



US008652038B2

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

(10) **Patent No.:** **US 8,652,038 B2**
(45) **Date of Patent:** ***Feb. 18, 2014**

(54) **HEALTH MONITORING APPLIANCE**

(56) **References Cited**

(71) Applicant: **Bao Tran**, Saratoga, CA (US)

U.S. PATENT DOCUMENTS

(72) Inventors: **Bao Tran**, Saratoga, CA (US); **Ha Tran**, Saratoga, CA (US)

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

This patent is subject to a terminal disclaimer.

(21) Appl. No.: **13/775,055**

(22) Filed: **Feb. 22, 2013**

(65) **Prior Publication Data**

US 2013/0178718 A1 Jul. 11, 2013

Related U.S. Application Data

(63) Continuation of application No. 13/488,351, filed on Jun. 4, 2012, now Pat. No. 8,323,189, and a continuation-in-part of application No. 13/337,217, filed on Dec. 26, 2011, now Pat. No. 8,328,718, and a continuation of application No. 12/426,232, filed on Apr. 18, 2009, now Pat. No. 8,121,673, and a continuation of application No. 11/433,900, filed on May 12, 2006, now Pat. No. 7,539,532, and a continuation of application No. 13/676,138, filed on Nov. 14, 2012, now Pat. No. 8,475,368.

(51) **Int. Cl.**
A61B 5/00 (2006.01)

(52) **U.S. Cl.**
USPC **600/300**

(58) **Field of Classification Search**
USPC 600/300, 485, 509, 513; 128/903
See application file for complete search history.

4,234,926 A	11/1980	Wallace et al.
4,275,742 A	6/1981	Faisandier
4,320,767 A	3/1982	Villa-Real
4,412,545 A	11/1983	Okino et al.
4,420,000 A	12/1983	Bailey
4,537,202 A	8/1985	Mancini et al.
4,557,270 A	12/1985	John
4,595,018 A	6/1986	Rantala
4,656,319 A	4/1987	Bially
4,679,144 A	7/1987	Cox et al.
4,778,449 A	10/1988	Weber et al.
4,807,642 A	2/1989	Brown
4,819,648 A	4/1989	Ko
4,862,359 A	8/1989	Trivedi et al.
4,869,261 A	9/1989	Penaz
4,880,013 A	11/1989	Chio
4,907,597 A	3/1990	Chamonu
4,922,229 A	5/1990	Guenst
4,964,411 A	10/1990	Johnson et al.
5,090,418 A	2/1992	Squires et al.
RE34,015 E	8/1992	Duffy
5,140,990 A	8/1992	Jones et al.
5,161,529 A	11/1992	Stotts et al.
5,162,991 A	11/1992	Chio
5,271,411 A	12/1993	Ripley et al.
5,285,793 A	2/1994	Slovut et al.
5,287,859 A	2/1994	John

(Continued)

OTHER PUBLICATIONS

U.S. Appl. No. 10/938,783, Wen.

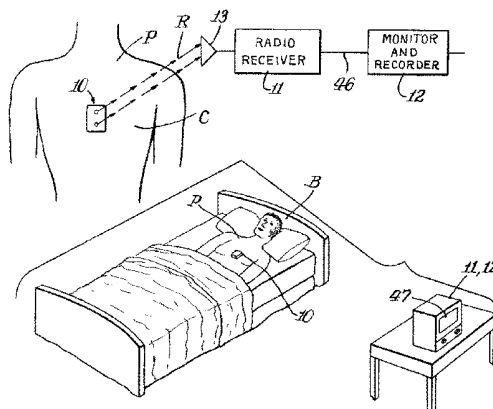
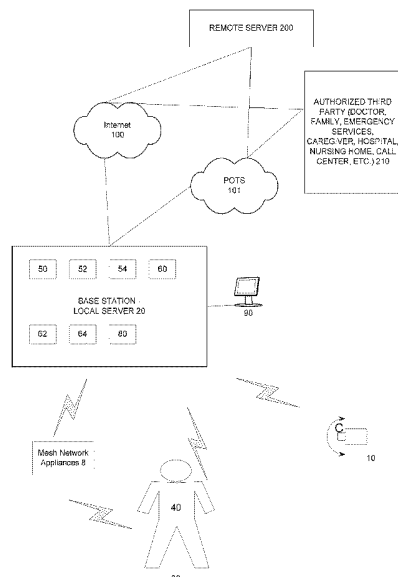
Primary Examiner — Brian T Gedeon

(74) *Attorney, Agent, or Firm* — Tran & Associates

(57) **ABSTRACT**

A heart monitoring system for a person includes one or more wireless nodes; and a wearable appliance in communication with the one or more wireless nodes, the appliance monitoring vital signs.

27 Claims, 29 Drawing Sheets



(56)

References Cited

U.S. PATENT DOCUMENTS

5,309,916	A	5/1994	Hatschek	6,230,056	B1	5/2001	Kroll
5,331,969	A	7/1994	Silberstein	6,246,894	B1	6/2001	Steuer et al.
5,361,775	A	11/1994	Remes et al.	6,280,393	B1	8/2001	Granger et al.
5,365,426	A	11/1994	Siegel et al.	6,287,252	B1	9/2001	Lugo
5,368,039	A	11/1994	Moses	6,299,583	B1	10/2001	Eggers et al.
5,370,126	A	12/1994	Clifford, Jr.	6,336,900	B1	1/2002	Alleckson et al.
5,375,610	A	12/1994	LaCourse et al.	6,364,834	B1	4/2002	Reuss et al.
5,434,419	A	7/1995	Decupper	6,364,842	B1	4/2002	Amano et al.
5,435,316	A	7/1995	Kruse	6,368,283	B1	4/2002	Xu et al.
5,438,607	A	8/1995	Przygoda, Jr. et al.	6,374,614	B2	4/2002	Prueitt
5,441,047	A	8/1995	David et al.	6,375,614	B1	4/2002	Braun et al.
5,458,620	A	10/1995	Adams et al.	6,398,727	B1	6/2002	Bui et al.
5,462,065	A	10/1995	Cusimano	6,406,426	B1	6/2002	Reuss et al.
5,497,781	A	3/1996	Chen et al.	6,413,223	B1	7/2002	Yang et al.
5,505,208	A	4/1996	Toomim et al.	6,416,480	B1	7/2002	Nenov
5,513,651	A	5/1996	Cusimano et al.	6,433,690	B2	8/2002	Petelenz et al.
5,544,649	A	8/1996	David et al.	6,440,067	B1	8/2002	DeLuca et al.
5,544,661	A	8/1996	Davis et al.	6,443,906	B1	9/2002	Ting et al.
5,546,953	A	8/1996	Garfield	6,445,299	B1	9/2002	Rojas, Jr.
5,551,435	A	9/1996	Sramek	6,466,816	B2	10/2002	Granger et al.
5,551,438	A	9/1996	Moses	6,473,640	B1	10/2002	Erlebacher
5,579,001	A	11/1996	Dempsey et al.	6,475,146	B1	11/2002	Frelburger et al.
5,579,775	A	12/1996	Dempsey et al.	6,475,153	B1	11/2002	Khair et al.
5,584,298	A	12/1996	Kabal	6,487,442	B1	11/2002	Wood
5,623,939	A	4/1997	Garfield	6,514,211	B1	2/2003	Baura
5,633,910	A	5/1997	Cohen	6,515,593	B1	2/2003	Stark et al.
5,652,570	A	7/1997	Lepkofker	6,533,729	B1	3/2003	Khair et al.
5,662,118	A	9/1997	Skubick	6,544,173	B2	4/2003	West et al.
5,671,741	A	9/1997	Lang et al.	6,544,174	B2	4/2003	West et al.
5,683,425	A	11/1997	Hauptmann	6,546,269	B1	4/2003	Kurnik
5,685,316	A	11/1997	Schookin	6,554,774	B1	4/2003	Miele
5,692,215	A	11/1997	Kutzik et al.	6,556,860	B1	4/2003	Groenewegen
5,694,940	A	12/1997	Unger et al.	6,595,316	B2	7/2003	Cybulski et al.
5,708,417	A	1/1998	Tallman et al.	6,597,944	B1	7/2003	Hadas
5,709,215	A	1/1998	Perttu et al.	6,605,038	B1	8/2003	Teller et al.
5,710,551	A	1/1998	Ridgeway	6,609,023	B1	8/2003	Fischer et al.
5,720,771	A	2/1998	Snell	6,612,984	B1	9/2003	Kerr, II
5,722,420	A	3/1998	Lee	6,615,074	B2	9/2003	Mickle et al.
5,733,230	A	3/1998	Sawchuck et al.	6,615,075	B2	9/2003	Mlynash et al.
5,738,104	A	4/1998	Lo et al.	6,616,613	B1	9/2003	Goodman
5,738,105	A	4/1998	Kroll	6,660,042	B1	12/2003	Curcie et al.
5,743,268	A	4/1998	Kabal	6,661,161	B1	12/2003	Lanzo et al.
5,772,603	A	6/1998	Ohlsson	6,676,600	B1	1/2004	Conero et al.
5,775,330	A	7/1998	Kangas et al.	6,678,549	B2	1/2004	Cusimano
5,776,073	A	7/1998	Garfield et al.	6,678,551	B2	1/2004	Maalouf et al.
5,785,960	A	7/1998	Rigg et al.	6,687,519	B2	2/2004	Steuer et al.
5,788,643	A	8/1998	Feldman	6,705,990	B1	3/2004	Gallant et al.
5,857,975	A	1/1999	Golub	6,720,984	B1	4/2004	Jorgensen et al.
5,865,755	A	2/1999	Golub	6,730,038	B2	5/2004	Gallant et al.
5,876,350	A	3/1999	Lo et al.	6,740,045	B2	5/2004	Amano
5,935,077	A	8/1999	Ogle	6,747,561	B1	6/2004	Reeves
5,945,112	A	8/1999	Flynn et al.	6,760,620	B2	7/2004	Sippens Groenewegen
5,964,719	A	10/1999	Costello et al.	6,783,492	B2	8/2004	Domiguez et al.
5,967,975	A	10/1999	Ridgeway	6,804,656	B1	10/2004	Rosenfeld et al.
5,987,519	A	11/1999	Peifer et al.	6,816,266	B2	11/2004	Varshaneya et al.
5,996,889	A	12/1999	Fuchs et al.	6,829,501	B2	12/2004	Nielsen et al.
6,002,957	A	12/1999	Finneran	6,832,251	B1	12/2004	Gelvin et al.
6,004,312	A	12/1999	Finneran et al.	6,834,436	B2	12/2004	Townsend et al.
6,010,454	A	1/2000	Arieff et al.	6,843,771	B2	1/2005	Lo et al.
6,014,626	A	1/2000	Cohen	6,856,291	B2	2/2005	Mickle et al.
6,032,035	A	2/2000	Webster et al.	6,865,409	B2	3/2005	Getsla et al.
6,038,469	A	3/2000	Karlsson et al.	6,871,211	B2	3/2005	Labounty et al.
6,047,202	A	4/2000	Finneran et al.	6,882,128	B1	4/2005	Rahmel et al.
6,050,940	A	4/2000	Braun et al.	6,887,202	B2	5/2005	Currie et al.
6,052,619	A	4/2000	John	6,898,460	B2	5/2005	Hector et al.
6,056,671	A	5/2000	Marmar	6,907,288	B2	6/2005	Daum
6,063,051	A	5/2000	Stern	6,912,420	B2	6/2005	Scheiner et al.
6,070,140	A	5/2000	Tran	6,944,496	B2	9/2005	Jeong et al.
6,076,011	A	6/2000	Hoover	6,961,285	B2	11/2005	Niemiec et al.
6,108,685	A	8/2000	Kutzik et al.	6,965,794	B2	11/2005	Brody
6,171,237	B1	1/2001	Avitall	6,973,344	B2	12/2005	Finneran et al.
6,175,308	B1	1/2001	Tallman et al.	6,974,419	B1	12/2005	Voss et al.
6,176,831	B1	1/2001	Voss et al.	6,975,206	B2	12/2005	Reining
6,184,797	B1	2/2001	Stark et al.	6,980,852	B2	12/2005	Jersey-Willuhn et al.
6,228,034	B1	5/2001	Voss et al.	6,989,751	B2	1/2006	Richards
				7,004,907	B2	2/2006	Banet et al.
				7,011,629	B2	3/2006	Bulat
				7,035,679	B2	4/2006	Addison et al.
				7,043,293	B1	5/2006	Baura

(56)

References Cited

U.S. PATENT DOCUMENTS

7,285,090	B2	10/2007	Stivoric	2005/0113630	A1	5/2005	Fox et al.
2002/0120184	A1	8/2002	Beck et al.	2005/0113710	A1	5/2005	Stahmann et al.
2003/0004421	A1	1/2003	Ting et al.	2005/0115561	A1	6/2005	Stahmann et al.
2003/0125635	A1	7/2003	Maalouf et al.	2005/0143667	A1	6/2005	Park et al.
2003/0135127	A1	7/2003	Sackner et al.	2005/0148882	A1	7/2005	Banet
2003/0216630	A1	11/2003	Jersey-Willuhn et al.	2005/0194926	A1	9/2005	Di Stefano et al.
2003/0216662	A1	11/2003	Jersey-Willuhn et al.	2005/0201585	A1	9/2005	Jannard et al.
2003/0216663	A1	11/2003	Jersey-Willuhn et al.	2005/0209521	A1	9/2005	Kettunen et al.
2003/0231115	A1	12/2003	Stanners et al.	2005/0222515	A1	10/2005	Polyshchuk et al.
2004/0044273	A1	3/2004	Keith et al.	2005/0228244	A1	10/2005	Banet et al.
2004/0102683	A1	5/2004	Khanuja et al.	2005/0228299	A1	10/2005	Banet et al.
2004/0122487	A1	6/2004	Hatlestad et al.	2005/0228316	A1	10/2005	Morgenstern et al.
2004/0138568	A1	7/2004	Lo et al.	2005/0240086	A1	10/2005	Akay et al.
2004/0167409	A1	8/2004	Lo et al.	2005/0240523	A1	10/2005	Richardson et al.
2004/0167417	A1	8/2004	Schulhauser et al.	2005/0245831	A1	11/2005	Banet et al.
2004/0220490	A1	11/2004	Appel et al.	2005/0261594	A1	11/2005	Banet et al.
2004/0254493	A1	12/2004	Chervin et al.	2005/0261598	A1	11/2005	Banet et al.
2004/0260188	A1	12/2004	Syed et al.	2005/0278409	A1	12/2005	Kutzik et al.
2004/0267148	A1	12/2004	Arand et al.	2006/0009697	A1	1/2006	Banet et al.
2005/0038345	A1	2/2005	Gorgenburg et al.	2006/0009698	A1	1/2006	Banet et al.
2005/0061568	A1	3/2005	Schondorf et al.	2006/0047215	A1	3/2006	Newman et al.
2005/0079871	A1	4/2005	Kirk	2006/0079801	A1	4/2006	DeLuca et al.
2005/0099387	A1	5/2005	Matsumoto	2006/0082727	A1	4/2006	Bolger et al.
				2006/0084878	A1	4/2006	Banet et al.
				2006/0094975	A1	5/2006	Manto
				2006/0105357	A1	5/2006	Benesch et al.

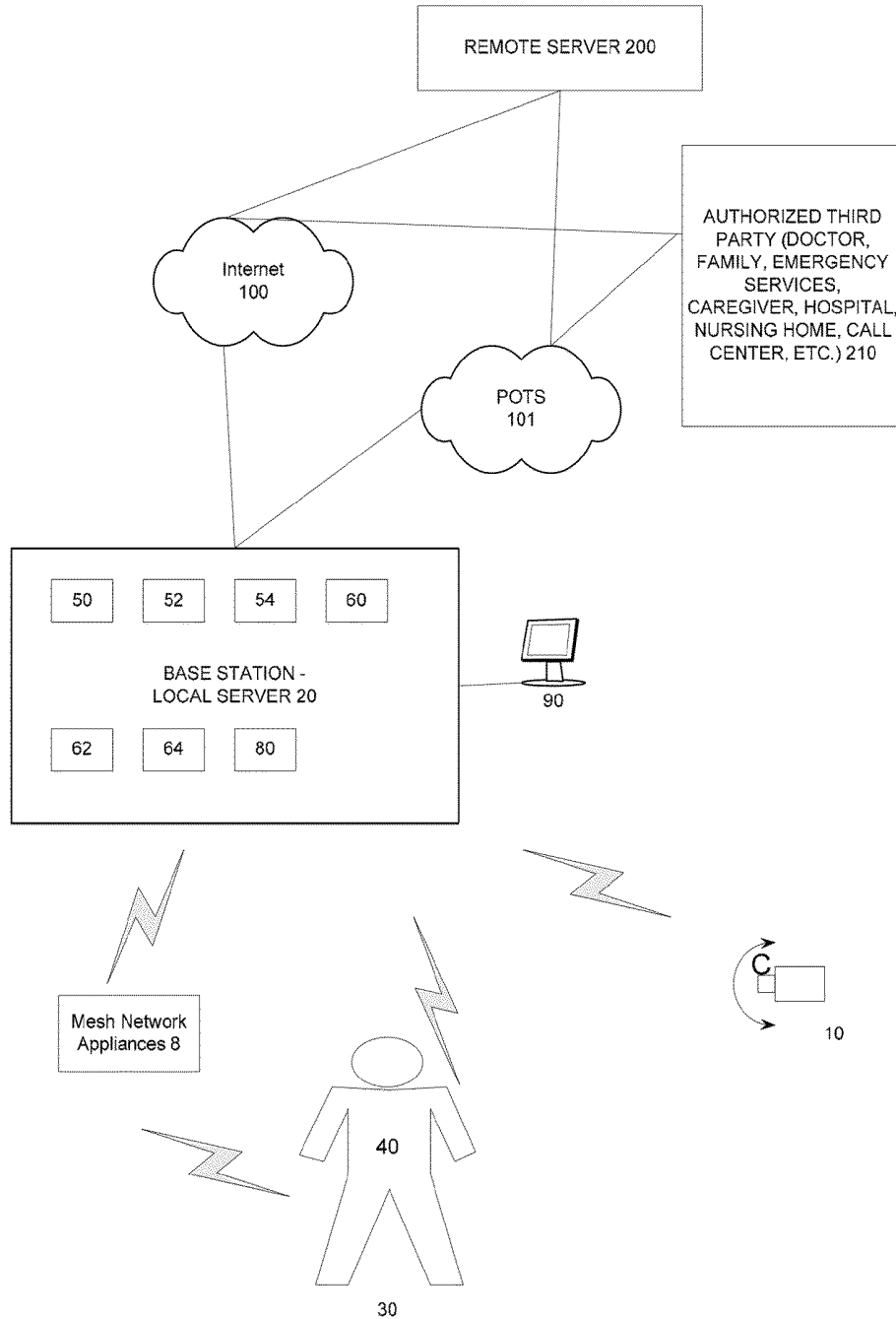


FIG. 1A

FIG. 1B

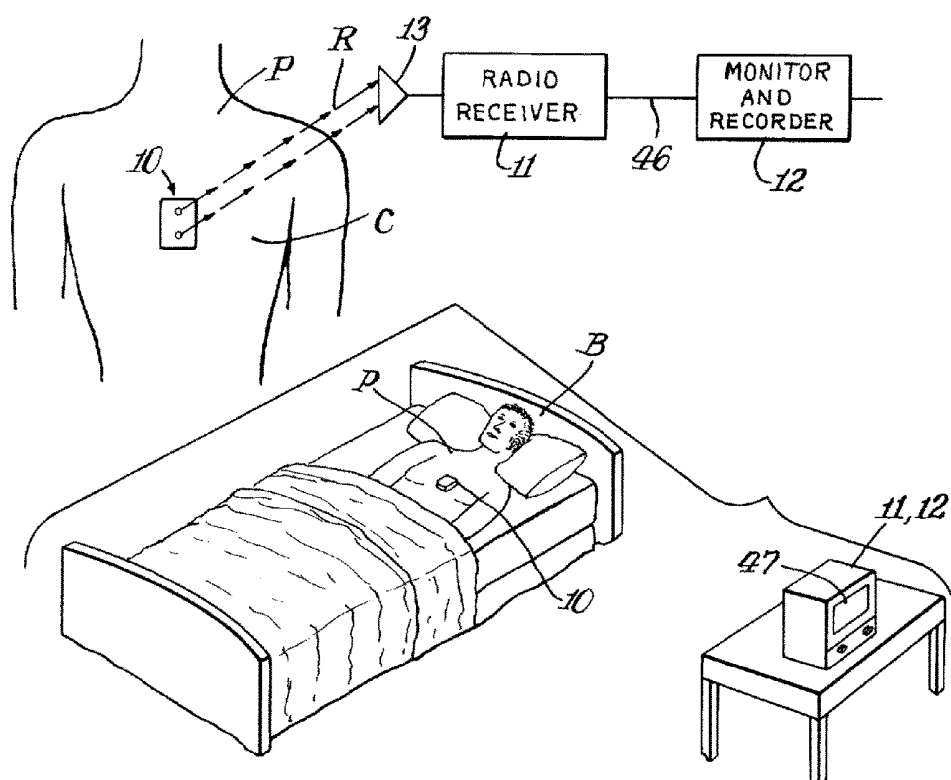


FIG. 1C

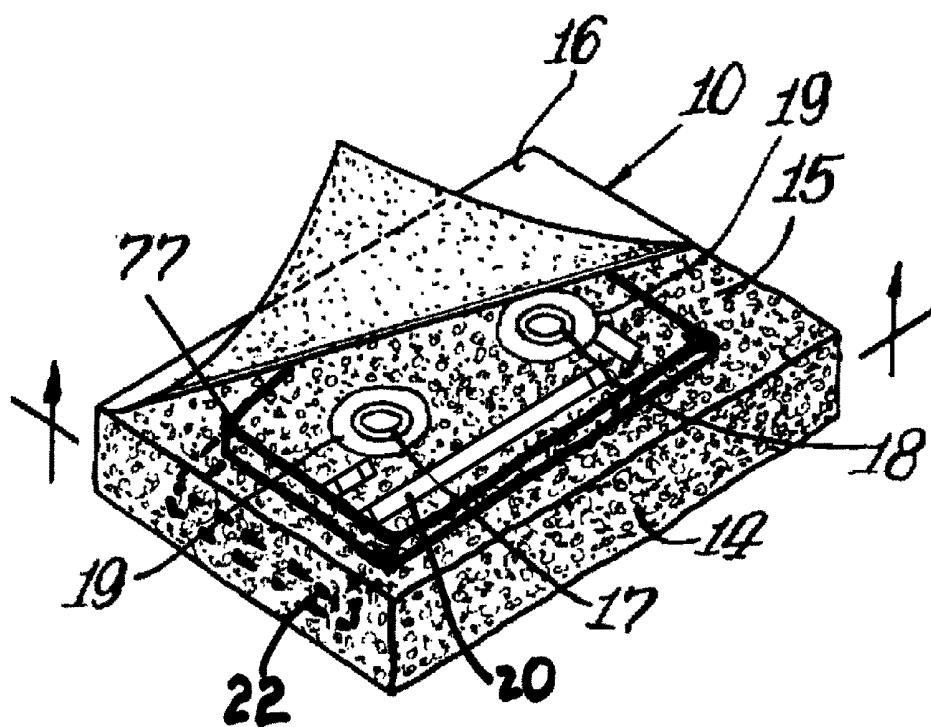


FIG. 1D

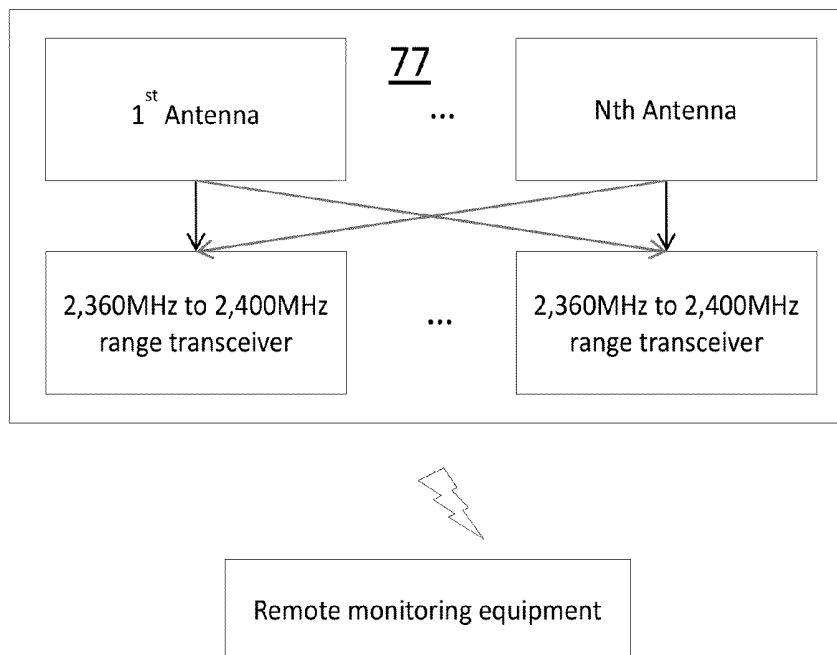


FIG. 1E

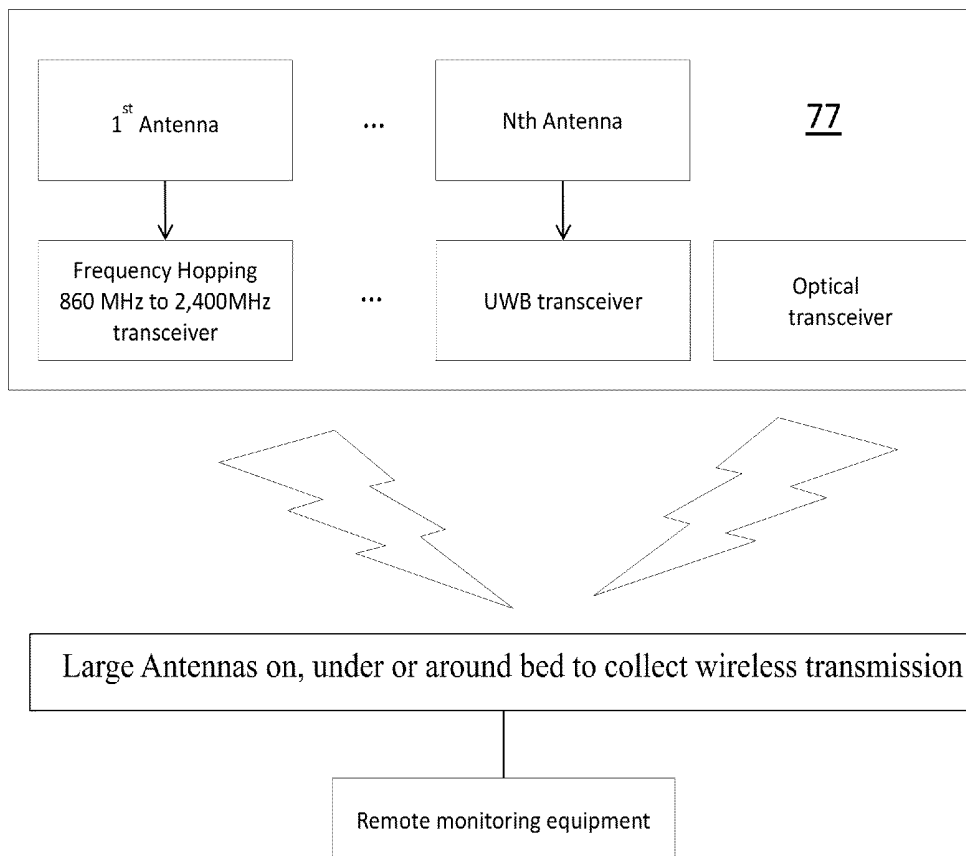


FIG. 1F

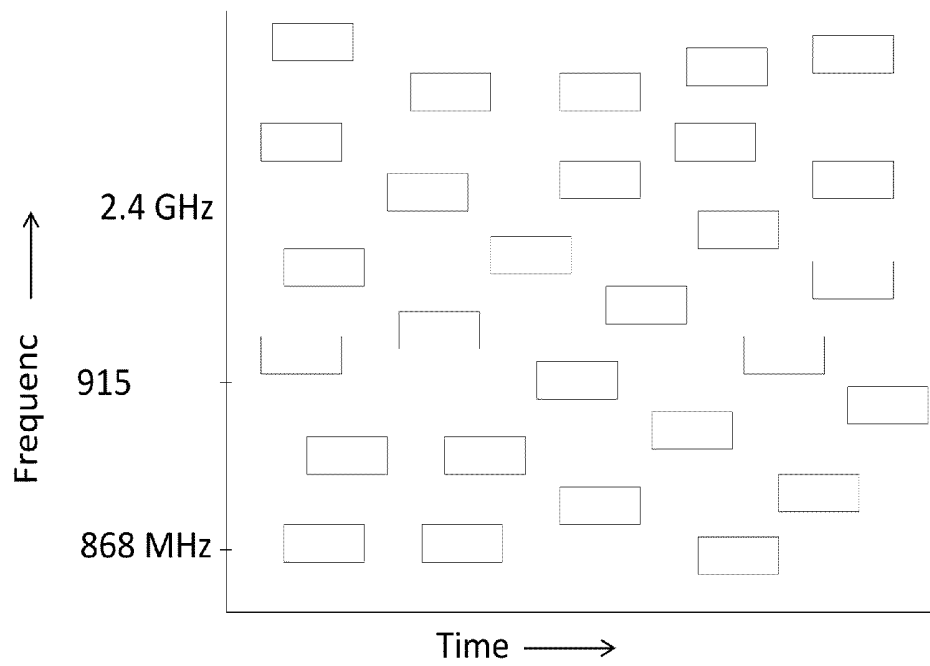


Fig. 1F

RF transceiver

FIG. 1G

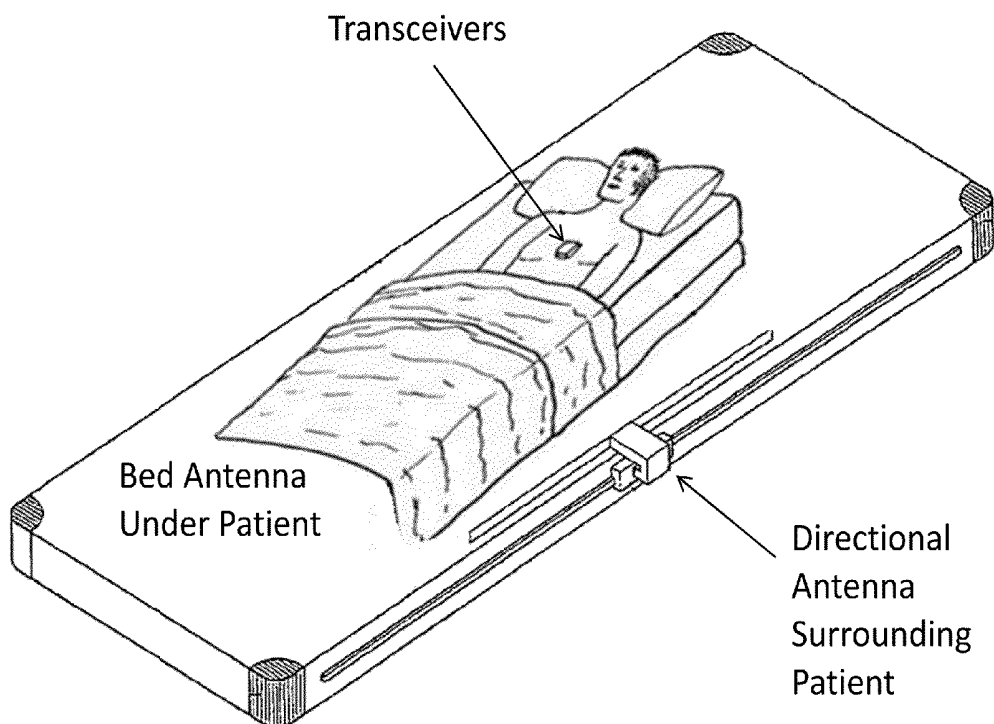


FIG. 1H

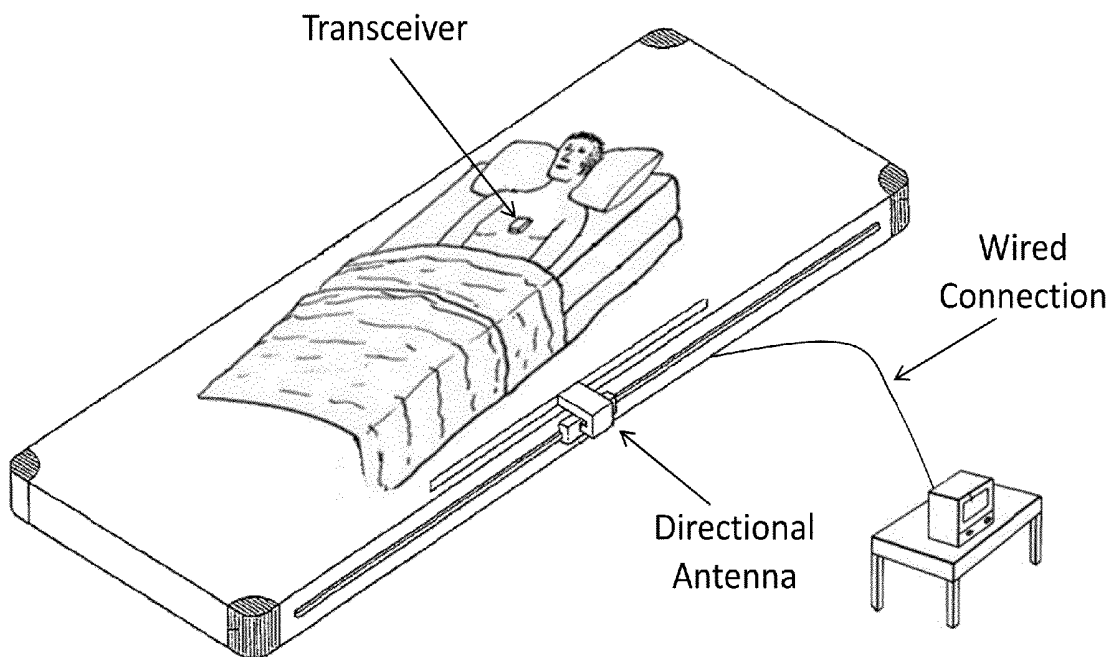


FIG. 11

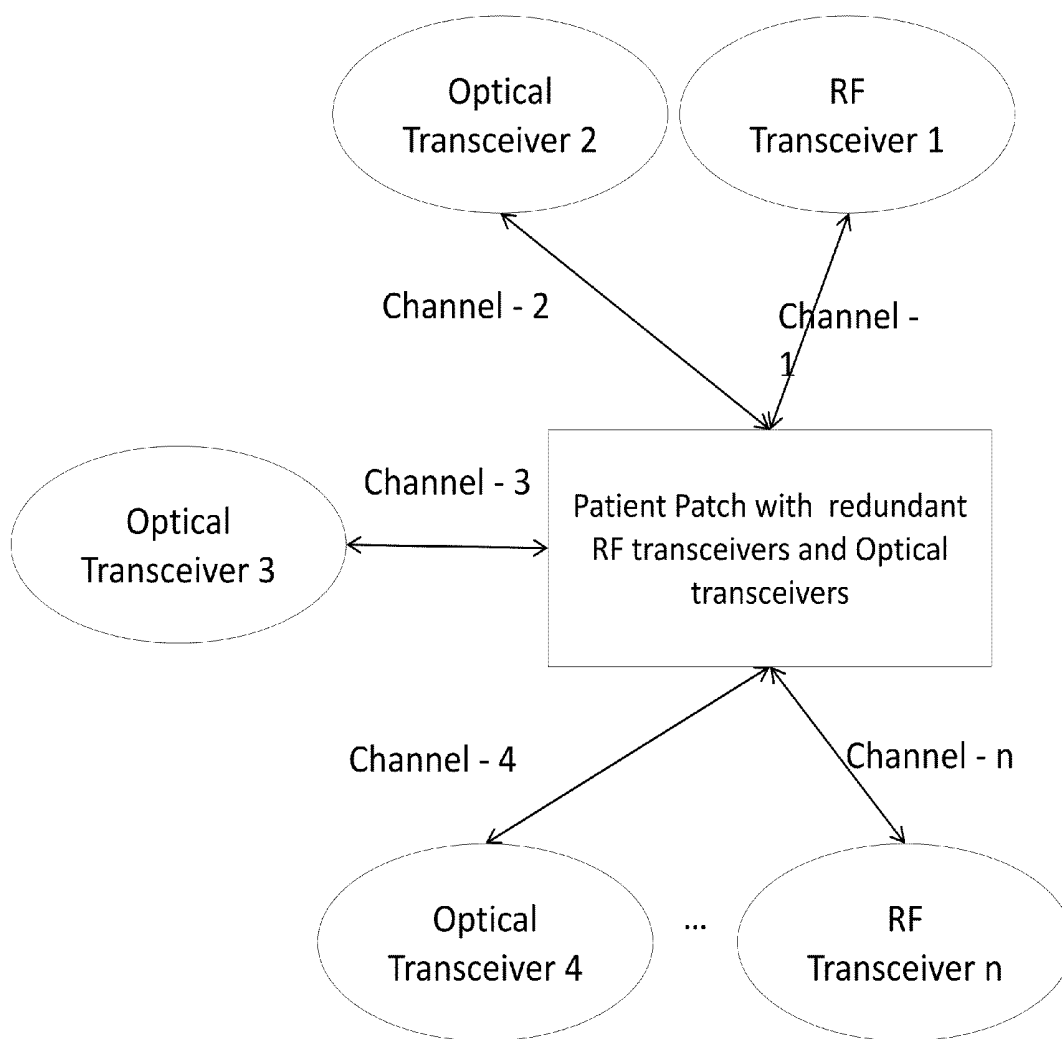


FIG. 1J

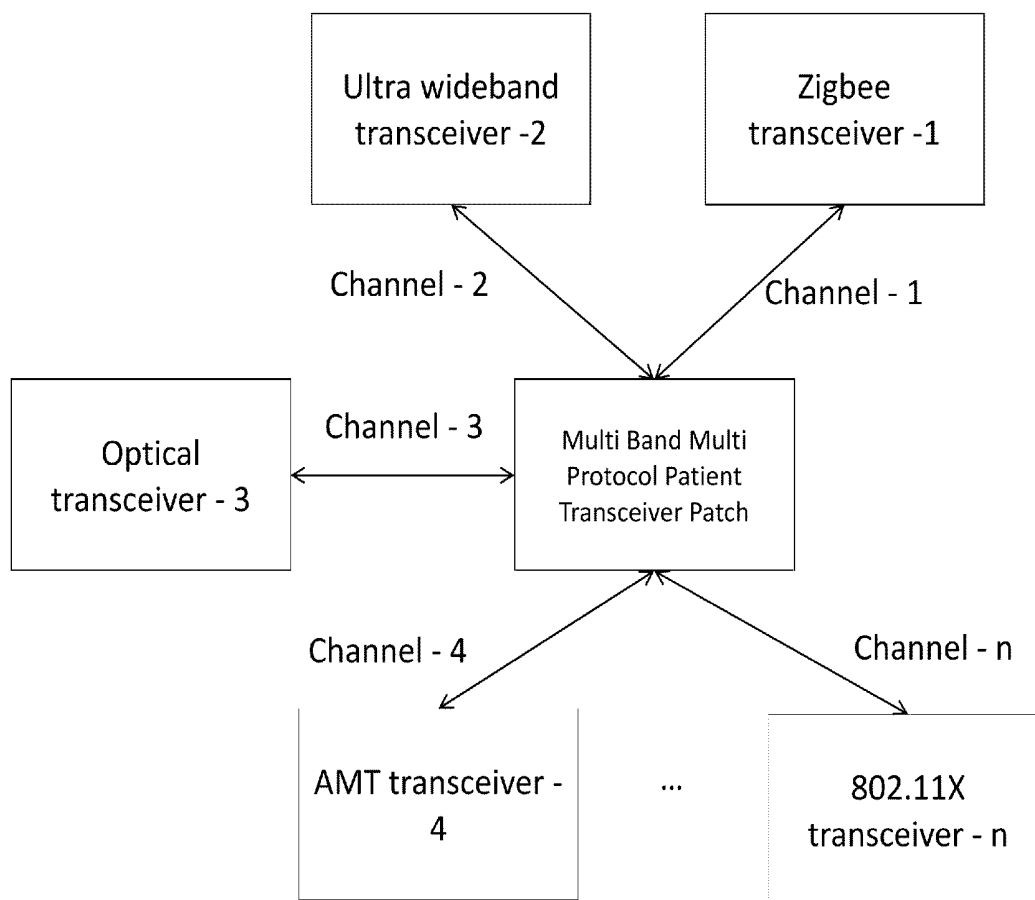


FIG. 1K

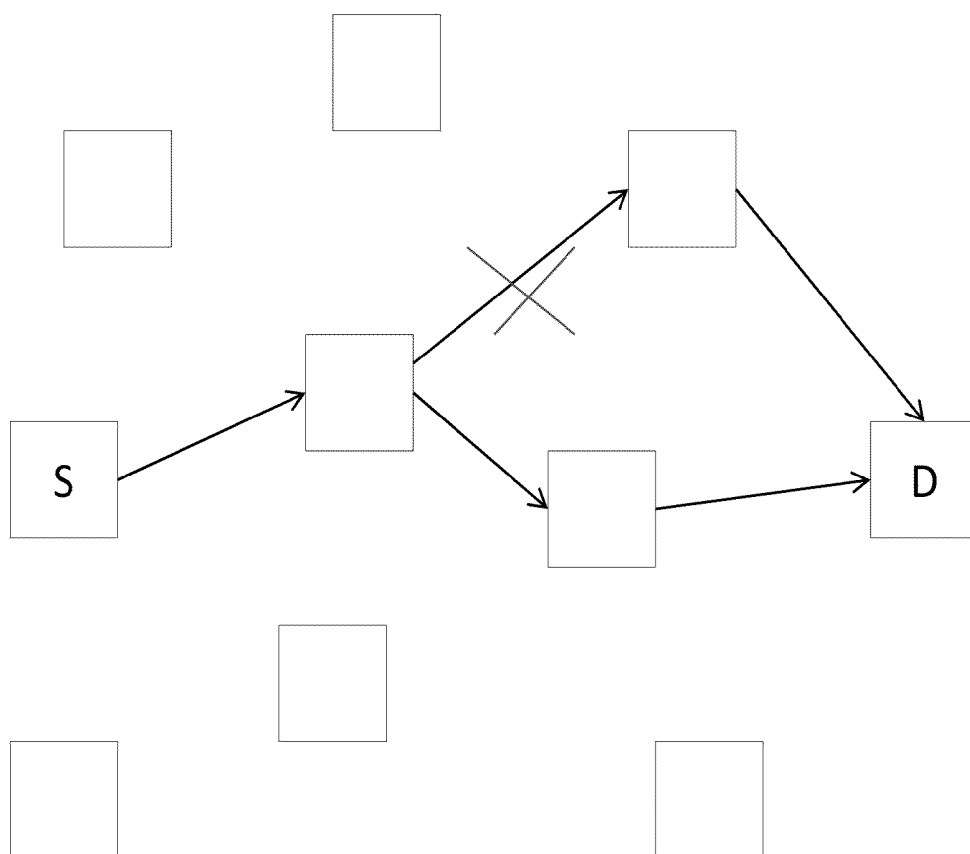
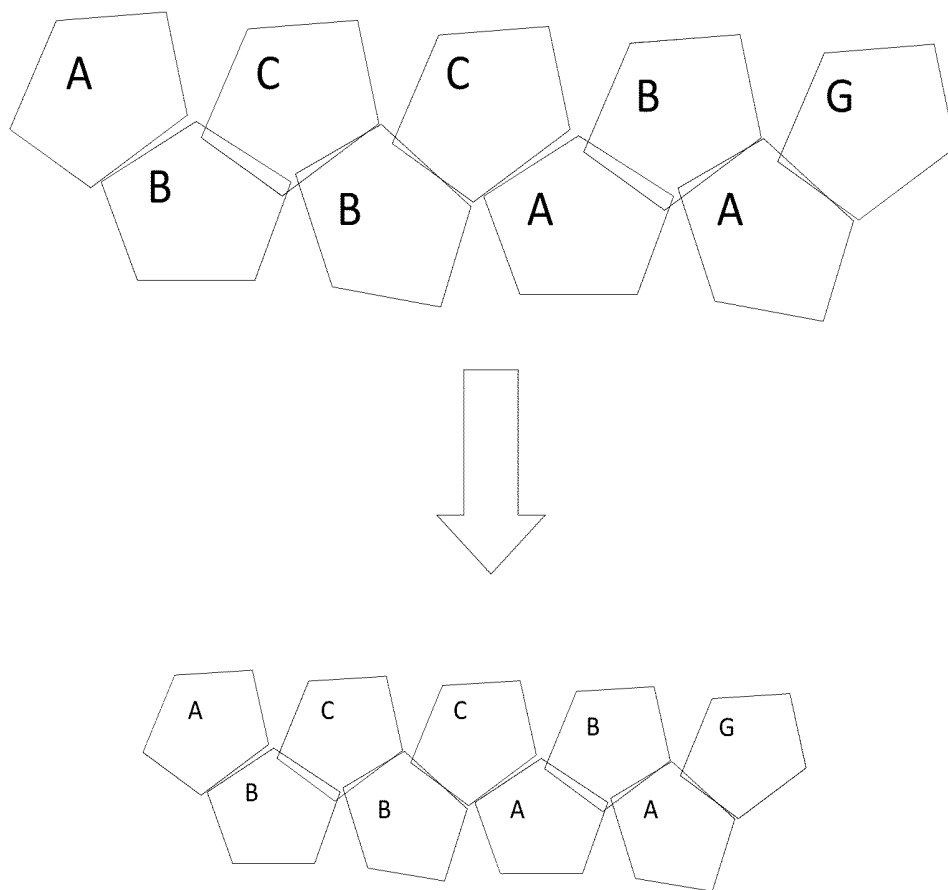


FIG. 1L



Reduced cell size

FIG. 1M

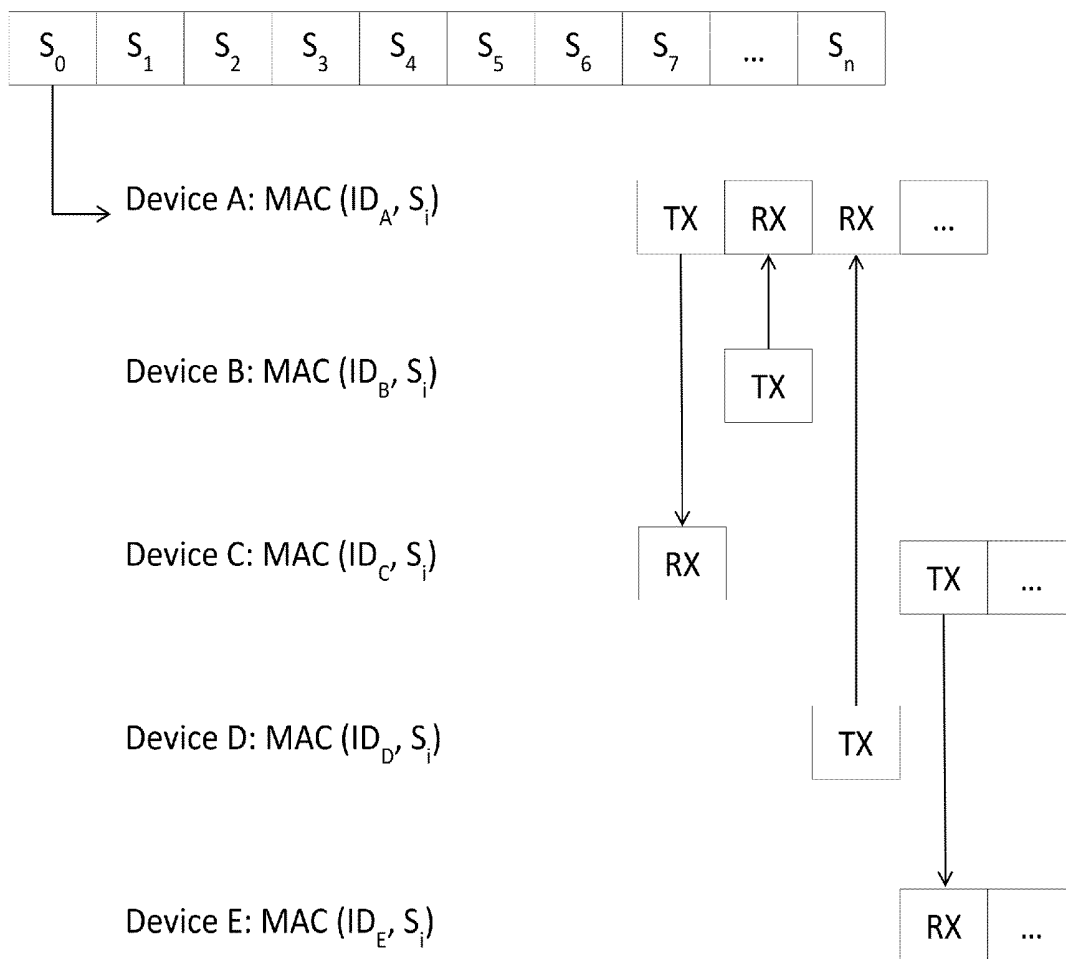
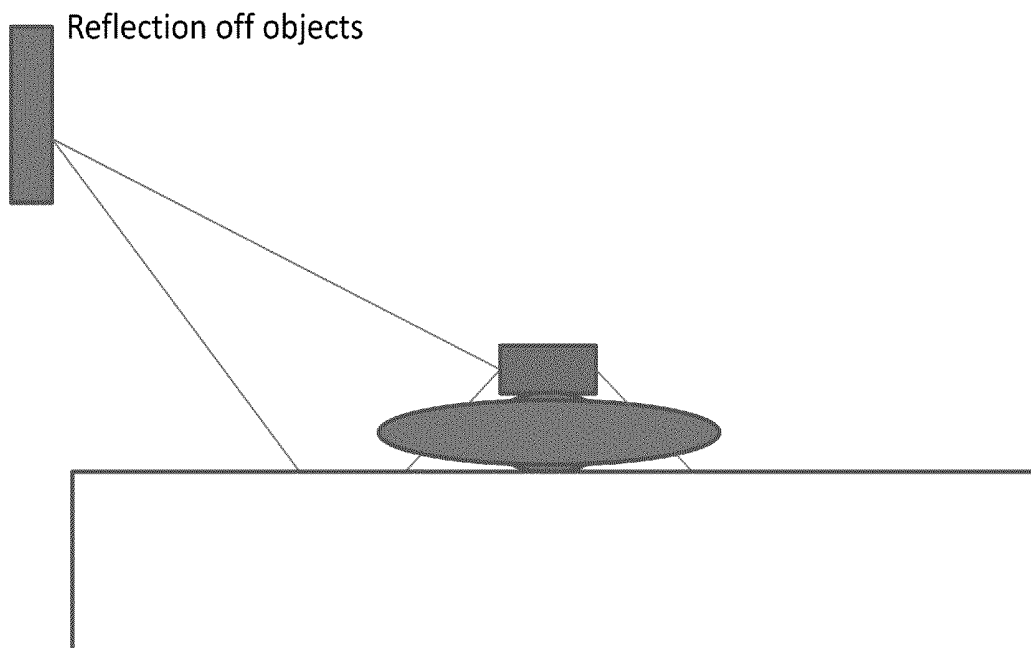


FIG. 1N



Place a calibration sheet with known dots at a known distance and perpendicular to a camera view
Take snap shot of the sheet, and correlate the position of the dots to the camera image
Place a different calibration sheet that contains known dots at another different known distance and perpendicular to camera view.
Take snap shot of the sheet and correlate the position of the dots to the camera image
Smooth the dots to the pixels to minimize digitization errors
For each pixel, draw a line from Dot1(x,y,z) to Dot2 (x, y, z) defining a cone center where the camera can view

FIG. 2A

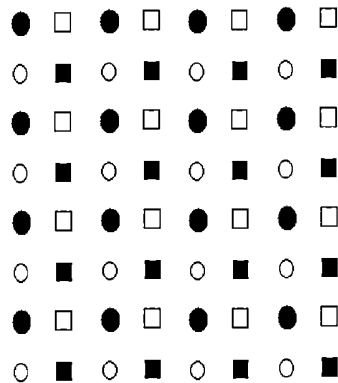


FIG. 2B

Find floor space area
Define camera view background 3D scene
Calculate patient's key features
Detect fall

FIG. 3

Find floor space area
Define camera view background 3D scene
Calculate patient's key features
Extract facial objects
Detect facial orientation
Detect facial expression

FIG. 4

Set up mesh network appliances (1000)
Determine patient position using in-door positioning system (1002)
Determine patient movement using accelerometer output (1004)
Determine vital parameter including patient heart rate (1006)
Determine if patient needs assistance based on in-door position, fall detection and vital parameter (1008)
Confirm prior to calling third party (1010)
If confirmed or non-responsive, make connection with third party and send voice over mesh network to appliance worn by the patient (1012)
If needed, call emergency personnel to get medical care (1014)

FIG. 5

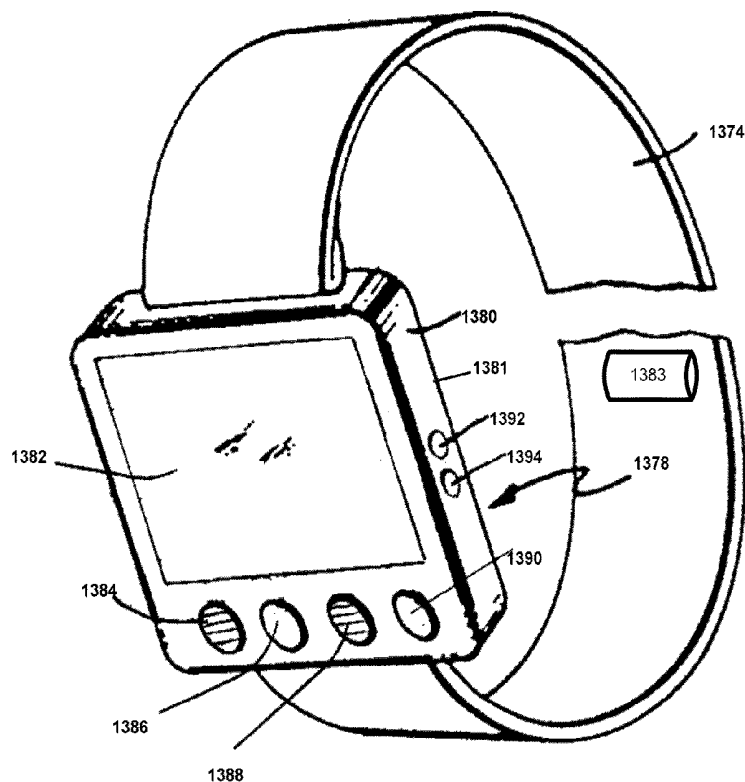


FIG. 6A

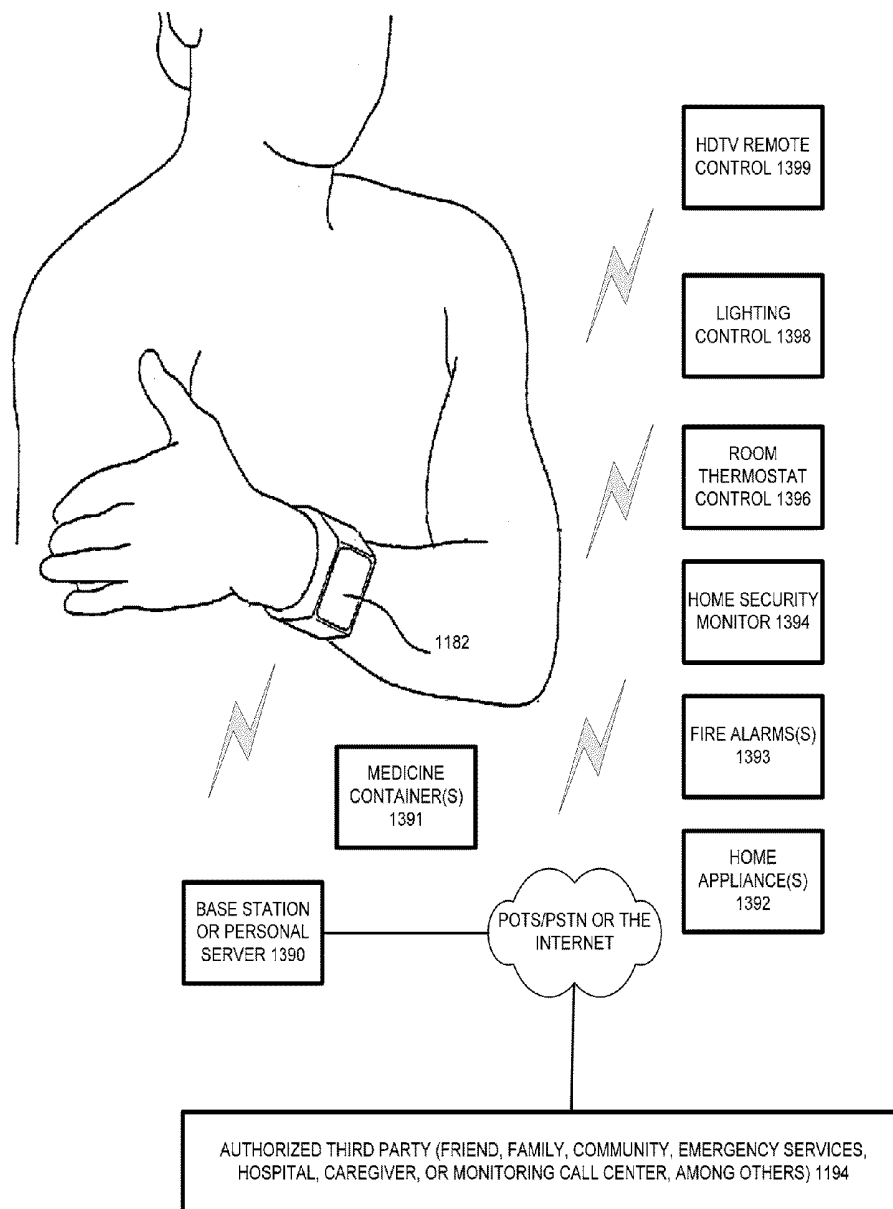


FIG. 6B

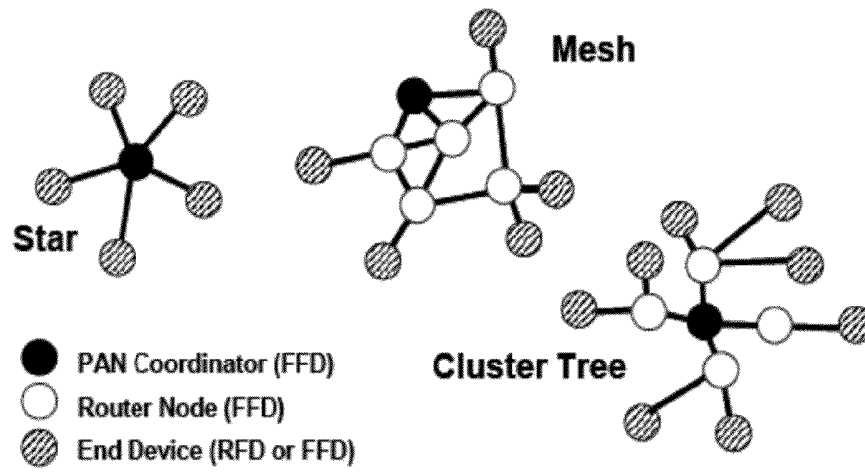


FIG. 7

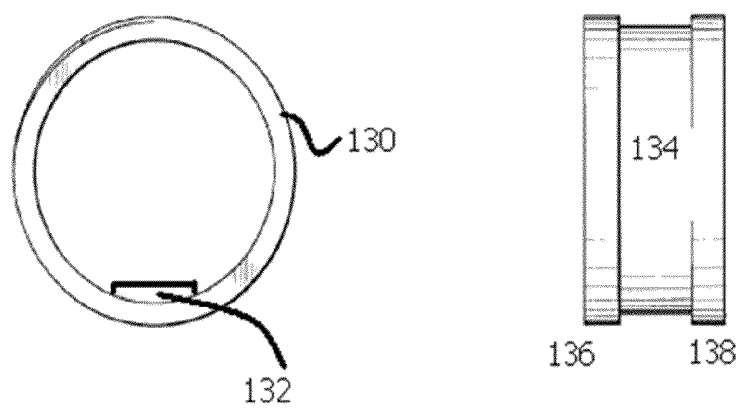


FIG. 8

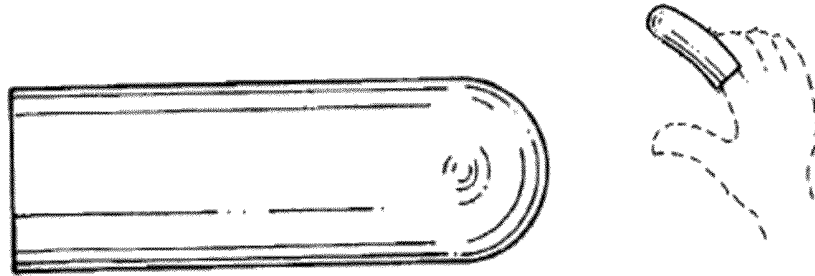


FIG. 9

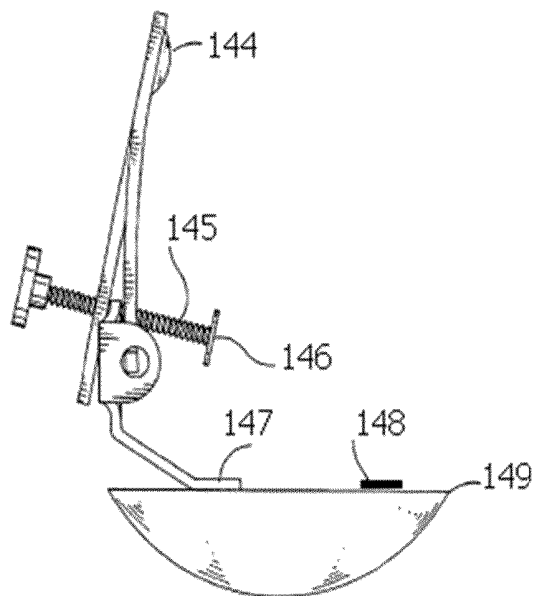


FIG. 10

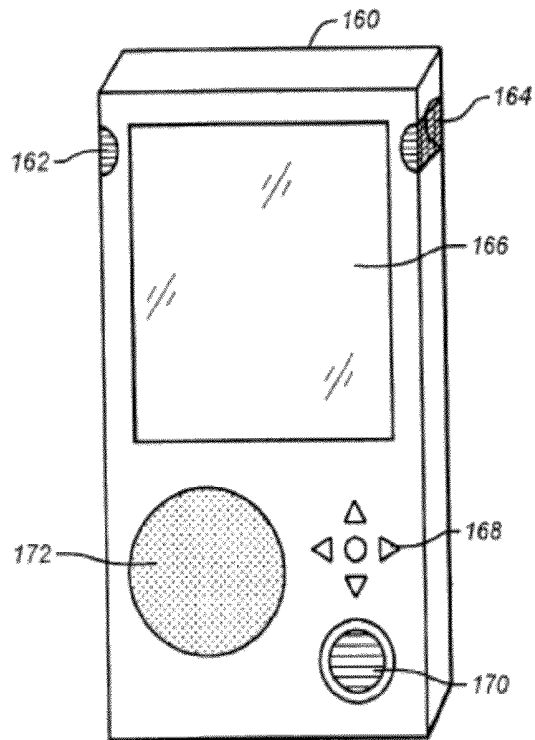


FIG. 11

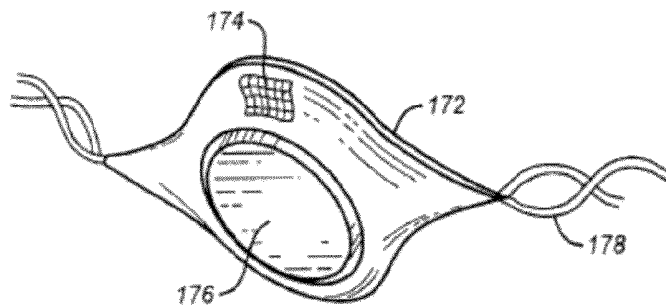


FIG. 12



FIG. 13

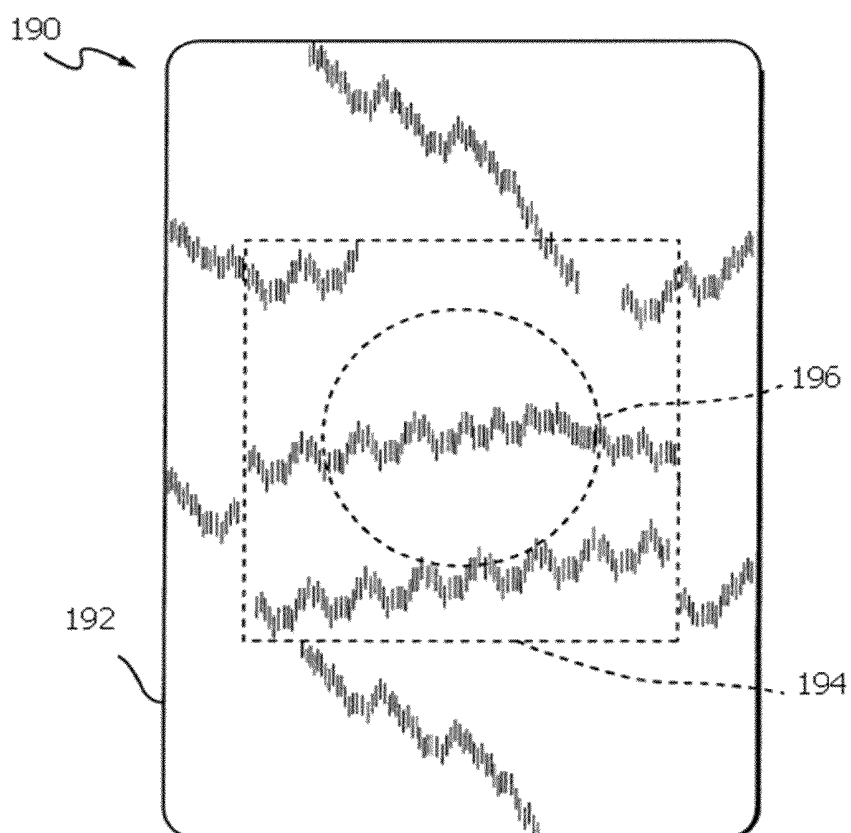


FIG. 14

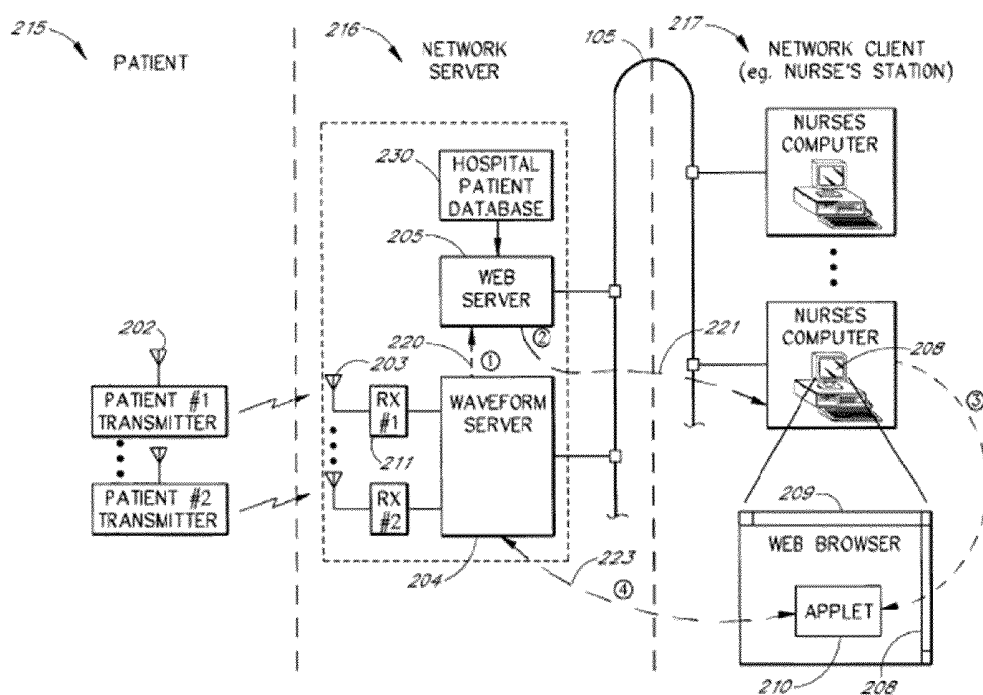


FIG. 15A

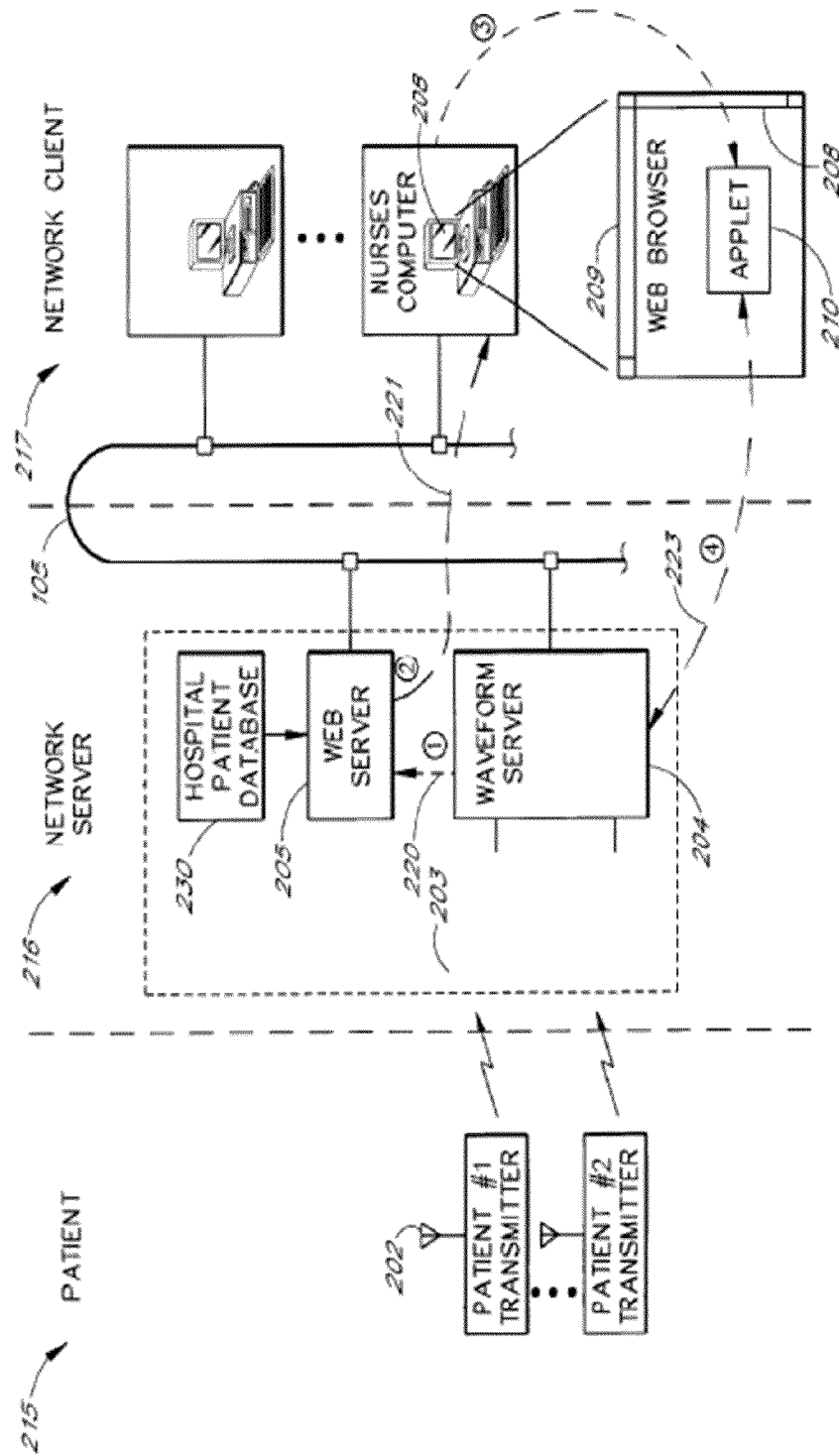


FIG. 15B

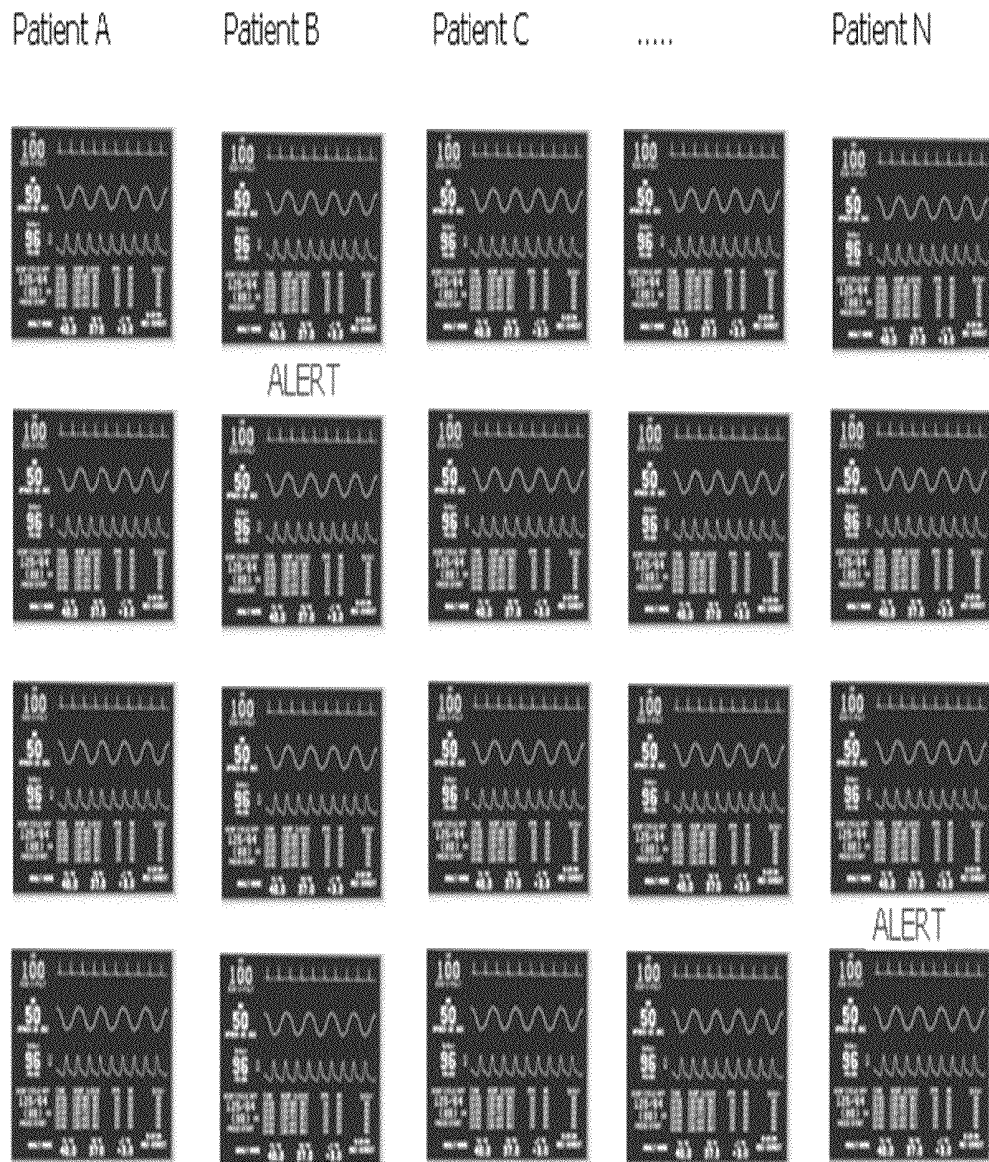
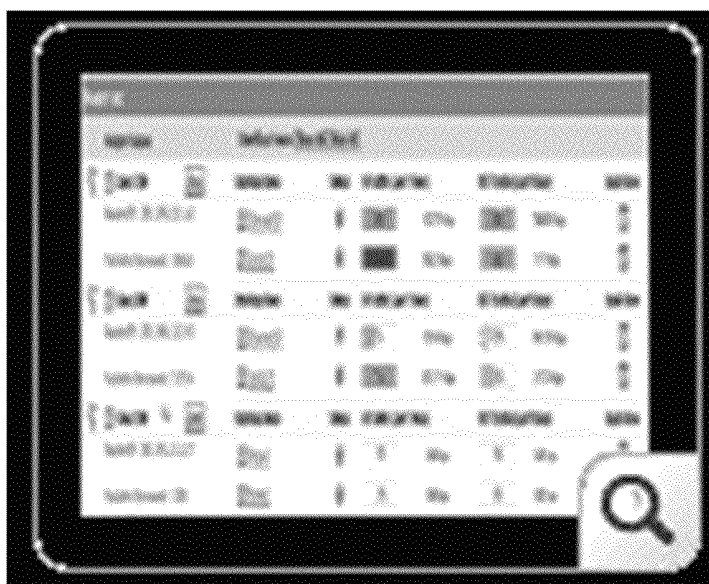


FIG. 15C



Name	Information	Value	Unit	Value	Unit
Group 1	Information	Value	Unit	Value	Unit
Item 1.1	Information	Value	Unit	Value	Unit
Item 1.2	Information	Value	Unit	Value	Unit
Group 2	Information	Value	Unit	Value	Unit
Item 2.1	Information	Value	Unit	Value	Unit
Item 2.2	Information	Value	Unit	Value	Unit
Group 3	Information	Value	Unit	Value	Unit
Item 3.1	Information	Value	Unit	Value	Unit
Item 3.2	Information	Value	Unit	Value	Unit

FIG. 15D



FIG. 15E

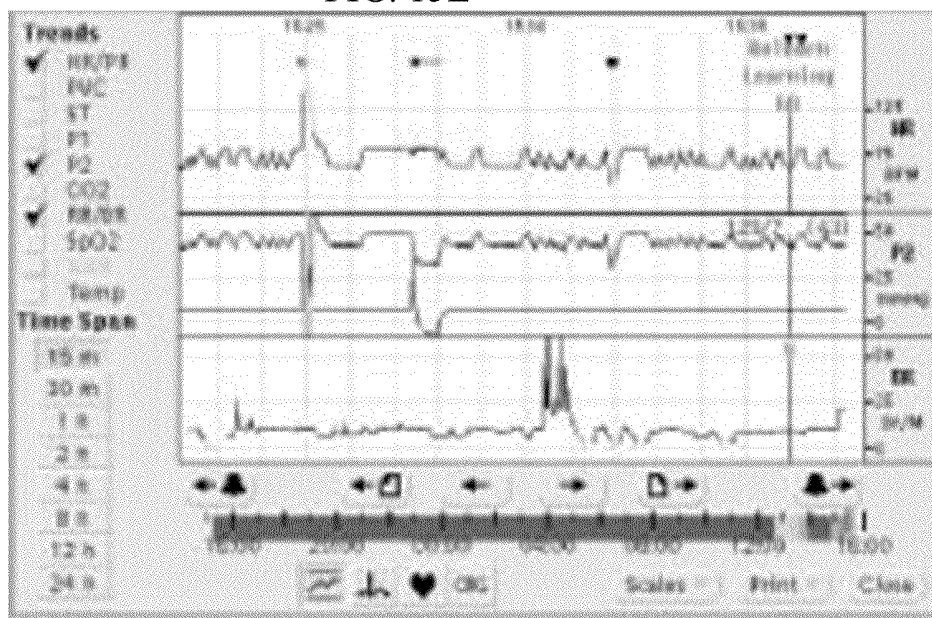


FIG. 15F

Generate a blood pressure model of a patient (2002)
Determine a blood flow velocity using a piezoelectric transducer (2004)
Provide the blood flow velocity to the blood pressure model to continuously estimate blood pressure (2006)

FIG. 16A

Attach monitoring device and calibration device to patient (2010)
Determine blood flow velocity from the monitoring device and actual blood pressure from the calibration device (2012)
Generate a blood pressure model based on the blood flow velocity and the actual blood pressure (2014)
Remove calibration device (2016)
Determine blood flow velocity (2018)
Provide blood flow velocity to the blood pressure model to estimate blood pressure (2020)

FIG. 16B

HEALTH MONITORING APPLIANCE

This application is a continuation of U.S. application Ser. No. 13/676,138, now U.S. Pat. No. 8,475,368, filed 14 Nov. 2012, which is a continuation of U.S. application Ser. No. 13/488,351, now U.S. Pat. No. 8,323,189, filed 4 Jun. 2012, which is a continuation-in-part of U.S. application Ser. No. 13/337,217, now U.S. Pat. No. 8,328,718, filed 26 Dec. 2011, which is a continuation of U.S. application Ser. No. 11/433,900, now U.S. Pat. No. 7,539,532, filed 12 May 2006 and a continuation of U.S. application Ser. No. 12/426,232, now U.S. Pat. No. 8,121,673, filed 4 Jun. 2012, the contents of which are incorporated by reference.

BACKGROUND

This invention relates generally to methods and apparatus for monitoring patients.

Conventionally, the monitoring of a patient such as in an intensive care unit of a hospital has required vigilant surveillance by one or more nurses. More recently, monitoring means have been utilized in the form of electrical monitoring and recording devices which are connected to the patient by suitable electrical wires. During surgical operations and during their stay in the ICU (Intensive Care Unit), patients are attached by cables to the monitoring or treatment equipment. Cabling, however, complicates the treatment process in many ways; for instance, cables obstruct nursing procedures and complicate transfers, as they have to be attached and detached. Such conventional devices have required that the patient's bed be made electrically shockproof. The use of such wiring limits the mobility of the patient and conventionally has required that the patient remain in the shockproof bed. Because of the high cost of the equipment and the installation, the use thereof has been primarily limited to intensive care units of hospitals and, thus, such monitoring has not been readily available in connection with less serious patient problems.

U.S. Pat. No. 3,212,496 discloses a molecular physiological monitoring system wherein the voltage produced by the heart in the functioning thereof is sensed and transmitted to an EKG receiver and recording or display device. In Preston, the transducer system is implanted subcutaneously or externally.

U.S. Pat. No. 3,943,918 discloses a throwaway, one-time use signal sensing and telemetric transmitting device for use such as in the care of medical patients requiring a monitoring of a physiological function such as the cardiac function of the patient. The device includes one-time use self-powering battery means, adhesive means for attachment of the device to the patient and electrodes for sensing the physiological functioning. A disposable cover is removed to expose the adhesive means and the battery means are actuated to power the device at the time of use. The radio frequency transmitted signal is received on a suitable radio telemetry receiver for monitoring and recording as desired.

SUMMARY

In one aspect, a monitoring system includes one or more wireless nodes communicating over an aeronautical mobile telemetry (AMT) band; and a wearable appliance in communication with the one or more wireless nodes to capture a patient vital sign.

In another aspect, a monitoring system for a person includes one or more wireless nodes; and a wearable appliance in communication with the one or more wireless nodes,

the appliance continuously monitoring vital sign. Other implementations can monitor heart rate, heart rate variability, respiratory rate, fluid status, posture and activity.

In yet another aspect, a monitoring system for a person includes one or more wireless nodes forming a wireless network and a wearable appliance having a sound transducer coupled to the wireless transceiver; and a heart disease recognizer coupled to the sound transducer to determine cardiovascular health and to transmit heart sound over the wireless network to a remote listener if the recognizer identifies a cardiovascular problem. The heart sound being transmitted may be compressed to save transmission bandwidth.

In a further aspect, a monitoring system for a person includes one or more wireless nodes; and a wristwatch having a wireless transceiver adapted to communicate with the one or more wireless nodes; and an accelerometer to detect a dangerous condition and to generate a warning when the dangerous condition is detected.

In another aspect, a monitoring system for a person includes one or more wireless nodes forming a wireless network; and a wearable appliance having a wireless transceiver adapted to communicate with the one or more wireless nodes; and a heartbeat detector coupled to the wireless transceiver. The system may also include an accelerometer to detect a dangerous condition such as a falling condition and to generate a warning when the dangerous condition is detected.

In another aspect, a monitoring system for a person includes one or more wireless bases; and a cellular telephone having a wireless transceiver adapted to communicate with the one or more wireless bases; and an accelerometer to detect a dangerous condition and to generate a warning when the dangerous condition is detected.

In yet another aspect, a monitoring system includes one or more cameras to determine a three dimensional (3D) model of a person; means to detect a dangerous condition based on the 3D model; and means to generate a warning when the dangerous condition is detected.

In another aspect, a method to detect a dangerous condition for an infant includes placing a pad with one or more sensors in the infant's diaper; collecting infant vital parameters; processing the vital parameter to detect SIDS onset; and generating a warning.

Implementations of the above aspect may include one or more of the following. The system wirelessly monitors parameters such as RR (respiratory rate), SpO₂ (oxygen saturation), ECG (electrocardiogram), HR (heart rate), core temperature (inside the heart) and peripheral temperature (on top of the instep), CI (cardiac output index, CO/m²), systematic pressures (SSAP, systematic systolic arterial pressure; SDAP, systematic diastolic arterial pressure; SMAP, systematic mean arterial pressure), CVP (central venous pressure), pulmonary pressures (PSAP, pulmonary systolic arterial pressure; PDAP, pulmonary diastolic arterial pressure; PMAP, pulmonary mean arterial pressure), svO₂ (oxygen saturation in the lung artery), ET/CO₂ (outcoming carbon dioxide), FIO₂ (ingoing oxygen), diuretics, the patient's weight, fluid balance (ingoing and outcoming fluids) and EEG. The device can be a wristwatch that determines position based on triangulation. The wristwatch determines position based on RF signal strength and RF signal angle. A switch detects a confirmatory signal from the person. The confirmatory signal includes a head movement, a hand movement, or a mouth movement. The confirmatory signal includes the person's voice. A processor in the system executes computer readable code to transmit a help request to a remote computer. The code can encrypt or scramble data for privacy. The processor can execute voice over IP (VOIP) code to allow a user and a

remote person to audibly communicate with each other. The voice communication system can include Zigbee VOIP or Bluetooth VOIP or 802.XX VOIP. The remote person can be a doctor, a nurse, a medical assistant, or a caregiver. The system includes code to store and analyze patient information. The patient information includes medicine taking habits, eating and drinking habits, sleeping habits, or exercise habits. A patient interface is provided on a user computer for accessing information and the patient interface includes in one implementation a touch screen; voice-activated text reading; and one touch telephone dialing. The processor can execute code to store and analyze information relating to the person's ambulation. A global positioning system (GPS) receiver can be used to detect movement and where the person falls. The system can include code to map the person's location onto an area for viewing. The system can include one or more cameras positioned to capture three dimensional (3D) video of the patient; and a server coupled to the one or more cameras, the server executing code to detect a dangerous condition for the patient based on the 3D video and allow a remote third party to view images of the patient when the dangerous condition is detected.

Advantages of the invention may include one or more of the following. The system eliminates cables attaching patient monitoring sensors to monitoring devices without disturbing and obstructing nursing staff. Vital parameters, such as blood pressure, electrocardiography (ECG), respiration rate, heart rate and temperature, from patient who is in critical condition are measured all the time. Wireless connections enable free redeployment of patient between separate units and allow free movement of the patient. Wireless connections will offer quick and easy way to redeploy the patient, e.g., from an operating room to an intensive care unit (ICU) and make the movement of patient more easier.

The system is highly reliable and is at least as reliable as wired techniques. Interference is reduced by operating at a different frequency than Bluetooth and 2.45 GHz WLAN. The system avoids interference from surgical knives, mobile phones and even microwave ovens through error correction and redundant transmission within a hospital. The absence of cables around patients will improve the working conditions of nursing staff, and enhance their efficiency. When patients arrive at the hospital, sensors will be attached to them and removed as they check out. All patient information, including personal data, laboratory test results, patient monitor signals etc., will be transferred to one single database. During their stay, patients remain connected to the hospital network and can either move freely or be transferred anywhere in the hospital. Relevant nursing staff will be able to examine patient information via a PDA, for example, anywhere in the hospital. Additional information can be sent outside the hospital to a consulting doctor through a mobile phone, PDA or via e-mail.

The device can be applied in the manner of a conventional bandage to the patient's body without complicated or extensive preparation of the patient. As the device is extremely simple and economical of construction, it may be utilized as a one-time use, throwaway device which permits high mobility of the patient while yet providing continuous monitoring of the sensed physiological function. The system for non-invasively and continually monitors a subject's arterial blood pressure, with reduced susceptibility to noise and subject movement, and relative insensitivity to placement of the apparatus on the subject. The system does not need frequent recalibration of the system while in use on the subject.

In particular, it allows patients to conduct a low-cost, comprehensive, real-time monitoring of their blood pressure.

Using the web services software interface, the invention then avails this information to hospitals, home-health care organizations, insurance companies, pharmaceutical agencies conducting clinical trials and other organizations. Information can be viewed using an Internet-based website, a personal computer, or simply by viewing a display on the monitor. Data measured several times each day provide a relatively comprehensive data set compared to that measured during medical appointments separated by several weeks or even months. This allows both the patient and medical professional to observe trends in the data, such as a gradual increase or decrease in blood pressure, which may indicate a medical condition. The invention also minimizes effects of white coat syndrome since the monitor automatically makes measurements with basically no discomfort; measurements are made at the patient's home or work, rather than in a medical office.

The wearable appliance is small, easily worn by the patient during periods of exercise or day-to-day activities, and non-invasively measures blood pressure can be done in a matter of seconds without affecting the patient. An on-board or remote processor can analyze the time-dependent measurements to generate statistics on a patient's blood pressure (e.g., average pressures, standard deviation, beat-to-beat pressure variations) that are not available with conventional devices that only measure systolic and diastolic blood pressure at isolated times.

The wearable appliance provides an in-depth, cost-effective mechanism to evaluate a patient's cardiac condition. Certain cardiac conditions can be controlled, and in some cases predicted, before they actually occur. Moreover, data from the patient can be collected and analyzed while the patient participates in their normal, day-to-day activities.

In cases where the device has fall detection in addition to blood pressure measurement, other advantages of the invention may include one or more of the following. The system provides timely assistance and enables elderly and disabled individuals to live relatively independent lives. The system monitors physical activity patterns, detects the occurrence of falls, and recognizes body motion patterns leading to falls. Continuous monitoring of patients is done in an accurate, convenient, unobtrusive, private and socially acceptable manner since a computer monitors the images and human involvement is allowed only under pre-designated events. The patient's privacy is preserved since human access to videos of the patient is restricted: the system only allows human viewing under emergency or other highly controlled conditions designated in advance by the user. When the patient is healthy, people cannot view the patient's video without the patient's consent. Only when the patient's safety is threatened would the system provide patient information to authorized medical providers to assist the patient. When an emergency occurs, images of the patient and related medical data can be compiled and sent to paramedics or hospital for proper preparation for pick up and check into emergency room.

The system allows certain designated people such as a family member, a friend, or a neighbor to informally check on the well-being of the patient. The system is also effective in containing the spiraling cost of healthcare and outpatient care as a treatment modality by providing remote diagnostic capability so that a remote healthcare provider (such as a doctor, nurse, therapist or caregiver) can visually communicate with the patient in performing remote diagnosis. The system allows skilled doctors, nurses, physical therapists, and other scarce resources to assist patients in a highly efficient manner since they can do the majority of their functions remotely.

Additionally, a sudden change of activity (or inactivity) can indicate a problem. The remote healthcare provider may

receive alerts over the Internet or urgent notifications over the phone in case of such sudden accident indicating changes. Reports of health/activity indicators and the overall well-being of the individual can be compiled for the remote healthcare provider. Feedback reports can be sent to monitored subjects, their designated informal caregiver and their remote healthcare provider. Feedback to the individual can encourage the individual to remain active. The content of the report may be tailored to the target recipient's needs, and can present the information in a format understandable by an elder person unfamiliar with computers, via an appealing patient interface. The remote healthcare provider will have access to the health and well-being status of their patients without being intrusive, having to call or visit to get such information interrogatively. Additionally, remote healthcare provider can receive a report on the health of the monitored subjects that will help them evaluate these individuals better during the short routine check-up visits. For example, the system can perform patient behavior analysis such as eating/drinking/smoke habits and medication compliance, among others.

The patient's home equipment is simple to use and modular to allow for the accommodation of the monitoring device to the specific needs of each patient. Moreover, the system is simple to install. Regular monitoring of the basic wellness parameters provides significant benefits in helping to capture adverse events sooner, reduce hospital admissions, and improve the effectiveness of medications, hence, lowering patient care costs and improving the overall quality of care. Suitable users for such systems are disease management companies, health insurance companies, self-insured employers, medical device manufacturers and pharmaceutical firms.

The system reduces costs by automating data collection and compliance monitoring, and hence reduce the cost of nurses for hospital and nursing home applications. At-home vital signs monitoring enables reduced hospital admissions and lower emergency room visits of chronic patients. Operators in the call centers or emergency response units get high quality information to identify patients that need urgent care so that they can be treated quickly, safely, and cost effectively. The Web based tools allow easy access to patient information for authorized parties such as family members, neighbors, physicians, nurses, pharmacists, caregivers, and other affiliated parties to improve the Quality of Care for the patient.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1A illustrates an exemplary system for monitoring a person.

FIG. 1B a schematic representation of a patient provided with a signal sensing and telemetric transmitting device embodying the invention in connection with the transmitting of physiologic signals to a suitable receiver, monitor, and recorder.

FIG. 1C is a perspective view of a monitoring patch, pad or band-aid ready to be worn by a patient for transmitting vital signs.

FIG. 1D shows an exemplary module with redundant transceivers and redundant antennas.

FIG. 1E shows another exemplary module with redundant transceivers and redundant antennas.

FIG. 1F shows a multi-band frequency hopping embodiment.

FIG. 1G shows a wideband bed antenna embodiment.

FIG. 1H shows a bed receiver with large antenna where the receiver is wired to hospital monitoring equipment.

FIG. 1I shows a multi-band system that transmits over RF and optical bands.

FIG. 1J shows a multi-band system that transmits over ultra-wideband, Zigbee, WiFi and optical bands.

FIG. 1K shows mesh networking that transmits over various wireless and optical protocols in a node to node basis to avoid a jammed node for enhanced reliability.

FIG. 1L shows exemplary low power cells for patient telemetry.

FIG. 1M shows an embodiment with scheduled transmissions.

FIG. 1N shows an exemplary high gain parabolic bed antenna.

FIG. 2A illustrates a process for determining three dimensional (3D) detection.

FIG. 2B shows an exemplary calibration sheet.

FIG. 3 illustrates a process for detecting falls.

FIG. 4 illustrates a process for detecting facial expressions.

FIG. 5 illustrates an exemplary process for determining and getting assistance for a patient or user.

FIG. 6A shows an exemplary wrist-watch based assistance device.

FIG. 6B shows an exemplary mesh network working with the wearable appliance of FIG. 6A.

FIG. 7 shows an exemplary mesh network in communication with the wrist-watch device of FIG. 6.

FIGS. 8-14 show various exemplary wearable appliances to monitor a patient.

FIGS. 15A-15B show exemplary systems for performing patient monitoring.

FIG. 15C shows an exemplary interface to monitor a plurality of persons.

FIG. 15D shows an exemplary dash-board that provides summary information on the status of a plurality of persons.

FIG. 15E shows an exemplary multi-station vital parameter user interface for a professional embodiment.

FIG. 15F shows an exemplary trending pattern display.

FIGS. 16A-16B show exemplary blood pressure determination processes.

DESCRIPTION

FIG. 1A shows an exemplary patient monitoring system. The system can operate in a home, a nursing home, or a hospital. In this system, one or more mesh network appliances 8 are provided to enable wireless communication in the home monitoring system. Appliances 8 in the mesh network can include home security monitoring devices, door alarm, window alarm, home temperature control devices, fire alarm devices, among others. Appliances 8 in the mesh network can be one of multiple portable physiological transducer, such as a blood pressure monitor, heart rate monitor, weight scale, thermometer, spirometer, single or multiple lead electrocardiograph (ECG), a pulse oximeter, a body fat monitor, a cholesterol monitor, a signal from a medicine cabinet, a signal from a drug container, a signal from a commonly used appliance such as a refrigerator/stove/oven/washer, or a signal from an exercise machine, such as a heart rate. As will be discussed in more detail below, one appliance is a patient monitoring device that can be worn by the patient and includes a single or bi-directional wireless communication link, generally identified by the bolt symbol in FIG. 1, for transmitting data from the appliances 8 to the local hub or receiving station or base station server 20 by way of a wireless radio frequency (RF) link using a proprietary or non-proprietary protocol. For example, within a house, a user may have mesh network appliances that detect window and door contacts, smoke detectors and motion sensors, video cameras, key chain control, temperature monitors, CO and other gas

detectors, vibration sensors, and others. A user may have flood sensors and other detectors on a boat. An individual, such as an ill or elderly grandparent, may have access to a panic transmitter or other alarm transmitter. Other sensors and/or detectors may also be included. The user may register these appliances on a central security network by entering the identification code for each registered appliance/device and/or system. The mesh network can be Zigbee network or 802.15 network. More details of the mesh network is shown in FIG. 7 and discussed in more detail below.

A plurality of monitoring cameras **10** may be placed in various predetermined positions in a home of a patient **30**. The cameras **10** can be wired or wireless. For example, the cameras can communicate over infrared links or over radio links conforming to the 802X (e.g. 802.11A, 802.11B, 802.11G, 802.15) standard or the Bluetooth standard to a base station/server **20** may communicate over various communication links, such as a direct connection, such a serial connection, USB connection, Firewire connection or may be optically based, such as infrared or wireless based, for example, home RF, IEEE standard 802.11a/b, Bluetooth or the like. In one embodiment, appliances **8** monitor the patient and activates the camera **10** to capture and transmit video to an authorized third party for providing assistance should the appliance **8** detects that the user needs assistance or that an emergency had occurred.

The base station/server **20** stores the patient's ambulation pattern and vital parameters and can be accessed by the patient's family members (sons/daughters), physicians, caretakers, nurses, hospitals, and elderly community. The base station/server **20** may communicate with the remote server **200** by DSL, T-1 connection over a private communication network or a public information network, such as the Internet **100**, among others.

The patient **30** may wear one or more wearable patient monitoring appliances such as wrist-watches or clip on devices or electronic jewelry to monitor the patient. One wearable appliance such as a wrist-watch includes sensors **40**, for example devices for sensing ECG, EKG, blood pressure, sugar level, among others. In one embodiment, the sensors **40** are mounted on the patient's wrist (such as a wristwatch sensor) and other convenient anatomical locations. Exemplary sensors **40** include standard medical diagnostics for detecting the body's electrical signals emanating from muscles (EMG and EOG) and brain (EEG) and cardiovascular system (ECG). Leg sensors can include piezoelectric accelerometers designed to give qualitative assessment of limb movement. Additionally, thoracic and abdominal bands used to measure expansion and contraction of the thorax and abdomen respectively. A small sensor can be mounted on the subject's finger in order to detect blood-oxygen levels and pulse rate. Additionally, a microphone can be attached to throat and used in sleep diagnostic recordings for detecting breathing and other noise. One or more position sensors can be used for detecting orientation of body (lying on left side, right side or back) during sleep diagnostic recordings. Each of sensors **40** can individually transmit data to the server **20** using wired or wireless transmission. Alternatively, all sensors **40** can be fed through a common bus into a single transceiver for wired or wireless transmission. The transmission can be done using a magnetic medium such as a floppy disk or a flash memory card, or can be done using infrared or radio network link, among others. The sensor **40** can also include an indoor positioning system or alternatively a global position system (GPS) receiver that relays the position and ambulatory patterns of the patient to the server **20** for mobility tracking.

In one embodiment, the sensors **40** for monitoring vital signs are enclosed in a wrist-watch sized case supported on a wrist band. The sensors can be attached to the back of the case. For example, in one embodiment, Cygnus' AutoSensor (Redwood City, Calif.) is used as a glucose sensor. A low electric current pulls glucose through the skin. Glucose is accumulated in two gel collection discs in the AutoSensor. The AutoSensor measures the glucose and a reading is displayed by the watch.

In another embodiment, EKG/ECG contact points are positioned on the back of the wrist-watch case. In yet another embodiment that provides continuous, beat-to-beat wrist arterial pulse rate measurements, a pressure sensor is housed in a casing with a 'free-floating' plunger as the sensor applanates the radial artery. A strap provides a constant force for effective applanation and ensuring the position of the sensor housing to remain constant after any wrist movements. The change in the electrical signals due to change in pressure is detected as a result of the piezoresistive nature of the sensor are then analyzed to arrive at various arterial pressure, systolic pressure, diastolic pressure, time indices, and other blood pressure parameters.

The case may be of a number of variations of shape but can be conveniently made a rectangular, approaching a box-like configuration. The wrist-band can be an expansion band or a wristwatch strap of plastic, leather or woven material. The wrist-band further contains an antenna for transmitting or receiving radio frequency signals. The wristband and the antenna inside the band are mechanically coupled to the top and bottom sides of the wrist-watch housing. Further, the antenna is electrically coupled to a radio frequency transmitter and receiver for wireless communications with another computer or another user. Although a wrist-band is disclosed, a number of substitutes may be used, including a belt, a ring holder, a brace, or a bracelet, among other suitable substitutes known to one skilled in the art. The housing contains the processor and associated peripherals to provide the human-machine interface. A display is located on the front section of the housing. A speaker, a microphone, and a plurality of push-button switches and are also located on the front section of housing. An infrared LED transmitter and an infrared LED receiver are positioned on the right side of housing to enable the watch to communicate with another computer using infrared transmission.

In another embodiment, the sensors **40** are mounted on the patient's clothing. For example, sensors can be woven into a single-piece garment (an undershirt) on a weaving machine. A plastic optical fiber can be integrated into the structure during the fabric production process without any discontinuities at the armhole or the seams. An interconnection technology transmits information from (and to) sensors mounted at any location on the body thus creating a flexible "bus" structure. T-Connectors—similar to "button clips" used in clothing—are attached to the fibers that serve as a data bus to carry the information from the sensors (e.g., EKG sensors) on the body. The sensors will plug into these connectors and at the other end similar T-Connectors will be used to transmit the information to monitoring equipment or personal status monitor. Since shapes and sizes of humans will be different, sensors can be positioned on the right locations for all patients and without any constraints being imposed by the clothing. Moreover, the clothing can be laundered without any damage to the sensors themselves. In addition to the fiber optic and specialty fibers that serve as sensors and data bus to carry sensory information from the wearer to the monitoring devices, sensors for monitoring the respiration rate can be integrated into the structure.

In another embodiment, instead of being mounted on the patient, the sensors can be mounted on fixed surfaces such as walls or tables, for example. One such sensor is a motion detector. Another sensor is a proximity sensor. The fixed sensors can operate alone or in conjunction with the cameras 10. In one embodiment where the motion detector operates with the cameras 10, the motion detector can be used to trigger camera recording. Thus, as long as motion is sensed, images from the cameras 10 are not saved. However, when motion is not detected, the images are stored and an alarm may be generated. In another embodiment where the motion detector operates stand alone, when no motion is sensed, the system generates an alarm.

The server 20 also executes one or more software modules to analyze data from the patient. A module 50 monitors the patient's vital signs such as ECG/EKG and generates warnings should problems occur. In this module, vital signs can be collected and communicated to the server 20 using wired or wireless transmitters. In one embodiment, the server 20 feeds the data to a statistical analyzer such as a neural network which has been trained to flag potentially dangerous conditions. The neural network can be a back-propagation neural network, for example. In this embodiment, the statistical analyzer is trained with training data where certain signals are determined to be undesirable for the patient, given his age, weight, and physical limitations, among others. For example, the patient's glucose level should be within a well-established range, and any value outside of this range is flagged by the statistical analyzer as a dangerous condition. As used herein, the dangerous condition can be specified as an event or a pattern that can cause physiological or psychological damage to the patient. Moreover, interactions between different vital signals can be accounted for so that the statistical analyzer can take into consideration instances where individually the vital signs are acceptable, but in certain combinations, the vital signs can indicate potentially dangerous conditions. Once trained, the data received by the server 20 can be appropriately scaled and processed by the statistical analyzer. In addition to statistical analyzers, the server 20 can process vital signs using rule-based inference engines, fuzzy logic, as well as conventional if-then logic. Additionally, the server can process vital signs using Hidden Markov Models (HMMs), dynamic time warping, or template matching, among others.

Through various software modules, the system reads video sequence and generates a 3D anatomy file out of the sequence. The proper bone and muscle scene structure are created for head and face. A based profile stock phase shape will be created by this scene structure. Every scene will then be normalized to a standardized viewpoint.

A module 52 monitors the patient ambulatory pattern and generates warnings should the patient's patterns indicate that the patient has fallen or is likely to fall.

FIG. 1B shows one exemplary embodiment hospital wireless monitoring system that forms a "medical body area network" (MBAN). In this embodiment, a physiologic signal sensing and transmitting telemetric device generally designated 10 is shown to comprise a device adapted to be affixed to the chest C of a patient for sensing a physiological function of the patient such as a cardiac function and transmitting suitable radio signals R corresponding thereto to a receiver 11 and a suitable monitor and recording device 12. The receiver 11 may be provided with a suitable antenna 13 for receiving the radio signals R at a location remote from the patient. The receiver may be installed in the same room with the patient, or at a central surveillance area as desired within the range of the transmitting device 10. The patient may utilize a conventional hospital bed B which need not be electrically shockproofed

and is free to move about within the range of the transmitter with the device 10 remaining attached to the patient's body in the manner of a small adhesive bandage which may preferably be of the nonallergenic type.

As shown in FIG. 1C, device 10 may comprise a block 14 of suitable nonallergenic foam plastic having a nonallergenic adhesive coated front surface 15 normally covered by a suitable protective sheet 16. Nonallergenic electrodes 17 project outwardly from the surface 15 and an optional body of suitable electrically conductive nonallergenic jelly of conventional form 18 is provided in association with each electrode also suitably covered by the sheet 16 prior to use of the device. Each electrode illustratively may be formed of silver-silver chloride and comprise a circular electrode of approximately $\frac{3}{4}$ inch diameter with the electrodes being spaced apart approximately 1 inch on center from the block 14. The device may be stored in a sterile pack for extended periods of time within the normal shelf life of the batteries and made available substantially instantaneously for use by the simple removal from the sterile pack and removal of the protective cover sheet 16 and manipulation of the battery actuator 20. An electronic unit 77 contains electronic conditioning circuits and wireless telemetry circuits and rechargeable battery. In one embodiment, the electronic unit 77 is inserted through slots 22 and then slides on rails 20 to make secure electrical contacts with the electrodes 17. In this manner, the electronic unit 77 can be securely coupled to the electrodes 17 to make good electrical contacts thereto. In one embodiment, the electronic unit 77 includes an analog front end chip that can sense bio-potential signals in; digitized signals out; acquisition (ECG) channels+1 driven lead; AC and DC Leads Off Detection; Internal Pace Detection Algorithm on 3 leads; and Thoracic Impedance Measurement (internal/external path).

The package cover device 10 is for one time use and is disposed of after use. To aid in recycling, the electronic unit 77 is removed prior to disposal of device 10 and is disinfected and recharged for a subsequent use. In this manner, the system is environmentally friendly while keeping cost down at a high level of performance. In one embodiment, prior to use, a protective tape electrically isolating the battery is removed, and the electronic module 77 is inserted into disposable device 10. The electronic module 77 cooperates with rails 20 that connect the unit 77 with body contacts 17-18 that make electrical contact with the patient.

The recyclable transceiver with the disposable shell can monitor an array of physiological data, such as temperature, pulse, blood glucose level, blood pressure, and respiratory function. One embodiment can use existing transceivers such as Zigbee, Bluetooth, or WiFi. The frequencies for the MBAN network (2360-2400 MHz) can be aeronautical mobile telemetry (AMT) frequency, for federal radio location tasks, and by amateur radio operators. The bandwidths cover 2360-2400 MHz; 2300-2305 MHz and 2395-2400 MHz; 2400-2483.5 MHz; or 5150-5250 MHz—and reside next those now used by Bluetooth devices. In addition, the modifications proposed by the industry representatives would use the 2390-2400 MHz range as a secondary MBAN network when the primary frequencies interfere with aeronautical industry communications. By allocating spectrum for medical sensors, patients will avoid having wireless dead zones interrupt their transmission of vital data to doctors. This spectrum, previously reserved for commercial test pilots, could be used in hospitals, clinics and doctors' offices. Before a health care facility could use the 2,360 to 2,390 MHz band, they go through a coordination process that considers its actual geographic location in the context of actual aeronautical telemetry receiver locations and actual existing use of the band. Since

there are relatively few aeronautical receive locations and they tend to be clustered-around certain military bases, for example—the majority of hospitals-around 96 percent-would have access to the entire band.

In one implementation, the 2360-2400 MHz for Medical Body Area Networks (MBAN) can be divided into two sub-bands:

In 2360-2390 MHz, devices would be limited to use inside hospitals, contain electronic means to prevent operation outside the hospital and be excluded from zones around AMT facilities using the band unless specifically coordinated with the AMT facility. Hospitals would be required to register with a coordinator when MBAN devices are first deployed.

In 2390-2400 MHz, devices would be permitted anywhere, such as in ambulances for monitoring on way to hospitals, in homes to permit remote monitoring, and anywhere else that electronic monitoring of patients' vital signs may be desired.

These short-range networks comprise small, low-power sensors that can be placed on the body to pick up vital data, such as body temperature and respiratory function. Sensors attach to the body and a local wireless hub. Hospitals will receive a unique key to activate their portion of the band. This key is automatically distributed to MBAN devices in the hospital by the beacon signal, which ensures that the MBAN devices operating according to the key are actually located in the hospital at the time.

One embodiment provides for an 11.5 mile radius exclusion zone to protect AMT sites. Frequency coordination are used to register all locations where devices are operating within hospitals in the 2360-2390 MHz sub-band. Operation of medical devices within an exclusion zone must be pre-coordinated. Conversely, new AMT requirements and mobile AMT operations would have access to accurate information on the location of hospitals with deployed MBAN devices so that coordination can be accomplished without any potential for harmful interference.

In one implementation, compliance with in-hospital and exclusion zone limits can be assured a software key with the controller. The controller could operate in the 2360-2390 MHz band only if loaded with the software key assigned it by the spectrum coordinator. The individual body-worn devices would be operable only as "slaves" and transmit only if they detect an authorized controller. Since controllers are professionally installed and configured, individual body-worn devices could not operate outside the immediate vicinity of a properly authorized controller.

Without the proper coding, both the controller and the body worn devices would default to the 2390-2400 MHz band where operation would be permitted anywhere. Specifically, MBAN devices operating in the 2360-2390 MHz sub-band worn outside the hospital setting would be automatically disabled or could revert to the generally available 2390-2400 MHz sub-band if within range of a controller operating within the same available sub-band. In another embodiment, cognitive technologies including listen-before-talk (LBT), DFS, etc. would permit devices to avoid channels already in use. This embodiment prevents interference to others and also prevents interference to the device itself from others. A hub device can control MBAN sensors and seamlessly move devices to another frequency when interference is encountered, protecting both the other user and the integrity of the MBAN communication. Automatic Power Control (APC) also is likely to be utilized to preserve and extend battery power for these small devices. Another advantage of APC is that the transmitted signal will decrease automatically to the

minimum required to maintain the communications link. This capability will further reduce the interference potential of MBAN devices wherever they are deployed.

Next, security issues will be addressed. In one embodiment, the system uses 64 as well as 128-bit AES encryption. The system implements multiple layers of security measures to control access to mission-critical systems and networks. These are often the targets that an attacker attempts to gain unauthorized access to by compromising a wireless network and using it as an attack path or vector in to an organizational network such as a hospital network where the target systems reside. In order to defend the target environment, multiple security measures are implemented so that if one measure is defeated by an attacker, additional measures and layers of security remain to protect the target environment. Measures such as separation of wireless and wired network segments, strong device and user authentication methods, filtering of traffic based on addresses and protocols, securing end-points/stations from unauthorized access, and monitoring and intrusion detection on the wireless and wired segments are examples of multiple layers of defense that can be employed to achieve a defense-in-depth design. The wireless networks and wired networks should not be directly connected if possible. For example, a wireless environmental sensing LR-WPAN or equipment monitoring LR-WPAN network should not have direct connectivity to the wired healthcare network, but instead be separated by a device such as a firewall, bastion host, or security gateway to establish a security perimeter that can more effectively isolate, segment, and control traffic flows between them. Security features at the upper layer standards and the IEEE lower layer standards are enabled. Both standards have security services defined in their specifications. Having security defined at both the higher and lower layers of the protocol stack creates a stronger security solution.

Source node authentication is implemented in order to cryptographically verify the identity of a transmitting node. Although a shared Network Key will provide a security check for packets utilizing the network, source node authentication can be used by the destination device to verify the identity of the source device. In order to authenticate a source device, a Link Key (end-to-end crypto key) must be generated and used. This key is unique to a pair of devices that are communicating with each other and is derived from their respective Master Keys. (This is equivalent to the concept of a shared secret or unique session key that is derived between two entities in order secure data transmitted between them.)

One implementation of the security architecture includes security mechanisms at three layers of the protocol stack—MAC, Network, and Application. Each layer has services defined for the secure transport of their respective frames. The MAC layer is responsible for its own security processing, but the upper layers determine which security level to use. When MAC layer integrity protection is employed, the entire MAC frame is protected, including the MAC header that contains the hardware source and destination addresses. By enabling MAC frame integrity, the MAC layer source address can be authenticated. This measure can counter address spoofing attacks and allow a device to more effectively process and compare a received MAC frame against an Access Control List (ACL). Cryptography is based on the use of 128-bit keys and the AES encryption standard. Encryption, integrity, and authentication can be applied at the Application, Network, and MAC layers to secure the frames at each of those levels. Master, Link, and Network keys secure transmitted frames. A Network Key is a common key shared among all nodes in a network. The standard also specifies an Alternate Network

Key as a form of key rotation that may be employed for key update purposes. At a minimum, a network should be secured with the use of a Network Key used by all the devices to protect all network frames (routing messages, network join requests, etc.) and prevent the unauthorized joining and use of the network by illegitimate devices. Link Keys, on the other hand, are secret session keys used between two communicating devices and are unique to those devices. Devices use their Master Key to generate the Link Key. The manner in which Master, Link and Network Keys are generated, stored, processed, and delivered to devices determines the effectiveness and degree of security of the overall network implementation.

All devices within a network recognize and trust exactly one Trust Center (TC). The TC stores and distributes keys to devices, but it is preferred to pre-load the keys into the devices directly. The functions performed by the TC are trust management, network management, and configuration management.

In addition to configuring a dedicated Coordinator for the network, a predetermined PAN Identifier is used by the Coordinator. The nodes are limited to joining only the network with the pre-assigned PAN Identifier. Also, the network policy is configured to use the permit join access control to restrict device connectivity. Preferably, an out-of-band method is used for loading the cryptographic key(s) onto the devices. The methods for key management (generation, distribution, updating, revoking, etc.) will vary. Generally, the initial generation and loading of cryptographic keys (e.g. the Master key) will be possible in three ways. One way is out-of-band: This method entails loading the key into the device using a mechanism other than through the normal wireless communication channels used for network operation. An example would be a serial port on the device through which a key could be loaded with a cable attachment to the key generation device (such as a laptop or the Trust Center host). Alternatively, in-band method can be used which delivers keys over-the-air through the normal wireless communication channels used for network operation. This is a less secure method of key delivery because the transmission of the key to a device joining the network that has not been pre-configured is unprotected (creating a potential short period of vulnerability). The least preferred is factory pre-loaded: This method of key deliver consists of the vendor generating and loading the key(s) into the devices at the manufacturing location prior to deliver to the customer. Key values must subsequently be conveyed to the customer when they receive the equipment. This approach is the least secure because the vendor has knowledge of the key values and must also successfully convey the information to the customer in a secure manner. Also, as there is exactly one TC in a network, if possible, the address of the TC should be pre-loaded into the node which can be combined with pre-loading of the crypto keys.

In an embodiment, the communication of health related information between sensors in a MBAN may be subjected to the security requirements such as data confidentiality, data authenticity, data integrity, and data freshness. The data confidentiality described herein may enable the access to the transmitted information by authorized persons (such as the doctor attending the patient). In an embodiment, the data confidentiality can be achieved by encrypting the information before sending it using a secret key and can be both symmetrically and asymmetrically. The data authenticity described herein may provide a means for making sure that the information is sent by the claimed sender (such as a doctor or a patient). In an embodiment, a Message Authentication Code (MAC3) may be calculated using a shared secret key. The data integrity described herein may enable to determine that the

information received has not been tampered with any other intermediate sources. In an embodiment, the system may implement security protocols such as by verifying the MAC. The data freshness described herein may guarantee most recent data can be accessed.

One implementation of the security architecture includes security mechanism that provides secure communication such as by encrypting data using one or more encryption keys. A configuration security server in communication with a security manager of the network may provide a temporary secure communication path between the security manager and a wireless device of the MBAN network. The cryptographic material and other configuration data can then be transferred between the security manager and the wireless devices in the MBAN network via the configuration security server.

One implementation of the security architecture includes techniques for secure communications among MBAN. In response to a packet received at a first access point of the wireless MBAN from a local end-user destined to another end-user associated, the packet can be routed via a tunnelling protocol such as to provide secure communication in the MBAN.

Next, reliable transmission of critical patient data is discussed. As discussed above, hospital environments can produce a significant amount of electromagnetic noise from surgical knives, mobile phones, microwave ovens, and various actuator devices. The resulting EMI (Electro Magnetic Interference) can interfere with the operation of LR-WPAN networks by increasing the white noise floor and reducing the signal-to-noise quality of transmissions. In one implementation, the MAC layer is based on the CSMA/CA (Carrier Sense Multiple Access with Collision Avoidance) channel access method in which a station will first listen for an open channel before transmitting. This is done by sensing the energy level in the frequency band corresponding to the channel. In an environment with significant levels of EMI, the noise floor in the operating frequency ranges of networks can prevent stations from transmitting because the RF energy threshold level for an open channel has been exceeded.

To address this issue, in one embodiment, a Guaranteed Time Slot (GTS) transmission mode accesses the wireless medium based on regular time slots assigned to devices by the LR-WPAN Coordinator. GTS mode can be employed to ensure that devices with critical data to transmit are guaranteed the opportunity to send during a specified time interval without risking a collision with other devices transmitting at the same time.

Another embodiment uses Frequency Hopping (FH) radio with configurable hopping channels and patterns to avoid RFI from surgical knives and similar devices. This type of temporal and frequency diversity approach can improve EMI immunity in an industrial environment as well as provide an additional measure of security if a non-default hopping pattern is used and also changed on a periodic basis.

In another embodiment, reliable multicast provides reliability for information multicast. Usually it is done by negative acknowledgements or repair requests. As one example, SRM (Scalable Reliable Multicast) lets each multicast recipient be responsible for information loss or error by requesting a repair from the whole multicast group (not necessarily from the sender) or by initiating a local recovery.

In yet another embodiment, broadcasting and flooding are used to reach multiple destinations with a best effort in just one session transmission. A recipient may receive more than one copy of exactly the same information, which inadvertently gives rise to some level of redundancy by heavy use of

bandwidth. The broadcast and flooding methods assume that all members are following the rules, and that delivery on a link level is assured.

In a further embodiment, a dynamic redundant transmission policy, with Markov decision theory, is used to find the optimal policy numerically. By relaxing the integer requirement on the number of packets transmitted, the system explicitly finds a real-time recursion for the optimal transmission function. The properties of this recursion are then analyzed to propose a simple numerical procedure for finding the minimum over all real valued transmission functions, by searching over a one-dimensional parameter space. This then yields an implementable, suboptimal policy when discretized to integer values.

In addition to software, the system uses redundant transmitters and antennas to improve reliability. One embodiment increases node density in order to reduce required transmitter power and enable shorter link distances. A mesh topology allows a device to have multiple next-hop neighbors to communicate with and therefore spatial diversity in terms of multiple transmission paths. Higher node density will also permit shorter distances between devices and can result in increased received signal strength and improved signal-to-noise ratios.

FIG. 1D describes generally, an example of transceiver for receiving data over the wireless network. One embodiment can use existing transceivers such as Zigbee, Bluetooth, WiFi, or the like. The frequencies for the MBAN network (2360-2400 MHz) can be aeronautical mobile telemetry (AMT) frequency. In an embodiment, the central station or base stations may be equipped with the different spatially separated antennas and switch the antennas in a round-robin fashion such as while receiving or transmitting data over the MBAN network. In an embodiment, the antennas described herein may directional antennas that may be employed on the wireless devices. In an example, Microstrip antennas may be used by the system that may have a patch area of $3.2 \times 3.2 \text{ cm}^2$, which may allow the placement of these antennas on front, back, or any other sides of the wireless devices. Unlike digital beamforming, the passive directional antennas described herein may produce a directional radiation pattern without extra circuitry or power. In an embodiment, with a directional radiation pattern pointed at the right direction, a wireless client (such as a doctor or a patient) may use reduced transmit power such as to deliver a required receiver signal strength (RSS), or it may increase the RSS with the same transmit power. Further, the client's interference to its peers may also be reduced. In an embodiment, the term "antenna" may refer to the passive antenna without the RF chain.

In an embodiment, the multiple directional antennas (such as antenna 1 to n) may be placed on the surface of the wireless devices such as to select one for directional transmission without adding RF circuitry, in contrast to the simultaneous use of multiple RF chains by beamforming. In an embodiment, the antennas and hospital wireless monitoring system may allow communication to and from the wireless devices in the MBAN. In an embodiment, the wireless devices receive communications covers bandwidth of 2360-2400 MHz, 2300-2305 MHz and 2395-2400 MHz, 2400-2483.5 MHz, 5150-5250 MHz, and the like. In an embodiment, the system may include one or more transceivers configured to transmit and receive communications using different frequency band.

In an embodiment, a challenge for the system is the rotation of the wireless devices during the wireless access. In an embodiment, the device orientation may be critical during the use of the directional antenna because a wireless device can rotate and the rotation may change the device direction much faster than mobility does. The system may be configured to

collect accelerometer and compass readings along with network usage information from different wireless devices in the field. From such field-collected traces, the system may be configured to estimate the orientation and rotation of the wireless devices during the wireless usage.

In an embodiment, the orientation and rotation may impact the directional channels. In an implementation, the system may use a computerized motor platform such as to measure the Received Signal Strength Indicator (RSSI) of directional antennas for indoor and non-line-of-sight (NLOS) environments, with not only controlled orientations, but also rotation according to the field collected traces. In an embodiment, the system may present the directional antennas outperform omni-directional ones for a considerable range of orientations even in NLOS indoor environments. In an embodiment, the directional channels may be highly reciprocal for 802.11 like frequency bands, and their performance may be predictable in short intervals even under realistic rotations.

In an embodiment, another challenge is to allow the wireless devices to dynamically select the best antenna. The system may use a multi-antenna system that may consist of an omni directional antenna, one or more directional antennas, and an antenna switch. In an embodiment, the system works with existing mobile devices that may include a single RF chain and may use only one antenna at a time. In an embodiment, two antenna selection methods such as a packet-based method and a symbol-based method are disclosed herein. The packet-based method may use one packet to assess an antenna without any changes to the network infrastructure. The symbol-based method may use the PHY training symbols such that all antennas are assessed with a single packet. In an embodiment, the use of antenna selection methods, the directional antennas may be effectively employed to improve the transmission gain of wireless devices by almost 3 dB under various propagation environments and realistic rotation. Such gain may be achieved without any change to the deployed network infrastructure such as MBAN.

In another embodiment, antenna redundancy is applied to a central station (base station, access point) with a number K of spatially separated antennas, and to switch the antennas in a round-robin fashion when performing retransmissions for frames directed from the base station to a wireless station. As an example, the first packet is transmitted on antenna 1, the first retransmission on antenna 2, the second retransmission on antenna 3 and so forth.

Alternatively, a number K of tightly synchronized and coupled base stations can be used to achieve the same effect. The system could also use different center frequencies to obtain different channels, but using multiple antennas and spatially diverse channels is more advantageous if obstacles move from time to time in the path between transmission. This approach leverages transmit diversity (and of receiver diversity in the case of packets sent from the wireless station to the access point) while keeping the complexity of the receiver low, as compared to true transmit diversity/MIMO systems. This makes antenna redundancy attractive for scenarios where the wireless stations are small and cheap field devices. For the case of independent (and rather bad) channels between the antennas and a wireless station the antenna redundancy approach decreases the failure probability (i.e. the probability of a request to miss its deadline) exponentially in the number of antennas. For the investigated scenario the difference is approximately one order of magnitude per additional antenna. In addition, already for the second antenna, the system achieves a significant reduction in the mean number of packets needed to transmit a request. This saves an enormous amount of bandwidth, which can be used to serve

other wireless stations. When additionally the antenna reuse strategy is used, then for scenarios with low request inter-arrival periods (as often found in hospital applications) further bandwidth savings can be achieved. This approach is highly effective with antenna redundancy for cases where the channels are not independent but correlated, e.g. for the case where the wireless station is located close to an interferer (surgical knives). In these situations, the best antenna is preferentially used for the cluster of transmission nodes.

In an embodiment, the MBAN can facilitate data redundancy and retransmission techniques such as to high availability data storage, file replication, and some fault tolerant techniques. The security mechanisms can have different characteristics that require different approaches to redundancy. The MBAN can use redundancy to increase assurance of security and data delivery. In an embodiment, different transmission mechanism can be used such as to wirelessly transfer the data among the devices of the system.

In an embodiment, the devices may transmit messages among each in the MBAN. However, in the face of interference such as surgical knives, the message may not be successfully received by the receivers. In an embodiment, the communications may be acknowledged to ensure the reliable delivery of data among the devices. In an embodiment, each device may be configured to include a brief time window in which it is required to send back a short message acknowledging the receipt of the data. In an embodiment, the transmitter or the transmitting device may be configured to wait for the acknowledgement receipt or response message from the receivers such as to ensure the reliable delivery of the data. In an embodiment, if the acknowledgement is received within brief time window then the transmitting device or the transmitter may assume that the message or the data is not received by the receiver and resend the message again. In an embodiment, the process may repeat until the message and acknowledgement are both received. In an embodiment, if even after a few tries the data is not acknowledged by the receiver then the transmitter may be configured to reports a failure notification to the system.

In an implementation, the module 77 may use mesh networking to make the wireless links as reliable as possible. In an embodiment, the devices of interest may configured to be placed close together to each other such that a robust network can be formed such as by simply allowing some of them to route messages on each other's behalf. In an embodiment, the better use of the channel may be made if devices limit their transmit power and communicate only with their near neighbours. Once a mesh network is in place, a number of possible paths may exist between the devices in the MBAN network. In an embodiment, the module 77 may exploit the path diversity such as by using a form of dynamic routing techniques. In an embodiment, the module 77 may allow the system to determine a path failure, as a result of interference or some other change in the environment, the network may pick a different path for the system to transfer the data. In an embodiment, a single-hop transmissions may acknowledged and retried if they fail, multi-hop transmissions through the mesh may also be acknowledged and retried.

In an embodiment, a robust rate adaptation scheme that can be resilient to jamming from surgical knives and other interference sources is disclosed. In an embodiment, the network devices jamming may be detected such as by measuring PDR (Packet Delivery Ratio) with SS (Signal Strength). In an embodiment, the rate adaptation scheme described herein may detect jamming from surgical knives or other equipment and select the data transmission mode, which may include the expected maximum throughput. Through the performance

evaluations, the rate adaptation scheme may improve the packet delivery ratio and the wireless link utilization in the MBAN network. In an embodiment, it may also improve the wireless link utilization by detecting jamming interference and adapting the data transmission mode (modulation and coding levels) to the successful transmission probability.

In another embodiment, an effective approach to diminish the impact of a statistical jammer on TDMA-based MAC protocols is to eliminate the possibility to extract patterns in the wireless network (such as MBAN). In an embodiment, the system may be configured to use these patterns appear as a result of the use of fixed schedules, which may be set when one or more nodes joins the wireless network and may be assumed to repeat till the network is disbanded. In an embodiment, such simple and repetitive patterns may be maintained with tight time synchronization and may result in minimal energy consumption, deterministic end-to-end delay and perhaps maximal transmission concurrency. In an embodiment, the system may be configured to maintain the time synchronization and change the schedule, transmission duration and routes in a randomized yet coordinated manner along small time scales such that impact of the jamming may be reduced in the wireless network.

Transfer Control Protocol (TCP) can provide reliable one-to-one information transmission on top of the IP layer, at which an IP packet is routed to the destination devices along a dynamically determined physical path. If a TCP packet is lost according to acknowledgement information from the receiver, or if its own retransmission timer times out, a TCP sender retransmits the TCP packet. In an embodiment if the data packet is blocked by one or more path interruption threats then the packet can be retransmitted by the system, since the system may get a report that the packet is missing. Further, the system can implement time out techniques before hearing any acknowledgement such that the system can retransmit the data packets if dropping or damaging of acknowledgement.

In an implementation, reliable multicast technique can provide reliability for information multicast. This can be done by negative acknowledgements or repair requests. For example Scalable Reliable Multicast (SRM) can let each multicast device be responsible for information loss or error by requesting a repair from the whole multicast group (not necessarily from the sender) or by initiating a local recovery.

In an embodiment, broadcasting and flooding techniques can be used such as to reach multiple destinations in one session transmission. In MBAN network the receiver may receive more than one copy of exactly the same information, which inadvertently gives rise to some level of redundancy of data. The standard broadcast and flooding techniques assume that all receivers are following the rules, and that delivery on a link level is assured.

In an implementation, high availability data storage techniques can be used that can include one or more one disks to store a copy of the data or the data is dispersed to more than one disk with built-in redundancy such as to deal with disk crashes and provide lower latency for data access.

In an implementation, file replication techniques can be used to make replicas to support easier access to the data among the devices. Establishing mirror web sites for lower latency is one such example. The data backup can be done periodically such as to help restore damaged or lost files from backed-up copies.

In hospital wireless monitoring system, replicated execution may be employed to run a program concurrently at multiple places. The computation can smoothly continue if one execution succeeds. Mapping one transceiver to several dif-

ferent server machines, in a round robin fashion or other more sophisticated way, can prevent one single server from being overloaded and ensure that the data can be accessible even if some server machines have crashed.

In an embodiment, redundancy may be improved by simply increasing the number of the sources or database of same information or the number of transmission paths, particularly when information corruption is detected.

In an embodiment, the system may suffer from data packet drops, transmitting multiple, redundant packets during each sampling interval can improve estimation performance, but at the expense of a higher communication rate. To avoid this, the system can implement techniques such as by relaxing the integer requirement on the number of packets transmitted and identifying real-time recursion for the optimal transmission function.

The technique may implement redundant transmission policies for state estimation of unstable stochastic linear systems over packet dropping channels such as by relaxing the integer requirement on the number of packets transmitted during each sampling interval. The system can determine a real-time recursion for the optimal transmission function. The system can analyze the properties of this recursion such as to propose a simple numerical procedure for finding the minimum over all real-valued transmission functions, which may yield an implementable policy when discretized.

In an embodiment, the system can implement antenna redundancy techniques, which can use different transmitter antennas for performing the data retransmissions. The antenna redundancy technique can equip the system (such as central station, base station, access point, and the like) with a number K of spatially separated antennas capable of switching the antennas in a round-robin fashion when performing retransmissions for frames directed in the MBAN network. For example, the first packet can be transmitted on antenna 1, the first retransmission on antenna 2, the second retransmission on antenna 3 and so forth. In an embodiment, different frequencies can be used to obtain different channels. The use of multiple antennas and spatially diverse channels can be advantageous if obstacles move from time to time in the path between transmitter and receiver, which may happen due to mobility.

In an embodiment, an example of the antenna redundancy technique with K antennas is disclosed. A packet directed from or to the MBAN network can be first transmitted over antenna 1. If there is need for a retransmission, then antenna 2 can be used. If another retransmission is needed, antenna 3 can be used and so forth, until the packet is successfully received or a prescribed deadline for transmitting the request expires. The antennas described herein can be used in round-robin fashion.

FIG. 1E shows another exemplary module with redundant transceivers and redundant antennas. The module includes a plurality of transceivers, for example transceivers that can be selectively operated between 860 MHz to 2.4 GHz, UWB transceiver, and optical transceivers. In this embodiment, the system may employ a simple "listen before you talk" strategy such that the device may be configured to listen such as to check if the channel is busy, and if it is, then the device may wait before checking again. In an embodiment, if the channel is busy and the device keeps on failing to find a clear channel then module 77 can perform exception handling during the jamming period in a number of ways, such as:

- 1) perform frequency hopping between major bands such as AMT 2.4 GHz with 16 channels, 915 MHz and 868 MHz. This can be done by providing programmable/selectable crystals as inputs to the RF transceiver to

periodically alter the frequency. The frequency hopping can be done at preprogrammed intervals or the mesh transceivers can tell each other to flock to new frequency if jamming is detected. In an embodiment, various spreading methods may be commonly used, but the essential idea behind all of them may be to use a bandwidth that may be several orders of magnitude greater than strictly required by the information that is sent. In an embodiment, the signal may be spread over a large bandwidth that may coexist with narrow-band signals, which generally appear to the spread-spectrum receiver as a slight reduction in the signal-to-noise ratio over the spectrum being used;

- 2) place directional antenna on patient bed or directly under patient so that the antenna is very close to transceivers worn by patient and can wirelessly receive data and transfer data back to monitoring devices using power-line transceiver networks;
- 3) place directional antenna on patient bed (or use patient bed as antenna) so that it is very close to transceivers worn by patient to wirelessly receive data and have a wired connection back to monitoring devices to minimize risk of jamming;
- 4) transmit through slower optical transceivers (infrared transceivers such as those on TV remote controls) to transmit data during period where the RF transceivers could not transmit due to jamming;
- 5) use ultrawideband (UWB) transceivers in short duration to transmit data during period where the transceivers could not transmit due to jamming.

FIG. 1F shows a multi-band frequency hopping embodiment. In this embodiment, technique for robust transmission of data in the MBAN network is disclosed. The traditional defenses against jamming may frequency hopping at the physical layer. In an embodiment, the frequency hopping technique may be implemented between major bands such as AMT 2.4 GHz with 16 channels, 915 MHz and 868 MHz, and the like. In an embodiment, the technique may allow the signal hops from channel to channel such as by transmitting short bursts of data at each channel for a set period of time. This may be implemented by providing programmable or selectable crystals as inputs to the RF transceiver to periodically alter the frequency. This may be done at preprogrammed intervals such that the meshed transceivers may inform each other to flock to new frequency if jamming is detected.

FIG. 1G shows a bed antenna embodiment. In this embodiment, distance may have a strong influence on the signal loss or jamming in the MBAN network. In an embodiment, the directional antenna may be placed on the patient bed or directly under the patient such that it may be very close to the transceivers worn by the patient. In an embodiment, the transceivers can wirelessly receive data and transfer data back to the monitoring devices (such as devices of the hospital monitoring system) using power line transceiver networks, there by substantially reducing the risk of jamming in the MBAN network.

FIG. 1H shows a bed receiver with large antenna where the receiver is wired to hospital monitoring equipment. In this embodiment, the directional antenna may be placed on the patient bed (or may use patient bed as antenna) such that it may be very close to transceivers worn by the patient. In an embodiment, the transceivers can wirelessly receive data and may include a wired connection back to the monitoring devices such as to minimize risk of jamming.

FIG. 1I shows a multi-band system that transmits over RF and optical bands. In this embodiment, one technique by which the system may reduce or avoid the jamming is by

using slower optical transceivers. In an embodiment, the slower optical transceivers described herein may include for example, infrared transceivers such as those on TV remote controls. In an embodiment, the system may be configured to detect the jamming such as by determining the RF energy threshold level for an open channel. For example, if the RF energy threshold level for an open is such that no data may be transmitted by the transceivers for a long time, then in an embodiment, slower optical infrared (IR) transceivers may be configured to transmit the data during such period where the wireless transceivers may not transmit the data due to RF interference or jamming. At the end of interference, the system switches back to the regular transceiver for speed and power optimization.

FIG. 1J shows a multi-band system that transmits over ultra-wideband, Zigbee, WiFi and optical bands. In an embodiment, power hungry ultra wideband transceivers may be used to for reliable transfer of data in the MBAN network. In an embodiment, the system may be configured to detect the jamming such as by determining the RF energy threshold level for an open channel. For example, if the RF energy threshold level for an open channel indicates that transmission cannot be done for an extended period (due to interference), where no data may be transmitted by the transceivers, then power hungry ultra wideband transceivers may be configured to transmit data during the short period where the transceivers may not transmit due to jamming. At the end of interference, the system switches back to the regular transceiver for speed and power optimization.

FIG. 1K shows mesh networking that transmits over various wireless and optical protocols in a node to node basis to avoid a jammed node for enhanced reliability. In an implementation, the system may uses mesh networking technology such as to make the wireless links as reliable as possible. In one implementation, the use of ultrawideband, Zigbee, WiFi, AMT, and optical transceivers, each of which connected to corresponding mesh network for ultrawideband, Zigbee, WiFi, AMT, and optical transceivers, renders the system super reliable for patient telemetry. In an embodiment, the devices of interest may be configured to be placed close together to each other such that a robust network can be formed such as by simply allowing some of them to route messages on each other's behalf. In an embodiment, the better use of the channel may be made if devices limit their transmit power and communicate only with their near neighbors. In an embodiment, the mesh network may include a number of possible paths between the devices in the network (such as MBAN). In an embodiment, if the path chosen by the transceivers may be failed, as a result of interference or some other change in the environment, the network may pick a different path for the system to transfer the data. Thus, the system avoids interference or jamming by allowing the mesh transceivers' to immediate switch to another path.

FIG. 1L shows exemplary low power cells for patient telemetry. In an embodiment, the effects of jamming may be reduced such by reducing the transmission range for each wireless node (cell size) in the network. In an embodiment, the cell sizes may be as small as possible such as to make full use of the limited maximum output power of the monitoring devices and to increase the number of possible call re-establishment cells. In an embodiment, the small cell size avoids or reduces the risk of jamming such as by handover to another cell for data transmission.

In an embodiment, the resistance to the jamming may be increased by slowing down the data rate of transmissions. In

an embodiment, as the data transmission rate decreases, the signal-to-noise ratio increases, thereby minimizing jamming in the MBAN network.

FIG. 1M shows an embodiment with scheduled transmissions. In an embodiment, the impact of jamming is diminished by using scheduled transmission techniques. In an embodiment, the system may implement tight time synchronization techniques and may result in minimal energy consumption, deterministic end-to-end delay and perhaps maximal transmission concurrency. In an embodiment, the systems may synchronization signal to be transmitted between the transceivers and the monitoring devices. In an embodiment, the system may be configured to maintain the time synchronization and change the schedule periods and transmission route in randomly on small time periods such that impact of the jamming may be reduced in the MBAN.

FIG. 1N shows an exemplary high gain parabolic bed antenna. In this embodiment, the parabolic antenna is positioned under the patient and the focus of the antenna is aimed at the RF transceiver on a wearable appliance mounted on the patient. The parabolic antenna captures signals from the RF transceiver or signals that bounce off objects such as wall or furniture, for example. The radiation pattern of the feed antenna has to be tailored to the shape of the dish, because it has a strong influence on the aperture efficiency, which determines the antenna gain. Radiation from the feed that falls outside the edge of the dish is called "spillover" and is wasted, reducing the gain and increasing the backlobes, possibly causing interference or (in receiving antennas) increasing susceptibility to ground noise. However, maximum gain is only achieved when the dish is uniformly "illuminated" with a constant field strength to its edges. So the ideal radiation pattern of a feed antenna would be a constant field strength throughout the solid angle of the dish, dropping abruptly to zero at the edges.

In one embodiment, a technique called "dual reflector shaping" may be used. This involves changing the shape of the sub-reflector (usually in a Cassegrain configuration) to map the known pattern of the feed into a uniform illumination of the primary, to maximize the gain. However, this results in a secondary that is no longer precisely hyperbolic (though it is still very close), so the constant phase property is lost. This phase error, however, can be compensated for by slightly tweaking the shape of the primary mirror. The result is a higher gain, or gain/spillover ratio.

Parabolic antennas can use the following type of feed, that is, how the radio waves are supplied to the antenna. One embodiment is an axial or front feed—This is the most common type of feed, with the feed antenna located in front of the dish at the focus, on the beam axis. A disadvantage of this type is that the feed and its supports block some of the beam, which limits the aperture efficiency. Another embodiment is an off-axis or offset feed—The reflector is an asymmetrical segment of a paraboloid, so the focus, and the feed antenna, are located to one side of the dish. The purpose of this design is to move the feed structure out of the beam path, so it doesn't block the beam. Offset feed is also used in multiple reflector designs such as the Cassegrain and Gregorian. In a Cassegrain embodiment, the Cassegrain antenna the feed is located on or behind the dish, and radiates forward, illuminating a convex hyperboloidal secondary reflector at the focus of the dish. The radio waves from the feed reflect back off the secondary reflector to the dish, which forms the outgoing beam. An advantage of this configuration is that the feed, with its waveguides and "front end" electronics does not have to be suspended in front of the dish, so it is used for antennas with complicated or bulky feeds, such as large satellite communi-

cation antennas and radio telescopes. In another embodiment, a Gregorian design is used which is similar to the Cassegrain design except that the secondary reflector is concave, (ellipsoidal) in shape. Aperture efficiency over 70% can be achieved.

Advantages of the wireless monitoring system can include one or more of the following. Patients will enjoy greater mobility when wires can be dispensed with. Immobilized patients are at higher risk for emboli, wasting, bed sores, pneumonias, and other problems. Since cables must be sterilized after use by one patient and before use by another, a wireless approach may decrease the risk of cross-infection. The system presents opportunities for cost savings as well. Earlier intervention is often cheaper, more effective intervention. Wireless monitoring requires fewer staff than more conventional approaches. Sometimes, patients are admitted to ICUs not so much for specialized nursing care as because they need monitoring. If the same monitoring could be carried out in less costly settings, the savings could be appreciable. While as presently envisioned wireless monitoring is intended for hospital use, these devices could also help protect soldiers in combat, and eventually the technology may become suitable for home use. The system can be used for remotely monitoring critically and chronically ill people via small wireless devices so that medical workers can track the person's health status as well as take swift action in emergencies.

By using the system, doctors and nurses in hospitals can avoid having to disconnect patients multiple times, whether it's in an ambulance or various areas of a hospital. The system can also speed up diagnoses, reduce readmissions and allow patients to remain in their homes.

In one embodiment, cameras with 3D detection are used to monitor the patient's ambulation. In the 3D detection process, by putting 3 or more known coordinate objects in a scene, camera origin, view direction and up vector can be calculated and the 3D space that each camera views can be defined.

In one embodiment with two or more cameras, camera parameters (e.g. field of view) are preset to fixed numbers. Each pixel from each camera maps to a cone space. The system identifies one or more 3D feature points (such as a birthmark or an identifiable body landmark) on the patient. The 3D feature point can be detected by identifying the same point from two or more different angles. By determining the intersection for the two or more cones, the system determines the position of the feature point. The above process can be extended to certain feature curves and surfaces, e.g. straight lines, arcs; flat surfaces, cylindrical surfaces. Thus, the system can detect curves if a feature curve is known as a straight line or arc. Additionally, the system can detect surfaces if a feature surface is known as a flat or cylindrical surface. The further the patient is from the camera, the lower the accuracy of the feature point determination. Also, the presence of more cameras would lead to more correlation data for increased accuracy in feature point determination. When correlated feature points, curves and surfaces are detected, the remaining surfaces are detected by texture matching and shading changes. Predetermined constraints are applied based on silhouette curves from different views. A different constraint can be applied when one part of the patient is occluded by another object. Further, as the system knows what basic organic shape it is detecting, the basic profile can be applied and adjusted in the process.

In a single camera embodiment, the 3D feature point (e.g. a birth mark) can be detected if the system can identify the same point from two frames. The relative motion from the two frames should be small but detectable. Other features curves and surfaces will be detected correspondingly, but can be

tessellated or sampled to generate more feature points. A transformation matrix is calculated between a set of feature points from the first frame to a set of feature points from the second frame. When correlated feature points, curves and surfaces are detected, the rest of the surfaces will be detected by texture matching and shading changes.

Each camera exists in a sphere coordinate system where the sphere origin (0,0,0) is defined as the position of the camera. The system detects theta and phi for each observed object, but not the radius or size of the object. The radius is approximated by detecting the size of known objects and scaling the size of known objects to the object whose size is to be determined. For example, to detect the position of a ball that is 10 cm in radius, the system detects the ball and scales other features based on the known ball size. For human, features that are known in advance include head size and leg length, among others. Surface texture can also be detected, but the light and shade information from different camera views is removed. In either single or multiple camera embodiments, depending on frame rate and picture resolution, certain undetected areas such as holes can exist. For example, if the patient yawns, the patient's mouth can appear as a hole in an image. For 3D modeling purposes, the hole can be filled by blending neighborhood surfaces. The blended surfaces are behind the visible line.

In one embodiment shown in FIG. 2A, each camera is calibrated before 3D detection is done. Pseudo-code for one implementation of a camera calibration process is as follows:

Place a calibration sheet with known dots at a known distance (e.g. 1 meter), and perpendicular to a camera view. Take snap shot of the sheet, and correlate the position of the dots to the camera image:

Dot1(x,y,1) ← pixel(x,y)

Place a different calibration sheet that contains known dots at another different known distance (e.g. 2 meters), and perpendicular to camera view.

Take another snapshot of the sheet, and correlate the position of the dots to the camera image:

Dot2(x,y,2) ← pixel(x,y)

Smooth the dots and pixels to minimize digitization errors. By smoothing the map using a global map function, step errors will be eliminated and each pixel will be mapped to a cone space.

For each pixel, draw a line from Dot1 (x,y,z) to Dot2 (x,y,z) defining a cone center where the camera can view.

One smoothing method is to apply a weighted filter for Dot1 and Dot2. A weight filter can be used. In one example, the following exemplary filter is applied.

1 2 1

2 4 2

1 2 1

Assuming Dot1_Left refers to the value of the dot on the left side of Dot1 and Dot1_Right refers to the value of the dot to the right of Dot1 and Dot1_Upper refers to the dot above Dot1, for example, the resulting smoothed Dot1 value is as follows:

$$\frac{1}{16}(\text{Dot1} * 4 + \text{Dot1_Left} * 2 + \text{Dot1_Right} * 2 + \text{Dot1_Upper} * 2 + \text{Dot1_Down} * 2 + \text{Dot1_UpperLeft} + \text{Dot1_UpperRight} + \text{Dot1_LowerLeft} + \text{Dot1_LowerRight})$$

25

Similarly, the resulting smoothed Dot2 value is as follows:

$$\frac{1}{16} * (\text{Dot2_4} + \text{Dot2_Left} * 2 + \text{Dot2_Right} * 2 + \text{Dot2_Upper} * 2 + \text{Dot2_Down} * 2 + \text{Dot2_UpperLeft} + \text{Dot2_UpperRight} + \text{Dot2_LowerLeft} + \text{Dot2_LowerRight})$$

In another smoothing method, features from Dot1 sheet are mapped to a sub pixel level and features of Dot2 sheet are mapped to a sub pixel level and smooth them. To illustrate, Dot1 dot center (5, 5, 1) are mapped to pixel (1.05, 2.86), and Dot2 dot center (10, 10, 2) are mapped to pixel (1.15, 2.76). A predetermined correlation function is then applied.

FIG. 2B shows an exemplary calibration sheet having a plurality of dots. In this embodiment, the dots can be circular dots and square dots which are interleaved among each others. The dots should be placed relatively close to each other and each dot size should not be too large, so we can have as many dots as possible in one snapshot. However, the dots should not be placed too close to each other and the dot size should not be too small, so they are not identifiable.

A module 54 monitors patient activity and generates a warning if the patient has fallen. In one implementation, the system detects the speed of center of mass movement. If the center of mass movement is zero for a predetermined period, the patient is either sleeping or unconscious. The system then attempts to signal the patient and receive confirmatory signals indicating that the patient is conscious. If patient does not confirm, then the system generates an alarm. For example, if the patient has fallen, the system would generate an alarm signal that can be sent to friends, relatives or neighbors of the patient. Alternatively, a third party such as a call center can monitor the alarm signal. Besides monitoring for falls, the system performs video analysis of the patient. For example, during a particular day, the system can determine the amount of time for exercise, sleep, and entertainment, among others. The network of sensors in a patient's home can recognize ordinary patterns—such as eating, sleeping, and greeting visitors—and to alert caretakers to out-of-the-ordinary ones—such as prolonged inactivity or absence. For instance, if the patient goes into the bathroom then disappears off the sensor for 13 minutes and don't show up anywhere else in the house, the system infers that patient had taken a bath or a shower. However, if a person falls and remains motionless for a predetermined period, the system would record the event and notify a designated person to get assistance.

A fall detection process (shown in FIG. 3) performs the following operations:

- Find floor space area
- Define camera view background 3D scene
- Calculate patient's key features
- Detect fall

In one implementation, pseudo-code for determining the floor space area is as follows:

2. Sample the camera view space by M by N, e.g. M=1000, N=500.
3. Calculate all sample points the 3D coordinates in room coordinate system; where Z axis is pointing up. Refer to the 3D detection for how to calculate 3D positions.
4. Find the lowest Z value point (Zmin)
5. Find all points whose Z values are less than Zmin+Ztol; where Ztol is a user adjustable value, e.g. 2 inches.
6. If rooms have different elevation levels, then excluding the lowest Z floor points, repeat step 2, 3 and 4 while keeping the lowest Z is within Ztol2 of previous Z. In this example Ztol2=2 feet, which means the floor level difference should be within 2 feet.
7. Detect stairs by finding approximate same flat area but within equal Z differences between them.

26

8. Optionally, additional information from the user can be used to define floor space more accurately, especially in single camera system where the coordinates are less accurate, e.g.:
 - a. Import the CAD file from constructors' blue prints.
 - b. Pick regions from the camera space to define the floor, then use software to calculate its room coordinates.
 - c. User software to find all flat surfaces, e.g. floors, counter tops, then user pick the ones, which are actually floors and/or stairs.

In the implementation, pseudo-code for determining the camera view background 3D scene is as follows:

1. With the same sample points, calculate x, y coordinates and the Z depth and calculate 3D positions.
2. Determine background scene using one the following methods, among others:
 - a. When there is nobody in the room.
 - b. Retrieve and update the previous calculated background scene.
 - c. Continuous updating every sample point when the furthest Z value was found, that is the background value.

In the implementation, pseudo-code for determining key features of the patient is as follows:

1. Foreground objects can be extracted by comparing each sample point's Z value to the background scene point's Z value, if it is smaller, then it is on the foreground.
2. In normal condition, the feet/shoe can be detected by finding the lowest Z point clouds close the floor in room space, its color will be extracted.
3. In normal condition, the hair/hat can be detected by finding the highest Z point clouds close the floor in room space, its color will be extracted.
4. The rest of the features can be determined by searching from either head or toe. E.g. hat, hair, face, eye, mouth, ear, earring, neck, lipstick, moustache, jacket, limbs, belt, ring, hand, etc.
5. The key dimension of features will be determined by retrieving the historic stored data or recalculated, e.g., head size, mouth width, arm length, leg length, waist, etc.
6. In abnormal conditions, features can be detected by detect individual features then correlated them to different body parts. E.g. if patient's skin is black, we can hardly get a yellow or white face, by detecting eye and nose, we know which part is the face, then we can detect other characteristics.

To detect fall, the pseudo-code for the embodiment is as follows:

1. The fall has to be detected in almost real time by tracking movements of key features very quickly. E.g. if patient has black hair/face, track the center of the black blob will know roughly where his head move to.
2. Then the center of mass will be tracked, center of mass is usually around belly button area, so the belt or borderline between upper and lower body closed will be good indications.
3. Patient's fall always coupled with rapid deceleration of center of mass. Software can adjust this threshold based on patient age, height and physical conditions.
4. Then if the fall is accidental and patient has difficult to get up, one or more of following will happen:
 - a. Patient will move very slowly to find support object to get up.
 - b. Patient will wave hand to camera ask for help. To detect this condition, the patient hand has to be detected first by finding a blob of points with his skin

color. Hand motion can be tracked by calculate the motion of the center of the points, if it swings left and right, it means patient is waving to camera.

- c. Patient is unconscious, motionless. To detect this condition, extract the foreground object, calculate its motion vectors, if it is within certain tolerance, it means patient is not moving. In the mean time, test how long it last, if it past a user defined time threshold, it means patient is in great danger.

In one embodiment for fall detection, the system determines a patient fall-down as when the patient's knee, butt or hand is on the floor. The fall action is defined a quick deceleration of center of mass, which is around belly button area. An accidental fall action is defined when the patient falls down with limited movement for a predetermined period.

The system monitors the patients' fall relative to a floor. In one embodiment, the plan of the floor is specified in advance by the patient. Alternatively, the system can automatically determine the floor layout by examining the movement of the patient's feet and estimated the surfaces touched by the feet as the floor.

In one embodiment with in door positioning, the user can create a facsimile of the floor plan during initialization by walking around the perimeter of each room and recording his/her movement through the in-door positioning system and when complete, press a button to indicate to the system the type of room such as living room, bed room, bath room, among others. Also, the user can calibrate the floor level by sitting down and then standing up (or vice versa) and allowing the accelerometer to sense the floor through the user motion. Periodically, the user can recalibrate the floor plan and/or the floor level.

The system detects a patient fall by detecting a center of mass of an exemplary feature. Thus, the software can monitor the center of one or more objects, for example the head and toe, the patient's belt, the bottom line of the shirt, or the top line of the pants.

The detection of the fall can be adjusted based on two thresholds:

- a. Speed of deceleration of the center of mass.
- b. The amount of time that the patient lies motionless on the floor after the fall.

In one example, once a stroke occurs, the system detects a slow motion of patient as the patient rests or a quick motion as the patient collapses. By adjust the sensitivity threshold, the system detects whether a patient is uncomfortable and ready to rest or collapse.

If the center of mass movement ceases to move for a predetermined period, the system can generate the warning. In another embodiment, before generating the warning, the system can request the patient to confirm that he or she does not need assistance. The confirmation can be in the form of a button that the user can press to override the warning. Alternatively, the confirmation can be in the form of a single utterance that is then detected by a speech recognizer.

In another embodiment, the confirmatory signal is a patient gesture. The patient can nod his or her head to request help and can shake the head to cancel the help request. Alternatively, the patient can use a plurality of hand gestures to signal to the server the actions that the patient desires.

By adding other detecting mechanism such as sweat detection, the system can know whether patient is uncomfortable or not. Other items that can be monitored include chest movement (frequency and amplitude) and rest length when the patient sits still in one area, among others.

Besides monitoring for falls, the system performs video analysis of the patient. For example, during a particular day,

the system can determine the amount of time for exercise, sleep, entertainment, among others. The network of sensors in a patient's home can recognize ordinary patterns—such as eating, sleeping, and greeting visitors—and to alert caretakers to out-of-the-ordinary ones—such as prolonged inactivity or absence. For instance, if the patient goes into the bathroom then disappears off the camera view for a predetermined period and does not show up anywhere else in the house, the system infers that patient had taken a bath or a shower. However, if a person falls and remains motionless for a predetermined period, the system would record the event and notify a designated person to get assistance.

In one embodiment, changes in the patient's skin color can be detected by measuring the current light environment, properly calibrating color space between two photos, and then determining global color change between two states. Thus, when the patient's face turn red, based on the redness, a severity level warning is generated.

In another embodiment, changes in the patient's face are detected by analyzing a texture distortion in the images. If the patient perspires heavily, the texture will show small glisters, make-up smudges, or sweat/tear drippings. Another example is, when long stretched face will be detected as texture distortion. Agony will show certain wrinkle texture patterns, among others.

The system can also utilize high light changes. Thus, when the patient sweats or changes facial appearance, different high light areas are shown, glisters reflect light and pop up geometry generates more high light areas.

A module analyzes facial changes such as facial asymmetries. The change will be detected by superimpose a newly acquired 3D anatomy structure to a historical (normal) 3D anatomy structure to detect face/eye sagging or excess stretch of facial muscles.

In one embodiment, the system determines a set of base 3D shapes, which are a set of shapes which can represent extremes of certain facial effects, e.g. frown, open mouth, smiling, among others. The rest of the 3D face shape can be generated by blending/interpolating these base shapes by applied different weight to each base shapes.

The base 3D shape can be captured using 1) a 3D camera such as cameras from Steinbichler, Genex Technology, Minolta 3D, Olympus 3D or 2) one or more 2D camera with preset camera field of view (FOV) parameters. To make it more accurate, one or more special markers can be placed on patient's face. For example, a known dimension square stick can be placed on the forehead for camera calibration purposes.

Using the above 3D detection method, facial shapes are then extracted. The proper features (e.g. a wrinkle) will be detected and attached to each base shape. These features can be animated or blended by changing the weight of different shape(s). The proper features change can be detected and determine what type of facial shape it will be.

Next, the system super-imposes two 3D facial shapes (historical or normal facial shapes and current facial shapes). By matching features and geometry of changing areas on the face, closely blended shapes can be matched and facial shape change detection can be performed. By overlaying the two shapes, the abnormal facial change such as sagging eyes or mouth can be detected.

The above processes are used to determine paralysis of specific regions of the face or disorders in the peripheral or central nervous system (trigeminal paralysis; CVA, among others). The software also detects eyelid positions for evidence of ptosis (incomplete opening of one or both eyelids) as a sign of innervation problems (CVA; Horner syndrome, for

example). The software also checks eye movements for pathological conditions, mainly of neurological origin are reflected in aberrations in eye movement. Pupil reaction is also checked for abnormal reaction of the pupil to light (pupil gets smaller the stronger the light) may indicate various pathological conditions mainly of the nervous system. In patients treated for glaucoma pupillary status and motion pattern may be important to the follow-up of adequate treatment. The software also checks for asymmetry in tongue movement, which is usually indicative of neurological problems. Another check is neck veins: Engorgement of the neck veins may be an indication of heart failure or obstruction of normal blood flow from the head and upper extremities to the heart. The software also analyzes the face, which is usually a mirror of the emotional state of the observed subject. Fear, joy, anger, apathy are only some of the emotions that can be readily detected, facial expressions of emotions are relatively uniform regardless of age, sex, race, etc. This relative uniformity allows for the creation of computer programs attempting to automatically diagnose people's emotional states.

Speech recognition is performed to determine a change in the form of speech (slurred speech, difficulties in the formation of words, for example) may indicated neurological problems, such an observation can also indicate some outward effects of various drugs or toxic agents.

In one embodiment shown in FIG. 4, a facial expression analysis process performs the following operations:

- Find floor space area
- Define camera view background 3D scene
- Calculate patient's key features
- Extract facial objects
- Detect facial orientation
- Detect facial expression

The first three steps are already discussed above. The patient's key features provide information on the location of the face, and once the face area has been determined, other features can be detected by detecting relative position to each other and special characteristics of the features:

- Eye: pupil can be detected by applying Chamfer matching algorithm, by using stock pupil objects.
- Hair: located on the top of the head, using previous stored hair color to locate the hair point clouds.
- Birthmarks, wrinkles and tattoos: pre store all these features then use Chamfer matching to locate them.
- Nose: nose bridge and nose holes usually show special characteristics for detection, sometime depend on the view angle, is side view, special silhouette will be shown.
- Eye browse, Lips and Moustache: All these features have special colors, e.g. red lipstick; and base shape, e.g. patient has no expression with mouth closed. Software will locate these features by color matching, then try to deform the base shape based on expression, and match shape with expression, we can detect objects and expression at the same time.
- Teeth, earring, necklace: All these features can be detected by color and style, which will give extra information.

In one implementation, pseudo-code for detecting facial orientation is as follows:

- Detect forehead area
- Use the previously determined features and superimpose them on the base face model to detect a patient face orientation.

Depends on where patient is facing, for a side facing view, silhouette edges will provide unique view information because there is a one to one correspondent between the view and silhouette shape.

Once the patient's face has been aligned to the right view, exemplary pseudo code to detect facial expression is as follows:

1. Detect shape change. The shape can be match by superimpose different expression shapes to current shape, and judge by minimum discrepancy. E.g. wide mouth open.
2. Detect occlusion. Sometime the expression can be detected by occlusal of another objects, e.g., teeth show up means mouth is open.
3. Detect texture map change. The expression can relate to certain texture changes, if patient smile, certain wrinkles patents will show up.
4. Detect highlight change. The expression can relate to certain high light changes, if patient sweats or cries, different highlight area will show up.

Speech recognition can be performed in one embodiment to determine a change in the form of speech (slurred speech, difficulties in the formation of words, for example) may indicated neurological problems, such an observation can also indicate some outward effects of various drugs or toxic agents.

A module communicates with a third party such as the police department, a security monitoring center, or a call center. The module operates with a POTS telephone and can use a broadband medium such as DSL or cable network if available. The module **80** requires that at least the telephone is available as a lifeline support. In this embodiment, duplex sound transmission is done using the POTS telephone network. The broadband network, if available, is optional for high resolution video and other advanced services transmission.

During operation, the module checks whether broadband network is available. If broadband network is available, the module **80** allows high resolution video, among others, to be broadcasted directly from the server **20** to the third party or indirectly from the server **20** to the remote server **200** to the third party. In parallel, the module **80** allows sound to be transmitted using the telephone circuit. In this manner, high resolution video can be transmitted since sound data is separately sent through the POTS network.

If broadband network is not available, the system relies on the POTS telephone network for transmission of voice and images. In this system, one or more images are compressed for burst transmission, and at the request of the third party or the remote server **200**, the telephone's sound system is placed on hold for a brief period to allow transmission of images over the POTS network. In this manner, existing POTS lifeline telephone can be used to monitor patients. The resolution and quantity of images are selectable by the third party. Thus, using only the lifeline as a communication medium, the person monitoring the patient can elect to only listen, to view one high resolution image with duplex telephone voice transmission, to view a few low resolution images, to view a compressed stream of low resolution video with digitized voice, among others.

During installation or while no live person in the scene, each camera will capture its own environment objects and store it as background images, the software then detect the live person in the scene, changes of the live person, so only the portion of live person will be send to the local server, other compression techniques will be applied, e.g. send changing file, balanced video streaming based on change.

The local server will control and schedule how the video/picture will be send, e.g. when the camera is view an empty room, no pictures will be sent, the local server will also determine which camera is at the right view, and request only the corresponding video be sent. The local server will deter-

mine which feature it is interested in looking at, e.g. face and request only that portion be sent.

With predetermined background images and local server controlled streaming, the system will enable higher resolution and more camera system by using narrower bandwidth.

Through this module, a police officer, a security agent, or a healthcare agent such as a physician at a remote location can engage, in interactive visual communication with the patient. The patient's health data or audio-visual signal can be remotely accessed. The patient also has access to a video transmission of the third party. Should the patient experience health symptoms requiring intervention and immediate care, the health care practitioner at the central station may summon help from an emergency services provider. The emergency services provider may send an ambulance, fire department personnel, family member, or other emergency personnel to the patient's remote location. The emergency services provider may, perhaps, be an ambulance facility, a police station, the local fire department, or any suitable support facility.

Communication between the patient's remote location and the central station can be initiated by a variety of techniques. One method is by manually or automatically placing a call on the telephone to the patient's home or from the patient's home to the central station.

Alternatively, the system can ask a confirmatory question to the patient through text to speech software. The patient can be orally instructed by the health practitioner to conduct specific physical activities such as specific arm movements, walking, bending, among others. The examination begins during the initial conversation with the monitored subject. Any changes in the spontaneous gestures of the body, arms and hands during speech as well as the fulfillment of nonspecific tasks are important signs of possible pathological events. The monitoring person can instruct the monitored subject to perform a series of simple tasks which can be used for diagnosis of neurological abnormalities. These observations may yield early indicators of the onset of a disease.

A network 100 such as the Internet receives images from the server 20 and passes the data to one or more remote servers 200. The images are transmitted from the server 200 over a secure communication link such as virtual private network (VPN) to the remote server(s) 200.

In one embodiment where cameras are deployed, the server 200 collects data from a plurality of cameras and uses the 3D images technology to determine if the patient needs help. The system can transmit video (live or archived) to the friend, relative, neighbor, or call center for human review. At each viewer site, after a viewer specifies the correct URL to the client browser computer, a connection with the server 200 is established and user identity authenticated using suitable password or other security mechanisms. The server 200 then retrieves the document from its local disk or cache memory storage and transmits the content over the network. In the typical scenario, the user of a Web browser requests that a media stream file be downloaded, such as sending, in particular, the URL of a media redirection file from a Web server. The media redirection file (MRF) is a type of specialized Hypertext Markup Language (HTML) file that contains instructions for how to locate the multimedia file and in what format the multimedia file is in. The Web server returns the MRF file to the user's browser program. The browser program then reads the MRF file to determine the location of the media server containing one or more multimedia content files. The browser then launches the associated media player application program and passes the MRF file to it. The media player reads the MRF file to obtain the information needed to open a connection to a media server, such as a URL, and the required

protocol information, depending upon the type of medial content is in the file. The streaming media content file is then routed from the media server down to the user.

In the camera embodiment, the transactions between the server 200 and one of the remote servers 200 are detailed. The server 200 compares one image frame to the next image frame. If no difference exists, the duplicate frame is deleted to minimize storage space. If a difference exists, only the difference information is stored as described in the JPEG standard. This operation effectively compresses video information so that the camera images can be transmitted even at telephone modem speed of 64 k or less. More aggressive compression techniques can be used. For example, patient movements can be clusterized into a group of known motion vectors, and patient movements can be described using a set of vectors. Only the vector data is saved. During view back, each vector is translated into a picture object which is suitably rasterized. The information can also be compressed as motion information.

Next, the server 200 transmits the compressed video to the remote server 200. The server 200 stores and caches the video data so that multiple viewers can view the images at once since the server 200 is connected to a network link such as telephone line modem, cable modem, DSL modem, and ATM transceiver, among others.

In one implementation, the servers 200 use RAID-5 striping and parity techniques to organize data in a fault tolerant and efficient manner. The RAID (Redundant Array of Inexpensive Disks) approach is well described in the literature and has various levels of operation, including RAID-5, and the data organization can achieve data storage in a fault tolerant and load balanced manner. RAID-5 provides that the stored data is spread among three or more disk drives, in a redundant manner, so that even if one of the disk drives fails, the data stored on the drive can be recovered in an efficient and error free manner from the other storage locations. This method also advantageously makes use of each of the disk drives in relatively equal and substantially parallel operations. Accordingly, if one has a six gigabyte cluster volume which spans three disk drives, each disk drive would be responsible for servicing two gigabytes of the cluster volume. Each two gigabyte drive would be comprised of one-third redundant information, to provide the redundant, and thus fault tolerant, operation required for the RAID-5 approach. For additional physical security, the server can be stored in a Fire Safe or other secured box, so there is no chance to erase the recorded data, this is very important for forensic analysis.

The system can also monitor the patient's gait pattern and generate warnings should the patient's gait patterns indicate that the patient is likely to fall. The system will detect patient skeleton structure, stride and frequency; and based on this information to judge whether patient has joint problem, asymmetrical bone structure, among others. The system can store historical gait information, and by overlaying current structure to the historical (normal) gait information, gait changes can be detected. In the camera embodiment, an estimate of the gait pattern is done using the camera. In a camera-less embodiment, the gait can be sensed by providing a sensor on the floor and a sensor near the head and the variance in the two sensor positions are used to estimate gait characteristics.

The system also provides a patient interface 90 to assist the patient in easily accessing information. In one embodiment, the patient interface includes a touch screen; voice-activated text reading; one touch telephone dialing; and video conferencing. The touch screen has large icons that are pre-selected to the patient's needs, such as his or her favorite web sites or application programs. The voice activated text reading allows

a user with poor eye-sight to get information from the patient interface **90**. Buttons with pre-designated dialing numbers, or video conferencing contact information allow the user to call a friend or a healthcare provider quickly.

In one embodiment, medicine for the patient is tracked using radio frequency identification (RFID) tags. In this embodiment, each drug container is tracked through an RFID tag that is also a drug label. The RF tag is an integrated circuit that is coupled with a mini-antenna to transmit data. The circuit contains memory that stores the identification Code and other pertinent data to be transmitted when the chip is activated or interrogated using radio energy from a reader. A reader consists of an RF antenna, transceiver and a micro-processor. The transceiver sends activation signals to and receives identification data from the tag. The antenna may be enclosed with the reader or located outside the reader as a separate piece. RFID readers communicate directly with the RFID tags and send encrypted usage data over the patient's network to the server **200** and eventually over the Internet **100**. The readers can be built directly into the walls or the cabinet doors.

In one embodiment, capacitively coupled RFID tags are used. The capacitive RFID tag includes a silicon microprocessor that can store 96 bits of information, including the pharmaceutical manufacturer, drug name, usage instruction and a 40-bit serial number. A conductive carbon ink acts as the tag's antenna and is applied to a paper substrate through conventional printing means. The silicon chip is attached to printed carbon-ink electrodes on the back of a paper label, creating a low-cost, disposable tag that can be integrated on the drug label. The information stored on the drug labels is written in a Medicine Markup Language (MML), which is based on the eXtensible Markup Language (XML). MML would allow all computers to communicate with any computer system in a similar way that Web servers read Hyper Text Markup Language (HTML), the common language used to create Web pages.

After receiving the medicine container, the patient places the medicine in a medicine cabinet, which is also equipped with a tag reader. This smart cabinet then tracks all medicine stored in it. It can track the medicine taken, how often the medicine is restocked and can let the patient know when a particular medication is about to expire. At this point, the server **200** can order these items automatically. The server **200** also monitors drug compliance, and if the patient does not remove the bottle to dispense medication as prescribed, the server **200** sends a warning to the healthcare provider.

The user's habits can be determined by the system. This is done by tracking location, ambulatory travel vectors and time in a database. Thus, if the user typically sleeps between 10 pm to 6 am, the location would reflect that the user's location maps to the bedroom between 10 pm and 6 am. In one exemplary system, the system builds a schedule of the user's activity as follows:

Location	Time Start	Time End	Heart Rate
Bed room	10 pm	6 am	60-80
Gym room	6 am	7 am	90-120
Bath room	7 am	7:30 am	85-120
Dining room	7:30 am	8:45 am	80-90
Home Office	8:45 am	11:30 am	85-100
...			
...			

The habit tracking is adaptive in that it gradually adjusts to the user's new habits. If there are sudden changes, the system flags these sudden changes for follow up. For instance, if the user spends three hours in the bathroom, the system prompts the third party (such as a call center) to follow up with the patient to make sure he or she does not need help.

In one embodiment, data driven analyzers may be used to track the patient's habits. These data driven analyzers may incorporate a number of models such as parametric statistical models, non-parametric statistical models, clustering models, nearest neighbor models, regression methods, and engineered (artificial) neural networks. Prior to operation, data driven analyzers or models of the patient's habits or ambulation patterns are built using one or more training sessions. The data used to build the analyzer or model in these sessions are typically referred to as training data. As data driven analyzers are developed by examining only training examples, the selection of the training data can significantly affect the accuracy and the learning speed of the data driven analyzer. One approach used heretofore generates a separate data set referred to as a test set for training purposes. The test set is used to avoid overfitting the model or analyzer to the training data. Overfitting refers to the situation where the analyzer has memorized the training data so well that it fails to fit or categorize unseen data. Typically, during the construction of the analyzer or model, the analyzer's performance is tested against the test set. The selection of the analyzer or model parameters is performed iteratively until the performance of the analyzer in classifying the test set reaches an optimal point. At this point, the training process is completed. An alternative to using an independent training and test set is to use a methodology called cross-validation. Cross-validation can be used to determine parameter values for a parametric analyzer or model for a non-parametric analyzer. In cross-validation, a single training data set is selected. Next, a number of different analyzers or models are built by presenting different parts of the training data as test sets to the analyzers in an iterative process. The parameter or model structure is then determined on the basis of the combined performance of all models or analyzers. Under the cross-validation approach, the analyzer or model is typically retrained with data using the determined optimal model structure.

In one embodiment, clustering operations are performed to detect patterns in the data. In another embodiment, a neural network is used to recognize each pattern as the neural network is quite robust at recognizing user habits or patterns. Once the treatment features have been characterized, the neural network then compares the input user information with stored templates of treatment vocabulary known by the neural network recognizer, among others. The recognition models can include a Hidden Markov Model (HMM), a dynamic programming model, a neural network, a fuzzy logic, or a template matcher, among others. These models may be used singly or in combination.

Dynamic programming considers all possible points within the permitted domain for each value of i . Because the best path from the current point to the next point is independent of what happens beyond that point. Thus, the total cost of $[i(k), j(k)]$ is the cost of the point itself plus the cost of the minimum path to it. Preferably, the values of the predecessors can be kept in an $M \times N$ array, and the accumulated cost kept in a $2 \times N$ array to contain the accumulated costs of the immediately preceding column and the current column. However, this method requires significant computing resources. For the recognizer to find the optimal time alignment between a sequence of frames and a sequence of node models, it must compare most frames against a plurality of node models. One

method of reducing the amount of computation required for dynamic programming is to use pruning. Pruning terminates the dynamic programming of a given portion of user habit information against a given treatment model if the partial probability score for that comparison drops below a given threshold. This greatly reduces computation.

Considered to be a generalization of dynamic programming, a hidden Markov model is used in the preferred embodiment to evaluate the probability of occurrence of a sequence of observations $O(1), O(2), \dots, O(t), \dots, O(T)$, where each observation $O(t)$ may be either a discrete symbol under the VQ approach or a continuous vector. The sequence of observations may be modeled as a probabilistic function of an underlying Markov chain having state transitions that are not directly observable. In one embodiment, the Markov network is used to model a number of user habits and activities. The transitions between states are represented by a transition matrix $A=[a(i,j)]$. Each $a(i,j)$ term of the transition matrix is the probability of making a transition to state j given that the model is in state i . The output symbol probability of the model is represented by a set of functions $B=[b(j) O(t)]$, where the $b(j) O(t)$ term of the output symbol matrix is the probability of outputting observation $O(t)$, given that the model is in state j . The first state is always constrained to be the initial state for the first time frame of the utterance, as only a prescribed set of left to right state transitions are possible. A predetermined final state is defined from which transitions to other states cannot occur. Transitions are restricted to reentry of a state or entry to one of the next two states. Such transitions are defined in the model as transition probabilities. Although the preferred embodiment restricts the flow graphs to the present state or to the next two states, one skilled in the art can build an HMM model without any transition restrictions, although the sum of all the probabilities of transitioning from any state must still add up to one. In each state of the model, the current feature frame may be identified with one of a set of predefined output symbols or may be labeled probabilistically. In this case, the output symbol probability $b(j) O(t)$ corresponds to the probability assigned by the model that the feature frame symbol is $O(t)$. The model arrangement is a matrix $A=[a(i,j)]$ of transition probabilities and a technique of computing $B=b(j) O(t)$, the feature frame symbol probability in state j . The Markov model is formed for a reference pattern from a plurality of sequences of training patterns and the output symbol probabilities are multivariate Gaussian function probability densities. The patient habit information is processed by a feature extractor. During learning, the resulting feature vector series is processed by a parameter estimator, whose output is provided to the hidden Markov model. The hidden Markov model is used to derive a set of reference pattern templates, each template representative of an identified pattern in a vocabulary set of reference treatment patterns. The Markov model reference templates are next utilized to classify a sequence of observations into one of the reference patterns based on the probability of generating the observations from each Markov model reference pattern template. During recognition, the unknown pattern can then be identified as the reference pattern with the highest probability in the likelihood calculator. The HMM template has a number of states, each having a discrete value. However, because treatment pattern features may have a dynamic pattern in contrast to a single value. The addition of a neural network at the front end of the HMM in an embodiment provides the capability of representing states with dynamic values. The input layer of the neural network comprises input neurons. The outputs of the input layer are distributed to all neurons in the middle layer. Similarly, the outputs of the middle layer are distributed

to all output states, which normally would be the output layer of the neuron. However, each output has transition probabilities to itself or to the next outputs, thus forming a modified HMM. Each state of the thus formed HMM is capable of responding to a particular dynamic signal, resulting in a more robust HMM. Alternatively, the neural network can be used alone without resorting to the transition probabilities of the HMM architecture.

The system allows patients to conduct a low-cost, comprehensive, real-time monitoring of their vital parameters such as ambulation and falls. Information can be viewed using an Internet-based website, a personal computer, or simply by viewing a display on the monitor. Data measured several times each day provide a relatively comprehensive data set compared to that measured during medical appointments separated by several weeks or even months. This allows both the patient and medical professional to observe trends in the data, such as a gradual increase or decrease in blood pressure, which may indicate a medical condition. The invention also minimizes effects of white coat syndrome since the monitor automatically makes measurements with basically no discomfort; measurements are made at the patient's home or work, rather than in a medical office.

The wearable appliance is small, easily worn by the patient during periods of exercise or day-to-day activities, and non-invasively measures blood pressure can be done in a matter of seconds without affecting the patient. An on-board or remote processor can analyze the time-dependent measurements to generate statistics on a patient's blood pressure (e.g., average pressures, standard deviation, beat-to-beat pressure variations) that are not available with conventional devices that only measure systolic and diastolic blood pressure at isolated times.

The wearable appliance provides an in-depth, cost-effective mechanism to evaluate a patient's health condition. Certain cardiac conditions can be controlled, and in some cases predicted, before they actually occur. Moreover, data from the patient can be collected and analyzed while the patient participates in their normal, day-to-day activities.

Software programs associated with the Internet-accessible website, secondary software system, and the personal computer analyze the blood pressure, and heart rate, and pulse oximetry values to characterize the patient's cardiac condition. These programs, for example, may provide a report that features statistical analysis of these data to determine averages, data displayed in a graphical format, trends, and comparisons to doctor-recommended values.

When the appliance cannot communicate with the mesh network, the appliance simply stores information in memory and continues to make measurements. The watch component automatically transmits all the stored information (along with a time/date stamp) when it comes in proximity to the wireless mesh network, which then transmits the information through the wireless network.

In one embodiment, the server provides a web services that communicate with third party software through an interface. To generate vital parameters such as blood pressure information for the web services software interface, the patient continuously wears the blood-pressure monitor for a short period of time, e.g. one to two weeks after visiting a medical professional during a typical 'check up' or after signing up for a short-term monitoring program through the website. In this case, the wearable device such as the watch measures mobility through the accelerometer and blood pressure in a near-continuous, periodic manner such as every fifteen minutes.

This information is then transmitted over the mesh network to a base station that communicates over the Internet to the server.

To view information sent from the blood-pressure monitor and fall detector on the wearable appliance, the patient or an authorized third party such as family members, emergency personnel, or medical professional accesses a patient user interface hosted on the web server **200** through the Internet **100** from a remote computer system. The patient interface displays vital information such as ambulation, blood pressure and related data measured from a single patient. The system may also include a call center, typically staffed with medical professionals such as doctors, nurses, or nurse practitioners, whom access a care-provider interface hosted on the same website on the server **200**. The care-provider interface displays vital data from multiple patients.

The wearable appliance has an indoor positioning system and processes these signals to determine a location (e.g., latitude, longitude, and altitude) of the monitor and, presumably, the patient. This location could be plotted on a map by the server, and used to locate a patient during an emergency, e.g. to dispatch an ambulance.

In one embodiment, the web page hosted by the server **200** includes a header field that lists general information about the patient (e.g. name, age, and ID number, general location, and information concerning recent measurements); a table that lists recently measured blood pressure data and suggested (i.e. doctor-recommended) values of these data; and graphs that plot the systolic and diastolic blood pressure data in a time-dependent manner. The header field additionally includes a series of tabs that each link to separate web pages that include, e.g., tables and graphs corresponding to a different data measured by the wearable device such as calorie consumption/dissipation, ambulation pattern, sleeping pattern, heart rate, pulse oximetry, and temperature. The table lists a series of data fields that show running average values of the patient's daily, monthly, and yearly vital parameters. The levels are compared to a series of corresponding 'suggested' values of vital parameters that are extracted from a database associated with the web site. The suggested values depend on, among other things, the patient's age, sex, and weight. The table then calculates the difference between the running average and suggested values to give the patient an idea of how their data compares to that of a healthy patient. The web software interface may also include security measures such as authentication, authorization, encryption, credential presentation, and digital signature resolution. The interface may also be modified to conform to industry-mandated, XML schema definitions, while being 'backwards compatible' with any existing XML schema definitions.

The system provides for self-registration of appliances by the user. Data can be synchronized between the Repository and appliance(s) via the base station **20**. The user can preview the readings received from the appliance(s) and reject erroneous readings. The user or treating professional can set up the system to generate alerts against received data, based on pre-defined parameters. The system can determine trends in received data, based on user defined parameters.

Appliance registration is the process by which a patient monitoring appliance is associated with one or more users of the system. This mechanism is also used when provisioning appliances for a user by a third party, such as a clinician (or their respective delegate). In one implementation, the user (or delegate) logs into the portal to select one or more appliances and available for registration. In turn, the base station server **20** broadcasts a query to all nodes in the mesh network to retrieve identification information for the appliance such as

manufacturer information, appliance model information, appliance serial number and optionally a hub number (available on hub packaging). The user may register more than one appliance at this point. The system optionally sets up a service subscription for appliance(s) usage. This includes selecting service plans and providing payment information. The appliance(s) are then associated with this user's account and a control file with appliance identification information is synchronized between the server **200** and the base station **20** and each appliance on initialization. In one embodiment, each appliance **8** transmits data to the base station **20** in an XML format for ease of interfacing and is either kept encrypted or in a non-readable format on the base station **20** for security reasons.

The base station **20** frequently collects and synchronizes data from the appliances **8**. The base station **20** may use one of various transportation methods to connect to the repository on the server **200** using a PC as conduit or through a connection established using an embedded modem (connected to a phone line), a wireless router (DSL or cable wireless router), a cellular modem, or another network-connected appliance (such as, but not limited to, a web-phone, video-phone, embedded computer, PDA or handheld computer).

In one embodiment, users may set up alerts or reminders that are triggered when one or more reading meet a certain set of conditions, depending on parameters defined by the user. The user chooses the condition that they would like to be alerted to and by providing the parameters (e.g. threshold value for the reading) for alert generation. Each alert may have an interval which may be either the number of data points or a time duration in units such as hours, days, weeks or months. The user chooses the destination where the alert may be sent. This destination may include the user's portal, e-mail, pager, voice-mail or any combination of the above.

Trends are determined by applying mathematical and statistical rules (e.g. moving average and deviation) over a set of reading values. Each rule is configurable by parameters that are either automatically calculated or are set by the user.

The user may give permission to others as needed to read or edit their personal data or receive alerts. The user or clinician could have a list of people that they want to monitor and have it show on their "My Account" page, which serves as a local central monitoring station in one embodiment. Each person may be assigned different access rights which may be more or less than the access rights that the patient has. For example, a doctor or clinician could be allowed to edit data for example to annotate it, while the patient would have read-only privileges for certain pages. An authorized person could set the reminders and alerts parameters with limited access to others. In one embodiment, the base station server **20** serves a web page customized by the user or the user's representative as the monitoring center that third parties such as family, physicians, or caregivers can log in and access information. In another embodiment, the base station **20** communicates with the server **200** at a call center so that the call center provides all services. In yet another embodiment, a hybrid solution where authorized representatives can log in to the base station server **20** access patient information while the call center logs into both the server **200** and the base station server **20** to provide complete care services to the patient.

The server **200** may communicate with a business process outsourcing (BPO) company or a call center to provide central monitoring in an environment where a small number of monitoring agents can cost effectively monitor multiple people 24 hours a day. A call center agent, a clinician or a nursing home manager may monitor a group or a number of users via a summary "dashboard" of their readings data, with

ability to drill-down into details for the collected data. A clinician administrator may monitor the data for and otherwise administer a number of users of the system. A summary “dashboard” of readings from all Patients assigned to the Administrator is displayed upon log in to the Portal by the Administrator. Readings may be color coded to visually distinguish normal vs. readings that have generated an alert, along with description of the alert generated. The Administrator may drill down into the details for each Patient to further examine the readings data, view charts etc. in a manner similar to the Patient’s own use of the system. The Administrator may also view a summary of all the appliances registered to all assigned Patients, including but not limited to all appliance identification information. The Administrator has access only to information about Patients that have been assigned to the Administrator by a Super Administrator. This allows for segmenting the entire population of monitored Patients amongst multiple Administrators. The Super Administrator may assign, remove and/or reassign Patients amongst a number of Administrators.

In one embodiment, a patient using an Internet-accessible computer and web browser, directs the browser to an appropriate URL and signs up for a service for a short-term (e.g., 1 month) period of time. The company providing the service completes an accompanying financial transaction (e.g. processes a credit card), registers the patient, and ships the patient a wearable appliance for the short period of time. The registration process involves recording the patient’s name and contact information, a number associated with the monitor (e.g. a serial number), and setting up a personalized website. The patient then uses the monitor throughout the monitoring period, e.g. while working, sleeping, and exercising. During this time the monitor measures data from the patient and wirelessly transmits it through the channel to a data center. There, the data are analyzed using software running on computer servers to generate a statistical report. The computer servers then automatically send the report to the patient using email, regular mail, or a facsimile machine at different times during the monitoring period. When the monitoring period is expired, the patient ships the wearable appliance back to the monitoring company.

Different web pages may be designed and accessed depending on the end-user. As described above, individual users have access to web pages that only their ambulation and blood pressure data (i.e., the patient interface), while organizations that support a large number of patients (nursing homes or hospitals) have access to web pages that contain data from a group of patients using a care-provider interface. Other interfaces can also be used with the web site, such as interfaces used for: insurance companies, members of a particular company, clinical trials for pharmaceutical companies, and e-commerce purposes. Vital patient data displayed on these web pages, for example, can be sorted and analyzed depending on the patient’s medical history, age, sex, medical condition, and geographic location. The web pages also support a wide range of algorithms that can be used to analyze data once they are extracted from the data packets. For example, an instant message or email can be sent out as an ‘alert’ in response to blood pressure indicating a medical condition that requires immediate attention. Alternatively, the message could be sent out when a data parameter (e.g. systolic blood pressure) exceeds a predetermined value. In some cases, multiple parameters (e.g., fall detection, positioning data, and blood pressure) can be analyzed simultaneously to generate an alert message. In general, an alert message can be sent out after analyzing one or more data parameters using any type of algorithm. These algorithms range from the rela-

tively simple (e.g., comparing blood pressure to a recommended value) to the complex (e.g., predictive medical diagnoses using ‘data mining’ techniques). In some cases data may be ‘fit’ using algorithms such as a linear or non-linear least-squares fitting algorithm.

In one embodiment, a physician, other health care practitioner, or emergency personnel is provided with access to patient medical information through the server 200. In one embodiment, if the wearable appliance detects that the patient needs help, or if the patient decides help is needed, the system can call his or her primary care physician. If the patient is unable to access his or her primary care physician (or another practicing physician providing care to the patient) a call from the patient is received, by an answering service or a call center associated with the patient or with the practicing physician. The call center determines whether the patient is exhibiting symptoms of an emergency condition by polling vital patient information generated by the wearable device, and if so, the answering service contacts 911 emergency service or some other emergency service. The call center can review falls information, blood pressure information, and other vital information to determine if the patient is in need of emergency assistance. If it is determined that the patient is not exhibiting symptoms of an emergent condition, the answering service may then determine if the patient is exhibiting symptoms of a non-urgent condition. If the patient is exhibiting symptoms of a non-urgent condition, the answering service will inform the patient that he or she may log into the server 200 for immediate information on treatment of the condition. If the answering service determines that the patient is exhibiting symptoms that are not related to a non-urgent condition, the answering service may refer the patient to an emergency room, a clinic, the practicing physician (when the practicing physician is available) for treatment.

In another embodiment, the wearable appliance permits direct access to the call center when the user pushes a switch or button on the appliance, for instance. In one implementation, telephones and switching systems in call centers are integrated with the home mesh network to provide for, among other things, better routing of telephone calls, faster delivery of telephone calls and associated information, and improved service with regard to client satisfaction through computer-telephony integration (CTI). CTI implementations of various design and purpose are implemented both within individual call-centers and, in some cases, at the telephone network level. For example, processors running CTI software applications may be linked to telephone switches, service control points (SCPs), and network entry points within a public or private telephone network. At the call-center level, CTI-enhanced processors, data servers, transaction servers, and the like, are linked to telephone switches and, in some cases, to similar CTI hardware at the network level, often by a dedicated digital link. CTI processors and other hardware within a call-center is commonly referred to as customer premises equipment (CPE). It is the CTI processor and application software in such centers that provides computer enhancement to a call center. In a CTI-enhanced call center, telephones at agent stations are connected to a central telephony switching apparatus, such as an automatic call distributor (ACD) switch or a private branch exchange (PBX). The agent stations may also be equipped with computer terminals such as personal computer/video display unit’s (PC/VDU’s) so that agents manning such stations may have access to stored data as well as being linked to incoming callers by telephone equipment. Such stations may be interconnected through the PC/VDUs by a local area network (LAN). One or more data or transaction servers may also be connected to the LAN that intercon-

nects agent stations. The LAN is, in turn, typically connected to the CTI processor, which is connected to the call switching apparatus of the call center.

When a call from a patient arrives at a call center, whether or not the call has been pre-processed at an SCP, the telephone number of the calling line and the medical record are made available to the receiving switch at the call center by the network provider. This service is available by most networks as caller-ID information in one of several formats such as Automatic Number Identification (ANI). Typically the number called is also available through a service such as Dialed Number Identification Service (DNIS). If the call center is computer-enhanced (CTI), the phone number of the calling party may be used as a key to access additional medical and/or historical information from a customer information system (CIS) database at a server on the network that connects the agent workstations. In this manner information pertinent to a call may be provided to an agent, often as a screen pop on the agent's PC/VDU.

The call center enables any of a first plurality of physician or health care practitioner terminals to be in audio communication over the network with any of a second plurality of patient wearable appliances. The call center will route the call to a physician or other health care practitioner at a physician or health care practitioner terminal and information related to the patient (such as an electronic medical record) will be received at the physician or health care practitioner terminal via the network. The information may be forwarded via a computer or database in the practicing physician's office or by a computer or database associated with the practicing physician, a health care management system or other health care facility or an insurance provider. The physician or health care practitioner is then permitted to assess the patient, to treat the patient accordingly, and to forward updated information related to the patient (such as examination, treatment and prescription details related to the patient's visit to the patient terminal) to the practicing physician via the network 200.

In one embodiment, the system informs a patient of a practicing physician of the availability of the web services and referring the patient to the web site upon agreement of the patient. A call from the patient is received at a call center. The call center enables physicians to be in audio communication over the network with any patient wearable appliances, and the call is routed to an available physician at one of the physician so that the available physician may carry on a two-way conversation with the patient. The available physician is permitted to make an assessment of the patient and to treat the patient. The system can forward information related to the patient to a health care management system associated with the physician. The health care management system may be a healthcare management organization, a point of service health care system, or a preferred provider organization. The health care practitioner may be a nurse practitioner or an internist.

The available health care practitioner can make an assessment of the patient and to conduct an examination of the patient over the network, including optionally by a visual study of the patient. The system can make an assessment in accordance with a protocol. The assessment can be made in accordance with a protocol stored in a database and/or making an assessment in accordance with the protocol may include displaying in real time a relevant segment of the protocol to the available physician. Similarly, permitting the physician to prescribe a treatment may include permitting the physician to refer the patient to a third party for treatment and/or referring the patient to a third party for treatment may include referring the patient to one or more of a primary care

physician, specialist, hospital, emergency room, ambulance service or clinic. Referring the patient to a third party may additionally include communicating with the third party via an electronic link included in a relevant segment of a protocol stored in a protocol database resident on a digital storage medium and the electronic link may be a hypertext link. When a treatment is being prescribed by a physician, the system can communicate a prescription over the network to a pharmacy and/or communicating the prescription over the network to the pharmacy may include communicating to the pharmacy instructions to be given to the patient pertaining to the treatment of the patient. Communicating the prescription over the network to the pharmacy may also include communicating the prescription to the pharmacy via a hypertext link included in a relevant segment of a protocol stored in a database resident on a digital storage medium. In accordance with another related embodiment, permitting the physician to conduct the examination may be accomplished under conditions such that the examination is conducted without medical instruments at the patient terminal where the patient is located.

In another embodiment, a system for delivering medical examination, diagnosis, and treatment services from a physician to a patient over a network includes a first plurality of health care practitioners at a plurality of terminals, each of the first plurality of health care practitioner terminals including a display device that shows information collected by the wearable appliances and a second plurality of patient terminals or wearable appliances in audiovisual communication over a network with any of the first plurality of health care practitioner terminals. A call center is in communication with the patient wearable appliances and the health care practitioner terminals, the call center routing a call from a patient at one of the patient terminals to an available health care practitioner at one of the health care practitioner terminals, so that the available health care practitioner may carry on a two-way conversation with the patient. A protocol database resident on a digital storage medium is accessible to each of the health care practitioner terminals. The protocol database contains a plurality of protocol segments such that a relevant segment of the protocol may be displayed in real time on the display device of the health care practitioner terminal of the available health care practitioner for use by the available health care practitioner in making an assessment of the patient. The relevant segment of the protocol displayed in real time on the display device of the health care practitioner terminal may include an electronic link that establishes communication between the available health care practitioner and a third party and the third party may be one or more of a primary care physician, specialist, hospital, emergency room, ambulance service, clinic or pharmacy.

In accordance with other related embodiment, the patient wearable appliance may include establish a direct connection to the call center by pushing a button on the appliance. Further, the protocol database may be resident on a server that is in communication with each of the health care practitioner terminals and each of the health care practitioner terminals may include a local storage device and the protocol database is replicated on the local storage device of one or more of the physician terminals.

In another embodiment, a system for delivering medical examination, diagnosis, and treatment services from a physician to a patient over a network includes a first plurality of health care practitioner terminals, each of the first plurality of health care practitioner terminals including a display device and a second plurality of patient terminals in audiovisual communication over a network with any of the first plurality of health care practitioner terminals. Each of the second plu-

reality of patient terminals includes a camera having pan, tilt and zoom modes, such modes being controlled from the first plurality of health care practitioner terminals. A call center is in communication with the patient terminals and the health care practitioner terminals and the call center routes a call from a patient at one of the patient terminals to an available health care practitioner at one of the health care practitioner terminals, so that the available health care practitioner may carry on a two-way conversation with the patient and visually observe the patient.

In one embodiment, the information is store in a secure environment, with security levels equal to those of online banking, social security number input, and other confidential information. Conforming to Health Insurance Portability and Accountability Act (HIPAA) requirements, the system creates audit trails, requires logins and passwords, and provides data encryption to ensure the patient information is private and secure. The HIPAA privacy regulations ensure a national floor of privacy protections for patients by limiting the ways that health plans, pharmacies, hospitals and other covered entities can use patients' personal medical information. The regulations protect medical records and other individually identifiable health information, whether it is on paper, in computers or communicated orally.

Due to its awareness of the patient's position, the server **200** can optionally control a mobility assistance device such as a smart cane or robot. The robotic smart cane sends video from its camera to the server **20**, which in turn coordinates the position of the robot, as determined by the cameras **10** mounted in the home as well as the robot camera. The robot position, as determined by the server **20**, is then transmitted to the robot for navigation. The robot has a frame with an extended handle. The handle includes handle sensors mounted thereon to detect the force places on each handle to receive as input the movement desired by the patient. In one embodiment, the robot has a control navigation system that accepts patient command as well as robot self-guidance command. The mobility is a result of give-and-take between the patient's self-propulsion and the walker's automated reactions. Thus, when the patient moves the handle to the right, the robot determines that the patient is interested in turning and actuates the drive systems appropriately. However, if the patient is turning into an obstacle, as determined by the cameras and the server **20**, the drive system provides gentle resistance that tells the patient of an impending collision.

If, for example, a patient does not see a coffee table ahead, the walker will detect it, override the patient's steering to avoid it, and thereby prevent a possible fall. Onboard software processes the data from 180 degrees of approaching terrain and steers the front wheel toward openings and away from obstacles.

The control module executes software that enables the robot to move around its environment safely. The software performs localization, mapping, path planning and obstacle avoidance. In one embodiment, images from a plurality of wall-mounted cameras **10** are transmitted to the server **20**. The server **20** collects images of the robot and triangulates the robot position by cross-referencing the images. The information is then correlated with the image from the robot-mounted camera and optical encoders that count the wheel rotations to calculate traveled distance for range measurement. In this process, a visual map of unique "landmarks" created as the robot moves along its path is annotated with the robot's position to indicate the position estimate of the landmark. The current image, seen from the robot, is compared with the images in the database to find matching landmarks. Such matches are used to update the position of the robot according

to the relative position of the matching landmark. By repeatedly updating the position of landmarks based on new data, the software incrementally improves the map by calculating more accurate estimates for the position of the landmarks. An improved map results in more accurate robot position estimates. Better position estimates contribute to better estimates for the landmark positions and so on. If the environment changes so much that the robot no longer recognizes previous landmarks, the robot automatically updates the map with new landmarks. Outdated landmarks that are no longer recognized can easily be deleted from the map by simply determining if they were seen or matched when expected.

Using the obstacle avoidance algorithm, the robot generates corrective movements to avoid obstacles not represented in the path planner such as open/closed doors, furniture, people, and more. The robot rapidly detects obstacles using its sensors and controls its speed and heading to avoid obstacles.

The hazard avoidance mechanisms provide a reflexive response to hazardous situations to insure the robot's safety and guarantee that it does not damage itself or the environment. Mechanisms for hazard avoidance include collision detection using not one but a complementary set of sensors and techniques. For instance, collision avoidance can be provided using contact sensing, motor load sensing, and vision. The combination of multiple sources for collision detection guarantees safe collision avoidance. Collision detection provides a last resort for negotiating obstacles in case obstacle avoidance fails to do so in the first place, which can be caused by moving objects or software and hardware failures.

If the walker is in motion (as determined by the wheel encoder), the force applied to the brake pads is inversely proportional to the distance to obstacles. If the walker is stopped, the brakes should be fully applied to provide a stable base on which the patient can rest. When the walker is stopped and the patient wishes to move again, the brakes should come off slowly to prevent the walker from lurching forward.

The walker should mostly follow the patient's commands, as this is crucial for patient acceptance. For the safety braking and the safety braking and steering control systems, the control system only influences the motion when obstacles or cliffs are near the patient. In other words, the walker is, typically, fully patient controlled. For all other situations, the control system submits to the patient's desire. This does not mean that the control system shuts down, or does not provide the usual safety features. In fact, all of the control systems fall back on their emergency braking to keep the patient safe. When the control system has had to brake to avoid an obstacle or has given up trying to lead the patient on a particular path, the patient must disengage the brakes (via a pushbutton) or re-engage the path following (again via a pushbutton) to regain control or allow collaboration again. This lets the patient select the walker's mode manually when they disagree with the control system's choices.

FIG. **5** shows an exemplary process to monitor patient. First, the process sets up mesh network appliances (**1000**). Next, the process determines patient position using in-door positioning system (**1002**). The process then determines patient movement using accelerometer output (**1004**). Sharp accelerations may be used to indicate fall. Further, the z axis accelerometer changes can indicate the height of the appliance from the floor and if the height is near zero, the system infers that the patient had fallen. The system can also determine vital parameter including patient heart rate (**1006**). The system determines if patient needs assistance based on in-door position, fall detection and vital parameter (**1008**). If a fall is suspected, the system confirms the fall by communi-

cating with the patient prior to calling a third party such as the patient's physician, nurse, family member, 911, 511, 411, or a paid call center to get assistance for the patient (1010). If confirmed or if the patient is non-responsive, the system contacts the third party and sends voice over mesh network to appliance on the patient to allow one or more third parties to talk with the patient (1012). If needed, the system calls and/or conferences emergency personnel into the call (1014).

In one embodiment, if the patient is outside of the mesh network range such as when the user is traveling away from his/her home, the system continuously records information into memory until the home mesh network is reached or until the monitoring appliance reaches an internet access point. While the wearable appliance is outside of the mesh network range, the device searches for a cell phone with an expansion card plugged into a cell phone expansion slot such as the SDIO slot. If the wearable appliance detects a cell phone that is mesh network compatible, the wearable appliance communicates with the cell phone and provides information to the server 200 using the cellular connection. In one embodiment, a Zigbee SDIO card from C-guys, Inc., enables device-to-device communications for PDAs and smart phones. C-guys' ZigBee SDIO card includes the company's CG-100 SDIO application interface controller, which is designed to convert an application signal to an SD signal (or vice versa). The ZigBee card can provide signal ranges of up to 10 m in the 2.4 GHz band and data rates of up to 200 kbps. The card has peer-to-peer communications mode and supports direct application to PDAs or any SD supported hand-held cell phones. In this embodiment, the PDA or cell phone can provide a GPS position information instead of the indoor position information generated by the mesh network appliances 8. The cell phone GPS position information, accelerometer information and vital information such as heart rate information is transmitted using the cellular channel to the server 200 for processing as is normal. In another embodiment where the phone works through WiFi (802.11) or WiMAX (802.16) or ultra-wideband protocol instead of the cellular protocol, the wearable appliance can communicate over these protocols using a suitable mesh network interface to the phone. In instances where the wearable appliance is outside of its home base and a dangerous condition such as a fall is detected, the wearable appliance can initiate a distress call to the authorized third party using cellular, WiFi, WiMAX, or UWB protocols as is available.

FIG. 6A shows a portable embodiment of the present invention where the voice recognizer is housed in a wrist-watch. As shown in FIG. 6, the device includes a wrist-watch sized case 1380 supported on a wrist band 1374. The case 1380 may be of a number of variations of shape but can be conveniently made a rectangular, approaching a box-like configuration. The wrist-band 1374 can be an expansion band or a wristwatch strap of plastic, leather or woven material. The processor or CPU of the wearable appliance is connected to a radio frequency (RF) transmitter/receiver (such as a Bluetooth device, a Zigbee device, a WiFi device, a WiMAX device, or an 802.X transceiver, among others).

In one embodiment, the back of the device is a conductive metal electrode 1381 that in conjunction with a second electrode 1383 mounted on the wrist band 1374, enables differential EKG or ECG to be measured. The electrical signal derived from the electrodes is typically 1 mV peak-peak. In one embodiment where only one electrode 1381 or 1383 is available, an amplification of about 1000 is necessary to render this signal usable for heart rate detection. In the embodiment with electrodes 1381 and 1383 available, a differential amplifier is used to take advantage of the identical

common mode signals from the EKG contact points, the common mode noise is automatically cancelled out using a matched differential amplifier. In one embodiment, the differential amplifier is a Texas Instruments INA321 instrumentation amplifier that has matched and balanced integrated gain resistors. This device is specified to operate with a minimum of 2.7V single rail power supply. The INA321 provides a fixed amplification of 5× for the EKG signal. With its CMRR specification of 94 dB extended up to 3 KHz the INA321 rejects the common mode noise signals including the line frequency and its harmonics. The quiescent current of the INA321 is 40 mA and the shut down mode current is less than 1 mA. The amplified EKG signal is internally fed to the on chip analog to digital converter. The ADC samples the EKG signal with a sampling frequency of 512 Hz. Precise sampling period is achieved by triggering the ADC conversions with a timer that is clocked from a 32.768 kHz low frequency crystal oscillator. The sampled EKG waveform contains some amount of super imposed line frequency content. This line frequency noise is removed by digitally filtering the samples. In one implementation, a 17-tap low pass FIR filter with pass band upper frequency of 6 Hz and stop band lower frequency of 30 Hz is implemented in this application. The filter coefficients are scaled to compensate the filter attenuation and provide additional gain for the EKG signal at the filter output. This adds up to a total amplification factor of greater than 1000× for the EKG signal.

The wrist band 1374 can also contain other electrical devices such as ultrasound transducer, optical transducer or electromagnetic sensors, among others. In one embodiment, the transducer is an ultrasonic transducer that generates and transmits an acoustic wave upon command from the CPU during one period and listens to the echo returns during a subsequent period. In use, the transmitted bursts of sonic energy are scattered by red blood cells flowing through the subject's radial artery, and a portion of the scattered energy is directed back toward the ultrasonic transducer 84. The time required for the return energy to reach the ultrasonic transducer varies according to the speed of sound in the tissue and according to the depth of the artery. Typical transit times are in the range of 6 to 7 microseconds. The ultrasonic transducer is used to receive the reflected ultrasound energy during the dead times between the successive transmitted bursts. The frequency of the ultrasonic transducer's transmit signal will differ from that of the return signal, because the scattering red blood cells within the radial artery are moving. Thus, the return signal, effectively, is frequency modulated by the blood flow velocity.

A driving and receiving circuit generates electrical pulses which, when applied to the transducer, produce acoustic energy having a frequency on the order of 8 MHz, a pulse width or duration of approximately 8 microseconds, and a pulse repetition interval (PRI) of approximately 16 μs, although other values of frequency, pulse width, and PRI may be used. In one embodiment, the transducer 84 emits an 8 microsecond pulse, which is followed by an 8 microsecond "listen" period, every 16 microseconds. The echoes from these pulses are received by the ultrasonic transducer 84 during the listen period. The ultrasonic transducer can be a ceramic piezoelectric device of the type well known in the art, although other types may be substituted.

An analog signal representative of the Doppler frequency of the echo is received by the transducer and converted to a digital representation by the ADC, and supplied to the CPU for signal processing. Within the CPU, the digitized Doppler frequency is scaled to compute the blood flow velocity within the artery based on the Doppler frequency. Based on the real

time the blood flow velocity, the CPU applies the vital model to the corresponding blood flow velocity to produce the estimated blood pressure value.

Prior to operation, calibration is done using a calibration device and the monitoring device to simultaneously collect blood pressure values (systolic, diastolic pressures) and a corresponding blood flow velocity generated by the monitoring device. The calibration device is attached to the base station and measures systolic and diastolic blood pressure using a cuff-based blood pressure monitoring device that includes a motor-controlled pump and data-processing electronics. While the cuff-based blood pressure monitoring device collects patient data, the transducer collects patient data in parallel and through the watch's radio transmitter, blood flow velocity is sent to the base station for generating a computer model that converts the blood flow velocity information into systolic and diastolic blood pressure values and this information is sent wirelessly from the base station to the watch for display and to a remote server if needed. This process is repeated at a later time (e.g., 15 minutes later) to collect a second set of calibration parameters. In one embodiment, the computer model fits the blood flow velocity to the systolic/diastolic values. In another embodiment, the computer trains a neural network or HMM to recognize the systolic and diastolic blood pressure values.

After the computer model has been generated, the system is ready for real-time blood pressure monitoring. In an acoustic embodiment, the transducer directs ultrasound at the patient's artery and subsequently listens to the echos therefrom. The echoes are used to determine blood flow, which is fed to the computer model to generate the systolic and diastolic pressure values as well as heart rate value. The CPU's output signal is then converted to a form useful to the user such as a digital or analog display, computer data file, or audible indicator. The output signal can drive a speaker to enable an operator to hear a representation of the Doppler signals and thereby to determine when the transducer is located approximately over the radial artery. The output signal can also be wirelessly sent to a base station for subsequent analysis by a physician, nurse, caregiver, or treating professional. The output signal can also be analyzed for medical attention and medical treatment.

It is noted that while the above embodiment utilizes a preselected pulse duration of 8 microseconds and pulse repetition interval of 16 microseconds, other acoustic sampling techniques may be used in conjunction with the invention. For example, in a second embodiment of the ultrasonic driver and receiver circuit (not shown), the acoustic pulses are range-gated with a more complex implementation of the gate logic. As is well known in the signal processing arts, range-gating is a technique by which the pulse-to-pulse interval is varied based on the receipt of range information from earlier emitted and reflected pulses. Using this technique, the system may be "tuned" to receive echoes falling within a specific temporal window which is chosen based on the range of the echo-producing entity in relation to the acoustic source. The delay time before the gate is turned on determines the depth of the sample volume. The amount of time the gate is activated establishes the axial length of the sample volume. Thus, as the acoustic source (in this case the ultrasonic transducer 84) is tuned to the echo-producing entity (red blood cells, or arterial walls), the pulse repetition interval is shortened such that the system may obtain more samples per unit time, thereby increasing its resolution. It will be recognized that other acoustic processing techniques may also be used, all of which are considered to be equivalent.

In one optical embodiment, the transducer can be an optical transducer. The optical transducer can be a light source and a photo-detector embedded in the wrist band portions 1374. The light source can be light-emitting diodes that generate red ($\lambda 630$ nm) and infrared ($\lambda 900$ nm) radiation, for example. The light source and the photo-detector are slidably adjustable and can be moved along the wrist band to optimize beam transmission and pick up. As the heart pumps blood through the patient's finger, blood cells absorb and transmit varying amounts of the red and infrared radiation depending on how much oxygen binds to the cells' hemoglobin. The photo-detector detects transmission at the predetermined wavelengths, for example red and infrared wavelengths, and provides the detected transmission to a pulse-oximetry circuit embedded within the wrist-watch. The output of the pulse-oximetry circuit is digitized into a time-dependent optical waveform, which is then sent back to the pulse-oximetry circuit and analyzed to determine the user's vital signs.

In the electromagnetic sensor embodiment, the wrist band 1374 is a flexible plastic material incorporated with a flexible magnet. The magnet provides a magnetic field, and one or more electrodes similar to electrode 1383 are positioned on the wrist band to measure voltage drops which are proportional to the blood velocity. The electromagnetic embodiment may be mounted on the upper arm of the patient, on the ankle or on the neck where peripheral blood vessels pass through and their blood velocity may be measured with minimal interruptions. The flexible magnet produces a pseudo-uniform (non-gradient) magnetic field. The magnetic field can be normal to the blood flow direction when wrist band 1374 is mounted on the user's wrist or may be a rotative pseudo-uniform magnetic field so that the magnetic field is in a transversal direction in respect to the blood flow direction. The electrode output signals are processed to obtain a differential measurement enhancing the signal to noise ratio. The flow information is derived based on the periodicity of the signals. The decoded signal is filtered over several periods and then analyzed for changes used to estimate artery and vein blood flow. Systemic stroke volume and cardiac output may be calculated from the peripheral SV index value.

The wrist-band 1374 further contains an antenna 1376 for transmitting or receiving radio frequency signals. The wrist-band 1374 and the antenna 1376 inside the band are mechanically coupled to the top and bottom sides of the wrist-watch housing 1380. Further, the antenna 1376 is electrically coupled to a radio frequency transmitter and receiver for wireless communications with another computer or another user. Although a wrist-band is disclosed, a number of substitutes may be used, including a belt, a ring holder, a brace, or a bracelet, among other suitable substitutes known to one skilled in the art. The housing 1380 contains the processor and associated peripherals to provide the human-machine interface. A display 1382 is located on the front section of the housing 1380. A speaker 1384, a microphone 1388, and a plurality of push-button switches 1386 and 1390 are also located on the front section of housing 1380.

The electronic circuitry housed in the watch case 1380 detects adverse conditions such as falls or seizures. In one implementation, the circuitry can recognize speech, namely utterances of spoken words by the user, and converting the utterances into digital signals. The circuitry for detecting and processing speech to be sent from the wristwatch to the base station 20 over the mesh network includes a central processing unit (CPU) connected to a ROM/RAM memory via a bus. The CPU is a preferably low power 16-bit or 32-bit microprocessor and the memory is preferably a high density, low-power RAM. The CPU is coupled via the bus to processor

wake-up logic, one or more accelerometers to detect sudden movement in a patient, an ADC **102** which receives speech input from the microphone. The ADC converts the analog signal produced by the microphone into a sequence of digital values representing the amplitude of the signal produced by the microphone at a sequence of evenly spaced times. The CPU is also coupled to a digital to analog (D/A) converter, which drives the speaker to communicate with the user. Speech signals from the microphone are first amplified, pass through an antialiasing filter before being sampled. The front-end processing includes an amplifier, a bandpass filter to avoid antialiasing, and an analog-to-digital (A/D) converter or a CODEC. To minimize space, the ADC, the DAC and the interface for wireless transceiver and switches may be integrated into one integrated circuit to save space. In one embodiment, the wrist watch acts as a walkie-talkie so that voice is received over the mesh network by the base station **20** and then delivered to a call center over the POTS or PSTN network. In another embodiment, voice is provided to the call center using the Internet through suitable VOIP techniques. In one embodiment, speech recognition such as a speech recognizer is discussed in U.S. Pat. No. 6,070,140 by the inventor of the instant invention, the content of which is incorporated by reference.

FIG. **6B** shows an exemplary mesh network working with the wearable appliance of FIG. **6A**. Data collected and communicated on the display **1382** of the watch as well as voice is transmitted to a base station **1390** for communicating over a network to an authorized party **1394**. The watch and the base station is part of a mesh network that may communicate with a medicine cabinet to detect opening or to each medicine container **1391** to detect medication compliance. Other devices include mesh network thermometers, scales, or exercise devices. The mesh network also includes a plurality of home/room appliances **1392-1399**. The ability to transmit voice is useful in the case the patient has fallen down and cannot walk to the base station **1390** to request help. Hence, in one embodiment, the watch captures voice from the user and transmits the voice over the Zigbee mesh network to the base station **1390**. The base station **1390** in turn dials out to an authorized third party to allow voice communication and at the same time transmits the collected patient vital parameter data and identifying information so that help can be dispatched quickly, efficiently and error-free. In one embodiment, the base station **1390** is a POTS telephone base station connected to the wired phone network. In a second embodiment, the base station **1390** can be a cellular telephone connected to a cellular network for voice and data transmission. In a third embodiment, the base station **1390** can be a WiMAX or 802.16 standard base station that can communicate VOIP and data over a wide area network. Alternatively, the base station can communicate over POTS and a wireless network such as cellular or WiMAX or both.

In one embodiment, the processor and transceiver on the watch and the base station conform to the Zigbee protocol. ZigBee is a cost-effective, standards-based wireless networking solution that supports low data-rates, low-power consumption, security, and reliability. Single chip Zigbee controllers with wireless transceivers built-in include the Chipcon/Ember CC2420: Single-chip 802.15.4 radio transceiver and the FreeScale single chip Zigbee and microcontroller. In various embodiments, the processor communicates with a Z axis accelerometer measures the patient's up and down motion and/or an X and Y axis accelerometer measures the patient's forward and side movements. In one embodiment, EKG and/or blood pressure parameters can be captured

by the processor. The controllers upload the captured data when the memory is full or while in wireless contact with other Zigbee nodes.

The wristwatch device can also be used to control home automation. The user can have flexible management of lighting, heating and cooling systems from anywhere in the home. The watch automates control of multiple home systems to improve conservation, convenience and safety. The watch can capture highly detailed electric, water and gas utility usage data and embed intelligence to optimize consumption of natural resources. The system is convenient in that it can be installed, upgraded and networked without wires. The patient can receive automatic notification upon detection of unusual events in his or her home. For example, if smoke or carbon monoxide detectors detect a problem, the wrist-watch can buzz or vibrate to alert the user and the central hub triggers selected lights to illuminate the safest exit route.

In another embodiment, the watch serves a key fob allowing the user to wirelessly unlock doors controlled by Zigbee wireless receiver. In this embodiment, when the user is within range, the door Zigbee transceiver receives a request to unlock the door, and the Zigbee transceiver on the door transmits an authentication request using suitable security mechanism. Upon entry, the Zigbee doorlock device sends access signals to the lighting, air-conditioning and entertainment systems, among others. The lights and temperature are automatically set to pre-programmed preferences when the user's presence is detected.

Although Zigbee is mentioned as an exemplary protocol, other protocols such as UWB, Bluetooth, WiFi and WiMAX can be used as well.

While the foregoing addresses the needs of the elderly, the system can assist infants as well. Much attention has been given to ways to reduce a risk of dying from Sudden Infant Death Syndrome (SIDS), an affliction which threatens infants who have died in their sleep for heretofore unknown reasons. Many different explanations for this syndrome and ways to prevent the syndrome are found in the literature. It is thought that infants which sleep on their backs may be at risk of death because of the danger of formula regurgitation and liquid aspiration into the lungs. It has been thought that infants of six (6) months or less do not have the motor skills or body muscular development to regulate movements responsive to correcting breathing problems that may occur during sleep.

In an exemplary system to detect and minimize SIDS problem in an infant patient, a diaper pad is used to hold an array of integrated sensors and the pad can be placed over a diaper, clothing, or blanket. The integrated sensors can provide data for measuring position, temperature, sound, vibration, movement, and optionally other physical properties through additional sensors. Each pad can have sensors that provide one or more of the above data. The sensors can be added or removed as necessary depending on the type of data being collected.

The sensor should be water proof and disposable. The sensor can be switch on/off locally or remotely. The sensor can be removable or clip on easily. The sensor can store or beam out information for analysis purpose, e.g. store body temperature every 5 seconds. The sensor can be turn-on for other purposed, e.g. diaper wet, it will beep and allow a baby care provider to take care of the business in time. The array of sensors can be self selective, e.g., when one sensor can detect strong heart beat, it will turn off others to do so.

The sensor can be used for drug delivery system, e.g. when patient has abdomen pain, soothing drug can be applied, based on the level of pain the sensor detects, different dose of drugs will be applied.

The array of sensors may allow the selection and analysis of zones of sensors in the areas of interest such as the abdomen area. Each sensor array has a low spatial resolution: approximately 10 cm between each sensor. In addition to lower cost due to the low number of sensors, it is also possible to modify the data collection rate from certain sensors that are providing high-quality data. Other sensors may include those worn on the body, such as in watch bands, finger rings, or adhesive sensors, but telemetry, not wires, would be used to communicate with the controller.

The sensor can be passive device such as a reader, which mounted near the crib can active it from time to time. In any emergency situation, the sensor automatically signals a different state which the reader can detect.

The sensor can be active and powered by body motion or body heat. The sensor can detect low battery situation and warn the user to provide a replacement battery. In one embodiment, a plurality of sensors attached to the infant collects the vital parameters. For example, the sensors can be attached to the infant's clothing (shirt or pant), diaper, undergarment or bed sheet, bed linen, or bed spread.

The patient may wear one or more sensors, for example devices for sensing ECG, EKG, blood pressure, sugar level, weight, temperature and pressure, among others. In one embodiment, an optical temperature sensor can be used. In another embodiment, a temperature thermistor can be used to sense patient temperature. In another embodiment, a fat scale sensor can be used to detect the patient's fat content. In yet another embodiment, a pressure sensor such as a MEMS sensor can be used to sense pressure on the patient.

In one embodiment, the sensors are mounted on the patient's wrist (such as a wristwatch sensor) and other convenient anatomical locations. Exemplary sensors include standard medical diagnostics for detecting the body's electrical signals emanating from muscles (EMG and EOG) and brain (EEG) and cardiovascular system (ECG). Leg sensors can include piezoelectric accelerometers designed to give qualitative assessment of limb movement. Additionally, thoracic and abdominal bands used to measure expansion and contraction of the thorax and abdomen respectively. A small sensor can be mounted on the subject's finger in order to detect blood-oxygen levels and pulse rate. Additionally, a microphone can be attached to throat and used in sleep diagnostic recordings for detecting breathing and other noise. One or more position sensors can be used for detecting orientation of body (lying on left side, right side or back) during sleep diagnostic recordings. Each of sensors can individually transmit data to the server 20 using wired or wireless transmission. Alternatively, all sensors can be fed through a common bus into a single transceiver for wired or wireless transmission. The transmission can be done using a magnetic medium such as a floppy disk or a flash memory card, or can be done using infrared or radio network link, among others.

In one embodiment, the sensors for monitoring vital signs are enclosed in a wrist-watch sized case supported on a wrist band. The sensors can be attached to the back of the case. For example, in one embodiment, Cygnus' AutoSensor (Redwood City, Calif.) is used as a glucose sensor. A low electric current pulls glucose through the skin. Glucose is accumulated in two gel collection discs in the AutoSensor. The AutoSensor measures the glucose and a reading is displayed by the watch.

In another embodiment, EKG/ECG contact points are positioned on the back of the wrist-watch case. In yet another embodiment that provides continuous, beat-to-beat wrist arterial pulse rate measurements, a pressure sensor is housed in a casing with a 'free-floating' plunger as the sensor applan-

ates the radial artery. A strap provides a constant force for effective applanation and ensuring the position of the sensor housing to remain constant after any wrist movements. The change in the electrical signals due to change in pressure is detected as a result of the piezoresistive nature of the sensor are then analyzed to arrive at various arterial pressure, systolic pressure, diastolic pressure, time indices, and other blood pressure parameters.

The heartbeat detector can be one of: EKG detector, ECG detector, optical detector, ultrasonic detector, or microphone/digital stethoscope for picking up heart sound. In one embodiment, one EKG/ECG contact point is provided on the back of the wrist watch case and one or more EKG/ECG contact points are provided on the surface of the watch so that when a user's finger or skin touches the contact points, an electrical signal indicative of heartbeat activity is generated. An electrocardiogram (ECG) or EKG is a graphic tracing of the voltage generated by the cardiac or heart muscle during a heartbeat. It provides very accurate evaluation of the performance of the heart. The heart generates an electrochemical impulse that spreads out in the heart in such a fashion as to cause the cells to contract and relax in a timely order and thus give the heart a pumping characteristic. This sequence is initiated by a group of nerve cells called the sinoatrial (SA) node resulting in a polarization and depolarization of the cells of the heart. Because this action is electrical in nature and because the body is conductive with its fluid content, this electrochemical action can be measured at the surface of the body. An actual voltage potential of approximately 1 mV develops between various body points. This can be measured by placing electrode contacts on the body. The four extremities and the chest wall have become standard sites for applying the electrodes. Standardizing electrocardiograms makes it possible to compare them as taken from person to person and from time to time from the same person. The normal electrocardiogram shows typical upward and downward deflections that reflect the alternate contraction of the atria (the two upper chambers) and of the ventricles (the two lower chambers) of the heart. The voltages produced represent pressures exerted by the heart muscles in one pumping cycle. The first upward deflection, P, is due to atria contraction and is known as the atrial complex. The other deflections, Q, R, S, and T, are all due to the action of the ventricles and are known as the ventricular complexes. Any deviation from the norm in a particular electrocardiogram is indicative of a possible heart disorder.

The CPU measures the time duration between the sequential pulses and converts each such measurement into a corresponding timing measurement indicative of heart rate. The CPU also processes a predetermined number of most recently occurring timing measurements in a prescribed fashion, to produce an estimate of heartbeat rate for display on a display device on the watch and/or for transmission over the wireless network. This estimate is updated with the occurrence of each successive pulse.

In one embodiment, the CPU produces the estimate of heartbeat rate by first averaging a plurality of measurements, then adjusting the particular one of the measurements that differs most from the average to be equal to that average, and finally computing an adjusted average based on the adjusted set of measurements. The process may repeat the foregoing operations a number of times so that the estimate of heartbeat rate is substantially unaffected by the occurrence of heartbeat artifacts.

In one EKG or ECG detector, the heartbeat detection circuitry includes a differential amplifier for amplifying the signal transmitted from the EKG/ECG electrodes and for

converting it into single-ended form, and a bandpass filter and a 60 Hz notch filter for removing background noise. The CPU measures the time durations between the successive pulses and estimates the heartbeat rate. The time durations between the successive pulses of the pulse sequence signal provides an estimate of heartbeat rate. Each time duration measurement is first converted to a corresponding rate, preferably expressed in beats per minute (bpm), and then stored in a file, taking the place of the earliest measurement previously stored. After a new measurement is entered into the file, the stored measurements are averaged, to produce an average rate measurement. The CPU optionally determines which of the stored measurements differs most from the average, and replaces that measurement with the average.

Upon initiation, the CPU increments a period timer used in measuring the time duration between successive pulses. This timer is incremented in steps of about two milliseconds in one embodiment. It is then determined whether or not a pulse has occurred during the previous two milliseconds. If it has not, the CPU returns to the initial step of incrementing the period timer. If a heartbeat has occurred, on the other hand, the CPU converts the time duration measurement currently stored in the period timer to a corresponding heartbeat rate, preferably expressed in bpm. After the heartbeat rate measurement is computed, the CPU determines whether or not the computed rate is intermediate prescribed thresholds of 20 bpm and 240 bpm. If it is not, it is assumed that the detected pulse was not in fact a heartbeat and the period timer is cleared.

In an optical heartbeat detector embodiment, an optical transducer is positioned on a finger, wrist, or ear lobe. The ear, wrist or finger pulse oximeter waveform is then analyzed to extract the beat-to-beat amplitude, area, and width (half height) measurements. The oximeter waveform is used to generate heartbeat rate in this embodiment. In one implementation, a reflective sensor such as the Honeywell HLC1395 can be used. The device emits lights from a window in the infrared spectrum and receives reflected light in a second window. When the heart beats, blood flow increases temporarily and more red blood cells flow through the windows, which increases the light reflected back to the detector. The light can be reflected, refracted, scattered, and absorbed by one or more detectors. Suitable noise reduction is done, and the resulting optical waveform is captured by the CPU.

In another optical embodiment, blood pressure is estimated from the optical reading using a mathematical model such as a linear correlation with a known blood pressure reading. In this embodiment, the pulse oximeter readings are compared to the blood-pressure readings from a known working blood pressure measurement device during calibration. Using these measurements the linear equation is developed relating oximeter output waveform such as width to blood-pressure (systolic, mean and pulse pressure). In one embodiment, a transform (such as a Fourier analysis or a Wavelet transform) of the oximeter output can be used to generate a model to relate the oximeter output waveform to the blood pressure. Other non-linear math model or relationship can be determined to relate the oximeter waveform to the blood pressure.

In one implementation, the pulse oximeter probe and a blood pressure cuff are placed on the corresponding contralateral limb to the oscillometric (Dinamap 8100; Critikon, Inc, Tampa, Fla., USA) cuff site. The pulse oximeter captures data on plethysmographic waveform, heart rate, and oxygen saturation. Simultaneous blood pressure measurements were obtained from the oscillometric device, and the pulse oximeter. Systolic, diastolic, and mean blood pressures are recorded from the oscillometric device. This information is used derive calibration parameters relating the pulse oximeter output to

the expected blood pressure. During real time operation, the calibration parameters are applied to the oximeter output to predict blood pressure in a continuous or in a periodic fashion. In yet another embodiment, the device includes an accelerometer or alternative motion-detecting device to determine when the patient's hand is at rest, thereby reducing motion-related artifacts introduced to the measurement during calibration and/or operation. The accelerometer can also function as a falls detection device.

In an ultrasonic embodiment, a piezo film sensor element is placed on the wristwatch band. The sensor can be the SDT1-028K made by Measurement Specialties, Inc. The sensor should have features such as: (a) it is sensitive to low level mechanical movements, (b) it has an electrostatic shield located on both sides of the element (to minimize 50/60 Hz AC line interference), (c) it is responsive to low frequency movements in the 0.7-12 Hz range of interest. A filter/amplifier circuit has a three-pole low pass filter with a lower (-3 dB) cutoff frequency at about 12-13 Hz. The low-pass filter prevents unwanted 50/60 Hz AC line interference from entering the sensor. However, the piezo film element has a wide band frequency response so the filter also attenuates any extraneous sound waves or vibrations that get into the piezo element. The DC gain is about +30 dB.

Waveform averaging can be used to reduce noise. It reinforces the waveform of interest by minimizing the effect of any random noise. These pulses were obtained when the arm was motionless. If the arm was moved while capturing the data the waveform did not look nearly as clean. That's because motion of the arm causes the sonic vibrations to enter the piezo film through the arm or by way of the cable. An accelerometer is used to detect arm movement and used to remove inappropriate data capture.

In one embodiment that can determine blood pressure, two piezo film sensors and filter/amplifier circuits can be configured as a non-invasive velocity type blood pressure monitor. One sensor can be on the wrist and the other can be located on the inner left elbow at the same location where Korotkoff sounds are monitored during traditional blood pressure measurements with a sphygmometer. The correlation between pulse delay and blood pressure is well known in the art of non-invasive blood pressure monitors.

In yet another embodiment, an ultrasonic transducer generates and transmits an acoustic wave into the user's body such as the wrist or finger. The transducer subsequently receives pressure waves in the form of echoes resulting from the transmitted acoustic waves. In one embodiment, an ultrasonic driving and receiving circuit generates electrical pulses which, when applied to the transducer produce acoustic energy having a frequency on the order of 8 MHz, a pulse width or duration of approximately 8 microseconds, and a pulse repetition interval (PRI) of approximately 16 microseconds, although other values of frequency, pulse width, and PRI may be used. Hence, the transducer emits an 8 microsecond ultrasonic pulse, which is followed by an 8 microsecond "listen" period, every 16 microseconds. The echoes from these pulses are received by the ultrasonic transducer during the listen period. The ultrasonic transducer can be a ceramic piezoelectric device of the type well known in the art, although other types may be substituted. The transducer converts the received acoustic signal to an electrical signal, which is then supplied to the receiving section of the ultrasonic driver and receiver circuit 616, which contains two receiver circuits. The output of the first receiver circuit is an analog signal representative of the Doppler frequency of the echo received by the transducer which is digitized and supplied to the CPU. Within the CPU, the digitized Doppler frequency is

scaled to compute the blood velocity within the artery based on the Doppler frequency. The time-frequency distribution of the blood velocity is then computed. Finally, the CPU maps in time the peak of the time-frequency distribution to the corresponding pressure waveform to produce the estimated mean arterial pressure (MAP). The output of the ultrasonic receiver circuit is an analog echo signal proportional to absorption of the transmitted frequencies by blood or tissue. This analog signal is digitized and process so that each group of echoes, generated for a different transversal position, is integrated to determine a mean value. The mean echo values are compared to determine the minimum value, which is caused by direct positioning over the artery. In one embodiment, the device includes an accelerometer or alternative motion-detecting device to determine when the patient's hand is at rest, thereby reducing motion-related artifacts introduced to the measurement.

In yet another ultrasonic embodiment, a transducer includes a first and a second piezoelectric crystal, wherein the crystals are positioned at an angle to each other, and wherein the angle is determined based on the distance of the transducer to the living subject. The first piezoelectric crystal is energized by an original ultrasonic frequency signal, wherein the original ultrasonic frequency signal is reflected off the living subject and received by the second piezoelectric crystal. More specifically, the system includes a pair of piezoelectric crystals at an angle to each other, wherein the angle is determined by the depth of the object being monitored. If the object is the radial artery of a human subject (e.g., adult, infant), the angle of the two crystals with respect to the direction of the blood flow would be about 5 to about 20 degrees. One of the crystals is energized at an ultrasonic frequency. The signal is then reflected back by the user's wrist and picked up by the second crystal. The frequency received is either higher or lower than the original frequency depending upon the direction and the speed of the fluidic mass flow. For example, when blood flow is monitored, the direction of flow is fixed. Thus, the Doppler frequency which is the difference between the original and the reflected frequency depends only upon the speed of the blood flow. Ultrasonic energy is delivered to one of the two piezoelectric elements in the module by the power amplifier. The other element picks up the reflected ultrasonic signal as Doppler frequencies.

In a digital stethoscope embodiment, a microphone or a piezoelectric transducer is placed near the wrist artery to pick up heart rate information. In one embodiment, the microphone sensor and optionally the EKG sensor are placed on the wrist band 1374 of the watch to analyze the acoustic signal or signals emanating from the cardiovascular system and, optionally can combine the sound with an electric signal (EKG) emanating from the cardiovascular system and/or an acoustic signal emanating from the respiratory system. The system can perform automated auscultation of the cardiovascular system, the respiratory system, or both. For example, the system can differentiate pathological from benign heart murmurs, detect cardiovascular diseases or conditions that might otherwise escape attention, recommend that the patient go through for a diagnostic study such as an echocardiography or to a specialist, monitor the course of a disease and the effects of therapy, decide when additional therapy or intervention is necessary, and providing a more objective basis for the decision(s) made. In one embodiment, the analysis includes selecting one or more beats for analysis, wherein each beat comprises an acoustic signal emanating from the cardiovascular system; performing a time-frequency analysis of beats selected for analysis so as to provide information regarding the distribution of energy, the relative distribution

of energy, or both, over different frequency ranges at one or more points in the cardiac cycle; and processing the information to reach a clinically relevant conclusion or recommendation. In another implementation, the system selects one or more beats for analysis, wherein each beat comprises an acoustic signal emanating from the cardiovascular system; performs a time-frequency analysis of beats selected for analysis so as to provide information regarding the distribution of energy, the relative distribution of energy, or both, over different frequency ranges at one or more points in the cardiac cycle; and present information derived at least in part from the acoustic signal, wherein the information comprises one or more items selected from the group consisting of: a visual or audio presentation of a prototypical beat, a display of the time-frequency decomposition of one or more beats or prototypical beats, and a playback of the acoustic signal at a reduced rate with preservation of frequency content.

In an electromagnetic embodiment where the wrist band incorporates a flexible magnet to provide a magnetic field and one or more electrodes positioned on the wrist band to measure voltage drops which are proportional to the blood velocity, instantaneously variation of the flow can be detected but not artery flow by itself. To estimate the flow of blood in the artery, the user or an actuator such as motorized cuff temporarily stops the blood flow in the vein by applying external pressure or by any other method. During the period of time in which the vein flow is occluded, the decay of the artery flow is measured. This measurement may be used for zeroing the sensor and may be used in a model for estimating the steady artery flow. The decay in artery flow due to occlusion of veins is measured to arrive at a model the rate of artery decay. The system then estimates an average artery flow before occlusion. The blood flow can then be related to the blood pressure.

In another embodiment, an ionic flow sensor is used with a driving electrode that produces a pulsatile current. The pulsatile current causes a separation of positive and negative charges that flows in the blood of the arteries and veins passing in the wrist area. Using electrophoresis principle, the resistance of the volume surrounded by the source first decreases and then increases. The difference in resistance in the blood acts as a mark that moves according to the flow of blood so that marks are flowing in opposite directions by arteries and veins.

In the above embodiments, accelerometer information is used to detect that the patient is at rest prior to making a blood pressure measurement and estimation. Further, a temperature sensor may be incorporated so that the temperature is known at any minute. The processor correlates the temperature measurement to the blood flow measurement for calibration purposes.

In another embodiment, the automatic identification of the first, second, third and fourth heart sounds (S1, S2, S3, S4) is done. In yet another embodiment, based on the heart sound, the system analyzes the patient for mitral valve prolapse. The system performs a time-frequency analysis of an acoustic signal emanating from the subject's cardiovascular system and examines the energy content of the signal in one or more frequency bands, particularly higher frequency bands, in order to determine whether a subject suffers from mitral valve prolapse.

FIG. 7 shows an exemplary mesh network that includes the wrist-watch of FIG. 6 in communication with a mesh network including a telephone such as a wired telephone as well as a cordless telephone. In one embodiment, the mesh network is an IEEE 802.15.4 (ZigBee) network. IEEE 802.15.4 defines two device types; the reduced function device (RFD) and the full function device (FFD). In ZigBee these are referred to as

the ZigBee Physical Device types. In a ZigBee network a node can have three roles: ZigBee Coordinator, ZigBee Router, and ZigBee End Device. These are the ZigBee Logical Device types. The main responsibility of a ZigBee Coordinator is to establish a network and to define its main parameters (e.g. choosing a radio-frequency channel and defining a unique network identifier). One can extend the communication range of a network by using ZigBee Routers. These can act as relays between devices that are too far apart to communicate directly. ZigBee End Devices do not participate in routing. An FFD can talk to RFDs or other FFDs, while an RFD can talk only to an FFD. An RFD is intended for applications that are extremely simple, such as a light switch or a passive infrared sensor; they do not have the need to send large amounts of data and may only associate with a single FFD at a time. Consequently, the RFD can be implemented using minimal resources and memory capacity and have lower cost than an FFD. An FFD can be used to implement all three ZigBee Logical Device types, while an RFD can take the role as an End Device.

One embodiment supports a multicluster-multihop network assembly to enable communication among every node in a distribution of nodes. The algorithm should ensure total connectivity, given a network distribution that will allow total connectivity. One such algorithm of an embodiment is described in U.S. Pat. No. 6,832,251, the content of which is incorporated by referenced. The '251 algorithm runs on each node independently. Consequently, the algorithm does not have global knowledge of network topology, only local knowledge of its immediate neighborhood. This makes it well suited to a wide variety of applications in which the topology may be time-varying, and the number of nodes may be unknown. Initially, all nodes consider themselves remotes on cluster zero. The assembly algorithm floods one packet (called an assembly packet) throughout the network. As the packet is flooded, each node modifies it slightly to indicate what the next node should do. The assembly packet tells a node whether it is a base or a remote, and to what cluster it belongs. If a node has seen an assembly packet before, it will ignore all further assembly packets.

The algorithm starts by selecting (manually or automatically) a start node. For example, this could be the first node to wake up. This start node becomes a base on cluster 1, and floods an assembly packet to all of its neighbors, telling them to be remotes on cluster 1. These remotes in turn tell all their neighbors to be bases on cluster 2. Only nodes that have not seen an assembly packet before will respond to this request, so nodes that already have decided what to be will not change their status. The packet continues on, oscillating back and forth between "become base/become remote", and increasing the cluster number each time. Since the packet is flooded to all neighbors at every step, it will reach every node in the network. Because of the oscillating nature of the "become base/become remote" instructions, no two bases will be adjacent. The basic algorithm establishes a multi-cluster network with all gateways between clusters, but self-assembly time is proportional with the size of the network. Further, it includes only single hop clusters. Many generalizations are possible, however. If many nodes can begin the network nucleation, all that is required to harmonize the clusters is a mechanism that recognizes precedence (e.g., time of nucleation, size of sub-network), so that conflicts in boundary clusters are resolved. Multiple-hop clusters can be enabled by means of establishing new clusters from nodes that are N hops distant from the master.

Having established a network in this fashion, the masters can be optimized either based on number of neighbors, or

other criteria such as minimum energy per neighbor communication. Thus, the basic algorithm is at the heart of a number of variations that lead to a scalable multi-cluster network that establishes itself in time, and that is nearly independent of the number of nodes, with clusters arranged according to any of a wide range of optimality criteria. Network synchronism is established at the same time as the network connections, since the assembly packet(s) convey timing information outwards from connected nodes.

The network nodes can be mesh network appliances to provide voice communications, home security, door access control, lighting control, power outlet control, dimmer control, switch control, temperature control, humidity control, carbon monoxide control, fire alarm control, blind control, shade control, window control, oven control, cooking range control, personal computer control, entertainment console control, television control, projector control, garage door control, car control, pool temperature control, water pump control, furnace control, heater control, thermostat control, electricity meter monitor, water meter monitor, gas meter monitor, or remote diagnostics. The telephone can be connected to a cellular telephone to answer calls directed at the cellular telephone. The connection can be wired or wireless using Bluetooth or ZigBee. The telephone synchronizes calendar, contact, emails, blogs, or instant messaging with the cellular telephone. Similarly, the telephone synchronizes calendar, contact, emails, blogs, or instant messaging with a personal computer. A web server can communicate with the Internet through the POTS to provide information to an authorized remote user who logs into the server. A wireless router such as 802.11 router, 802.16 router, WiFi router, WiMAX router, Bluetooth router, X10 router can be connected to the mesh network.

A mesh network appliance can be connected to a power line to communicate X10 data to and from the mesh network. X10 is a communication protocol that allows up to 256 X10 products to talk to each other using the existing electrical wiring in the home. Typically, the installation is simple, a transmitter plugs (or wires) in at one location in the home and sends its control signal (on, off, dim, bright, etc.) to a receiver which plugs (or wires) into another location in the home. The mesh network appliance translates messages intended for X10 device to be relayed over the ZigBee wireless network, and then transmitted over the power line using a ZigBee to X10 converter appliance.

An in-door positioning system links one or more mesh network appliances to provide location information. Inside the home or office, the radio frequency signals have negligible multipath delay spread (for timing purposes) over short distances. Hence, radio strength can be used as a basis for determining position. Alternatively, time of arrival can be used to determine position, or a combination of radio signal strength and time of arrival can be used. Position estimates can also be achieved in an embodiment by beamforming, a method that exchanges time-stamped raw data among the nodes. While the processing is relatively more costly, it yields processed data with a higher signal to noise ratio (SNR) for subsequent classification decisions, and enables estimates of angles of arrival for targets that are outside the convex hull of the participating sensors. Two such clusters of ZigBee nodes can then provide for triangulation of distant targets. Further, beamforming enables suppression of interfering sources, by placing nulls in the synthetic beam pattern in their directions. Another use of beamforming is in self-location of nodes when the positions of only a very small number of nodes or appliances are known such as those sensors nearest the wireless stations. In one implementation where each node knows the

distances to its neighbors due to their positions, and some small fraction of the nodes (such as those nearest a PC with GPS) of the network know their true locations. As part of the network-building procedure, estimates of the locations of the nodes that lie within or near the convex hull of the nodes with known position can be quickly generated. To start, the shortest distance (multihop) paths are determined between each reference node. All nodes on this path are assigned a location that is the simple linear average of the two reference locations, as if the path were a straight line. A node which lies on the intersection of two such paths is assigned the average of the two indicated locations. All nodes that have been assigned locations now serve as references. The shortest paths among these new reference nodes are computed, assigning locations to all intermediate nodes as before, and continuing these iterations until no further nodes get assigned locations. This will not assign initial position estimates to all sensors. The remainder can be assigned locations based on pairwise averages of distances to the nearest four original reference nodes. Some consistency checks on location can be made using trigonometry and one further reference node to determine whether or not the node likely lies within the convex hull of the original four reference sensors.

In two dimensions, if two nodes have known locations, and the distances to a third node are known from the two nodes, then trigonometry can be used to precisely determine the location of the third node. Distances from another node can resolve any ambiguity. Similarly, simple geometry produces precise calculations in three dimensions given four reference nodes. But since the references may also have uncertainty, an alternative procedure is to perform a series of iterations where successive trigonometric calculations result only in a delta of movement in the position of the node. This process can determine locations of nodes outside the convex hull of the reference sensors. It is also amenable to averaging over the positions of all neighbors, since there will often be more neighbors than are strictly required to determine location. This will reduce the effects of distance measurement errors. Alternatively, the network can solve the complete set of equations of intersections of hyperbola as a least squares optimization problem.

In yet another embodiment, any or all of the nodes may include transducers for acoustic, infrared (IR), and radio frequency (RF) ranging. Therefore, the nodes have heterogeneous capabilities for ranging. The heterogeneous capabilities further include different margins of ranging error. Furthermore, the ranging system is re-used for sensing and communication functions. For example, wideband acoustic functionality is available for use in communicating, bistatic sensing, and ranging. Such heterogeneous capability of the sensors **40** can provide for ranging functionality in addition to communications functions. As one example, repeated use of the communications function improves position determination accuracy over time. Also, when the ranging and the timing are conducted together, they can be integrated in a self-organization protocol in order to reduce energy consumption. Moreover, information from several ranging sources is capable of being fused to provide improved accuracy and resistance to environmental variability. Each ranging means is exploited as a communication means, thereby providing improved robustness in the presence of noise and interference. Those skilled in the art will realize that there are many architectural possibilities, but allowing for heterogeneity from the outset is a component in many of the architectures.

Turning now to FIGS. **8-13**, various exemplary monitoring devices are shown. In FIG. **8**, a ring **130** has an opening **132**

for transmitting and receiving acoustic energy to and from the sensor **84** in an acoustic implementation. In an optical implementation, a second opening (not shown) is provided to emit an optical signal from an LED, for example, and an optical detector can be located at the opening **132** to receive the optical signal passing through the finger wearing the ring **130**. In another implementation, the ring has an electrically movable portion **134** and rigid portions **136-138** connected thereto. The electrically movable portion **134** can squeeze the finger as directed by the CPU during an applanation sweep to determine the arterial blood pressure.

FIG. **9** shows an alternate finger cover embodiment where a finger-mounted module housing the photo-detector and light source. The finger mounted module can be used to measure information that is processed to determine the user's blood pressure by measuring blood flow in the user's finger and sending the information through a wireless connection to the base station. In one implementation, the housing is made from a flexible polymer material.

In an embodiment to be worn on the patient's ear lobe, the monitoring device can be part of an earring jewelry clipped to the ear lobe. In the implementation of FIG. **10**, the monitoring device has a jewelry body **149** that contains the monitoring electronics and power source. The surface of the body **149** is an ornamental surface such as jade, ivory, pearl, silver, or gold, among others. The body **149** has an opening **148** that transmits energy such as optical or acoustic energy through the ear lobe to be detected by a sensor **144** mounted on a clamp portion that is secured to the body **149** at a base **147**. The energy detected through the sensor **144** is communicated through an electrical connector to the electronics in the jewelry body **149** for processing the received energy and for performing wireless communication with a base station. In FIG. **2E**, a bolt **145** having a stop end **146** allows the user to adjust the pressure of the clamp against the ear lobe. In other implementations, a spring biased clip is employed to retain the clip on the wearer's ear lobe. A pair of members, which snap together under pressure, are commonly used and the spring pressure employed should be strong enough to suit different thicknesses of the ear lobe.

FIGS. **11** and **12** show two additional embodiments of the monitoring device. In FIG. **11**, a wearable monitoring device is shown. The monitoring device has a body **160** comprising microphone ports **162**, **164** and **170** arranged in a first order noise cancelling microphone arrangement. The microphones **162** and **164** are configured to optimally receive distant noises, while the microphone **170** is optimized for capturing the user's speech. A touch sensitive display **166** and a plurality of keys **168** are provided to capture hand inputs. Further, a speaker **172** is provided to generate a verbal feedback to the user.

Turning now to FIG. **12**, a jewelry-sized monitoring device is illustrated. In this embodiment, a body **172** houses a microphone port **174** and a speaker port **176**. The body **172** is coupled to the user via the necklace **178** so as to provide a personal, highly accessible personal computer. Due to space limitations, voice input/output is an important user interface of the jewelry-sized computer. Although a necklace is disclosed, one skilled in the art can use a number of other substitutes such as a belt, a brace, a ring, or a band to secure the jewelry-sized computer to the user.

FIG. **13** shows an exemplary ear phone embodiment **180**. The ear phone **180** has an optical transmitter **182** which emits LED wavelengths that are received by the optical receiver **184**. The blood oximetry information is generated and used to determine blood pulse or blood pressure. Additionally, a module **186** contains mesh network communication electron-

61

ics, accelerometer, and physiological sensors such as EKG/ECG sensors or temperature sensors or ultrasonic sensors. In addition, a speaker (not shown) is provided to enable voice communication over the mesh network, and a microphone 188 is provided to pick up voice during verbal communication and pick up heart sound when the user is not using the microphone for voice communication. The ear phone optionally has an ear canal temperature sensor for sensing temperature in a human.

FIG. 14 shows an exemplary adhesive patch embodiment. The patch may be applied to a person's skin by anyone including the person themselves or an authorized person such as a family member or physician. The adhesive patch is shown generally at 190 having a gauze pad 194 attached to one side of a backing 192, preferably of plastic, and wherein the pad can have an impermeable side 194 coating with backing 192 and a module 196 which contains electronics for communicating with the mesh network and for sensing acceleration and EKG/ECG, heart sound, microphone, optical sensor, or ultrasonic sensor in contact with a wearer's skin. In one embodiment, the module 196 has a skin side that may be coated with a conductive electrode lotion or gel to improve the contact. The entire patch described above may be covered with a plastic or foil strip to retain moisture and retard evaporation by a conductive electrode lotion or gel provided to improve the electrode contact. In one embodiment, an acoustic sensor (microphone or piezoelectric sensor) and an electrical sensor such as EKG sensor contact the patient with a conductive gel material. The conductive gel material provides transmission characteristics so as to provide an effective acoustic impedance match to the skin in addition to providing electrical conductivity for the electrical sensor. The acoustic transducer can be directed mounted on the conductive gel material substantially with or without an intermediate air buffer. The entire patch is then packaged as sterile as are other over-the-counter adhesive bandages. When the patch is worn out, the module 196 may be removed and a new patch backing 192 may be used in place of the old patch. One or more patches may be applied to the patient's body and these patches may communicate wirelessly using the mesh network or alternatively they may communicate through a personal area network using the patient's body as a communication medium.

The term "positional measurement," as that term is used herein, is not limited to longitude and latitude measurements, or to metes and bounds, but includes information in any form from which geophysical positions can be derived. These include, but are not limited to, the distance and direction from a known benchmark, measurements of the time required for certain signals to travel from a known source to the geophysical location where the signals may be electromagnetic or other forms, or measured in terms of phase, range, Doppler or other units.

FIG. 15A shows a system block diagram of the network-based patient monitoring system in a hospital or nursing home setting. The system has a patient component 215, a server component 216, and a client component 217. The patient component 215 has one or more mesh network patient transmitters 202 for transmitting data to the central station. The central server comprises one or more Web servers 205, one or more waveform servers 204 and one or more mesh network receivers 211. The output of each mesh network receiver 211 is connected to at least one of the waveform servers 204. The waveform servers 204 and Web servers 205 are connected to the network 105. The Web servers 205 are also connected to a hospital database 230. The hospital database 230 contains patient records. In the embodiment of

62

FIG. 15A, a plurality of nurse stations provide a plurality of nurse computer user interface 208. The user interface 208 receives data from an applet 210 that communicates with the waveform server 204 and updates the display of the nurse computers for treating patients.

The network client component 217 comprises a series of workstations 106 connected to the network 105. Each workstation 106 runs a World Wide Web (WWW or Web) browser application 208. Each Web browser can open a page that includes one or more media player applets 210. The waveform servers 204 use the network 105 to send a series of messages 220 to the Web servers 205. The Web servers 205 use the network 105 to communicate messages, shown as a path 221, to the workstations 106. The media player applets running on the workstations 106 use the network 105 to send messages over a path 223 directly to the waveform servers 204.

FIG. 15B shows a variation of the system of FIG. 15A for call center monitoring. In this embodiment, the patient appliances 202 wirelessly communicate to home base stations (not shown) which are connected to the POTS or PSTN network for voice as well as data transmission. The data is captured by the waveform server 204 and the voice is passed through to the call center agent computer 207 where the agent can communicate by voice with the patient. The call center agent can forward the call to a professional such as a nurse or doctor or emergency service personnel if necessary. Hence, the system can include a patient monitoring appliance coupled to the POTS or PSTN through the mesh network. The patient monitoring appliance monitors drug usage and patient falls. The patient monitoring appliance monitors patient movement. A call center can call to the telephone to provide a human response.

In one exemplary monitoring service providing system, such as an emergency service providing system, the system includes a communication network (e.g., the Public Switch Telephone Network or PSTN or POTS), a wide area communication network (e.g., TCP/IP network) in call centers. The communication network receives calls destined for one of the call centers. In this regard, each call destined for one of the call centers is preferably associated with a particular patient, a call identifier or a call identifier of a particular set of identifiers. A call identifier associated with an incoming call may be an identifier dialed or otherwise input by the caller. For example, the call centers may be locations for receiving calls from a particular hospital or nursing home.

To network may analyze the automatic number information (ANI) and/or automatic location information (ALI) associated with the call. In this regard, well known techniques exist for analyzing the ANI and ALI of an incoming call to identify the call as originating from a particular calling device or a particular calling area. Such techniques may be employed by the network to determine whether an incoming call originated from a calling device within an area serviced by the call centers. Moreover, if an incoming call originated from such an area and if the incoming call is associated with the particular call identifier referred to above, then the network preferably routes the call to a designated facility.

When a call is routed to the facility, a central data manager, which may be implemented in software, hardware, or a combination thereof, processes the call according to techniques that will be described in more detail hereafter and routes the call, over the wide area network, to one of the call centers depending on the ANI and/or ALI associated with the call. In processing the call, the central data manager may convert the call from one communication protocol to another communication protocol, such as voice over internet protocol (VoIP),

for example, in order to increase the performance and/or efficiency of the system. The central data manager may also gather information to help the call centers in processing the call. There are various techniques that may be employed by the central data manager to enhance the performance and/or efficiency of the system, and examples of such techniques will be described in more detail hereafter.

Various benefits may be realized by utilizing a central facility to intercept or otherwise receive a call from the network and to then route the call to one of the call centers via WAN. For example, serving multiple call centers with a central data manager, may help to reduce total equipment costs. In this regard, it is not generally necessary to duplicate the processing performed by the central data manager at each of the call centers. Thus, equipment at each of the call centers may be reduced. As more call centers are added, the equipment savings enabled by implementing equipment at the central data manager instead of the call centers generally increases. Furthermore, the system is not dependent on any telephone company's switch for controlling the manner in which data is communicated to the call centers. In this regard, the central data manager may receive a call from the network and communicate the call to the destination call centers via any desirable communication technique, such as VoIP, for example. Data security is another possible benefit of the exemplary system 10 as the central data manager is able to store the data for different network providers associated with network on different partitions.

While the patient interface 90 (FIG. 1A) can provide information for a single person, FIG. 15C shows an exemplary interface to monitor a plurality of persons, while FIG. 15D shows an exemplary dash-board that provides summary information on the status of a plurality of persons. As shown in FIG. 1C, for professional use such as in hospitals, nursing homes, or retirement homes, a display can track a plurality of patients. In FIG. 15C, a warning (such as sound or visual warning in the form of light or red flashing text) can be generated to point out the particular patient that may need help or attention. In FIG. 15D, a magnifier glass can be dragged over a particular individual icon to expand and show detailed vital parameters of the individual and if available, images from the camera 10 trained on the individual for real time video feedback. The user can initiate voice communication with the user for confirmation purposes by clicking on a button provided on the interface and speaking into a microphone on the professional's workstation.

In one embodiment for professional users such as hospitals and nursing homes, a Central Monitoring Station provides alarm and vital sign oversight for a plurality of patients from a single computer workstation. FIG. 15E shows an exemplary multi-station vital parameter user interface for a professional embodiment, while FIG. 15F shows an exemplary trending pattern display. The clinician interface uses simple point and click actions with a computer mouse or trackball. The clinician can initiate or change monitoring functions from either the Central Station or the bedside monitor. One skilled in the art will recognize that patient data such as EKG data can be shown either by a scrolling waveform that moves along the screen display, or by a moving bar where the waveform is essentially stationary and the bar moves across the screen.

In one embodiment, software for the professional monitoring system provides a login screen to enter user name and password, together with database credentials. In Select Record function, the user can select a person, based on either entered or pre-selected criteria. From here navigate to their demographics, medical record, etc. The system can show a persons demographics, includes aliases, people involved in

their care, friends and family, previous addresses, home and work locations, alternative numbers and custom fields. The system can show all data elements of a person's medical record. These data elements are not 'hard wired', but may be configured in the data dictionary to suit your particular requirements. It is possible to create views of the record that filter it to show (for instance) just the medications or diagnosis, etc. Any data element can be designated 'plan able' in the data dictionary and then scheduled. A Summary Report can be done. Example of a report displayed in simple format, selecting particular elements and dates. As many of these reports as required can be created, going across all data in the system based on some criteria, with a particular selection of fields and sorting, grouping and totaling criteria. Reports can be created that can format and analyze any data stored on the server. The system supports OLE controls and can include graphs, bar codes, etc. These can be previewed on screen, printed out or exported in a wide variety of formats. The system also maintains a directory of all organizations the administrator wishes to record as well as your own. These locations are then used to record the location for elements of the medical record (where applicable), work addresses for people involved in the care and for residential addresses for people in residential care. The data elements that form the medical record are not 'hard wired' (ie predefined) but may be customized by the users to suit current and future requirements.

In one embodiment, the wearable appliance can store patient data in its data storage device such as flash memory. The data can include Immunizations and dates; medications (prescriptions and supplements); physician names, addresses, phone numbers, email addresses; location and details of advance directives; insurance company, billing address, phone number, policy number; emergency contacts, addresses, home/business/pager phone numbers, email addresses. The data can include color or black and white photo of the wearer of the device; a thumb print, iris print of other distinguishing physical characteristic; dental records; sample ECG or Cardiac Echo Scan.; blood type; present medication being taken; drug interaction precautions; drug and/or allergic reaction precautions; a description of serious preexisting medical conditions; Emergency Medical Instructions, which could include: administering of certain suggested drugs or physical treatments; calling emergency physician numbers listed; bringing the patient to a certain type of clinic or facility based on religious beliefs; and living will instructions in the case of seriously ill patients; Organ Donor instructions; Living Will instructions which could include: instructions for life support or termination of treatment; notification of next of kin and/or friends including addresses and telephone numbers; ECG trace; Cardiac Echo Scan; EEG trace; diabetes test results; x-ray scans, among others. The wearable appliance stores the wearer's medical records and ID information. In one embodiment, to start the process new/original medical information is organized and edited to fit into the BWD page format either in physicians office or by a third party with access to a patient's medical records using the base unit storage and encrypting software which can be stored in a normal pc or other compatible computer device. The system can encrypt the records so as to be secure and confidential and only accessible to authorized individuals with compatible de-encrypting software. In the event the wearer is stricken with an emergency illness a Paramedic, EMT or Emergency Room Technician can use a wireless interrogator to rapidly retrieve and display the stored medical records in the wearable appliance and send the medical records via wireless telemetry to a remote emergency room or physician's office

for rapid and life-saving medical intervention in a crisis situation. In a Non-emergency Situation, the personal health information service is also helpful as it eliminates the hassle of repeatedly filling out forms when changing health plans or seeing a new physician; stores vaccination records to schools or organizations without calling the pediatrician; or enlists the doctor's or pharmacist's advice about multiple medications without carrying all the bottles to a personal visit.

In one embodiment, a plurality of body worn sensors with in-door positioning can be used as an Emergency Department and Urgent Care Center Tracking System. The system tracks time from triage to MD assessment, identifies patients that have not yet been registered, records room usage, average wait time, and average length of stay. The system allows user defined "activities" so that hospitals can track times and assist in improving patient flow and satisfaction. The system can set custom alerts and send email/pager notifications to better identify long patient wait times and record the number of these alert occurrences. The system can manage room usage by identifying those rooms which are under/over utilized. The hospital administrator can set manual or automatic alerts and generate custom reports for analysis of patient flow. The system maximizes revenue by streamlining processes and improving throughput; improves charge capture by ensuring compliance with regulatory standards; increases accountability by collecting clear, meaningful data; enhances risk management and QA; and decreases liability.

FIG. 16A shows an exemplary process to continuously determine blood pressure of a patient. The process generates a blood pressure model of a patient (2002); determines a blood flow velocity using a piezoelectric transducer (2004); and provides the blood flow velocity to the blood pressure model to continuously estimate blood pressure (2006).

FIG. 16B shows another exemplary process to continuously determine blood pressure of a patient. First, during an initialization mode, a monitoring device and calibration device are attached to patient (2010). The monitoring device generates patient blood flow velocity, while actual blood pressure is measured by a calibration device (2012). Next, the process generates a blood pressure model based on the blood flow velocity and the actual blood pressure (2014). Once this is done, the calibration device can be removed (2016). Next, during an operation mode, the process periodically samples blood flow velocity from the monitoring device on a real-time basis (18) and provides the blood flow velocity as input information to the blood pressure model to estimate blood pressure (20). This process can be done in continuously or periodically as specified by a user.

In one embodiment, to determine blood flow velocity, acoustic pulses are generated and transmitted into the artery using an ultrasonic transducer positioned near a wrist artery. These pulses are reflected by various structures or entities within the artery (such as the artery walls, and the red blood cells within the subject's blood), and subsequently received as frequency shifts by the ultrasonic transducer. Next, the blood flow velocity is determined. In this process, the frequencies of those echoes reflected by blood cells within the blood flowing in the artery differ from that of the transmitted acoustic pulses due to the motion of the blood cells. This well known "Doppler shift" in frequency is used to calculate the blood flow velocity. In one embodiment for determining blood flow velocity, the Doppler frequency is used to determine mean blood velocity. For example, U.S. Pat. No. 6,514, 211, the content of which is incorporated by reference, discusses blood flow velocity using a time-frequency representation.

In one implementation, the system can obtain one or more numerical calibration curves describing the patient's vital signs such as blood pressure. The system can then direct energy such as infrared or ultrasound at the patient's artery and detecting reflections thereof to determine blood flow velocity from the detected reflections. The system can numerically fit or map the blood flow velocity to one or more calibration parameters describing a vital-sign value. The calibration parameters can then be compared with one or more numerical calibration curves to determine the blood pressure.

Additionally, the system can analyze blood pressure, and heart rate, and pulse oximetry values to characterize the user's cardiac condition. These programs, for example, may provide a report that features statistical analysis of these data to determine averages, data displayed in a graphical format, trends, and comparisons to doctor-recommended values.

In one embodiment, feed forward artificial neural networks (NNs) are used to classify valve-related heart disorders. The heart sounds are captured using the microphone or piezoelectric transducer. Relevant features were extracted using several signal processing tools, discrete wavelet transfer, fast fourier transform, and linear prediction coding. The heart beat sounds are processed to extract the necessary features by: a) denoising using wavelet analysis, b) separating one beat out of each record c) identifying each of the first heart sound (FHS) and the second heart sound (SHS). Valve problems are classified according to the time separation between the FHS and th SHS relative to cardiac cycle time, namely whether it is greater or smaller than 20% of cardiac cycle time. In one embodiment, the NN comprises 6 nodes at both ends, with one hidden layer containing 10 nodes. In another embodiment, linear predictive code (LPC) coefficients for each event were fed to two separate neural networks containing hidden neurons.

In another embodiment, a normalized energy spectrum of the sound data is obtained by applying a Fast Fourier Transform. The various spectral resolutions and frequency ranges were used as inputs into the NN to optimize these parameters to obtain the most favorable results.

In another embodiment, the heart sounds are denoised using six-stage wavelet decomposition, thresholding, and then reconstruction. Three feature extraction techniques were used: the Decimation method, and the wavelet method. Classification of the heart diseases is done using Hidden Markov Models (HMMs).

In yet another embodiment, a wavelet transform is applied to a window of two periods of heart sounds. Two analyses are realized for the signals in the window: segmentation of first and second heart sounds, and the extraction of the features. After segmentation, feature vectors are formed by using the wavelet detail coefficients at the sixth decomposition level. The best feature elements are analyzed by using dynamic programming.

In another embodiment, the wavelet decomposition and reconstruction method extract features from the heart sound recordings. An artificial neural network classification method classifies the heart sound signals into physiological and pathological murmurs. The heart sounds are segmented into four parts: the first heart sound, the systolic period, the second heart sound, and the diastolic period. The following features can be extracted and used in the classification algorithm: a) Peak intensity, peak timing, and the duration of the first heart sound b) the duration of the second heart sound c) peak intensity of the aortic component of S2(A2) and the pulmonic component of S2 (P2), the splitting interval and the reverse flag of A2 and P2, and the timing of A2 d) the duration, the three largest frequency components of the systolic signal and

the shape of the envelope of systolic murmur e) the duration the three largest frequency components of the diastolic signal and the shape of the envelope of the diastolic murmur.

In one embodiment, the time intervals between the ECG R-waves are detected using an envelope detection process. The intervals between R and T waves are also determined. The Fourier transform is applied to the sound to detect S1 and S2. To expedite processing, the system applies Fourier transform to detect S1 in the interval 0.1-0.5 R-R. The system looks for S2 the intervals R-T and 0.6 R-R. S2 has an aortic component A2 and a pulmonary component P2. The interval between these two components and its changes with respiration has clinical significance. A2 sound occurs before P2, and the intensity of each component depends on the closing pressure and hence A2 is louder than P2. The third heard sound S3 results from the sudden halt in the movement of the ventricle in response to filling in early diastole after the AV valves and is normally observed in children and young adults. The fourth heart sound S4 is caused by the sudden halt of the ventricle in response to filling in presystole due to atrial contraction.

In yet another embodiment, the S2 is identified and a normalized splitting interval between A2 and P2 is determined. If there is no overlap, A2 and P2 are determined from the heart sound. When overlap exists between A2 and P2, the sound is dechirped for identification and extraction of A2 and P2 from S2. The A2-P2 splitting interval (SI) is calculated by computing the cross-correlation function between A2 and P2 and measuring the time of occurrence of its maximum amplitude. SI is then normalized (NSI) for heart rate as follows: $NSI = SI / \text{cardiac cycle time}$. The duration of the cardiac cycle can be the average interval of QRS waves of the ECG. It could also be estimated by computing the mean interval between a series of consecutive S1 and S2 from the heart sound data. A non linear regressive analysis maps the relationship between the normalized NSI and PAP. A mapping process such as a curve-fitting procedure determines the curve that provides the best fit with the patient data. Once the mathematical relationship is determined, NSI can be used to provide an accurate quantitative estimate of the systolic and mean PAP relatively independent of heart rate and systemic arterial pressure.

In another embodiment, the first heart sound (S1) is detected using a time-delayed neural network (TDNN). The network consists of a single hidden layer, with time-delayed links connecting the hidden units to the time-frequency energy coefficients of a Morlet wavelet decomposition of the input phonocardiogram (PCG) signal. The neural network operates on a 200 msec sliding window with each time-delay hidden unit spanning 100 msec of wavelet data.

In yet another embodiment, a local signal analysis is used with a classifier to detect, characterize, and interpret sounds corresponding to symptoms important for cardiac diagnosis. The system detects a plurality of different heart conditions. Heart sounds are automatically segmented into a segment of a single heart beat cycle. Each segment are then transformed using 7 level wavelet decomposition, based on Coifman 4th order wavelet kernel. The resulting vectors 4096 values, are reduced to 256 element feature vectors, this simplified the neural network and reduced noise.

In another embodiment, feature vectors are formed by using the wavelet detail and approximation coefficients at the second and sixth decomposition levels. The classification (decision making) is performed in 4 steps: segmentation of the first and second heart sounds, normalization process, feature extraction, and classification by the artificial neural network.

In another embodiment using decision trees, the system distinguishes (1) the Aortic Stenosis (AS) from the Mitral

Regurgitation (MR) and (2) the Opening Snap (OS), the Second Heart Sound Split (A2_P2) and the Third Heart Sound (S3). The heart sound signals are processed to detect the first and second heart sounds in the following steps: a) wavelet decomposition, b) calculation of normalized average Shannon Energy, c) a morphological transform action that amplifies the sharp peaks and attenuates the broad ones d) a method that selects and recovers the peaks corresponding to S1 and S2 and rejects others e) algorithm that determines the boundaries of S1 and S2 in each heart cycle f) a method that distinguishes S1 from S2.

In one embodiment, once the heart sound signal has been digitized and captured into the memory, the digitized heart sound signal is parameterized into acoustic features by a feature extractor. The output of the feature extractor is delivered to a sound recognizer. The feature extractor can include the short time energy, the zero crossing rates, the level crossing rates, the filter-bank spectrum, the linear predictive coding (LPC), and the fractal method of analysis. In addition, vector quantization may be utilized in combination with any representation techniques. Further, one skilled in the art may use an auditory signal-processing model in place of the spectral models to enhance the system's robustness to noise and reverberation.

In one embodiment of the feature extractor, the digitized heart sound signal series $s(n)$ is put through a low-order filter, typically a first-order finite impulse response filter, to spectrally flatten the signal and to make the signal less susceptible to finite precision effects encountered later in the signal processing. The signal is pre-emphasized preferably using a fixed pre-emphasis network, or preemphasizer. The signal can also be passed through a slowly adaptive pre-emphasizer. The preemphasized heart sound signal is next presented to a frame blocker to be blocked into frames of N samples with adjacent frames being separated by M samples. In one implementation, frame 1 contains the first 400 samples. The frame 2 also contains 400 samples, but begins at the 300th sample and continues until the 700th sample. Because the adjacent frames overlap, the resulting LPC spectral analysis will be correlated from frame to frame. Each frame is windowed to minimize signal discontinuities at the beginning and end of each frame. The windower tapers the signal to zero at the beginning and end of each frame. Preferably, the window used for the autocorrelation method of LPC is the Hamming window. A noise canceller operates in conjunction with the autocorrelator to minimize noise. Noise in the heart sound pattern is estimated during quiet periods, and the temporally stationary noise sources are damped by means of spectral subtraction, where the autocorrelation of a clean heart sound signal is obtained by subtracting the autocorrelation of noise from that of corrupted heart sound. In the noise cancellation unit, if the energy of the current frame exceeds a reference threshold level, the heart is generating sound and the autocorrelation of coefficients representing noise is not updated. However, if the energy of the current frame is below the reference threshold level, the effect of noise on the correlation coefficients is subtracted off in the spectral domain. The result is half-wave rectified with proper threshold setting and then converted to the desired autocorrelation coefficients. The output of the autocorrelator and the noise canceller are presented to one or more parameterization units, including an LPC parameter unit, an FFT parameter unit, an auditory model parameter unit, a fractal parameter unit, or a wavelet parameter unit, among others. The LPC parameter is then converted into cepstral coefficients. The cepstral coefficients are the coefficients of the Fourier transform representation of the log magnitude spectrum. A filter bank spectral analysis, which

uses the short-time Fourier transformation (STFT) may also be used alone or in conjunction with other parameter blocks. FFT is well known in the art of digital signal processing. Such a transform converts a time domain signal, measured as amplitude over time, into a frequency domain spectrum, which expresses the frequency content of the time domain signal as a number of different frequency bands. The FFT thus produces a vector of values corresponding to the energy amplitude in each of the frequency bands. The FFT converts the energy amplitude values into a logarithmic value which reduces subsequent computation since the logarithmic values are more simple to perform calculations on than the longer linear energy amplitude values produced by the FFT, while representing the same dynamic range. Ways for improving logarithmic conversions are well known in the art, one of the simplest being use of a look-up table. In addition, the FFT modifies its output to simplify computations based on the amplitude of a given frame. This modification is made by deriving an average value of the logarithms of the amplitudes for all bands. This average value is then subtracted from each of a predetermined group of logarithms, representative of a predetermined group of frequencies. The predetermined group consists of the logarithmic values, representing each of the frequency bands. Thus, utterances are converted from acoustic data to a sequence of vectors of k dimensions, each sequence of vectors identified as an acoustic frame, each frame represents a portion of the utterance. Alternatively, auditory modeling parameter unit can be used alone or in conjunction with others to improve the parameterization of heart sound signals in noisy and reverberant environments. In this approach, the filtering section may be represented by a plurality of filters equally spaced on a log-frequency scale from 0 Hz to about 3000 Hz and having a prescribed response corresponding to the cochlea. The nerve fiber firing mechanism is simulated by a multilevel crossing detector at the output of each cochlear filter. The ensemble of the multilevel crossing intervals corresponding to the firing activity at the auditory nerve fiber-array. The interval between each successive pair of same direction, either positive or negative going, crossings of each predetermined sound intensity level is determined and a count of the inverse of these interspike intervals of the multilevel detectors for each spectral portion is stored as a function of frequency. The resulting histogram of the ensemble of inverse interspike intervals forms a spectral pattern that is representative of the spectral distribution of the auditory neural response to the input sound and is relatively insensitive to noise. The use of a plurality of logarithmically related sound intensity levels accounts for the intensity of the input signal in a particular frequency range. Thus, a signal of a particular frequency having high intensity peaks results in a much larger count for that frequency than a low intensity signal of the same frequency. The multiple level histograms of the type described herein readily indicate the intensity levels of the nerve firing spectral distribution and cancel noise effects in the individual intensity level histograms. Alternatively, the fractal parameter block can further be used alone or in conjunction with others to represent spectral information. Fractals have the property of self similarity as the spatial scale is changed over many orders of magnitude. A fractal function includes both the basic form inherent in a shape and the statistical or random properties of the replacement of that shape in space. As is known in the art, a fractal generator employs mathematical operations known as local affine transformations. These transformations are employed in the process of encoding digital data representing spectral data. The encoded output constitutes a "fractal transform" of

the spectral data and consists of coefficients of the affine transformations. Different fractal transforms correspond to different images or sounds.

Alternatively, a wavelet parameterization block can be used alone or in conjunction with others to generate the parameters. Like the FFT, the discrete wavelet transform (DWT) can be viewed as a rotation in function space, from the input space, or time domain, to a different domain. The DWT consists of applying a wavelet coefficient matrix hierarchically, first to the full data vector of length N , then to a smooth vector of length $N/2$, then to the smooth-smooth vector of length $N/4$, and so on. Most of the usefulness of wavelets rests on the fact that wavelet transforms can usefully be severely truncated, or turned into sparse expansions. In the DWT parameterization block, the wavelet transform of the heart sound signal is performed. The wavelet coefficients are allocated in a non-uniform, optimized manner. In general, large wavelet coefficients are quantized accurately, while small coefficients are quantized coarsely or even truncated completely to achieve the parameterization. Due to the sensitivity of the low-order cepstral coefficients to the overall spectral slope and the sensitivity of the high-order cepstral coefficients to noise variations, the parameters generated may be weighted by a parameter weighing block, which is a tapered window, so as to minimize these sensitivities. Next, a temporal derivator measures the dynamic changes in the spectra. Power features are also generated to enable the system to distinguish heart sound from silence.

After the feature extraction has been performed, the heart sound parameters are next assembled into a multidimensional vector and a large collection of such feature signal vectors can be used to generate a much smaller set of vector quantized (VQ) feature signals by a vector quantizer that cover the range of the larger collection. In addition to reducing the storage space, the VQ representation simplifies the computation for determining the similarity of spectral analysis vectors and reduces the similarity computation to a look-up table of similarities between pairs of codebook vectors. To reduce the quantization error and to increase the dynamic range and the precision of the vector quantizer, the preferred embodiment partitions the feature parameters into separate codebooks, preferably three. In the preferred embodiment, the first, second and third codebooks correspond to the cepstral coefficients, the differenced cepstral coefficients, and the differenced power coefficients.

With conventional vector quantization, an input vector is represented by the codeword closest to the input vector in terms of distortion. In conventional set theory, an object either belongs to or does not belong to a set. This is in contrast to fuzzy sets where the membership of an object to a set is not so clearly defined so that the object can be a part member of a set. Data are assigned to fuzzy sets based upon the degree of membership therein, which ranges from 0 (no membership) to 1.0 (full membership). A fuzzy set theory uses membership functions to determine the fuzzy set or sets to which a particular data value belongs and its degree of membership therein.

To handle the variance of heart sound patterns of individuals over time and to perform speaker adaptation in an automatic, self-organizing manner, an adaptive clustering technique called hierarchical spectral clustering is used. Such speaker changes can result from temporary or permanent changes in vocal tract characteristics or from environmental effects. Thus, the codebook performance is improved by collecting heart sound patterns over a long period of time to account for natural variations in speaker behavior. In one embodiment, data from the vector quantizer is presented to

one or more recognition models, including an HMM model, a dynamic time warping model, a neural network, a fuzzy logic, or a template matcher, among others. These models may be used singly or in combination.

In dynamic processing, at the time of recognition, dynamic programming slides, or expands and contracts, an operating region, or window, relative to the frames of heart sound so as to align those frames with the node models of each S1-S4 pattern to find a relatively optimal time alignment between those frames and those nodes. The dynamic processing in effect calculates the probability that a given sequence of frames matches a given word model as a function of how well each such frame matches the node model with which it has been time-aligned. The word model which has the highest probability score is selected as corresponding to the heart sound.

Dynamic programming obtains a relatively optimal time alignment between the heart sound to be recognized and the nodes of each word model, which compensates for the unavoidable differences in speaking rates which occur in different utterances of the same word. In addition, since dynamic programming scores words as a function of the fit between word models and the heart sound over many frames, it usually gives the correct word the best score, even if the word has been slightly misspoken or obscured by background sound. This is important, because humans often mispronounce words either by deleting or mispronouncing proper sounds, or by inserting sounds which do not belong.

In dynamic time warping (DTW), the input heart sound A, defined as the sampled time values $A=a(1) \dots a(n)$, and the vocabulary candidate B, defined as the sampled time values $B=b(1) \dots b(n)$, are matched up to minimize the discrepancy in each matched pair of samples. Computing the warping function can be viewed as the process of finding the minimum cost path from the beginning to the end of the words, where the cost is a function of the discrepancy between the corresponding points of the two words to be compared. Dynamic programming considers all possible points within the permitted domain for each value of i . Because the best path from the current point to the next point is independent of what happens beyond that point. Thus, the total cost of $[i(k), j(k)]$ is the cost of the point itself plus the cost of the minimum path to it. Preferably, the values of the predecessors can be kept in an $M \times N$ array, and the accumulated cost kept in a $2 \times \text{times} \cdot N$ array to contain the accumulated costs of the immediately preceding column and the current column. However, this method requires significant computing resources. For the heart sound recognizer to find the optimal time alignment between a sequence of frames and a sequence of node models, it must compare most frames against a plurality of node models. One method of reducing the amount of computation required for dynamic programming is to use pruning. Pruning terminates the dynamic programming of a given portion of heart sound against a given word model if the partial probability score for that comparison drops below a given threshold. This greatly reduces computation, since the dynamic programming of a given portion of heart sound against most words produces poor dynamic programming scores rather quickly, enabling most words to be pruned after only a small percent of their comparison has been performed. To reduce the computations involved, one embodiment limits the search to that within a legal path of the warping.

A Hidden Markov model can be used in one embodiment to evaluate the probability of occurrence of a sequence of observations $O(1), O(2), \dots O(t), \dots, O(T)$, where each observation $O(t)$ may be either a discrete symbol under the VQ approach or a continuous vector. The sequence of observa-

tions may be modeled as a probabilistic function of an underlying Markov chain having state transitions that are not directly observable. The transitions between states are represented by a transition matrix $A=[a(i,j)]$. Each $a(i,j)$ term of the transition matrix is the probability of making a transition to state j given that the model is in state i . The output symbol probability of the model is represented by a set of functions $B=[b(j)(O(t))]$, where the $b(j)(O(t))$ term of the output symbol matrix is the probability of outputting observation $O(t)$, given that the model is in state j . The first state is always constrained to be the initial state for the first time frame of the utterance, as only a prescribed set of left-to-right state transitions are possible. A predetermined final state is defined from which transitions to other states cannot occur. Transitions are restricted to reentry of a state or entry to one of the next two states. Such transitions are defined in the model as transition probabilities. For example, a heart sound pattern currently having a frame of feature signals in state 2 has a probability of reentering state 2 of $a(2,2)$, a probability $a(2,3)$ of entering state 3 and a probability of $a(2,4)=1-a(2,1)-a(2,2)$ of entering state 4. The probability $a(2,1)$ of entering state 1 or the probability $a(2,5)$ of entering state 5 is zero and the sum of the probabilities $a(2,1)$ through $a(2,5)$ is one. Although the preferred embodiment restricts the flow graphs to the present state or to the next two states, one skilled in the art can build an HMM model without any transition restrictions.

The Markov model is formed for a reference pattern from a plurality of sequences of training patterns and the output symbol probabilities are multivariate Gaussian function probability densities. The heart sound traverses through the feature extractor. During learning, the resulting feature vector series is processed by a parameter estimator, whose output is provided to the hidden Markov model. The hidden Markov model is used to derive a set of reference pattern templates, each template representative of an identified S1-S4 pattern in a vocabulary set of reference patterns. The Markov model reference templates are next utilized to classify a sequence of observations into one of the reference patterns based on the probability of generating the observations from each Markov model reference pattern template. During recognition, the unknown pattern can then be identified as the reference pattern with the highest probability in the likelihood calculator.

In one embodiment, a wireless monitoring system includes one or more wireless nodes communicating over aeronautical mobile telemetry frequency; and a wearable appliance in communication with the one or more wireless nodes, the appliance monitoring one or more vital signs.

In addition to hospital patient and equipment monitoring, the system can be used in other applications. In an employee access application, the system enables an employer to selectively grant access to specific rooms of a facility. Additionally, when an employee is in an area where he is not normally present, the system can flag a warning to the facility administrator. Further, the computer of the employee can be access locked so that only the employee with the proper wireless authorization can work on a particular computer. All employee accesses, physical as well as electronic, are tracked for regulatory requirements such as HIPAA requirements so that the administrator knows that only authorized personnel are present. Additionally, the employee can be paged in case he or she is needed through the voice walkie-talkie over the Zigbee network.

In a vending machine monitoring application, the system can monitor vending machines remotely located to a central monitoring system. For example, transmitters can be placed within soda machines to monitor the depletion of soda machines. When a "sold out" indication is present in the

vending machine, the system can transmit refill/reorder requests to a supplier using the wireless network or the POTS network. Also, the status of the vending machine can be monitored (e.g., the temperature of a ice cream machine or soda machine) to notify a supplier or the maintenance department when maintenance is required.

In a prisoner monitoring embodiment, people who are subject to incarceration need to be monitored. The system can constantly monitor the prisoners to ensure they are present. A prisoner has a wireless appliance that is secured or unremovably attached to his person. If the wireless appliance is forcibly removed it immediately transmits a notification to the prisoner monitor. In a Home Confinement Monitoring embodiment, a convict can be required not to leave their home. They are monitored by the wireless appliance attached to the "home prisoners" which are then monitored by a central monitoring center or station which can be a sheriff's office. In the home prisoner monitoring system, the wireless appliance is secured to or unremovably attached to the home prisoner and if they move outside of the range of the network (i.e., the leave the house), no transmission will be received by the wireless transceiver and an alarm is issued by the remote home prisoner monitor. In one embodiment, the alarm can be a phone call or an email message or fax message to the monitoring center or station.

In an Animal Monitoring embodiment, the system can monitor the status and presence of animals in a stock yard or on a farm by the similar methodologies discussed above. In this embodiment a plurality of animals can be monitored for presence as well as condition. For example, the system can ensure that animals have not wandered off as well as determine conditions such as temperature or heart rate of an animal, this can be accomplished by placing a wireless appliance on the animal.

In a utility monitoring embodiment, a wireless appliance is interfaced with a utility meter and thereafter transmits the current meter reading at predetermined intervals. Because of the low power requirements of Zigbee and the low duty cycle and low data rate required for transmitting the information, the battery for powering the Zigbee radio transmitter can last many months or more.

"Computer readable media" can be any available media that can be accessed by client/server devices. By way of example, and not limitation, computer readable media may comprise computer storage media and communication media. Computer storage media includes volatile and non-volatile, removable and non-removable media implemented in any method or technology for storage of information such as computer readable instructions, data structures, program modules or other data. Computer storage media includes, but is not limited to, RAM, ROM, EEPROM, flash memory or other memory technology, CD-ROM, digital versatile disks (DVD) or other optical storage, magnetic cassettes, magnetic tape, magnetic disk storage or other magnetic storage devices, or any other medium which can be used to store the desired information and which can be accessed by client/server devices. Communication media typically embodies computer readable instructions, data structures, program modules or other data in a modulated data signal such as a carrier wave or other transport mechanism and includes any information delivery media.

All references including patent applications and publications cited herein are incorporated herein by reference in their entirety and for all purposes to the same extent as if each individual publication or patent or patent application was specifically and individually indicated to be incorporated by reference in its entirety for all purposes. Many modifications

and variations of this invention can be made without departing from its spirit and scope, as will be apparent to those skilled in the art. The specific embodiments described herein are offered by way of example only. The above specification, examples and data provide a complete description of the manufacture and use of the composition of the invention. Since many embodiments of the invention can be made without departing from the spirit and scope of the invention, the invention resides in the claims hereinafter appended.

While this invention has been particularly shown and described with references to preferred embodiments thereof, it will be understood by those skilled in the art that various changes in form and details may be made therein without departing from the spirit and scope of the invention as defined by the appended claims.

What is claimed is:

1. A monitoring system, comprising:

one or more wireless nodes communicating over an aeronautical mobile telemetry (AMT) band from about 2360 to 2400 MHz; and

a wearable appliance in communication with the one or more wireless nodes to capture a patient vital sign.

2. The system of claim 1, comprising patient monitoring equipment coupled to the wireless nodes to display patient vital signs without attaching cables to a patient.

3. The system of claim 1, wherein the wearable appliance includes a high reliability transmission link with a redundant communication protocol to transmit patient data.

4. The system of claim 1, comprising a set of wireless nodes restricted to indoor uses and a set of wireless nodes without location restriction.

5. The system of claim 1, wherein wireless nodes have predetermined quality of service, coexistence with other wireless devices, data integrity and security.

6. The system of claim 1, wherein the wireless nodes comprise adaptive spectrum aware wireless nodes.

7. The system of claim 1, wherein the wearable appliance is implanted inside a patient.

8. The system of claim 1, comprising a wireless coordinator coupled to the wireless nodes.

9. The system of claim 1, comprising a call center coupled to the appliance to provide a human response.

10. The system of claim 1, comprising a database to store medicine taking habits, eating and drinking habits, sleeping habits, or exercise habits.

11. The system of claim 1, comprising hospital or operating room patient monitoring equipment coupled to the wireless node.

12. The system of claim 1, wherein the appliance comprises a sensor to capture at least one of: RR (respiratory rate), SpO2 (oxygen saturation), ECG (electrocardiogram), HR (heart rate), core temperature (inside heart) and peripheral temperature (on top of instep), CI (cardiac output index), systematic pressure, systematic systolic arterial pressure; systematic diastolic arterial pressure; systematic mean arterial pressure, CVP (central venous pressure), pulmonary pressures, pulmonary systolic arterial pressure, pulmonary diastolic arterial pressure, pulmonary mean arterial pressure, svO2 (oxygen saturation in the lung artery), ET/CO2 (outcoming carbon dioxide), FIO (ingoing oxygen), diuretics, patient weight, patient fluid balance (ingoing and outcoming fluids), EEG, intracranial pressure (ICP).

13. The system of claim 1, wherein the wireless nodes are in an ambulance, comprising a long range transceiver coupled to the wireless nodes to send patient data from the ambulance to the hospital.

75

14. The system of claim 1, comprising a cellular transceiver, optical transceiver, or body area network coupled to the wireless nodes.

15. The system of claim 1, comprising a plurality of directional antennas placed around the patient or under a patient, wherein each directional antenna is aimed at the wearable appliance to capture transmission from the wearable appliance.

16. A wireless system for a patient, comprising:

one or more wireless nodes; and

a wearable appliance monitoring clinical information from the patient and communicating with the one or more wireless nodes over an aeronautical mobile telemetry (AMT) band between about 2360 to 2400 MHz.

17. The system of claim 16, comprising a master transmitter or hub controlling the transmissions of the wearable appliance, where in the hub aggregate patient data and transmit clinical information for viewing by a healthcare professional.

18. The system of claim 16, wherein appliance is used for self-management of diabetes, wound healing, hypertension, or heart failure.

19. The system of claim 16, wherein the wearable appliance communicates with a receiver either in the patient's pocket or in a hospital room.

20. The system of claim 16, wherein the wearable appliance is outside a hospital, the information aggregated locally from at least one sensor coupled to the appliance is relayed into a cellular network to provide doctors or hospitals with patient monitoring.

21. The system of claim 16, comprising a processor with analytical software to process the clinical information and determine medical patterns in the data.

22. The system of claim 16, wherein the wearable appliance is made by a first manufacturer, comprising a second wearable appliance made by a second manufacturer coupled to the one or more wireless nodes, wherein the two wearable appliances are interoperable, or wherein the two wearable

76

appliances are compatible, or wherein the two wearable appliances communicate without interference from each other.

23. The system of claim 16, wherein the wearable appliance is made by a first manufacturer, comprising a second wearable appliance made by a second manufacturer coupled to the one or more wireless nodes, wherein the two wearable appliances do not interfere with each other's wireless communication.

24. The system of claim 16, wherein the wearable appliance minimizes electrical cable on the patient for ECG, EKG, EEG, or vital sign monitoring.

25. A wireless system for a patient, comprising:

one or more wireless nodes; and

a wearable appliance monitoring clinical information from the patient and communicating with the one or more wireless nodes over an aeronautical mobile telemetry (AMT) band from about 2360 to 2400 MHz, wherein the clinical information is aggregated at a nearby device for local processing and forwarded to one or more centralized displays and electronic medical records.

26. The system of claim 25, comprising an accelerometer to detect a dangerous condition and to generate a warning when the dangerous condition is detected.

27. The system of claim 25, wherein the appliance comprises a sensor to capture at least one of: RR (respiratory rate), SpO₂ (oxygen saturation), ECG (electrocardiogram), HR (heart rate), core temperature (inside heart) and peripheral temperature (on top of instep), CI (cardiac output index), systematic pressure, systematic systolic arterial pressure; systematic diastolic arterial pressure; systematic mean arterial pressure, CVP (central venous pressure), pulmonary pressures, pulmonary systolic arterial pressure, pulmonary diastolic arterial pressure, pulmonary mean arterial pressure, svO₂ (oxygen saturation in the lung artery), ETCO₂ (outcoming carbon dioxide), FIO (ingoing oxygen), diuretics, patient weight, patient fluid balance (ingoing and outcoming fluids), EEG, intracranial pressure (ICP).

* * * * *

专利名称(译)	健康监测设备		
公开(公告)号	US8652038	公开(公告)日	2014-02-18
申请号	US13/775055	申请日	2013-02-22
[标]申请(专利权)人(译)	TRAN包		
申请(专利权)人(译)	TRAN, 包		
当前申请(专利权)人(译)	TRAN, 包		
[标]发明人	TRAN BAO TRAN HA		
发明人	TRAN, BAO TRAN, HA		
IPC分类号	A61B5/00		
CPC分类号	A61B5/0402 A61B2560/0233 A61B5/1128 G08B21/02 A61B5/7225 A61B5/1117 A61B5/7214 A61B5/686 A61B5/0492 A61B2560/0412 A61B5/411 A61B7/04 A61B5/0205 A61B5/0022 A61B5/7267 A61B5/0488 A61B5/1112 A61B5/0476 A61B5/6891 A61B5/742 A61B5/0024 A61B5/0026 A61B5/7257 A61B5/6826 A61B5/681 A61B5/14542 A61B5/021 A61B5/14532 A61B5/726 A61B5/02055 A61B5/7465 A61B5/7275 A61B5/022 A61B5/4875 A61B5/201 A61B5/0006 A61B5/01 A61B5/14551 A61B5/743 A61B5/0816 A61B5/0008 A61B5/6846 A61B5/024 A61B5/031 A61B5/0215 A61B7/00 G16H40/67 G16H50/20		
其他公开文献	US20130178718A1		
外部链接	Espacenet USPTO		

摘要(译)

用于人的心脏监测系统包括一个或多个无线节点;以及与一个或多个无线节点通信的可穿戴设备, 该设备监视生命体征。

