

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization

International Bureau



(10) International Publication Number

WO 2015/173679 A1

(43) International Publication Date
19 November 2015 (19.11.2015)

WIPO | PCT

(51) International Patent Classification:
A61B 5/0205 (2006.01) *A61B 5/11* (2006.01)

HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(21) International Application Number:
PCT/IB2015/052964

(22) International Filing Date:
23 April 2015 (23.04.2015)

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
61/991,640 12 May 2014 (12.05.2014) US

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(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT,

Declarations under Rule 4.17:

- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))

Published:

- with international search report (Art. 21(3))

(54) Title: MOTION TRIGGERED VITAL SIGN MEASUREMENT

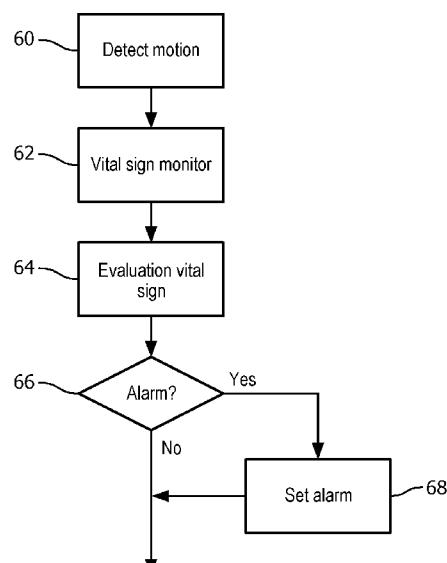


FIG. 5

(57) Abstract: A medical monitoring device (12) includes a motion detection device (20, 50) and a vital sign monitoring device (26, 52) communicatively connected to the motion detection device. The motion detection device (20, 50) detects whole body motion of a person. The vital sign monitoring device (26, 52) monitors, either continuously or intermittently, at least one vital sign for a predetermined time interval in response to the detected whole body motion. The vital sign is evaluated, e.g. for orthostatic intolerance, and in response to an affirmative evaluation an alarm signal is issued.

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MOTION TRIGGERED VITAL SIGN MEASUREMENT

The following relates generally to medical monitoring. It finds particular application in conjunction with vital sign monitoring and motion detection, and will be described with particular reference thereto. However, it will be understood that it also finds application in other usage scenarios and is not necessarily limited to the aforementioned 5 application.

People suffer from hemodynamic problems such as orthostatic hypotension, which can result in a patient falling and injuring themselves. For example, a person moving from a sitting position to a standing position can suffer from syncope or fainting, which can occur from 0-45 minutes after standing according to current medical guidelines.

10 Current methods to diagnosis orthostatic hypotension typically include a tilt table test in a hospital setting. The person is placed on a horizontal table surface, which rotates to a vertical or near vertical orientation, and the person is monitored for symptoms of syncope.

15 Patients are monitored non-invasively for vital signs, such as ECG, blood pressure, pulse, etc. Devices which monitor vital signs typically monitor either continuously or when manually initiated. Furthermore, there is a demand for mobility of vital sign monitoring devices. Vital sign devices are typically limited in their mobility by storage capacity and power generation capability. For example, a continuously monitored vital sign can quickly fill computer storage of a portable device and quickly consume battery power.

20 Monitoring a person for dizziness or fainting continuously, including during inactive periods, e.g. sitting, unnecessarily consumes both storage and power.

25 Accelerometers which detect motion have been included with some vital sign devices, such as athletic fitness devices which monitor pulse rate and motion rate. However, such devices do not discriminate body motion, such as changing from a sitting position to a standing position, and consume power, to measure and track motion.

The following discloses a new and improved motion triggered vital sign measurement device and method which addresses the above referenced issues, and others.

In accordance with one aspect, a medical monitoring device includes a motion detection device and a vital sign monitoring device communicatively connected to the motion

detection device. The motion detection device is adapted to detect whole body motion of a person. The vital sign monitoring device is adapted to monitor at least one vital sign for a predetermined time interval, in response to the detected whole body motion.

In accordance with another aspect, a method of monitoring vital signs includes 5 detecting whole body motion of a person. At least one vital sign is monitored for a predetermined time interval in response to the detected whole body motion.

In accordance with another aspect, a medical monitoring device includes a motion detection device, a vital sign monitoring device, a vital sign evaluation device, and an alarm device. The motion detection device is adapted to detect body motion of a person 10 indicative of a change in position, e.g. from sitting to standing. The vital sign monitoring device is communicatively connected to the motion detection device and adapted to monitor e.g. blood pressure for a predetermined time interval in response to the detected body motion. The vital sign evaluation device is adapted to evaluate the monitored blood pressure for conditions indicative of orthostatic intolerance. The alarm device is configured to generate a 15 signal in response to the evaluated conditions

One advantage is ability to monitor persons for syncope in a non-hospital setting.

Another advantage resides in alarms for possible falls due to fainting, 20 dizziness, or syncope.

Another advantage resides in reduced power and memory requirements.

Still further advantages will be appreciated to those of ordinary skill in the art upon reading and understanding the following detailed description.

The invention may take form in various components and arrangements of 25 components, and in various steps and arrangement of steps. The drawings are only for purposes of illustrating the preferred embodiments and are not to be construed as limiting the invention.

FIGURE 1 diagrammatically illustrates an embodiment of the motion triggered vital sign measurement system in a time sequence.

30 FIGURE 2 schematically illustrates one embodiment of the motion triggered medical monitoring device.

FIGURE 3 diagrammatically illustrates another embodiment of the motion triggered vital sign measurement system in a time sequence.

FIGURE 4 schematically illustrates another embodiment of the motion triggered medical monitoring device with an optical motion detection device.

FIGURE 5 flowcharts one method of motion triggered vital sign measurement.

5 With reference to FIGURE 1, an embodiment of the motion triggered vital sign measurement system **10** in a time sequence is diagrammatically illustrated. The system includes a medical monitoring device **12**, adapted to affix to a person **14** and monitor the person. The device **12** operates in two states, a resting state **16**, and a motion triggered state **18**. In the resting state **16**, the device does not monitor patient vital signs, or if intermittent 10 monitoring is required in resting state, the intermittent monitoring frequency is e.g. lower compared to the intermittent monitoring frequency in the motion triggered state. In the motion triggered state **18**, the device actively monitors vital signs of the person for a predetermined interval triggered by detected whole body motion at a motion triggered state **18**, either continuously or intermittent. For example, as a person moves from a sitting 15 position to a standing position, orthostatic stress is applied to the person. The system is ideally suited for use as a fall prediction and prevention system.

A motion detection device or means **20** detects whole body motion. For example, as the person moves to a standing position **18**, whole body movement **22**, such as an change in elevation **24** relative to the center of gravity, is detected. The change in 20 elevation creates orthostatic stress. The detected motion exceeds a predetermined threshold for whole body motion. For example, the person at rest generates motion, and the motion detection device discriminates between whole body motion indicative of standing and other motions such as respiration. In one embodiment, the motion detection device separates the acceleration vectors for elevation and other whole body movement. The motion detection 25 device is located and/or affixed to the person in a position to discriminate whole body motion from fine motor motion. The motion detection device determines acceleration vectors that separate elevation changes from other body motion vectors. For example, as a person stands, the motion includes forward movement, not to be confused with a leaning movement, e.g. forward motion vector, or a twisting movement, e.g. left-right motion vector without standing. 30 The motion detection device can alternatively detect motion with accelerometers, a multi-axis accelerometer, barometric pressure sensor, video monitoring, ultrasound monitoring, gyroscope, GPS receiver, or the like.

A vital sign monitoring device or means **26**, is triggered by the detected whole body movement to monitor one or more vital signs, such as blood pressure, pulse, ECG,

SpO₂ and the like. The vital sign monitoring device monitors the vital signs for a predetermined period of time, such as x minutes. For example, the person sits at a time t_0 and stands at a time t_1 , and the vital sign monitoring device **26** monitors blood pressure, continuously or intermittently, in a cuff-less configuration for a time interval of 45 minutes 5 beginning at t_1 . The predetermined time interval can be adjusted according to medical guidelines in which syncope presents. In one embodiment, the intermittent monitoring includes an adjustable sampling rate. For example, the rate is adjusted to sample frequently in an initial period after standing, and steps down the sampling rate according to medical guidelines and/or patient population statistics after the initial period. Inactive monitoring 10 during a resting state, and active monitoring limited to the predetermined interval, reduces power and memory or computer storage consumption. Blood pressure can also be detected with a cuff.

The medical monitoring device **12** evaluates the monitored vital sign, e.g. the signals generated by the vital sign monitoring device **24**, such as blood pressure. The vital 15 sign evaluation includes a determination of symptoms of syncope, such as a minimum blood pressure or a threshold change in blood pressure. For example, the blood pressure measured over time after the detected motion is compared to blood pressure versus time curves indicative or predictive of syncope. Such curves can be determined by data mining patient records of patient's that have suffered syncope for preceding blood pressure variations. As 20 another option, the monitored blood pressure can be compared with a preselected low blood pressure value, e.g. indicative of low blood flow to the brain.

The medical monitoring device **12** generates an alert signal in response to a minimum blood pressure and/or threshold change in blood pressure. The medical monitoring device **12** can audibly sound and/or visually indicate an alert **28**. A caregiver **30** responds to 25 the alert to prevent the person experiencing symptoms of syncope, e.g. dizziness, fainting, etc., from falling. The alert can be sent to a computing device **32** of the caregiver. The alert can also alert the patient to sit or recline before becoming unconscious.

The device **12** includes a mechanism or means **34** to affix to the patient, such as a strap, clip, band, adhesive, and/or the like. For example, the motion detector device is 30 affixed to the torso, or around the neck with a strap.

The device **12** can be configured with physically connected pieces with wired communication or physically separated pieces communicatively connected **36** via a wireless communication device, such as through a radio frequency (RF) transceiver or near body communication device. For example, the motion device communicates with an

electrocardiogram (ECG) monitoring device and a photoplethysmogram (PPG) communication through skin contact and the near body communication. In another example, the pieces communicate using Bluetooth or 802.x protocols and RF communications.

With reference to FIGURE 2 one embodiment of the motion triggered medical monitoring device **12** is schematically illustrated. The device **12** includes the motion detection device or means **20** to detect whole body motion indicative of orthostatic stress. For example, the motion detection device **20** includes an accelerometer, gyroscope, barometer, GPS receiver and/or the like to detect body motion and a processor or other software device which analyzes the body motion to determine whether whole body motion has occurred. The analysis can be based on changes in elevation, relative vertical motion vectors, horizontal motion vectors, and the like.

In response to the detected body motion being determined to be whole body motion, e.g. a vertical motion vector exceeding the threshold, a signal is communicated to the vital sign monitoring device **26**. The vital sign monitoring device, in one embodiment, includes a cuff-less blood pressure (BP) measuring device such as two pulse sensing devices, e.g. ECG, PPG, and the like, which are used to determine the pulse wave velocity between the pulse sensing devices. The blood pressure is proportional to the pulse wave velocity. The cuff-less BP measuring device uses the signals of the ECG and PPG to measure or determine BP, e.g. systolic/diastolic blood pressure. The measures can be continuous or repeated measures over the time interval.

A vital sign evaluation device or means **38** is communicatively connected to the vital sign monitoring device **26**, receives the monitored vital signs, and evaluates the received vital signs for symptoms of dizziness, fainting, and/or syncope which can cause the person to fall. For example, with the systolic blood pressure less than 90 mm Hg, or the diastolic blood pressure less than 60 mm Hg, the vital sign evaluation device generates an alarm signal in response. In another example, with a change in systolic or diastolic greater than 20 mm Hg, the vital sign evaluation device generates an alarm signal in response. The vital sign evaluation device can also compare other physiological values, such as heart rate or SpO₂, to thresholds. In another embodiment, time evolution patterns of physiological parameters are compared with patterns indicative of syncope or other potential events.

The vital sign evaluation device is communicatively connected to one or more alarm devices **40**. The alarm device includes an audible and/or visual indicator, such as sounded alarm and/or flashing light. The alarm device includes the computing device of the caregiver, which can indicate the alarm, the person's identity, and/or location. For example,

as the person stands, the motion detection device **20** detects an elevated whole body motion of 20 cm which triggers monitoring blood pressure by the vital sign monitoring device for a 45 minute period. During the 45-minute period, the vital sign evaluation device evaluates the monitored blood pressure, determines whether blood pressure drops more than 20 mm Hg
5 and, in response to the evaluated drop in blood pressure, generates the alarm signal. The alarm signal received by the alarm device sets the audible indicator and flashing light attached to the medical monitoring device **12**, indicating to the caregiver that the person is in danger of falling. Simultaneously, the alarm signal is communicated to the portable computing device **32**, which records the alarm signal, and/or indicates with a flashing
10 message display and sound that the person is in danger of falling.

The motion detection device **20**, vital sign monitoring device **26**, vital sign evaluation device **38**, and alarm device **40** are communicatively connected with a communication device or means **42**. The communication device **42** includes a wired connection in a wired configuration with devices physically connected. The communication
15 device **42** includes a wireless connection in a wireless configuration with devices physically separated. The wireless connection includes RF, infrared, or near body communication. The medical monitoring device can include different configurations between component devices. For example, the motion detection device can be in the wired configuration with the ECG device and in a wireless configuration with the PPG device.

With reference to FIGURE 3, another embodiment of the motion triggered vital sign measurement system **10** in a time sequence is diagrammatically illustrated. The person **14** moves from the resting state **16** at time t_0 to standing **18** at time t_1 . The motion detection device includes an optical motion detector **50**, such as one or more cameras and a processor configured to view body motion of the person, e.g. identify and track movement
25 indicative of orthostatic stress. For example, the camera receives images of the person in the field of view of the camera moving from the sitting position to the standing position. The processor communicatively connected to receive the images uses object recognition to identify the person and analyze the motion. For example, initial objects can include facial recognition, a badge or device worn by the patient, and the like, and then a growing algorithm to identify and associate other body parts. Alternatively, the person wears an
30 energy emitter, such as an LED, ultrasound emitter, infrared LED, RF transmitter, or the like. The emitted energy is received by appropriate stationary mounted receivers which use triangulation to monitor motion and distance and duration moved.

The medical monitoring device **12**, in communication with the optical motion detector, receives a signal indicating whole body motion. The medical monitoring device **12**, in one embodiment, includes a cuff-based vital sign monitoring device **52** adapted to monitor blood pressure of the person. The medical monitoring device **12** includes the components 5 integrated as a single physical device adapted to fit over the upper arm. For example, the medical monitoring device includes a pressure cuff to measure blood pressure, and the pressure cuff slides over the upper arm of the person and is adjustable, to be held in place by an attachment mechanism, such as elastic band, hook and loop, etc. The vital sign evaluation device and alarm device are physically attached to the pressure cuff. The cuff can include a 10 region of a characteristic color to assist in discriminating between monitored persons and bystanders.

With reference to FIGURE 4, another embodiment of the medical monitoring device **12** with an optical motion detection device **50** or means is schematically illustrated. The optical motion detection device **50**, such as the camera and configured or programmed 15 processor, views the person at time t_0 person in the sitting position, and at time t_1 person moving to standing position. The optical motion detection device **50** uses object recognition with motion separation to determine a change in position, e.g. a change in altitude of a center of mass, a characteristic point, or the like, giving rise to orthostatic stress. The motion detection device signals the vital sign monitoring device **52**, e.g. pressure cuff measuring 20 blood pressure for the predetermine time interval. The vital sign evaluation device **38** in response to blood pressure indicative of a minimum blood pressure, or a threshold change in blood pressure, signals the alarm device **54**, which includes a speaker that sounds the audible signal.

In one embodiment, the system is a person worn fall prevention system which 25 alerts the person or caregiver or relative of a potential fall before the fall occurs. In another embodiment, the system is a fall detection system that signals for help before or when a fall occurs.

The various devices **20**, **26**, **38**, **40**, **50**, **52** include or share one or more electronic data processing devices, such as a processor or electronic processing device 30 powered by either solar and/or battery power. The optical motion detection device **50** can include a AC/DC power source. Moreover, the disclosed vital sign evaluation and communication techniques are suitably implemented using a non-transitory storage medium storing instructions (e.g., software) readable by an electronic data processing device and executable by the electronic data processing device to perform the disclosed techniques.

With reference to FIGURE 5, one method of motion triggered vital sign measurement is flowcharted. In a step or by a module **60**, body motion of a person is detected, visually and/or physically. The motion is analyzed to determine whether the motion is whole body motion indicative of orthostatic stress, such as a change in elevation by moving from 5 the sitting to the standing position.

At least one vital sign is monitored by the vital sign monitoring device for a predetermined time interval in response to the detected whole body motion in a step or by a module **62**. For example, in response to the person standing, the blood pressure of the person is monitored using a cuff-less **26** or pressure cuff **42** blood pressure monitor.

10 In a step or by a module **64**, the at least one measured monitored vital sign is evaluated for orthostatic stress. For example, blood pressure is monitored for a minimum blood pressure, or a threshold change in blood pressure.

15 The alarm is signaled in response to an evaluated measure in a decision step or by a module **66**. For example, in response to monitored blood pressure dropping below a minimum value or changing more than a threshold value, the alarm condition is signaled.

20 The alarm is sent in response to the alarm signal in a step or by a module **68**. For example, the alarm signal causes an audible sound, tone, noise, or voice message indicative of the person in danger of falling. In another example, the alarm signal sets both the audible indicator and a visual indicator such as a light signal or a message on a caregiver display.

25 The communicating of conditions between modules or devices includes the detected motion, the monitored vital signs, the evaluated measures, and the alarm signal. The communicating can include devices physically distributed or separated. The devices or modules can include attachment mechanisms which attached the device or component to the person. A non-transitory computer-readable storage medium carrying software which controls one or more electronic data processing devices to perform the steps of communicating, detecting and/or analyzing motion, monitoring vital signs, evaluating vital signs, and signaling an alarm.

30 It is to be appreciated that in connection with the particular illustrative embodiments presented herein certain structural and/or functional features are described as being incorporated in defined elements and/or components. However, it is contemplated that these features may, to the same or similar benefit, also likewise be incorporated in other elements and/or components where appropriate. It is also to be appreciated that different aspects of the exemplary embodiments may be selectively employed as appropriate to

achieve other alternate embodiments suited for desired applications, the other alternate embodiments thereby realizing the respective advantages of the aspects incorporated therein.

It is also to be appreciated that particular elements or components described herein may have their functionality suitably implemented via hardware, software, firmware or a combination thereof. Additionally, it is to be appreciated that certain elements described herein as incorporated together may, under suitable circumstances, be stand-alone elements or otherwise divided. Similarly, a plurality of particular functions described as being carried out by one particular element may be carried out by a plurality of distinct elements acting independently to carry out individual functions, or certain individual functions may be split up and carried out by a plurality of distinct elements acting in concert. Alternately, some elements or components otherwise described and/or shown herein as distinct from one another may be physically or functionally combined where appropriate.

In short, the present specification has been set forth with reference to preferred embodiments. Obviously, modifications and alterations will occur to others upon reading and understanding the present specification. It is intended that the invention be construed as including all such modifications and alterations insofar as they come within the scope of the appended claims or the equivalents thereof. That is to say, it will be appreciated that various of the above-disclosed and other features and functions, or alternatives thereof, may be desirably combined into many other different systems or applications, and also that various presently unforeseen or unanticipated alternatives, modifications, variations or improvements therein may be subsequently made by those skilled in the art, which are similarly intended to be encompassed by the following claims.

CLAIMS:

What is claimed is:

1. A medical monitoring device (12), comprising:
 - a motion detection device (20, 50) adapted to detect whole body motion of a person; and
 - a vital sign monitoring device (26, 52) communicatively connected to the motion detection device and adapted to monitor at least one vital sign for a predetermined time interval, either continuously or intermittently, in response to the detected whole body motion.
2. The device according to claim 1, wherein the detected whole body motion (22) includes a determined change in elevation (24).
3. The device according to either one of claims 1 and 2, wherein the determined change in elevation (24) corresponds to a position of the person changing from a sitting position (16) to a standing position (18).
4. The device according to any one of claims 1-3, wherein the motion detection device (20, 50) includes at least one of:
 - an optical monitoring device adapted to monitor movements of the person;
 - an ultrasound monitoring device adapted to monitor movements of the person;
 - an accelerometer;
 - a gyroscope;
 - a barometer; or
 - a GPS receiver.
5. The device according to any one of claims 1-4, wherein the vital sign monitoring device (26, 52) is configured to measure blood pressure of a person, and includes either a cuff-less (26) or a cuff-based (52) blood pressure measuring device.

6. The device according to claim 5, wherein the cuff-less blood pressure measuring device **(26)** includes a plurality of pulse sensing devices.
7. The device according to any one of claims 1-6, further including:
 - a vital sign evaluation device **(38)** adapted to evaluate measures of the at least one monitored vital sign for orthostatic intolerance.
8. The device according to claim 7, wherein the evaluated measures include at least one of:
 - a minimum systolic or diastolic blood pressure; or
 - a threshold reduction in systolic or diastolic blood pressure.
9. The device according to either one of claims 7 and 8, wherein the vital sign evaluation device **(38)** is further configured to signal an alarm in response to an evaluated measure.
10. The device according to any one of claims 1-9, further including:
 - a communications device **(42)** adapted to communicate at least one of:
 - a prediction that a fall is imminent;
 - the detected motion;
 - the monitored at least one vital sign;
 - the evaluated measures; or
 - the alarm signal.
11. A method of monitoring vital signs, comprising:
 - detecting **(60)** whole body motion of a person; and
 - monitoring **(62)** at least one vital sign for a predetermined time interval in response to the detected whole body motion.
12. The method according to claim 11, wherein detecting whole body motion includes determining a change in elevation.
13. The method according to either one of claims 11 and 12, wherein monitoring **(62)** includes:
 - measuring blood pressure of the person.

14. The method according to either one of claims 11 and 12, further including:

- evaluating **(64)** measures of the at least one monitored vital sign for orthostatic stress; and
- signaling **(68)** an alarm in response to an evaluated measure.

15. The method according to any one of claims 11-14, further including:

- communicating at least one of:
 - a prediction that a fall is imminent;
 - the detected motion;
 - the monitored at least one vital sign;
 - the evaluated measures; or
 - the alarm signal.

16. A non-transitory computer-readable storage medium carrying software which controls one or more electronic data processing devices to perform the method according to any one of claims 11-15.

17. An electronic data processing device configured to perform the method according to any one of claims 11-15.

18. A medical monitoring device **(12)**, comprising:

- a motion detection device **(20, 50)** adapted to detect body motion of a person indicative of a change in position from sitting to standing;
- a vital sign monitoring device **(26, 52)** communicatively connected to the motion detection device and adapted to monitor blood pressure, either continuously or intermittently, for a predetermined time interval in response to the detected body motion;
- a vital sign evaluation device **(38)** adapted to evaluate the monitored blood pressure for conditions indicative of orthostatic intolerance; and
- an alarm device **(40, 54)** configured to generate a signal in response to the evaluated conditions.

19. The device according to claim 18, further including:

- an attachment mechanism **(34)** adapted to attach the motion detection device to the person to at least one of: around the neck, to the torso, or upper arm.

20. The device according to either one of claims 18 and 19, wherein the signal predicts or indicates the person falling.

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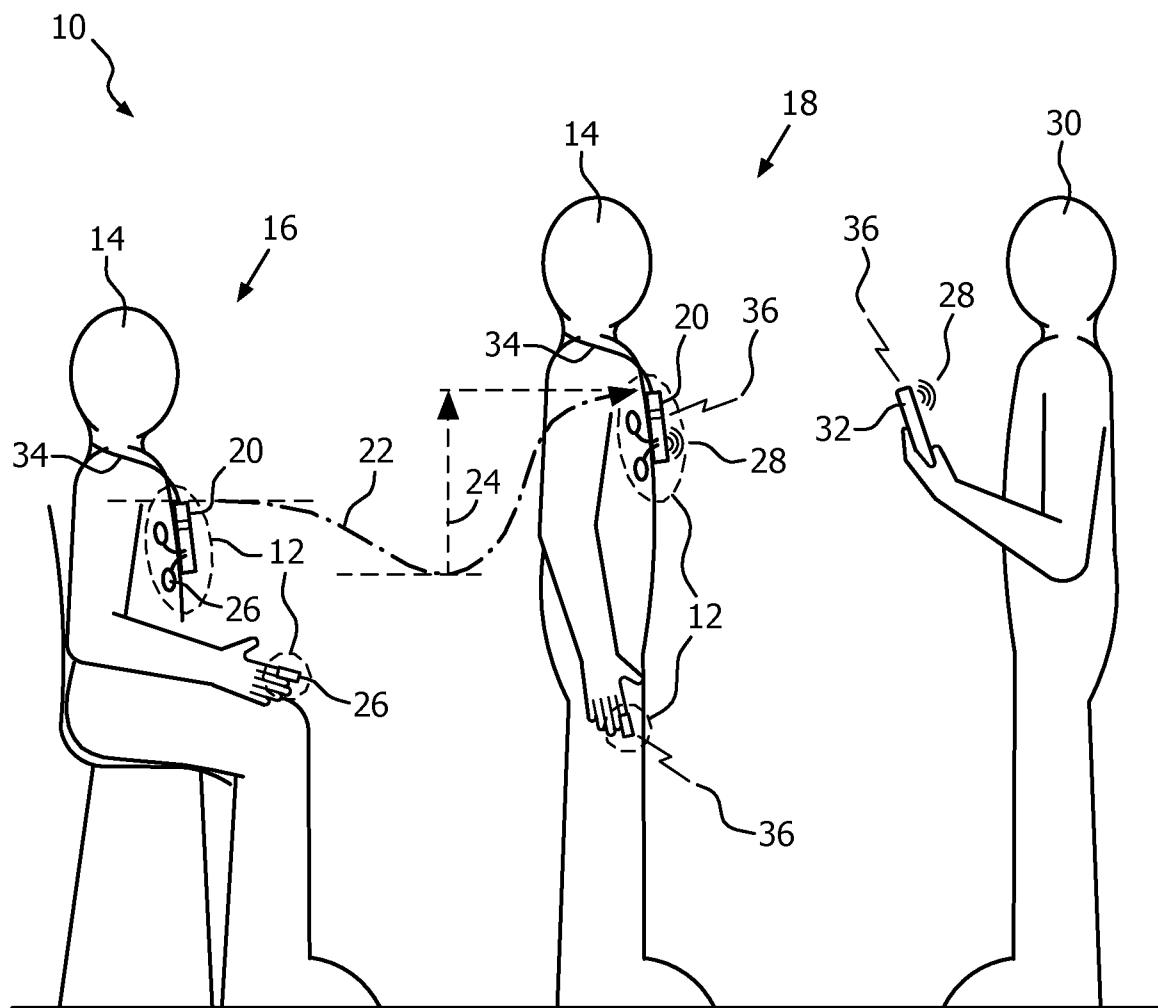


FIG. 1

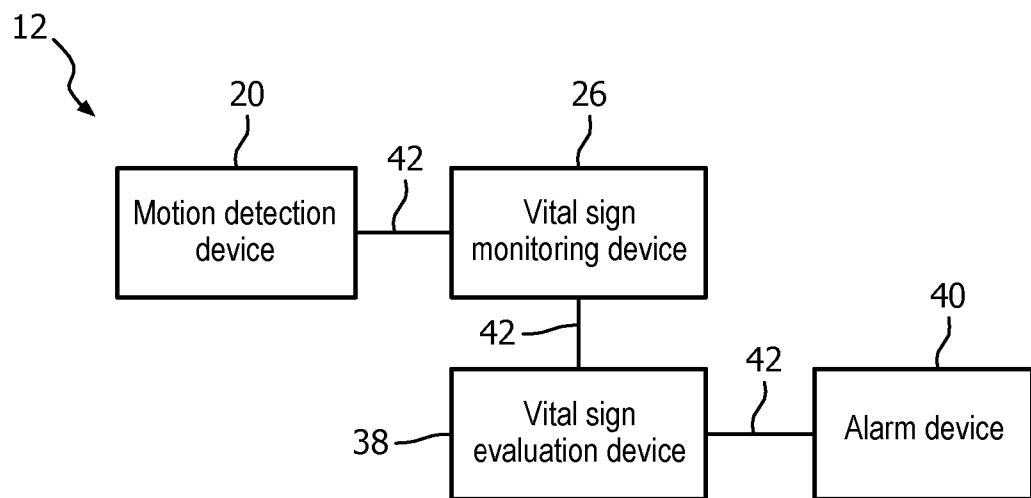


FIG. 2

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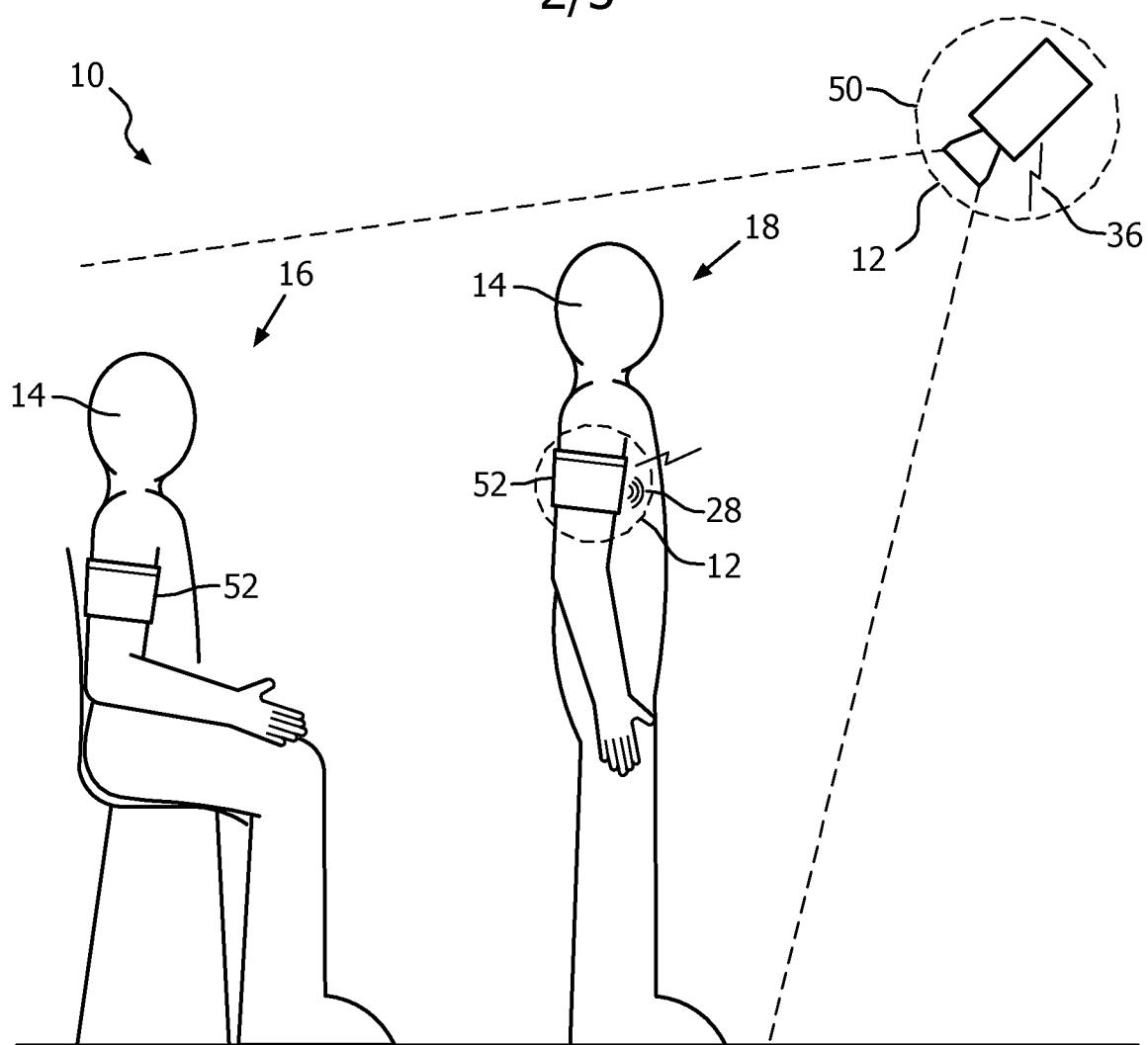


FIG. 3

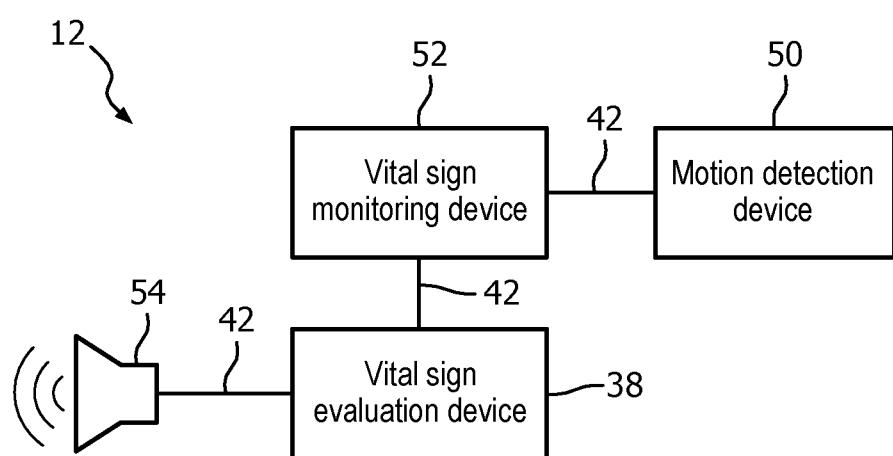


FIG. 4

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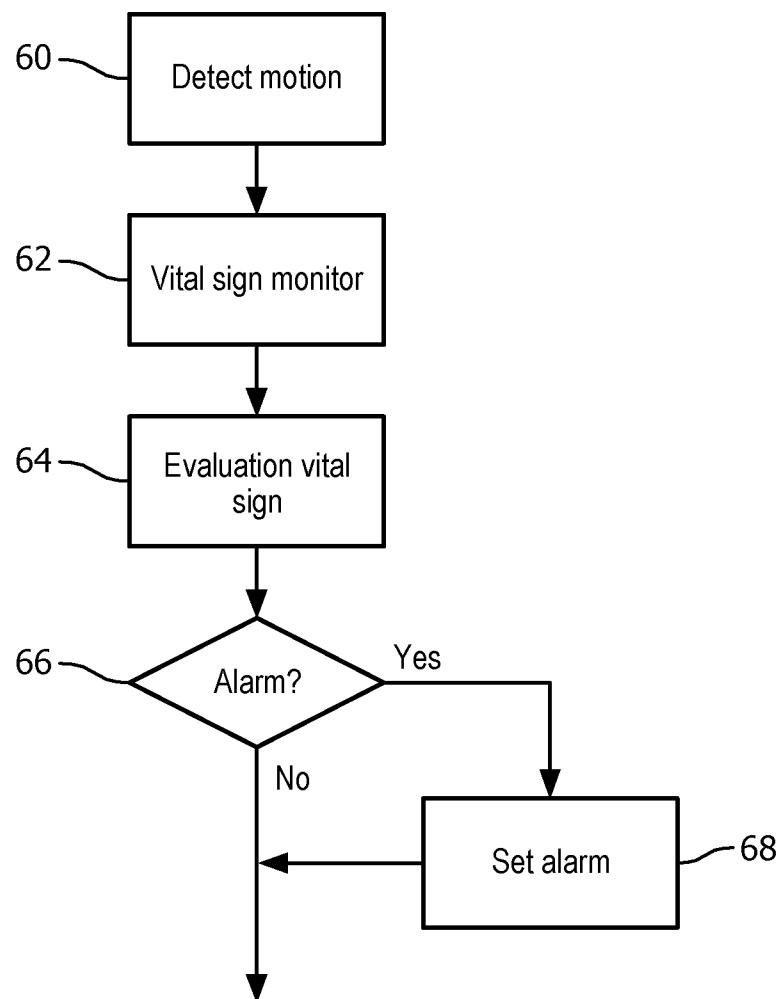


FIG. 5

INTERNATIONAL SEARCH REPORT

International application No

PCT/IB2015/052964

A. CLASSIFICATION OF SUBJECT MATTER
 INV. A61B5/0205 A61B5/11
 ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2004/215263 A1 (VIRAG NATHALIE [CH] ET AL) 28 October 2004 (2004-10-28) paragraph [0029] - paragraph [0030] paragraph [0041] paragraph [0048] - paragraph [0054]; figure 5 paragraph [0064] - paragraph [0068]; figure 7 ----- -/-	1-5,7, 9-14, 16-20

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

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- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

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Date of the actual completion of the international search

14 July 2015

Date of mailing of the international search report

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Name and mailing address of the ISA/

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 Fax: (+31-70) 340-3016

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INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2015/052964

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2010/113889 A1 (GHANEM RAJA N [US]) 6 May 2010 (2010-05-06) paragraph [0030] - paragraph [0035] paragraph [0100] paragraph [0113] - paragraph [0117] paragraph [0121] paragraph [0147] paragraph [0151] paragraph [0160] - paragraph [0161] -----	1-19
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