a progressive definition, refined over time as the number of monitored patients grows. Measures representing different types of patient information, such as measures X_0 , Y_0 , and Z_0 , are sibling measures. These are measures which are also measured over time, but which might have medically significant meaning when compared to each other within a set for an individual patient.

[**0045**] The comparison module **130** performs two basic forms of comparisons. First, individual measures for a given patient can be compared to other individual measures for that same patient (self-referencing). These comparisons might be peer-to-peer measures, that is, measures relating to a one specific type of patient information, projected over time, for instance, X_n , X_{n-1} , X_{n-2} , . . . X_0 , or sibling-to-sibling measures, that is, measures relating to multiple types of patient information measured during the same time period, for a single snapshot, for instance, X_n , Y_n , and Z_n , or projected over time, for instance, X_n , Y_n , Z_n , X_{n-1} , Y_{n-1} , Z_{n-1} , X_{n-2} , Y_{n-2} , Z_{n-2} , . . . X_0 , Y_0 , Z_0 . Second, individual measures for a given patient can be compared to other individual measures for a group of other patients sharing the same disorder- or disease-specific characteristics (peer group referencing) or to the patient population in general (population referencing). Again, these comparisons might be peer-to-peer measures projected over time, for instance, X_n , X_n'' , X_{n-1} , X_{n-1}' , X_{n-1}'' , X_{n-2} , X_{n-2}' , X_{n-2}'' , . . . X_0 , X_0' , X_0'' , or comparing the individual patient's measures to an average from the group. Similarly, these comparisons might be sibling-to-sibling measures for single snapshots, for instance, X_n , X_n'' , Y_n , Y_n'' , Z_n , Z_n'' , or projected over time, for instance, X_n , X_n'' , X_{n-1} , X_{n-1}' , X_{n-1}'' , Y_n , Y_n'' , Y_{n-1} , Y_{n-1}' , Y_{n-1}'' , Z_n , Z_n'' , Z_{n-1} , Z_{n-1}' , Z_{n-1}'' , X_{n-2} , X_{n-2}' , X_{n-2}'' , Y_{n-2} , Y_{n-2}' , Y_{n-2}'' , Z_{n-2} , Z_{n-2}' , Z_{n-2}'' , X_0 , X_0' , X_0'' , Y_0 , Y_0' , Y_0'' , and Z_0 , Z_0' , Z_0'' . Other forms of

comparisons are feasible, including multiple disease diagnoses for diseases exhibiting similar abnormalities in physiological measures that result from a second disease but manifest in different combinations or onset in different temporal sequences.

[0046] FIG. 7 is a Venn diagram showing, by way of example, peer group overlap between the partial patient care records 23 of FIG. 1. Each patient care record 23 includes characteristics data 350, 351, 352, including personal traits, demographics, medical history, and related personal data, for patients 1, 2 and 3, respectively. For example, the characteristics data 350 for patient 1 might include personal traits which include gender and age, such as male and an age between 40-45; a demographic of resident of New York City; and a medical history consisting of chronic bronchitis, recurrent pneumonia, a history of an inferior myocardial infarction and diabetes. Similarly, the characteristics data 351 for patient 2 might include identical personal traits, thereby resulting in partial overlap 353 of characteristics data 350 and 351. Similar characteristics overlap 354, 355, 356 can exist between each respective patient. The overall patient population 357 would include the universe of all characteristics data. As the monitoring population grows, the number of patients with personal traits matching those of the monitored patient will grow, increasing the value of peer group referencing. Large peer groups, well matched across all monitored measures, will result in a well known natural history of disease and will allow for more accurate prediction of the clinical course of the patient being monitored. If the population of patients is relatively small, only some traits 356 will be uniformly present in any particular peer group. Eventually, peer groups, for instance, composed of 100 or more patients each, would evolve under conditions in which there would be complete overlap of substantially all salient data, thereby forming a powerful core reference group for any new patient being monitored.

[0047] Referring back to FIG. 5, the analysis module 131 analyzes the results from the comparison module 130, which are stored as a combined measures set 95 (not shown), to a set of indicator thresholds 129, as further described below with reference to FIGS. 8A-8B. Similarly, the quality of life module 132 compares quality of life and symptom measures 25a, 25b from the reference baseline 26 and monitoring sets 27, the results of which are incorporated into the comparisons performed by the analysis module 131, in part, to either refute or support the findings based on physiological "hard" data. Finally, the feedback module 128 provides automated feedback to the individual patient based, in part, on the patient status indicator 127 generated by the diagnostic module 126. As described above, the feedback could be by electronic mail or by automated voice mail or facsimile. The feedback can also include normalized voice feedback, such as described in the related, commonly-owned U.S. Pat. No. 6,203,495, issued Mar. 20, 2001, the disclosure of which is incorporated herein by reference. In addition, the feedback module 128 determines whether any changes to interventive measures are appropriate based on threshold stickiness ("hysteresis") 133, as further described below with reference to FIG. 16. The threshold stickiness 133 can prevent fickleness in the diagnostic routines resulting from transient, non-trending and non-significant fluctuations in the various collected and derived measures in favor of more certainty in diagnosis. However, in the case of some of the parameters being followed, such as activity and pulmonary artery sys-

tolic and diastolic pressures, abrupt spikes in these measures can be indicative of coughing and therefore helpful in indicating the onset of pulmonary insufficiency. In a further embodiment of the present invention, the feedback module 128 includes a patient query engine 134 that enables the individual patient 11 to interactively query the server system 16 regarding the diagnosis, therapeutic maneuvers, and treatment regimen. Conversely, the patient query engines 134, found in interactive expert systems for diagnosing medical conditions, can interactively query the patient. Using the personal computer 18 (shown in FIG. 1), the patient can have an interactive dialogue with the automated server system 16, as well as human experts as necessary, to self assess his or her medical condition. Such expert systems are well known in the art, an example of which is the MYCIN expert system developed at Stanford University and described in Buchanan, B. & Shortliffe, E., "RULE-BASED EXPERT SYSTEMS. The MYCIN Experiments of the Stanford Heuristic Programming Project," Addison-Wesley (1984). The various forms of feedback described above help to increase the accuracy and specificity of the reporting of the quality of life and symptomatic measures.

[0048] FIGS. 8A-8B are flow diagrams showing a method for diagnosing and monitoring respiratory insufficiency and outcomes thereof 135 using an automated collection and analysis patient care system 10 in accordance with the present invention. First, the indicator thresholds 129 (shown in FIG. 5) are set (block 136) by defining a quantifiable physiological measure of a pathophysiology indicative of respiratory insufficiency and relating to the each type of patient information in the combined device and derived measures set 951 (shown in FIG. 4). The actual values of each indicator threshold can be finite cutoff values, weighted values, or statistical ranges, as discussed below with reference to FIGS. 11A-11F. Next, the reference baseline 26 (block 137) and monitoring sets 27 (block 138) are retrieved from the database 17, as further described below with reference to FIGS. 9 and 10, respectively. Each measure in the combined device and derived measures set 95 is tested against the threshold limits defined for each indicator threshold 129 (block 139), as further described below with reference to FIGS. 11A-11F. The potential onset, progression, regression, or status quo of respiratory insufficiency is then evaluated (block 140) based upon the findings of the threshold limits tests (block 139), as further described below with reference to FIGS. 13A-13C, 14A-14C, 15A-15C.

[0049] In a further embodiment, multiple near-simultaneous disorders are considered in addition to primary respiratory insufficiency. Primary respiratory insufficiency is defined as the onset or progression of respiratory insufficiency without obvious inciting cause. Secondary respiratory insufficiency is defined as the onset or progression of respiratory insufficiency (in a patient with or without pre-existing respiratory insufficiency) from another disease process, such as congestive heart failure, coronary insufficiency, atrial fibrillation, and so forth. Other health disorders and diseases can potentially share the same forms of symptomatology as respiratory insufficiency, such as congestive heart failure, myocardial ischemia, pneumonia, exacerbation of chronic bronchitis, renal failure, sleep-apnea, stroke, anemia, atrial fibrillation, other cardiac arrhythmias, and so forth. If more than one abnormality is present, the relative sequence and magnitude of onset of abnormalities in the

monitored measures becomes most important in sorting and prioritizing disease diagnosis and treatment.

[0050] Thus, if other disorders or diseases are being cross-referenced and diagnosed (block 141), their status is determined (block 142). In the described embodiment, the operations of ordering and prioritizing multiple near-simultaneous disorders (box 151) by the testing of threshold limits and analysis in a manner similar to congestive heart failure as described above, preferably in parallel to the present determination, is described in the related, commonly-owned U.S. Pat. No. 6,440,066, issued Aug. 27, 2002, the disclosure of which is incorporated herein by reference. If respiratory insufficiency is due to an obvious inciting cause, i.e., secondary respiratory insufficiency, (block 143), an appropriate treatment regimen for respiratory insufficiency as exacerbated by other disorders is adopted that includes treatment of secondary disorders, e.g., congestive heart failure, myocardial ischemia, atrial fibrillation, and so forth (block 144) and a suitable patient status indicator 127 for respiratory insufficiency is provided (block 146) to the patient. Suitable devices and approaches to diagnosing and treating congestive heart failure, myocardial infarction, and atrial fibrillation are described in related, commonly-owned U.S. Pat. No. 6,336,903, issued Jan. 8, 2002; U.S. Pat. No. 6,368,284, issued Apr. 9, 2002; and U.S. Pat. No. 6,411,840, issued Jun. 25, 2002, pending, the disclosures of which are incorporated herein by reference.

[0051] Otherwise, if primary respiratory insufficiency is indicated (block 143), a primary treatment regimen is followed (block 145). A patient status indicator 127 for respiratory insufficiency is provided (block 146) to the patient regarding physical well-being, disease prognosis, including any determinations of disease onset, progression, regression, or status quo, and other pertinent medical and general information of potential interest to the patient.

[0052] Finally, in a further embodiment, if the patient submits a query to the server system 16 (block 147), the patient query is interactively processed by the patient query engine (block 148). Similarly, if the server elects to query the patient (block 149), the server query is interactively processed by the server query engine (block 150). The method then terminates if no further patient or server queries are submitted.

[0053] FIG. 9 is a flow diagram showing the routine for retrieving reference baseline sets 137 for use in the method of FIGS. 8A-8B. The purpose of this routine is to retrieve the appropriate reference baseline sets 26, if used, from the database 17 based on the types of comparisons being performed. First, if the comparisons are self referencing with respect to the measures stored in the individual patient care record 23 (block 152), the reference device and derived measures set 24a and reference quality of life and symptom measures set 25a, if used, are retrieved for the individual patient from the database 17 (block 153). Next, if the comparisons are peer group referencing with respect to measures stored in the patient care records 23 for a health disorder- or disease-specific peer group (block 154), the reference device and derived measures set 24a and reference quality of life and symptom measures set 25a, if used, are retrieved from each patient care record 23 for the peer group from the database 17 (block 155). Data for each measure (e.g., minimum, maximum, averaged, standard deviation

(SD), and trending data) from the reference baseline 26 for the peer group is then calculated (block 156). Finally, if the comparisons are population referencing with respect to measures stored in the patient care records 23 for the overall patient population (block 157), the reference device and derived measures set 24a and reference quality of life and symptom measures set 25a, if used, are retrieved from each patient care record 23 from the database 17 (block 158). Minimum, maximum, averaged, standard deviation, and trending data and other numerical processes using the data, as is known in the art, for each measure from the reference baseline 26 for the peer group is then calculated (block 159). The routine then returns.

[0054] FIG. 10 is a flow diagram showing the routine for retrieving monitoring sets 138 for use in the method of FIGS. 8A-8B. The purpose of this routine is to retrieve the appropriate monitoring sets 27 from the database 17 based on the types of comparisons being performed. First, if the comparisons are self referencing with respect to the measures stored in the individual patient care record 23 (block 160), the device and derived measures set 24b and quality of life and symptom measures set 25b, if used, are retrieved for the individual patient from the database 17 (block 161). Next, if the comparisons are peer group referencing with respect to measures stored in the patient care records 23 for a health disorder- or disease-specific peer group (block 162), the device and derived measures set 24b and quality of life and symptom measures set 25b, if used, are retrieved from each patient care record 23 for the peer group from the database 17 (block 163). Data for each measure (e.g., minimum, maximum, averaged, standard deviation, and trending data) from the monitoring sets 27 for the peer group is then calculated (block 164). Finally, if the comparisons are population referencing with respect to measures stored in the patient care records 23 for the overall patient population (block 165), the device and derived measures set 24b and quality of life and symptom measures set 25b, if used, are retrieved from each patient care record 23 from the database 17 (block 166). Minimum, maximum, averaged, standard deviation, and trending data and other numerical processes using the data, as is known in the art, for each measure from the monitoring sets 27 for the peer group is then calculated (block 167). The routine then returns.

[0055] FIGS. 11A-11F are flow diagrams showing the routine for testing threshold limits 139 for use in the method of FIGS. 8A and 8B. The purpose of this routine is to analyze, compare, and log any differences between the observed, objective measures stored in the reference baseline 26, if used, and the monitoring sets 27 to the indicator thresholds 129. Briefly, the routine consists of tests pertaining to each of the indicators relevant to diagnosing and monitoring respiratory insufficiency. The threshold tests focus primarily on: (1) changes to and rates of change for the indicators themselves, as stored in the combined device and derived measures set 95 (shown in FIG. 4) or similar data structure; and (2) violations of absolute threshold limits which trigger an alert. The timing and degree of change may vary with each measure and with the natural fluctuations noted in that measure during the reference baseline period. In addition, the timing and degree of change might also vary with the individual and the natural history of a measure for that patient.

[0056] One suitable approach to performing the threshold tests uses a standard statistical linear regression technique using a least squares error fit. The least squares error fit can be calculated as follows:

$$y = \beta_0 + \beta_1 x \quad (1)$$

$$\beta = \frac{SS_{xy}}{SS_{xx}} \quad (2)$$

$$SS_{xy} = \sum_{i=1}^n x_i y_i - \frac{\left(\sum_{i=1}^n x_i \right) \left(\sum_{i=1}^n y_i \right)}{n} \quad (3)$$

$$SS_{xx} = \sum_{i=1}^n x_i^2 - \frac{\left(\sum_{i=1}^n x_i \right)^2}{n} \quad (4)$$

where n is the total number of measures, x_i is the time of day for measure i, and y_i is the value of measure i, β_1 is the slope, and β_0 is the y-intercept of the least squares error line. A positive slope β_1 indicates an increasing trend, a negative slope β_1 indicates a decreasing trend, and no slope indicates no change in patient condition for that particular measure. A predicted measure value can be calculated and compared to the appropriate indicator threshold 129 for determining whether the particular measure has either exceeded an acceptable threshold rate of change or the absolute threshold limit.

[0057] For any given patient, three basic types of comparisons between individual measures stored in the monitoring sets 27 are possible: self referencing, peer group, and general population, as explained above with reference to FIG. 6. In addition, each of these comparisons can include comparisons to individual measures stored in the pertinent reference baselines 26.

[0058] The indicator thresholds 129 for detecting a trend indicating progression into a state of respiratory insufficiency or a state of imminent or likely respiratory insufficiency, for example, over a one week time period, can be as follows:

[0059] (1) Respiratory rate (block 170): If the respiratory rate has increased over 1.0 SD from the mean respiratory rate in the reference baseline 26 (block 171), the increased respiratory rate and time span over which it occurs are logged in the combined measures set 95 (block 172).

[0060] (2) Heart rate (block 173): If the heart rate has increased over 1.0 SD from the mean heart rate in the reference baseline 26 (block 174), the increased heart rate and time span over which it occurs are logged in the combined measures set 95 (block 175).

[0061] (3) Transthoracic impedance (block 176): If the transthoracic impedance has increased over 1.0 SD from the mean transthoracic impedance in the reference baseline 26 (block 177), the increased transthoracic impedance and time span are logged in the combined measures set 95 (block 178).

[0062] (4) The ventilatory tidal volume (block 179): If the tidal volume has increased over 1.0 SD from the

tidal volume score in the reference baseline 26 (block 180), the increased tidal volume score and time span are logged in the combined measures set 95 (block 181).

[0063] (5) Arterial oxygen score (block 182): If the arterial oxygen score has decreased over 1.0 SD from the arterial oxygen score in the reference baseline 26 (block 183), the decreased arterial oxygen score and time span are logged in the combined measures set 95 (block 184).

[0064] (6) Arterial carbon dioxide score (block 185): If the arterial carbon dioxide score has decreased over 1.0 SD from the arterial carbon dioxide score in the reference baseline 26 (block 186), the decreased arterial carbon dioxide score and time span are logged in the combined measures set 95 (block 187).

[0065] (7) Patient activity score (block 188): If the mean patient activity score has decreased over 1.0 SD from the mean patient activity score in the reference baseline 26 (block 189), the decreased patient activity score and time span are logged in the combined measures set 95 (block 190).

[0066] (8) Temperature (block 191): If the patient temperature score has increased over 1.0 SD from the mean patient temperature score in the reference baseline 26 (block 192), the increased patient temperature score and the time span are logged in the combined measures set 95 (block 193).

[0067] (9) Spikes in patient activity (block 194): If short-lived spikes in the patient activity score occur over time periods under 5 minutes compared to the reference baseline 26 (block 195), the spike in patient activity score and time span are logged in the combined measures set 95 (block 196).

[0068] (10) Spikes in pulmonary arterial pressure (PAP) (block 197): If short-lived spikes in the pulmonary arterial pressure score occur over time periods under 5 minutes compared to the reference baseline 26 (block 198), the spike in the pulmonary arterial pressure score and time span are logged in the combined measures set 95 (block 199). In the described embodiment, the mean arterial pressure on any spike in the arterial pressure tracing could be utilized.

[0069] (11) Exercise tolerance quality of life (QOL) measures (block 200): If the exercise tolerance QOL has decreased over 1.0 SD from the mean exercise tolerance in the reference baseline 26 (block 201), the decrease in exercise tolerance and the time span over which it occurs are logged in the combined measures set 95 (block 202).

[0070] (12) Respiratory distress quality of life (QOL) measures (block 203): If the respiratory distress QOL measure has deteriorated by more than 1.0 SD from the mean respiratory distress QOL measure in the reference baseline 26 (block 204), the increase in respiratory distress and the time span over which it occurs are logged in the combined measures set 95 (block 205).

[0071] (13) Spikes in right ventricular (RV) pressure (block 206): If short-lived spikes in the right ventricular pressure occur over time periods under 5 minutes

compared to the reference baseline **26** (block **207**), the spike in the right ventricular pressure and time span are logged in the combined measures set **95** (block **208**).

[**0072**] (14) Spikes in transthoracic impedance (TTZ) (block **209**): If short-lived spikes in the transthoracic impedance occur over time periods under 5 minutes compared to the reference baseline **26** (block **210**), the spike in the transthoracic impedance and time span are logged in the combined measures set **95** (block **211**).

[**0073**] (15) Atrial fibrillation (block **212**): The presence or absence of atrial fibrillation (AF) is determined and, if present (block **213**), atrial fibrillation is logged (block **214**).

[**0074**] (16) Rhythm changes (block **215**): The type and sequence of rhythm changes is significant and is determined (block **216**) based on the timing of the relevant rhythm measure, such as sinus rhythm. For instance, a finding that a rhythm change to atrial fibrillation precipitated respiratory measures changes can indicate therapy directions against atrial fibrillation rather than the primary development of respiratory insufficiency. Thus, if there are rhythm changes (block **217**), the sequence of the rhythm changes and time span are logged (block **211**).

[**0075**] Note also that an inversion of the indicator thresholds **129** defined above could similarly be used for detecting a trend in disease regression. One skilled in the art would recognize that these measures would vary based on whether or not they were recorded during rest or during activity and that the measured activity score can be used to indicate the degree of patient rest or activity. The patient activity score can be determined via an implantable motion detector, for example, as described in U.S. Pat. No. 4,428,378, issued Jan. 31, 1984, to Anderson et al., the disclosure of which is incorporated herein by reference.

[**0076**] The indicator thresholds **129** for detecting a trend towards a state of respiratory insufficiency can also be used to declare, a priori, respiratory insufficiency present, regardless of pre-existing trend data when certain limits are established, such as:

[**0077**] (1) An absolute limit of arterial oxygen (block **182**) less than 85 mm Hg is an a priori definition of respiratory insufficiency from decreased oxygenation.

[**0078**] (2) An absolute limit of arterial carbon dioxide (block **185**) falling below 25 mm Hg (in the absence of marked exercise) or greater than 50 mm Hg are both a priori definitions of respiratory insufficiency as indicated by hyperventilation and hypoventilation, respectively.

[**0079**] FIG. **12** is a flow diagram showing the routine for evaluating the onset, progression, regression and status quo of respiratory insufficiency **140** for use in the method of FIGS. **8A** and **8B**. The purpose of this routine is to evaluate the presence of sufficient indicia to warrant a diagnosis of the onset, progression, regression, and status quo of respiratory insufficiency. Quality of life and symptom measures **25a**, **25b** can be included in the evaluation (block **230**) by determining whether any of the individual quality of life and symptom measures **25a**, **25b** have changed relative to the previously collected quality of life and symptom measures

from the monitoring sets **27** and the reference baseline **26**, if used. For example, an increase in the shortness of breath measure **87** and exercise tolerance measure **92** would corroborate a finding of respiratory insufficiency. Similarly, a transition from NYHA Class II to NYHA Class III would indicate a deterioration or, conversely, a transition from NYHA Class III to NYHA Class II status would indicate improvement or progress when adapting the NYHA classifications for their parallel in lung disorders. Incorporating the quality of life and symptom measures **25a**, **25b** into the evaluation can help, in part, to refute or support findings based on physiological data. Next, a determination as to whether any changes to interventive measures are appropriate based on threshold stickiness ("hysteresis") is made (block **231**), as further described below with reference to FIG. **16**.

[**0080**] The routine returns upon either the determination of a finding or elimination of all factors as follows. If a finding of respiratory insufficiency was not previously diagnosed (block **232**), a determination of disease onset is made (block **233**), as further described below with reference to FIGS. **13A-13C**. Otherwise, if respiratory insufficiency was previously diagnosed (block **232**), a further determination of either disease progression or worsening (block **234**) or regression or improving (block **235**) is made, as further described below with reference to FIGS. **14A-14C** and **15A-15C**, respectively. If, upon evaluation, neither disease onset (block **233**), worsening (block **234**) or improving (block **235**) is indicated, a finding of status quo is appropriate (block **236**) and noted (block **237**). Otherwise, respiratory insufficiency and the related outcomes are actively managed (block **238**) through the administration of, non-exclusively, antibiotic and antiviral therapies, bronchodilator therapies, oxygen therapies, antiinflammation therapies, electrical therapies, mechanical therapies, and other therapies as are known in the art. The management of respiratory insufficiency is described, by way of example, in A. S. Fauci et al. (Eds.), "Harrison's Principles of Internal Medicine," pp. 1407-1491, McGraw-Hill, 14th Ed. (1997), the disclosure of which is incorporated herein by reference. The routine then returns.

[**0081**] FIGS. **13A-13C** are flow diagrams showing the routine for determining an onset of respiratory insufficiency **232** for use in the routine of FIG. **12**. Respiratory insufficiency is possible based on two general symptom categories: reduced exercise capacity (block **244**) and respiratory distress (block **256**). An effort is made to diagnose respiratory insufficiency manifesting primarily as resulting in reduced exercise capacity (block **244**) and/or increased respiratory distress (block **256**). Reduced exercise capacity and respiratory distress can generally serve as markers of low systemic arterial oxygenation. The clinical aspects of respiratory insufficiency are described, by way of example, in A. S. Fauci et al. (Eds.), "Harrison's Principles of Internal Medicine," pp. 1410-1419, McGraw-Hill, 14th Ed. (1997), the disclosure of which is incorporated herein by reference.

[**0082**] As primary pulmonary disease considerations, multiple individual indications (blocks **240-243**, **245-255**) should be present for the two principal findings of respiratory insufficiency related reduced exercise capacity (block **244**), or respiratory insufficiency related respiratory distress (block **256**), to be indicated, both for disease onset or progression. The presence of primary key findings alone can

be sufficient to indicate an onset of respiratory insufficiency and secondary key findings serve to corroborate disease onset. Note the presence of any abnormality can trigger an analysis for the presence or absence of secondary disease processes, such as the presence of atrial fibrillation or congestive heart failure. Secondary disease considerations can be evaluated using the same indications (see, e.g., blocks 141-144 of FIGS. 8A-8B), but with adjusted indicator thresholds 129 (shown in FIG. 5) triggered at a change of 0.5 SD, for example, instead of 1.0 SD.

[0083] In the described embodiment, the reduced exercise capacity and respiratory distress findings (blocks 244, 256) can be established by consolidating the individual indications (blocks 240-243, 245-255) in several ways. First, in a preferred embodiment, each individual indication (blocks 240-243, 245-255) is assigned a scaled index value correlating with the relative severity of the indication. For example, decreased cardiac output (block 240) could be measured on a scale from '1' to '5' wherein a score of '1' indicates no change in cardiac output from the reference point, a score of '2' indicates a change exceeding 0.5 SD, a score of '3' indicates a change exceeding 1.0 SD, a score of '4' indicates a change exceeding 2.0 SD, and a score of '5' indicates a change exceeding 3.0 SD. The index value for each of the individual indications (blocks 240-243, 245-255) can then either be aggregated or averaged with a result exceeding the aggregate or average maximum indicating an appropriate respiratory insufficiency finding.

[0084] Preferably, all scores are weighted depending upon the assignments made from the measures in the reference baseline 26. For instance, arterial partial pressure of oxygen 102 could be weighted more importantly than respiratory rate 104 if the respiratory rate in the reference baseline 26 is particularly high at the outset, making the detection of further disease progression from increases in respiratory rate, less sensitive. In the described embodiment, arterial partial pressure of oxygen 102 receives the most weight in determining a reduced exercise capacity finding whereas arterial partial pressure of carbon dioxide 107 receives the most weight in determining a respiratory distress or dyspnea finding.

[0085] Alternatively, a simple binary decision tree can be utilized wherein each of the individual indications (blocks 240-243, 245-255) is either present or is not present. All or a majority of the individual indications (blocks 240-243, 245-255) should be present for the relevant respiratory insufficiency finding to be affirmed.

[0086] Other forms of consolidating the individual indications (blocks 240-243, 245-255) are feasible.

[0087] FIGS. 14A-14C are flow diagrams showing the routine for determining a progression or worsening of respiratory insufficiency 234 for use in the routine of FIG. 12. The primary difference between the determinations of disease onset, as described with reference to FIGS. 13A-13C, and disease progression is the evaluation of changes indicated in the same factors present in a disease onset finding. Thus, a revised respiratory insufficiency finding is possible based on the same two general symptom categories: reduced exercise capacity (block 274) and respiratory distress (block 286). The same factors which need be indicated to warrant a diagnosis of respiratory insufficiency onset are evaluated to determine disease progression.

[0088] Similarly, FIGS. 15A-15C are flow diagrams showing the routine for determining a regression or improving of respiratory distress 235 for use in the routine of FIG. 12. The same factors as described above with reference to FIGS. 13A-13C and 14A-14C, trending in opposite directions from disease onset or progression, are evaluated to determine disease regression. As primary cardiac disease considerations, multiple individual indications (blocks 300-303, 305-315) should be present for the two principal findings of respiratory insufficiency related reduced exercise capacity (block 304), or respiratory insufficiency related respiratory distress (block 316), to indicate disease regression.

[0089] FIG. 16 is a flow diagram showing the routine for determining threshold stickiness ("hysteresis") 231 for use in the method of FIG. 12. Stickiness, also known as hysteresis, is a medical practice doctrine whereby a diagnosis or therapy will not be changed based upon small or temporary changes in a patient reading, even though those changes might temporarily move into a new zone of concern. For example, if a patient measure can vary along a scale of '1' to '10' with '10' being worse, a transient reading of '6,' standing alone, on a patient who has consistently indicated a reading of '5' for weeks will not warrant a change in diagnosis without a definitive prolonged deterioration first being indicated. Stickiness dictates that small or temporary changes require more diagnostic certainty, as confirmed by the persistence of the changes, than large changes would require for any of the monitored (device) measures. Stickiness also makes reversal of important diagnostic decisions, particularly those regarding life-threatening disorders, more difficult than reversal of diagnoses of modest import. As an example, automatic external defibrillators (AEDs) manufactured by Heartstream, a subsidiary of Agilent Technologies, Seattle, Wash., monitor heart rhythms and provide interventional shock treatment for the diagnosis of ventricular fibrillation. Once diagnosis of ventricular fibrillation and a decision to shock the patient has been made, a pattern of no ventricular fibrillation must be indicated for a relatively prolonged period before the AED changes to a "no-shock" decision. As implemented in this AED example, stickiness mandates certainty before a decision to shock is disregarded. In practice, stickiness also dictates that acute deteriorations in disease state are treated aggressively while chronic, more slowly progressing disease states are treated in a more tempered fashion. However, in the case of some of the parameters being followed, such as activity and pulmonary artery systolic pressure, abrupt spikes in these measures can be indicative of coughing and therefore helpful in indicating the onset of a disorder that might lead to pulmonary insufficiency.

[0090] Thus, if the patient status indicates a status quo (block 330), no changes in treatment or diagnosis are indicated and the routine returns. Otherwise, if the patient status indicates a change away from status quo (block 330), the relative quantum of change and the length of time over which the change has occurred is determinative. If the change of approximately 0.5 SD has occurred over the course of about one month (block 331), a gradually deteriorating condition exists (block 332) and a very tempered diagnostic, and if appropriate, treatment program is undertaken. If the change of approximately 1.0 SD has occurred over the course of about one week (block 333), a more rapidly deteriorating condition exists (block 334) and a

slightly more aggressive diagnostic, and if appropriate, treatment program is undertaken. If the change of approximately 2.0 SD has occurred over the course of about one day (block 335), an urgently deteriorating condition exists (block 336) and a moderately aggressive diagnostic, and if appropriate, treatment program is undertaken. If the change of approximately 3.0 SD has occurred over the course of about one hour (block 337), an emergency condition exists (block 338) and an immediate diagnostic, and if appropriate, treatment program is undertaken as is practical. Finally, if the change and duration fall outside the aforementioned ranges (blocks 331-338), an exceptional condition exists (block 339) and the changes are reviewed manually, if necessary. The routine then returns. These threshold limits and time ranges may then be adapted depending upon patient history and peer-group guidelines.

[0091] The present invention provides several benefits. One benefit is improved predictive accuracy from the outset of patient care when a reference baseline is incorporated into the automated diagnosis. Another benefit is an expanded knowledge base created by expanding the methodologies applied to a single patient to include patient peer groups and the overall patient population. Collaterally, the information maintained in the database could also be utilized for the development of further predictive techniques and for medical research purposes. Yet a further benefit is the ability to hone and improve the predictive techniques employed through a continual reassessment of patient therapy outcomes and morbidity rates.

[0092] Other benefits include an automated, expert system approach to the cross-referral, consideration, and potential finding or elimination of other diseases and health disorders with similar or related etiological indicators and for those other disorders that may have an impact on respiratory insufficiency. Although disease specific markers will prove very useful in discriminating the underlying cause of symptoms, many diseases, other than respiratory insufficiency, will alter some of the same physiological measures indicative of respiratory insufficiency. Consequently, an important aspect of considering the potential impact of other disorders will be, not only the monitoring of disease specific markers, but the sequencing of change and the temporal evolution of more general physiological measures, for example respiratory rate, pulmonary artery diastolic pressure, and cardiac output, to reflect disease onset, progression or regression in more than one type of disease process, especially congestive heart failure from whatever cause.

[0093] While the invention has been particularly shown and described as referenced to the embodiments thereof, those skilled in the art will understand that the foregoing and other changes in form and detail may be made therein without departing from the spirit and scope of the invention.

What is claimed is:

1. A programmer for use in determining edema through remote patient monitoring, comprising:

an interface through which to interrogate and program a medical device for a patient; and

storage to accumulate device measures regularly recorded by the medical device pending upload,

wherein derived measures are determined at least in part from the device measures and a clinical indication of

one of an absence, onset, progression, regression, and status quo of edema is evaluated based on the device and derived measures.

2. A programmer according to claim 1, further comprising:

programming parameters for download by the programmer to effect the therapy modification for the medical device selected from the group comprising device parameters, drug infusion parameters, and neural stimulation parameters.

3. A programmer according to claim 2, wherein the programming parameters further comprise at least one of:

monitoring parameters to monitor the patient while asleep; and

exclusion parameters to exclude one or more of the device and derived measures when the patient exhibits an increased respiratory rate due to exercise or activity level.

4. A programmer according to claim 1, wherein the device and derived measures are obtained at least in part through one or more of transthoracic impedance and respiration rate.

5. A system for determining edema through remote patient monitoring, comprising:

storage to accumulate device measures regularly recorded by a medical device as raw physiometry for a patient; and

a processor, comprising:

an analysis module to determine derived measures at least in part from the device measures; and

an evaluation module to evaluate a clinical indication of one of an absence, onset, progression, regression, and status quo of edema based on the device and derived measures.

6. A system according to claim 5, further comprising:

a programmer to periodically monitor the device and derived measures; and

a diagnostic module to evaluate therapy modification for managing the patient.

7. A system according to claim 6, further comprising:

programming parameters to effect the therapy modification for the medical device selected from the group comprising device parameters, drug infusion parameters, and neural stimulation parameters.

8. A system according to claim 7, wherein the programming parameters further comprise at least one of:

monitoring parameters to monitor the patient while asleep; and

exclusion parameters to exclude one or more of the device and derived measures when the patient exhibits an increased respiratory rate due to exercise or activity level.

9. A system according to claim 5, wherein the device and derived measures are obtained at least in part through one or more of transthoracic impedance and respiration rate.

10. A system according to claim 5, further comprising:

a monitoring module to monitor well being of the patient in conjunction with conditions selected from the group comprising congestive heart failure, myocardial

ischemia, bronchitis, emphysema, pneumonia, pulmonary emboli, tumor infiltration of the lungs, abnormalities of the interstitium of the lungs potentially infectious in origin, immunological abnormalities, exposure to environmental pathogens, and high blood pressure.

11. A system according to claim 5, wherein the evaluation module processes the device and derived measures, comprising at least one of:

a recordation module to record, determine, and store the device and derived measures for edema evaluation;

an alert module to generate alerts in response to significant events characterized by the device and derived measures selected from the group comprising absolute measures, long term average measures, and short term average measures; and

an execution module to execute actions in response to indications selected from the group comprising the significant events, ECG signals, activity level, temperature, pressure, and oxygen saturation.

12. A system according to claim 5, further comprising:

trigger conditions characterized by the device and derived measures selected from the group comprising absolute measures, pattern matching, absolute differences between averaged measures, and comparisons of a current measure to another current measure or a previous measure; and

an execution monitor to executed the trigger conditions to cause activation of one or more of a therapy readout, alarm, or change in raw physiometry measured.

13. A method for determining edema through remote patient monitoring, comprising:

accumulating device measures regularly recorded by a medical device as raw physiometry for a patient and determining derived measures at least in part from the device measures; and

evaluating a clinical indication of one of an absence, onset, progression, regression, and status quo of edema based on the device and derived measures.

14. A method according to claim 13, further comprising: periodically monitoring the device and derived measures; and

evaluating therapy modification for managing the patient.

15. A method according to claim 14, further comprising:

effecting the therapy modification through programming parameters for the medical device selected from the group comprising device parameters, drug infusion parameters, and neural stimulation parameters.

16. A method according to claim 15, further comprising: generating the programming parameters, further comprising at least one of:

defining monitoring parameters to monitor the patient while asleep; and

defining exclusion parameters to exclude one or more of the device and derived measures when the patient exhibits an increased respiratory rate due to exercise or activity level.

17. A method according to claim 13, further comprising: obtaining the device and derived measures at least in part through one or more of transthoracic impedance and respiration rate.

18. A method according to claim 13, further comprising: monitoring well being of the patient in conjunction with conditions selected from the group comprising congestive heart failure, myocardial ischemia, bronchitis, emphysema, pneumonia, pulmonary emboli, tumor infiltration of the lungs, abnormalities of the interstitium of the lungs potentially infectious in origin, immunological abnormalities, exposure to environmental pathogens, and high blood pressure.

19. A method according to claim 13, further comprising: processing the device and derived measures, comprising at least one of:

recording, determining, and storing the device and derived measures for edema evaluation;

generating alerts in response to significant events characterized by the device and derived measures selected from the group comprising absolute measures, long term average measures, and short term average measures; and

executing actions in response to indications selected from the group comprising the significant events, ECG signals, activity level, temperature, pressure, and oxygen saturation.

20. A method according to claim 13, further comprising: defining trigger conditions characterized by the device and derived measures selected from the group comprising absolute measures, pattern matching, absolute differences between averaged measures, and comparisons of a current measure to another current measure or a previous measure; and

executing the trigger conditions to cause activation of one or more of a therapy readout, alarm, or change in raw physiometry measured.

21. A computer-readable storage medium holding code for performing the method according to claim 13.

* * * * *

专利名称(译)	用于通过远程患者监测确定水肿的系统和方法		
公开(公告)号	US20070203415A1	公开(公告)日	2007-08-30
申请号	US11/789388	申请日	2007-04-23
[标]申请(专利权)人(译)	BARDY GUST ^ h		
申请(专利权)人(译)	BARDY GUST ^ h		
当前申请(专利权)人(译)	BARDY GUST ^ h		
[标]发明人	BARDY GUST H		
发明人	BARDY, GUST H.		
IPC分类号	A61B5/00 A61B5/02 A61B5/08 G06F19/00		
CPC分类号	A61B5/0031 A61B2505/03 A61B5/0215 A61B5/024 A61B5/0452 A61B5/0535 A61B5/0816 A61B5/1112 A61B5/1116 A61B5/145 A61B5/14532 A61B5/14539 A61B5/14546 A61N1/3624 G06F19/322 G06F19/324 G06F19/3418 G06F19/3443 G06F19/345 G06F19/3456 G06F19/3481 G06F19/3487 Y10S128/92 Y10S128/923 A61B5/02055 G16H10/60 G16H15/00 G16H20/30 G16H20/40 G16H40/67 G16H50/20 G16H50/70 G16H70/60		
外部链接	Espacenet USPTO		

摘要(译)

提出了一种用于通过远程患者监测来确定水肿的系统和方法。由医疗设备定期记录的设备测量被累积为患者的原始生理测量。派生度量至少部分地根据设备度量来确定。基于装置和衍生的测量来评估水肿的缺失，发作，进展，退化和现状之一的临床指征。

