



(11)

**EP 1 558 193 B1**

(12)

## EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention of the grant of the patent:

**04.09.2019 Bulletin 2019/36**

(21) Application number: **03768544.3**

(22) Date of filing: **23.10.2003**

(51) Int Cl.:

**A61H 31/00** <sup>(2006.01)</sup> **A61B 5/00** <sup>(2006.01)</sup>  
**A61B 5/04** <sup>(2006.01)</sup> **A61B 5/053** <sup>(2006.01)</sup>  
**A61B 5/044** <sup>(2006.01)</sup> **A61M 16/00** <sup>(2006.01)</sup>  
**A61N 1/39** <sup>(2006.01)</sup> **G06K 9/00** <sup>(2006.01)</sup>

(86) International application number:

**PCT/US2003/034035**

(87) International publication number:

**WO 2004/037154 (06.05.2004 Gazette 2004/19)**

(54) **DEVICE FOR MEASURING DEPTH OF CHEST COMPRESSIONS DURING CARDIO-PULMONARY RESUSCITATION**

VORRICHTUNG ZUR BESTIMMUNG DER TIEFE VON KOMPRESSIENEN BEI DER KARDIOPULMONALEN WIEDERBELEBUNG

DISPOSITIF DE DETERMINATION DE LA PROFONDEUR DE COMPRESSIONS LORS DE REANIMATION CARDIO-PULMONAIRE

(84) Designated Contracting States:  
**AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LI LU MC NL PT RO SE SI SK TR**

(30) Priority: **25.10.2002 US 280220**

(43) Date of publication of application:  
**03.08.2005 Bulletin 2005/31**

(60) Divisional application:  
**12003127.3 / 2 532 338**  
**12003128.1 / 2 532 339**  
**12003129.9 / 2 532 340**  
**19186577.3**

(73) Proprietor: **ZOLL Circulation, Inc.**  
**San Jose, CA 95131 (US)**

(72) Inventors:

- **PALAZZOLO, James, A.**  
**Sunnyvale, CA 94089 (US)**
- **BERGER, Ronald D.**  
**Baltimore, MD 21287 (US)**
- **HALPERIN, Henry R.**  
**Baltimore, MD 21287 (US)**
- **SHERMAN, Darren, R.**  
**Sunnyvale, CA 94089 (US)**

(74) Representative: **Potter Clarkson**  
**The Belgrave Centre**  
**Talbot Street**  
**Nottingham NG1 5GG (GB)**

(56) References cited:

**WO-A2-02/15836** **US-A- 6 016 445**  
**US-A1- 2001 047 140** **US-A1- 2002 055 694**  
**US-B1- 6 171 267** **US-B1- 6 263 238**  
**US-B1- 6 390 996**

- "Citation Classics" CC, CURRENT CONTENTS, vol. 44, 30 October 1989 (1989-10-30), page 22, XP002560863
- SARAH MCCABE ET AL: "On the Use of Nonlinear Autoregressive Moving Average Models for Simulation and System Identification", PROCEEDINGS OF THE AMERICAN CONTROL CONFERENCE. BOSTON, JUNE 26 - 28, 1991; [PROCEEDINGS OF THE AMERICAN CONTROL CONFERENCE], NEW YORK, IEEE, US, 26 June 1991 (1991-06-26), pages 1758-1763, XP031430329, ISBN: 978-0-87942-565-4
- Anonymous: "Autoregressive-moving-average model - Wikipedia", , 12 July 2017 (2017-07-12), XP055406812, Retrieved from the Internet: URL: [https://en.wikipedia.org/wiki/Autoregressive-moving-average\\_model](https://en.wikipedia.org/wiki/Autoregressive-moving-average_model) [retrieved on 2017-09-14]

Note: Within nine months of the publication of the mention of the grant of the European patent in the European Patent Bulletin, any person may give notice to the European Patent Office of opposition to that patent, in accordance with the Implementing Regulations. Notice of opposition shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

**EP 1 558 193 B1**

- Anonymous: "ARMA-Modell - Wikipedia", , 8 August 2017 (2017-08-08), XP055406814, Retrieved from the Internet:  
URL:<https://de.wikipedia.org/wiki/ARMA-Modell> [retrieved on 2017-09-14]
- Anonymous: "Box-Jenkins method - Wikipedia", , 26 December 2016 (2016-12-26), XP055406820, Retrieved from the Internet:  
URL:[https://en.wikipedia.org/wiki/Box-Jenkins\\_method](https://en.wikipedia.org/wiki/Box-Jenkins_method) [retrieved on 2017-09-14]
- Anonymous: "Box-Jenkins-Methode - Wikipedia", , 15 February 2015 (2015-02-15), XP055406816, Retrieved from the Internet:  
URL:<https://de.wikipedia.org/wiki/Box-Jenkins-Methode> [retrieved on 2017-09-14]
- Anonymous: "Autoregressive integrated moving average - Wikipedia", , 13 September 2017 (2017-09-13), XP055406822, Retrieved from the Internet:  
URL:[https://en.wikipedia.org/wiki/Autoregressive\\_integrated\\_moving\\_average](https://en.wikipedia.org/wiki/Autoregressive_integrated_moving_average) [retrieved on 2017-09-14]

**Description**Field of the Inventions

**[0001]** The methods and devices described below relate to the field of cardio-pulmonary resuscitation (CPR).

Background of the Inventions

**[0002]** The American Heart Association guidelines for the correct application of cardio-pulmonary resuscitation (CPR) specify that chest compressions be performed at the rate of 80 to 100 per minute and at a depth, relative to the spine, of 3.81 to 5.08 cm (1.5 to 2.0 inches). (Guidelines 2000 for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care, 102 Circulation Supp. I (2000).) However, CPR is physically and emotionally challenging, even for trained professionals.

**[0003]** Research has shown that manual chest compressions rarely meet the guidelines. See, for example, Ochoa et al., The Effect of Rescuer Fatigue on the Quality of Chest Compressions, Resuscitation, vol. 37, p.149-52. See also Hightower et al., Decay in Quality of Closed-Chest Compressions over Time, Ann Emerg. Med, 26(3):300-333, Sept. 1995. One of the difficulties of performing correct chest compressions is that the rescuer imprecisely judges the timing and depth of compressions, particularly when the rescuer becomes tired. Thus, if accurate and timely user feedback could be provided to the rescuer then the rescuer would be more likely to perform CPR correctly.

**[0004]** Various devices have been proposed to assist a rescuer in properly applying CPR. For example, Kelley, Apparatus for Assisting in the Application of Cardiopulmonary Resuscitation, U.S. Patent 5,496,257 (Mar. 5, 1996) shows a device that uses a pressure sensor to monitor compression forces and timing. Groenke et al., AED with Force Sensor, U.S. Patent 6,125,299 (Sep. 26, 2000) shows a device that uses a force sensor to measure the compression force applied to a patient's chest. However, these devices only measure the force applied to the chest and do not measure the actual depth of compressions. A given force can compress the chests of different patients by different amounts, so measuring only force will not provide sufficient or consistent feedback to the rescuer. In addition, force-based measurements may also be inaccurate because of intra-patient variation in thoracic morphology and compliance (stiffness).

**[0005]** CPR devices that use only accelerometers to measure depth of compressions, other than our own patented device shown in Halperin et al., CPR Chest Compression Monitor, U.S. Patent 6,390,996 (May 21, 2002), do not fully or accurately account for errors in the measured acceleration; nor do they account for drift in the starting points of compressions. In addition, the integration process necessary to derive the depth of compressions greatly compounds any errors in the measured acceleration.

US 6390996 B1, discussed above, discloses a device for aiding in the administration of cardiopulmonary resuscitation (CPR). A hand-held CPR chest compression monitor comprises a displacement detector comprising an accelerometer coupled to a microprocessor via an interface. The monitor illustrated in Figure 1 of US 6390996 B1 determines chest displacement from a double integration of an acceleration signal produced by accelerometer. A signaling mechanism comprises an audible indicator (i.e., a loud speaker), which has an input connected to microprocessor via the interface. Interface, in addition to a serial or parallel interface, may further comprise A/D and D/A converters, including a D/A converter for driving audio transducer to indicate the amount of displacement and to prompt CPR at the correct rate (80--100 compressions per minute). The output from accelerometer is routed through an A/D converter provided as part of interface for digitization and subsequent analysis by microprocessor. Similarly, the output from each of first and second gyros is routed to microprocessor via an A/D converter provided as part of interface. In accordance with one aspect of the disclosure of US 6390996 B1, the device signals the rescuer to prompt correct compressions.

**[0006]** It is important to correct for errors in the measured acceleration since the total depth of compressions should be within the relatively narrow range of 3.81 to 5.08 cm (1.5 to 2.0 inches). Numerical simulations have shown that a total error in acceleration as small as 0.05 cm/sec<sup>2</sup> (0.02 in/sec<sup>2</sup>) results in an error of 0.64 cm (0.25 inches) in displacement. Given the narrow depth range of optimal compressions, an error of 0.64 cm (0.25 inches) is unacceptable.

**[0007]** For example, Freeman, Integrated Resuscitation, U. S. Publication 2001/0047140 (Nov. 29, 2001) shows a device that uses an accelerometer as a compression sensor and mentions gauging chest depth with the accelerometer.

However, Freeman

enables no method to account for the errors inherent in using an accelerometer alone. Thus any measurement Freeman makes of chest compression depth is inaccurate.

**[0008]** Myklebust et al., System for Measuring and Using Parameters During Chest Compression in a Life-Saving Situation or a Practice Situation and Also Application Thereof, U.S. Patent 6,306,107 (Oct. 23, 2001) describes a device which uses a pressure pad, containing an accelerometer and a force activated switch, to determine the depth of depressions. However, Myklebust does not provide a means to measure compression depth using an accelerometer alone, nor does Myklebust account for some kinds of error in the measured value of chest compression depth (such as drift).

**[0009]** The problems inherent in the above devices show the difficulty of solving the problem of measuring chest

compression depth using only an accelerometer. Nevertheless, the basic concept of determining displacement from a measured acceleration is straightforward (in a system with a known starting position). Displacement is determined by double integrating the measured acceleration.

**[0010]** However, this method of measuring chest compression depth is complicated by at least three major sources of error: signal error, external acceleration error, and drift in the actual or measured starting points of compressions from the initial starting point of compressions. Signal error comprises errors in the measured acceleration due to electronic noise, the shaking of wires or cables, errors inherent in the accelerometer, and other sources of noise in the acceleration itself.

**[0011]** External acceleration error comprises errors introduced by accelerations applied to the patient and/or the accelerometer other than accelerations caused by CPR. For example, if the patient is being transported in an ambulance and a rescuer is applying manual CPR with a compression monitor, then the accelerometer will measure accelerations caused by road vibrations as well as accelerations caused by CPR. (If the ambulance hits a pot hole then a large spike may appear in the compression waveform.) The accelerometer, by itself, cannot distinguish between the accelerations caused by road noise and the accelerations caused by compressions. In other words, the accelerometer measures a combined acceleration and not just the accelerations caused by compressions. Accordingly, the compression monitor will report a displacement value different from the actual chest displacement.

**[0012]** Another source of error, drift, comprises systematic shifts in the actual or reported starting points of each compression over an entire series of compressions. The accelerometer has no "memory" of the initial starting position. Thus, as the rescuer applies compressions the reported depth waveform can start to drift. The compression monitor may indicate that the reported depth waveform is increasingly deeper than the actual waveform. This form of drift is referred to as positive drift. On the other hand, drift can also cause the compression monitor to report a depth waveform that is increasingly more shallow than the actual waveform. In other words, actual compression starting points are becoming increasingly deeper, but the compression monitor instead reports each starting point as close to the initial starting point. This form of drift is referred to as negative drift.

**[0013]** One cause of negative drift is a failure to allow the chest to return to a fully relaxed position. Absent correction, the accelerometer will begin measuring displacement from the new "initial" position. Thus, the compression monitor erroneously informs the rescuer that the current starting point is at the initial starting point. However, the actual depth of the current starting point is more than the depth reported by the compression monitor. As a result, the rescuer may compress the chest harder than he should to achieve the erroneous depth suggested by the compression monitor.

**[0014]** Another source of both types of drift is a change in the overall position of the accelerometer with respect to the patient. For example, if the accelerometer is not fully secured then the accelerometer may systematically slip. (This may also cause external acceleration error.) Yet another source of drift is expansion and contraction of the chest due to ventilation performed simultaneously with compressions. Other sources of drift may also exist. Each source of drift may be independent of the others and may not cancel each other out, so the compression monitor should be able to account for both positive and negative drift.

**[0015]** Notwithstanding drift resulting from erroneous operation, changes in the actual starting point of compressions do occur. For example, if one or more ribs break during CPR then the actual starting point of each compression may be closer to the spine (a phenomena known as chest remodeling). Other types of chest injury or disease that affect the structure and strength of the rib cage can also cause chest remodeling. Chest remodeling can be gradual, in which case a gradual shift occurs in the actual initial starting point of compressions. A compression monitor should be able to account for the difference between erroneous drift and an actual shift in the starting points of compressions.

**[0016]** These and other sources of error are compounded by integrating the acceleration. The errors caused by signal noise and drift cause the constants of integration to have a value other than zero. The non-zero constants of integration compound the errors already present in the acceleration.

**[0017]** Thus, the total compression depth reported by the compression monitor can be very inaccurate. Accordingly, methods are needed to accurately and precisely derive the depth of chest compressions from a measured acceleration.

**[0018]** US 6,390,996 (B1) and US 2002/055694 (A1) discloses a method in which chest compressions are measured and prompted to facilitate the effective administration of CPR.

**[0019]** WO 02/15836 (A2) discloses a method for determining displacement of a region of a patient's chest.

**[0020]** "Citation Classics", CC, CURRENT CONTENTS, (19891030), vol. 44, page 22 provides a citation for Box, G E P & Jenkins G M, Times series analysis: forecasting and control, San Francisco, CA, Holden-Day, (1970) 1976, 575 p.

**[0021]** SARAH MCCABE ET AL: "On the Use of Nonlinear Autoregressive Moving Average Models for Simulation and System Identification", PROCEEDINGS OF THE AMERICAN CONTROL CONFERENCE. BOSTON, JUNE 26 - 28, 1991; [PROCEEDINGS OF THE AMERICAN CONTROL CONFERENCE], NEW YORK, IEEE, US, 26 June 1991 (1991-06-26), pages 1758-1763, ISBN: 978-0-87942-565-4 describes the Use of Nonlinear Autoregressive Moving Average Models for Simulation and System Identification.

**[0022]** US 6,171,267 (B1) relates to a cardiopulmonary resuscitation method and apparatus that is adapted to performing high-impulse CPR includes providing a chamber having an expandable volume and a patient-contacting pad

that moves as a function of volume of the chamber and supplying a controlled quantity of a fluid to the chamber.

**[0023]** US 6,016,445 (A) relates to a method and apparatus for electrode and transthoracic impedance estimation

**[0024]** US 6,263,238 (B1) relates to an automatic external defibrillator having a ventricular fibrillation detector

## 5 Summary

**[0025]** The invention defined by the independent claim relates to a device for measuring depth of chest compressions, said device comprising:

- 10 an accelerometer capable of producing an acceleration signal corresponding to the acceleration of chest compressions;
- a first filter capable of producing a filtered acceleration signal from the acceleration signal;
- a means for integrating the filtered acceleration signal, capable of producing a velocity signal;
- 15 a second filter (capable of producing a filtered velocity signal from the velocity signal;
- a means for integrating the filtered velocity signal and capable of producing a measured depth signal corresponding to the measured depth of chest compressions;
- a processor, operably connected to the means for integrating the filtered velocity signal, programmed to produce an estimated actual depth signal corresponding to the estimated actual depth of chest compressions by calculating an autoregressive moving average of the measured depth signal; and
- 20 a user feedback system configured to:

- receive the estimated actual depth signal;
- provide, to a rescuer or automatic CPR device, information corresponding to the estimated actual depth of a current chest compression and information concerning compression phase quality and decompression phase quality, wherein the information concerning compression phase quality and decompression phase quality is provided by combining information gained from the acceleration, velocity and estimated actual depth signals.
- 25

**[0026]** The methods and devices described below provide for signal processing techniques that precisely and accurately derive the depth of chest compressions from a measured acceleration of chest compressions. Specifically, the methods and devices provided below provide for a means to correct chest displacement errors caused by signal error, external acceleration error, and drift. A moving average technique is used to produce an accurate measurement of compression depth. There is also disclosed a method according to which a change in the patient's ECG (electrocardiogram) may be used to determine the starting points of compressions. These methods may be combined together to further increase the accuracy of chest depth measurement.

**[0027]** In broad terms, a moving average technique disclosed herein averages a plurality of compression cycles together, but weights recent compressions more heavily than compressions further in the past. One moving average technique begins with filtering a raw acceleration signal to eliminate as much signal noise as practicable. The filtered acceleration signal is then integrated to derive the velocity of compressions. The velocity is filtered to remove accumulated low frequency variations. The filtered velocity measurement is integrated again to derive chest displacement. Chest displacement is then processed through a baseline limiter and a peak limiter; the baseline limiter may comprise a moving average processor and the peak limiter may comprise a moving average processor.

**[0028]** The baseline limiter estimates the actual starting point of the current compression and the peak limiter estimates the actual peak depth of the current compression. A baseline detector then identifies the starting point of the current compression. A peak detector then identifies the peak depth of the current compression. A means for combining signals then combines the estimated starting point and the estimated peak depth to derive the estimated actual depth of the current compression. Finally, the estimated actual depth of the current compression is provided to one or more devices which provide intelligible feedback to a manual CPR provider, to an automated CPR device, or to an ECG operator.

**[0029]** In another method disclosed herein, a change in the noise component of the patient's ECG is correlated to the start of a chest compression. When the noise component of the patient's ECG signal exceeds a pre-determined threshold then the accelerometer begins to measure acceleration. Thus, the actual starting point of the current compression is established. This method reduces some forms of external acceleration error and reduced drift. The method also helps to set the constants of integration to zero.

## 55 Brief Description of The Drawings

**[0030]**

Figure 1 shows a patient and an accelerometer-based compression monitor in place on a patient.

Figure 2 shows a graph of compression depth over time before signal processing, where compression depth is derived from a measured acceleration.

Figure 3 shows a graph of compression velocity over time before signal processing, where compression velocity is derived from a measured acceleration.

Figure 4 shows a graph of compression acceleration over time before signal processing, where compression acceleration is measured by an accelerometer.

Figure 5 is a flow chart of a signal processing technique that converts a raw compression acceleration into an estimated actual compression depth.

Figure 6 is a flow chart of an alternate signal processing technique that converts a raw compression acceleration into an estimated actual compression depth.

Figure 7 shows the graph of compression depth over time after filtering the raw acceleration.

Figure 8 shows the graph of compression velocity over time after filtering the raw acceleration.

Figure 9 shows the graph of compression acceleration over time after filtering the raw acceleration.

Figure 10 shows the graph of compression depth over time after filtering both the raw acceleration and the derived velocity.

Figure 11 shows the graph of compression velocity over time after filtering both the raw acceleration and the derived velocity.

Figure 12 shows the graph of compression acceleration over time after filtering the raw acceleration.

Figure 13 shows the graph of compression depth over time after filtering both the raw acceleration and the derived velocity, and after applying a baseline limiter to the compression depth waveform.

Figure 14 shows the graph of compression velocity over time after filtering both the raw acceleration and the derived velocity, and after applying the baseline limiter to the compression velocity waveform.

Figure 15 shows the graph of compression acceleration over time after filtering the raw acceleration and after applying the baseline limiter to the compression acceleration waveform.

Figure 16 shows the graph of compression depth over time after filtering both the raw acceleration and the derived velocity, and after applying the baseline limiter and the peak limiter to the compression depth waveform.

Figure 17 shows the graph of compression velocity over time after filtering both the raw acceleration and the derived velocity, and after applying the baseline limiter and the peak limiter to the compression velocity waveform.

Figure 18 shows the graph of compression acceleration over time after filtering the raw acceleration and after applying the baseline limiter and the peak limiter to the compression acceleration waveform.

Figure 19 is a flow chart of a signal processing technique that uses a change in ECG noise to activate a switch which, in turn, controls when an accelerometer begins to measure acceleration.

Figure 20 shows a graph of compression depth over time before signal processing and with a negative drift in the reported compression depth waveform.

Figure 21 shows a graph of compression velocity over time before signal processing and with a negative drift in the reported compression velocity waveform.

Figure 22 shows a graph of compression acceleration over time before signal processing and with a negative drift in the reported compression acceleration waveform.

Figure 23 shows the graph of Figure 20 corrected by using a change in ECG noise to establish the actual starting points of compressions.

Figure 24 shows the graph of Figure 21 corrected by using a change in ECG noise to establish the actual starting points of compressions.

Figure 25 shows the graph of Figure 22 corrected by using a change in ECG noise to establish the actual starting points of compressions.

Figure 26 shows an accelerometer-based compression monitor in place on a patient and a system of reference sensors comprising a reference accelerometer, a switch, and a load sensor disposed such that each sensor may measure various parameters related to chest compressions.

Figure 27 illustrates a compression waveform that a user feedback system may prompt the rescuer to perform.

Figure 28 is a block diagram of how an actual chest compression acceleration is converted into a corrupted value for chest position.

Figure 29 is a block diagram of a general solution for converting a corrupted chest compression acceleration into an estimated actual depth of chest compressions.

Figure 30 is a block diagram of how an actual ECG signal is converted into a corrupted ECG signal.

Figure 31 is a block diagram of a general solution for converting a motion corrupted ECG signal into an estimated actual ECG signal.

Figure 32 is a graph of a pig's ECG signal that is corrupted by noise caused by chest compressions.

Figure 33 is a graph of CPR motion where CPR is performed on a pig.

Figure 34 is a graph of the pig's estimated ECG noise signal.

Figure 35 is a graph of the pig's estimated actual ECG signal.

#### Detailed Description

**[0031]** Figure 1 shows a patient 1 and an accelerometer-based compression monitor 2 in place on the patient. An accelerometer-based compression monitor uses one or more accelerometers to determine the depth of compressions. An example of an accelerometer-based compression monitor may be found in our own patent, Halperin et al., CPR Chest Compression Monitor, U. S. Patent 6,390,996 (May 21, 2002).

**[0032]** The compression monitor 2 is placed on the sternum 3 of the patient 1, on the rescuer's hands or arms, or on an automatic CPR device. The chest is then compressed. The accelerometer measures the acceleration of compressions and a processor 4 estimates the actual displacement of the accelerometer based on the measured acceleration. The signal processing techniques described below ensure that the estimated actual displacement is accurate and precise.

**[0033]** The estimated actual displacement may be provided to a displacement display 5 that provides intelligible feedback to a manual CPR provider or to an automated CPR device. Likewise, other CPR-related parameters may be provided to one or more compression device displays 6 (or other means for user feedback). CPR-related parameters include the depth of chest compressions, the velocity of chest compressions, the acceleration of chest compressions, and the patient's ECG.

**[0034]** In the case of the patient's ECG, the compression monitor may be provided with one or more electrodes. The processor may process the patient's ECG during compressions to produce an estimated actual ECG. The estimated actual ECG may then be provided to an ECG display 7 (or other means for user feedback) that provides intelligible feedback to the manual CPR provider, to an automated CPR device, or to other individuals or devices that monitor the patient's ECG.

**[0035]** The following terms are used throughout the specification and are defined as follows:

Actual compression depth: the actual depth of a compression at any given time.

**[0036]** Actual starting point of a compression: the actual place or point at which a chest compression begins.

**[0037]** Autoregressive moving average: a function that uses past data samples to modify the current data sample.

**[0038]** Baseline portion of the compression depth waveform: that portion of depth waveform where the set of actual starting points is most likely to be found.

**[0039]** Baseline limiter: a processor or function that operates on the baseline portion of the compression depth waveform.

**[0040]** Compression Peak: the place or point where maximum compression depth occurs.

**[0041]** Current compression depth: the depth of a compression at any given time.

**[0042]** Current starting points: the starting point of the current compression.

**[0043]** Depth of compressions: the depth the chest is compressed at any instant in time, where depth is measured relative to the relaxed position of the chest.

**[0044]** Estimated actual starting point of a compression: the estimated value of the actual place or point at which a chest compression begins.

**[0045]** Initial starting point of compressions: the place or point at which a series of compressions begins.

**[0046]** Measured starting point of a compression: the measured value of the place or point at which a chest compression begins.

**[0047]** Moving average: a function that uses past data samples to modify the current data sample.

**[0048]** Past starting points: the starting points of compressions that have already occurred.

**[0049]** Peak portion of the compression depth waveform: that portion of depth waveform where the set of actual peaks are most likely to be found.

**[0050]** Starting point of a compression: the place or point at which a chest compression is begun.

**[0051]** Figures 2 through 4 show graphs of compression depth, velocity, and acceleration over time for four hypothetical compressions. No signal processing has been applied to any of waveforms shown in Figures 2 through 4. Compression depth in Figure 2 is shown as a positive value-the higher the value, the deeper the chest has been compressed. The phantom waveforms 12 represent the actual waveforms for compression depth, velocity, and acceleration (measured independently of the accelerometer). The solid waveforms 13 represents the waveforms derived from the acceleration measured by the compression monitor accelerometer. The waveforms 13 are also the waveforms reported by the compression monitor to the signal processing system 4. Compression depth is measured in inches, marked at 2.54 cm (1 inch) intervals, compression velocity is measured in cm (or inches) per second, cm/s (in/s), marked at 2.54 cm/s (1 in/s) intervals, and compression acceleration is measured in cm (or inches) per second per second, cm/s<sup>2</sup> (in/s<sup>2</sup>), marked at 2.54 cm/s<sup>2</sup> (1 in/s<sup>2</sup>) intervals.

**[0052]** For all three Figures time is measured in seconds, marked at 1 second intervals. The start of compressions is at time equal to zero. The initial depth of compressions is at depth equal to zero.

**[0053]** Phantom lines 14 and 15 intersect all three graphs.

**[0054]** Phantom line 14 corresponds to the time at which maximum compression depth is obtained. Phantom line 15 corresponds to the time at which minimum compression depth is obtained. In addition, phantom line 14 indicates that a compression depth maximum 16 corresponds to a compression velocity of zero.

**[0055]** Phantom line 14 also indicates that an acceleration maximum 17 is slightly offset from the compression depth maximum 16.

**[0056]** Likewise, phantom line 15 indicates that a compression minimum 18 (or starting point or zero point) corresponds to a compression velocity of zero. Phantom line 15 also indicates that an acceleration minimum 19 is slightly offset from the compression depth minimum 18. A compression velocity maximum 20 and minimum 21 occur around the middle of a compression.

**[0057]** The solid waveforms show the effects of three major types of error: signal error, external acceleration error, and drift. Signal error is primarily represented by the "noisy" (rough) nature of the solid waveforms; however, external acceleration error can also form a portion of the "noise." Although the acceleration waveform is less noisy, integrating the acceleration increases the effect of the noise in the velocity waveform. Integrating the velocity waveform increases the effect of the noise yet again. Thus, the compression depth noise Figure 2 is higher than the compression velocity noise in Figure 3, which is in turn higher than the compression acceleration noise in Figure 4. Accordingly, the compression monitor will report a very noisy compression depth waveform.

**[0058]** External acceleration error is primarily represented by the large, positive spike 22 in the solid waveforms of Figures 2 through 4. (Although the spike in Figures 2 through 4 occurs at a maximum, spikes can occur anywhere in the compression cycle and can affect the measured acceleration both positively and negatively). The spike is caused by a large acceleration unrelated to compressions, but nevertheless measured by the accelerometer. Thus, the actual waveform 12 in all three figures shows a corresponding peak 23 significantly below spike 22. Accordingly, absent the correction suggested here, the compression monitor will report for that compression cycle a compression depth much higher than the actual compression depth.



**[0059]** Drift is primarily represented by the increasing distance between the respective minimums of the actual and reported waveforms of Figures 2 through 4, as shown by arrows **24** and **25**. The drift is causing the compression monitor to erroneously report a compression waveform that is becoming increasingly deeper (positive drift). However, the actual waveform is more closely returning to the initial starting point, and is thus the drift shown in Figures 2 through 4 is considered a positive drift. Likewise, arrows **24** and **25** in Figures 3 and 4 illustrate that drift has an increasing affect on the reported velocity and the reported acceleration. The effects of drift mean that the initial starting point of compressions cannot be used as a reliable starting point for all compressions. Accordingly, the starting point of compressions must be determined for every compression cycle. In addition, the other sources of noise must be either eliminated or greatly reduced.

**[0060]** Figure 5 is a flow chart of a signal processing technique that converts a raw acceleration into an estimated actual value for total compression depth. The raw acceleration **34** is filtered by a first filter in step **35** to produce a filtered acceleration. The first filter comprises a high-pass filter and greatly reduces most forms of signal noise. (In other embodiments the first filter may comprise a band pass filter, a moving average filter, an infinite impulse response filter, an autoregressive filter, or an autoregressive moving average filter.) The effects of the other steps shown in Figure 5 are described in the context of Figures 7 through 18.

**[0061]** Figure 6 is a flow chart of an alternate signal processing technique that converts a raw compression acceleration into an estimated actual compression depth. This flowchart is described after the description for Figures 7 through 18.

**[0062]** The effect of the filter operation **35** is seen in figures 7 through 9, which show the graphs of compression depth, velocity, and acceleration over time for four hypothetical compressions after the first filtering step **35**. (Figures 7 through 9 show the output of the first filtering step). The measured acceleration waveform **13** of Figure 9 is much less noisy than the corresponding unfiltered waveform **13** of Figure 4. Since the velocity and depth waveforms of Figures 8 and 9 are derived from the acceleration waveform they, too, are less noisy. Nevertheless, the integration process still causes the velocity waveform to be more noisy than the acceleration waveform and the depth waveform to be more noisy than the velocity waveform. In addition, the external acceleration spike **22** still remains, as do the errors caused by drift (as shown by arrows **24** and **25**).

**[0063]** Returning to Figure 5, the filtered acceleration is integrated in a first integration step **36** to derive the compression velocity. However, as shown in Figure 8, without further processing the velocity waveform is still noisy. Thus, the velocity is filtered by a second filter in step **37** to produce a filtered velocity. The second filter comprises a high pass filter and further reduces most signal noise in the velocity and depth waveforms. (In other embodiments the second filter may comprise a band pass filter, a moving average filter, an infinite impulse response filter, an autoregressive filter, or an autoregressive moving average filter.)

**[0064]** The effects of the filter operation **37** is seen in figures 10 through 12, which show the graphs of compression depth, velocity, and acceleration over time for four hypothetical compressions after the second filtering step **37**. (Figures 10 through 12 show the output of the second filtering step **37**.) The measured velocity waveform **13** of Figure 11 is less noisy than that of Figure 8 (the velocity waveform after the first filtering step). Since the depth waveform is derived from the velocity waveform it, too, is correspondingly less noisy. Nevertheless, the integration process still causes the depth waveform to be slightly more noisy than the acceleration and velocity waveforms. In addition, the external acceleration spike **22** still remains, as do the errors caused by drift (as shown by arrows **24** and **25**).

Returning to Figure 5, the filtered velocity is integrated in a second integration step **38** to calculate the chest compression depth. Signal noise has been substantially eliminated and thus a third filtering step is not required.

However, the noise in the depth waveform, as shown in Figure 10, is still slightly more than the noise in the velocity waveform, as shown in Figure 11. Thus in other embodiments a third filter, comprising a high pass, bandpass, or other filter may be used to further reduce signal noise in the depth waveform.

After the initial filtering steps (35 and 37) and integration steps (36 and 38), a baseline limiter estimates the actual starting point of a compression in step 39. The baseline limiter uses, among other techniques described below, the starting points from past compressions to estimate the current compression starting point. The baseline limiter itself comprises a digital or analog signal processor that operates on the baseline portion of the compression depth waveform of Figure 10. The baseline portion of the compression depth waveform comprises that portion of depth waveform where the set of actual starting points is most likely to be found. For example, the baseline may comprise the portion of the depth waveform that is equal to and below 2.79 cm (1.1 inches) compression depth. (Larger changes in the starting points of compressions are unlikely, and signals indicating large changes are probably wrong.) Thus, the limiter will disregard or arbitrarily assign a realistic depth value to any "starting point" above 2.79 cm (1.1 inches) depth. In one embodiment, past starting points above the baseline are disregarded and a current starting point above the baseline is reported or treated as an error. (Past starting points are the starting points of compressions that have already occurred. A current starting point is the starting point of the current compression.) In another embodiment a current starting point above the baseline is assigned a small probability and averaged with the past starting points.

**[0065]** In one embodiment the baseline limiter estimates the starting point of the current compression by applying a moving average to all starting points that fall within the baseline portion of the depth waveform. A moving average is a

function that uses past data samples to modify the current data sample. (Additional moving average techniques are described below.) In the case of the baseline limiter, the baseline limiter may weigh recent starting points more heavily than older starting points, meaning that the weight of a given starting point decays over time. Starting points that fall outside the baseline portion of the depth waveform are given an arbitrary weight or no weight. By applying a moving average to all starting points the baseline limiter reduces the effect of external acceleration error and drift on the current starting point. In other words, the moving average of all starting points will be statistically closer to the current actual starting point than the current measured starting point derived from the integration of the acceleration.

**[0066]** The following example shows an embodiment of a moving average technique. In this embodiment each compression starting point is given a weight of 1.25% of the previous compression starting point. In other embodiments the weighting may comprise a percentage in the range of about 0.1% to about 12.5% (which yields between about 0.3% to about 90% data weighting at the end of about 1 minute). In other words, the measured value of the current starting point (starting point 1) is weighted 100%, the most recent starting point (starting point 2) is weighted 98.75%, the next previous starting point (starting point 3) is weighted 97.5%, the next previous starting point (starting point 4) is weighted 96.25%, etc until all compressions are weighted. Eventually, compressions in the distant past are given no virtually no weight at all. The depth of all the weighted starting points is then averaged. The weighted average of all starting points is treated or reported as the current starting point.

In another embodiment, all compressions after a pre-determined time period (such as about 1 minute to about 15 minutes) are disregarded. Thus, only compressions within the last 1 to 15 minutes are averaged. In another embodiment, all compressions after a pre-determined number of compressions (such as about 5 to about 15) are disregarded.

Continuing the example, in one embodiment the measured values for starting point 1 = 1.27 cm (0.5 inches), starting point 2 = 2.79 cm (1.1 inches), starting point 3 = 10.16 cm (4.0 inches), and starting point 4 = 2.29 cm (0.9 inches). Starting point 3 is outside the baseline portion of the depth waveform (the baseline portion is 1.1 inches and below in this example). Starting points outside the baseline in this example are disregarded, so starting point 3 is disregarded. Thus, the current starting point would be reported as:  $[(1.27 \times 100\%) + (2.79 \times 98.75\%) + (2.29 \times 96.25\%)] / 3 = 2.17 \text{ cm}$   $[(0.5 \times 100\%) + (1.1 \times 98.75\%) + (0.9 \times 96.25\%)] / 3 = 0.853 \text{ inches}$  relative to the initial starting point.

Had starting point 3 been included in the moving average, then the current starting point would have been reported as:  $[(1.27 \times 100\%) + (2.79 \times 98.75\%) + (10.16 \times 97.5\%) + (2.29 \times 96.25\%)] / 4 = 4.10 \text{ cm}$   $[(0.5 \times 100\%) + (1.1 \times 98.75\%) + (4.0 \times 97.5\%) + (0.9 \times 96.25\%)] / 4 = 1.615 \text{ inches}$  relative to the initial starting point.

Stated differently, this value is the estimated actual starting point for the current compression.

**[0067]** Mathematically, the reported value of the current starting point is expressed as:

$$Ds = [\sum (DB_i \cdot \omega^{i-1})] / n_r$$

where  $DB_i = 0$  if  $DB_i > B$ ,

where  $Ds$  is the depth of the current starting point,  $n_r$  is the number of starting points remaining after all starting points that exceed the baseline have been disregarded,  $i$  is the starting point number (or sum index),  $DB_i$  is the measured depth of the  $i^{\text{th}}$  starting point,  $\omega$  is the weighting constant, and  $B$  is the baseline. Expressed differently,  $DB_i \cdot \omega^{i-1}$  is summed from  $i = 1$  to  $n_r$  and the sum is divided by  $n_r$ , but if a particular  $DB_i$  is greater than  $B$  then that  $DB_i$  is instead set to zero.

The baseline limiter may perform other functions to further increase the accuracy and precision of the estimated depth of the current starting point. For example, a probability can be assigned to a given change between the current starting point and the immediate previous starting point. (Likewise a probability can be assigned to a given change between the current starting point and the moving average of all previous starting points.) Large changes in starting point may be given less weight than smaller changes. This technique may be referred to as a "weighted moving average" technique.

Continuing the above example, measured depth 1 is treated as having a 100% probability of occurring. Then, the difference between the current starting point (depth 1) and the previous starting point (depth 2) is 2.78 cm - 1.27 cm = 1.52 cm (1.1 inches - 0.5 inches = 0.6 inches). The probability of a step of 1.52 cm (0.6 inches) occurring is assigned to be 97%, based on past experiments. Since the probability is not 100%, the current starting point is not treated as having jumped a full 1.52 cm (0.6 inches). Instead,

the current starting point is treated as having jumped  $0.25 \text{ cm} \times 0.97 = 0.582 \text{ cm}$  ( $0.6 \text{ inches} \times 0.97 = 0.582 \text{ inches}$ ). Accordingly when calculating the weighted moving average depth 2 is treated as being 2.75 cm (1.082 inches) and not 2.78 cm (1.1 inches). Starting point 3 is still disregarded. The difference between starting point 2 (2.78 cm (1.1 inches)) and starting point 4 (2.29 cm (0.9 inches)) is 0.51 cm (0.2 inches), which is assigned a 99% probability. Thus, the effective distance of the step between depth 2 and depth 4 is  $0.51 \text{ cm} \times 99\% = 0.50 \text{ cm}$  ( $0.2 \text{ inches} \times 99\% = 0.198 \text{ inches}$ ). Accordingly, depth 4 is treated as 2.291 cm (0.902 inches) instead of 2.286 cm (0.9 inches). Using the same moving average as above, the current starting point is now reported as:  $[(1.27 \times 100\%) + (2.748 \times 98.75\%) + (2.291 \times 96.25\%)] / 3 = 2.06 \text{ cm}$   $[(0.5 \times 100\%) + (1.082 \times 98.75\%) + (0.902 \times 96.25\%)] / 3 = 0.812 \text{ inches}$  relative to the initial starting point.

Stated differently, this value is the estimated actual starting point for the current compression.

**[0068]** Mathematically, the reported value of the current starting point is expressed as:

$$Ds = \{ \sum [DB_{i-j} + (DB_i - DB_{i-j}) * P_s] * \omega^{i-1} \} \div n_r$$

where  $DB_i = 0$  if  $DB_i > B$ ,

where  $Ds$  is the depth of the current starting point,  $n_r$  is the number of starting points remaining after all starting points that exceed the baseline have been disregarded,  $i$  is the starting point number,  $DB_i$  is the measured depth of the  $i^{\text{th}}$  starting point,  $j$  is the index for the most recent starting point that was still within the baseline,  $DB_{i-j}$  is the most recent starting point that was still within baseline,  $P_s$  is the probability that a step of size  $DB_i - DB_{i-j}$  will occur,  $\omega$  is the weighting constant, and  $B$  is the baseline. The result,  $Ds$ , is the reported depth of the current starting point. Expressed differently,  $[DB_{i-j} + (DB_i - DB_{i-j}) * P_s] * \omega^{i-1}$  is summed from  $i = 1$  to  $n$  and the sum is divided by  $n_r$ , but if a particular  $DB_i$  is greater than  $B$  (the baseline) then that  $DB_i$  is instead set to zero.

**[0069]** In another embodiment, a probability is assigned to the step size between the depth of the current starting point and the weighted average of all previous starting points. (In the above example, the probability is assigned to a step size between the current starting point and the immediate past starting point). This technique may be referred to as a "weighted moving average with memory" technique. In this technique the reported depth of the current starting point is expressed mathematically as:

$$Ds = \{ \sum [DB_{i-j} + (DB_i - Ds_{i-j}) * P_s] * \omega^{i-1} \} \div n_r$$

where  $Ds_{i-j} = [\sum (DB_{i-j} * \omega^{i-2})] \div n_r$  and  $DB_i = 0$  if  $DB_i > B$ ,

where the variables are defined above. Again, the value for  $Ds$  is also the estimated actual starting point for the current compression.

**[0070]** In another embodiment, an autoregressive moving average (ARMA) filter may be used as the baseline limiter. The ARMA filter is an exponentially decaying "forgetting" filter that weights more current data more heavily than past data. The ARMA operates on more than just the compression starting point or peak values. Instead, the ARMA filter operates on data samples of compression acceleration, velocity, or depth taken at rapid time intervals. Data samples may be taken at a rate of about 100 samples per second to about 2000 samples per second (with a rate of about 1000 samples per second preferred). Thus, the ARMA filter operates on the entire waveform and not just on the compression peaks and the starting points.

**[0071]** In low pass form (which eliminates high frequency variations in the baseline) the ARMA filter may be expressed mathematically as:

$$y[n] = (1-\alpha) * y[n-1] + \alpha * x[n].$$

**[0072]** In this case,  $n$  is the index of the current sample (the " $n^{\text{th}}$ " sample),  $y[n]$  is the output of the current sample,  $x[n]$  is the input of the current sample,  $y[n-1]$  is the output from the previous sample, and  $\alpha$  is an independent term that determines how fast the filter "forgets" past outputs and the amount of influence the current input has on the output. The value for  $\alpha$  may be in the range of about 0.02 to about 0.0002, with a value of about 0.002 being suitable for many CPR-related filter applications. Should it be desired to implement a high-pass ARMA filter for the baseline limiter, then the ARMA equation becomes:

$$y[n](\text{high pass}) = 1 - \{ (1-\alpha) * y[n-1] + \alpha * x[n] \},$$

where  $y[n](\text{high pass})$  is the high pass filter output and the other variables are defined in the context of the low pass ARMA filter. The high pass filter may be used to eliminate low-frequency variations in the depth, velocity, or acceleration signals.

**[0073]** The moving average techniques in the above examples have been described in the context of processing the compression depth waveform. However, the techniques can be used to process the velocity waveform and the acceleration waveform, should it be desired to report accurate values for the velocity and acceleration of compressions. The moving average techniques may be applied to each waveform separately. In other words, one does not necessarily apply a moving average technique to the acceleration waveform, then integrate the acceleration waveform, then apply a second moving average technique to the velocity waveform, then integrate the velocity waveform, and finally apply a third moving average technique to the depth waveform. However, in other embodiments this procedure may be used.

**[0074]** Other methods for analyzing the baseline signal may be used to determine the estimated actual starting point of compressions. Another embodiment of the baseline limiter comprises a signal processor that uses a transition probability map to identify the probability of particular shifts in the measured starting point. (The probability map may be pre-determined, such as by using a density estimator or kernel estimator, and then hard-coded into the compression monitor software.) A particular starting point measurement is compared to the probability map and the system determines by how much a given shift in the measured starting point is erroneous. The reported starting point is adjusted accordingly. (Likewise, a transition probability map may be used to estimate the actual peak and also the actual maximum depth for each compression.)

**[0075]** The effect of the baseline limiter **39** is seen in Figures 13 through 15, which show the graphs of compression depth, velocity, and acceleration over time for four hypothetical compressions. Figures 13 through 15 also show the output of steps **35** through **39** in Figure 5. The baseline limiter has been applied separately to the velocity waveform (Figure 14) in step **47** and to the acceleration waveform (Figure 15) in step **48**.

**[0076]** Figures 13 through 15 show that a moving average technique reduces the effect of drift in the reported starting point of each compression. (The moving average techniques also reduce the effect of external acceleration errors that appear in the baseline portion of the waveform). Before correction, the reported starting points were becoming increasingly deeper, though the actual starting points were returning to close to the actual initial starting point. By applying a moving average technique to the baseline of a measured waveform, the reported starting points of each compression are statistically closer to the actual starting points. Accordingly, the compression monitor will report an estimated actual compression depth that is closer to the actual compression depth. Arrows **49** and **50**, which are shorter than arrows **24** and **25** in Figures 2 through 4 and Figures 7 through 12, show the beneficial effect of applying a moving average technique to each waveform.

**[0077]** Returning to Figure 5, the compression depth waveform corrected by the baseline limiter may be passed through a third filter in step **51** to reduce any accumulated signal noise in the compression depth waveform. The third filter comprises a high pass filter, though in other embodiments the third filter may comprise a band pass filter.

**[0078]** Subsequently, the depth waveform (whether filtered or unfiltered) is provided to a starting point detector in step **52**. The starting point detector identifies the value of the current estimated starting point. The current estimated starting point is then provided to a means for combining signals **53** (as indicated by line **54**). The means for combining signals **53** will later use the current estimated starting point to calculate the estimated actual compression depth. The means for combining signals comprises a signal adder, a linear system model, a non-linear system model, or other means for combining signals.

**[0079]** Next, the compression waveform may be provided to a peak limiter in step **55**. The peak limiter is a signal processor that performs similar functions to the baseline limiter, but instead operates on the peak portion of a compression waveform. The peak portion of the waveform comprises that portion of the waveform in which a peak is most likely to occur. In one embodiment, the peak portion is the portion of the waveform above the baseline portion. Continuing the example given for the baseline limiter, the peak portion of the depth waveform would be the portion of the depth waveform that is above 2.79 cm (1.1 inches). The peak limiter thus will smooth the peak portion of a waveform in much the same way as the baseline limiter smoothes the baseline portion of a waveform.

In one embodiment the peak limiter sets an outside boundary on the size of the maximum compression depth. Thus, the peak limiter either disregards (throws out) or sets an arbitrary value to any peak that is greater than a known, improbable peak value (the depth of a large person's chest, for example, would not be a probable value for CPR compression depth). Thus, the peak limiter prevents the compression monitor from reporting a compression depth that is improbable.

The effect of the peak limiter is seen in Figures 16 through 18, which show the graphs of compression depth, velocity, and acceleration over time for four hypothetical compressions after the peak limiter step 55 in Figure 5.

(Figures 16 through 18 show the output of steps 35 through 55). A peak limiter has been applied separately to the velocity waveform in step 56 and to the acceleration waveform in step 57. By applying a moving average technique to the peak portion of the compression waveforms, the effect of the external acceleration spike 22 has been greatly reduced.

Combined with the techniques discussed in the previous processing steps, the reported waveforms are now close to the actual waveforms.

**[0080]** Returning to Figure 5, the estimated peak may optionally be provided to a fourth filter **58** to remove remaining signal noise. The fourth filter comprises a high pass filter, though in other embodiments the fourth filter may comprise a band pass or other filter.

**[0081]** Subsequently, the depth waveform is provided to a peak detector in step **59**. The peak detector identifies the value of the estimated peak (the estimated maximum depth of the current compression). The estimated peak is then provided to the means for combining signals **53**. The means for combining signals **53** combines the estimated starting point **52** with the estimated peak **59** to produce an estimated actual compression depth for the current compression **61**. The estimated actual depth is then provided to a means for user feedback **62** (a user feedback system). The means for user feedback may comprise a speaker, a visual display, one or more LEDs, a vibrator, radio, or other means for

communicating with the rescuer. The user feedback system in turn provides information corresponding to the estimated actual depth of the current compression to the rescuer.

**[0082]** In the technique of Figure 5, the baseline portion and the peak portion do not overlap. Thus, the compression depth waveform may be thought of as comprising two portions, the baseline portion and the peak portion. Each portion of the depth waveform is treated differently by two different procedures (the baseline limiter and the peak limiter) to extract different information. Thus, both the baseline limiter and the peak limiter operate on the same depth waveform. The effect of this is that the signal comprising the depth waveform is provided first to the baseline limiter and then to the peak limiter (the signal is not split).

The technique shown in Figure 6 may be used when the baseline portion and the peak portion overlap (though the technique may also be used when the baseline portion and peak portion do not overlap). For example, the technique of Figure 6 may be used when the baseline portion is set below 3.81 cm (1.5 inches) (relative to the chest's relaxed position) and the peak portion is set above 2.54 (1.0 inches) (relative to the chest's relaxed position). In this case the signal representing the depth waveform is split and is provided to two separate processors, a baseline limiter and a peak limiter. Each processor performs similar functions to the limiters already described. Thus, although the baseline limiter and the peak limiter act independently of each other, the technique of Figure 6 produces an estimated starting point and an estimated peak in much the same way as the technique shown in Figure 5.

The means for combining signals then combines the estimated starting point and estimated peak in step 53 to produce the estimated actual depth of the current compression. The estimated actual depth of the current compression is provided to the user feedback system in step 62. The user feedback system in turn provides the estimated actual depth of the current compression to the rescuer.

In addition to the signal processing techniques of Figures 5 and 6, other techniques can be used to correct for errors in the compression depth waveform. For example, Figure 19 is a flow chart of a signal processing technique that uses a change in ECG noise 63 to activate a switch 64 that, in turn, controls when an accelerometer begins to measure acceleration.

To implement this technique, the compression monitor is provided with one or more electrodes, or some other means for measuring the patient's ECG. As the rescuer performs compressions the patient's ECG becomes noisy. Even if the patient's actual ECG is flat (shows no activity) the reported ECG will still show the noise caused by chest compressions. Indeed, a motion artifact signal (an ECG noise component caused by chest compressions) will be superimposed on any ECG rhythm. Whatever the actual ECG rhythm, the ECG noise may be isolated and accounted for.

**[0083]** Since the bulk of ECG noise during compressions is caused by the act of compressing the chest, the starting point of a compression may be correlated to the point where the ECG noise exceeds a pre-determined threshold. However, there is some delay or lag between the onset of a compression and the onset of ECG noise. The time lag is on the order of milliseconds to tenths of a second. In order not to miss any part of a compression, a buffer (either digital or analog) may be employed to correct for the time lag. Thereafter, when the ECG noise exceeds the particular threshold then the switch is programmed to activate the accelerometer (which will begin to take acceleration measurements). Total compression depth is then determined by double integrating the measured acceleration.

**[0084]** The effect of using ECG noise as a reference sensor to establish the starting points of compressions is seen in Figures 20 through 25, which show compression depth, velocity, and acceleration over time for four hypothetical compressions. No signal processing is applied to any of waveforms shown in Figures 20 through 22. The phantom waveforms **12** represent the actual waveforms for compression depth, velocity, and acceleration (measured independently of the accelerometer). The solid waveforms **13** represent the waveforms derived from the acceleration measured by the accelerometer. The solid waveforms are also the waveforms reported by the compression monitor. The effects of signal noise are shown by the rough nature of the solid waveforms. The effects of external acceleration noise are shown by the two spikes, **65** and **66**, in the reported waveform. The effects of negative drift (increasingly shallow compressions) are shown by the increasing distance (represented by arrows **67** and **68**) between the minimums in the reported and the actual waveforms.

**[0085]** The effect of using ECG noise as a reference sensor to establish the starting points of compressions is seen in Figures 23 through 25, which show graphs of compression depth, velocity, and acceleration over time for hypothetical compressions. Using ECG noise as a reference sensor reduces certain external acceleration errors and reduces the effect of negative drift. (The ECG noise reference sensor can also reduce the effect of positive drift). Specifically, the ECG noise reference sensor reduces the effect of external acceleration noise that occurs near a compression minimum. Since the accelerometer is not "on," a portion of the external acceleration spike is "ignored". In practice the accelerometer is still taking data, but software or hardware is used to process out accelerometer data or signals that occur during a time period where ECG noise does not reach a predetermined level. In other methods, the estimated actual depth of compressions is calculated when the ECG noise falls within a predetermined threshold. In any case, the effect of spike **65** is reduced in the reported waveform. However, the accelerometer by itself still cannot tell the difference between a compression-related acceleration and an external acceleration. Thus, the reported waveform is still subject to external acceleration noise that occurs during a compression, as shown by spike **66**.

**[0086]** Nevertheless, the ECG noise reference sensor does reduce the effects of drift. Since the starting point of a compression is independently established, the waveform is much less subject to either positive or negative drift. In other words, the accelerometer will always measure acceleration after the actual start of compressions. Thus, the reported waveform of Figure 23 more accurately shows what the rescuer is actually doing - compressing the chest from starting points that are becoming increasingly deep. Thus, peaks **69** and **70** show that the measured waveform more closely matches the actual waveform.

**[0087]** Although the ECG noise reference sensor can reduce the effects of drift and reduce the effect of some forms of external acceleration noise, signal noise remains a problem. Thus, Figures 23 through 25 still show the same levels of signal noise as shown in Figures 20 through 22. To reduce all forms of noise the ECG noise reference sensor may be combined with the signal processing techniques of Figures 5 or 6. The combined techniques will produce a reported depth waveform that is close to the actual waveform.

**[0088]** Other reference sensors may be used to establish the actual starting point of a compression. Figure 26 shows an accelerometer-based compression monitor in place on a patient **1** who is lying on a surface **80**. A system of reference sensors comprising an accelerometer **81**, a load sensor **82**, and a switch **83** are disposed such that each sensor may measure various parameters related to chest compressions. In the case of reference accelerometers, the reference accelerometers may be disposed elsewhere on the patient, or upon any reference object that experiences the same external accelerations the patient experiences. The reference accelerometers may comprise a three-axis accelerometer, but may also comprise three orthogonal single-axis accelerometers or one single axis accelerometer (in which case the accelerations along the other two axes are assumed to be negligible).

**[0089]** The reference accelerometers **81** allow a signal processor to eliminate external acceleration error, such as those accelerations caused by transporting the patient. In one method, the acceleration sensed by the compression monitor or automatic CPR device (the device acceleration) is provided to a signal processor. The device acceleration contains the acceleration caused by compressions (the compression acceleration) and the acceleration caused by the external accelerations (the external acceleration). Next, the reference accelerometer or accelerometers provide a reference acceleration to the signal processor. The reference acceleration contains only the external acceleration of the patient. Then the reference acceleration is combined with the device acceleration to produce an estimated actual acceleration. (The effect of compression accelerations on the reference acceleration is negligible since the surface and patient are kept steady with respect to the compression monitor.)

**[0090]** Once obtained, the estimated actual acceleration may be double integrated to produce an estimated actual chest depth. Thus, the depth of compressions may be determined even in the presence of large external accelerations. Moreover, the position signal may be made more accurate and precise by combining the actual acceleration with the signal processing technique of Figures 5 or 6, or with other signal processing techniques.

**[0091]** In lieu of (or in addition to) the ECG noise sensor and reference accelerometers, other reference sensors may be used to set the actual starting point of a compression. Reference sensors may comprise a load sensor **82**, a switch **83**, a transthoracic impedance detector, an ECG noise detector (as described above), a voltage or current sensor in an automatic CPR device, a start signal in an automatic CPR device, an encoder in an automatic CPR device, or any other sensor capable of independently detecting the actual beginning of a compression. When the reference sensor detects the beginning of a compression then the starting point is set to zero. The acceleration is then processed to derive compression depth. The technique of setting the starting point to zero when a reference sensor detects the beginning of a compression may also be combined with the signal processing techniques of Figures 5 or 6.

**[0092]** In the case of a switch **83**, the switch is disposed such that when a compression begins the switch will be closed. For example, the switch may be disposed beneath or on the compression monitor, on the patient **1**, on the surface **80** upon which the patient lies, on the rescuer's hand, on a CPR machine, on the patient, or on some other location that allows the switch to register that a compression has begun.

**[0093]** The switch may comprise many different types of switches and sensors, including a contact switch, a motion sensor, a voltage sensor on an automatic CPR device, an optical, rotary, or other encoder on an automatic CPR device, the displacement of a shaft or other component on an automatic CPR device, a potentiometer, a strain gage, a piezoresistive transducer, a differential transformer, synchro and induction potentiometers, variable-inductance and variable-reluctance pickups, an eddy current non-conducting transducer, a capacitive transducer, an electro-optical transducer, a photographic switch, a video tape switch, a holographic switch, a switch that uses photoelastic techniques, translation encoders, an ultrasonic transducer, moving coil and moving magnet pickups, an AC or DC tachometer, an eddy-current drag-cup tachometer, additional accelerometers, or a gyroscopic displacement switch.

**[0094]** In the case of the load sensor **82**, the load sensor may be operatively connected to the rescuer, the patient, an automatic CPR device, beneath the patient, or elsewhere so long as the load sensor senses a load when compressions begin. When the load sensor measures a load that exceeds a pre-determined threshold, then the measured starting point is set to zero. The load sensor may also be operatively connected to a switch, which activates when the load sensor senses a load, or the load sensor may merely provide input to a signal processor system identifier (described in more detail below). Compression depth is then determined by integrating the acceleration twice. The technique of setting the

starting point to zero when a load sensor detects the beginning of a compression may also be combined with the signal processing techniques of Figures 5 or 6.

**[0095]** In another embodiment of the load sensor **82**, the load sensor may be disposed such that the sensor can sense both the weight of the patient and the force of compressions. The load sensor **82** may be disposed beneath the surface **80** upon which the patient **1** rests. During compressions the force of pressing on the patient causes the load sensor to report a total force greater than the patient's weight. Accordingly, a starting point is set to zero when the total force is about equal to the patient's weight.

**[0096]** Examples of force sensors that can be used with this technique include pressure sensors, elastic force transducers, shaft displacement on an automatic CPR device, a voltage or a current sensor on an automatic CPR device, an optical, rotary, or other encoder on an automatic CPR device, bonded strain gages, beam strain gages, differential transformers, piezoelectric transducers, variable reluctance/FM oscillators, gyroscopic force transducers, and vibrating wire force detectors. Examples of pressure sensors that can be used with this technique include deadweight gages, manometers, elastic transducers, piezoelectric transducers, and force-balance transducers.

In the case of a transthoracic impedance detector, one or more ECG, defibrillation, or other electrodes are disposed on the patient's thorax. When a compression begins the impedance of the thorax changes. The thoracic impedance comprises the impedance due to skin and thoracic contents between any two electrodes. The change in thoracic impedance may be measured by a small test current or by any other means for measuring impedance. When the impedance changes by a pre-determined amount then the starting point is set to zero. Total compression depth may then be determined by processing the measured acceleration.

Because the compression monitor can measure the compression waveform, the compression monitor can also prompt the rescuer or an automatic CPR device to perform a particular compression waveform. Figure 27 shows a compression waveform that the compression monitor may prompt the rescuer to perform. Depth is measured in cm (or inches) and time is measured in seconds. The scale shown in Figure 27 is marked in 0.5 second intervals and 2.54 cm (1.0 inch) intervals respectively. The compression phase of the cycle is indicated by the positively sloped curve **84**. The compression phase of the cycle ends at the maximum compression depth **85** (compression peak). The decompression phase of the cycle is indicated by the negatively sloped curve **86**. The decompression phase ends when the rescuer begins a new compression at the next starting point **87** (or baseline), which may or may not be at the initial starting point. Compressions are initiated at time = 0 and depth = 0, and the total depth of compressions is the distance represented by arrows **88**.

**[0097]** The compression waveform includes a compression hold **89**, where the rescuer maintains a hold at maximum compression depth for a short period of time, and an incomplete decompression hold **90**, where the rescuer maintains a short hold at a point deeper than the initial starting point. Each compression and decompression is performed quickly, at high acceleration and velocity, as indicated by the relatively steep slopes of the compression phase **84** and the decompression phase **86**. The duty cycle is slightly less than 50% (the ratio of compression and decompression time is 1), meaning that slightly less time is spent in the compression phase, as indicated by the distance between arrows **89**, than in the decompression phase, as indicated by the distance between arrows **90**.

**[0098]** Although the compression waveform of Figure 27 shows an example of a particular waveform that the compression monitor can instruct a rescuer to perform, other waveforms are also possible. For example, another waveform may lack a compression hold phase. Yet another varies the duty cycle and others increase the compression hold time. The exact waveform depends on the current state of the art of what kind of compression waveform comprises an optimal compression waveform for a particular kind of patient. In addition, the compression monitor may be provided with a switch, button, software, or other means for user input which allows the rescuer to enter the size or shape of the patient. The compression monitor may use this information to choose a particular waveform from a library of waveforms. The compression waveforms are thus adaptable to findings in future research, AHA guidelines, rescuer observations, and medical professional preferences. Accordingly, at various times different waveforms may be provided to the user feedback system, as described more fully below.

**[0099]** The prompted waveform may be provided by the user feedback system (step **62** in Figure 5). In addition, the user feedback system may provide the rescuer or automatic CPR device with other compression-related information. For example, the user feedback system may display information regarding the starting point of compressions, the compression depth waveform, the compression velocity waveform, and the compression acceleration waveform. Thus, the user feedback system may provide the rescuer or the automatic CPR device with all of the data needed to continuously track the position, velocity, and acceleration of the chest during all phases of CPR. This information may be used to evaluate the performance of a rescuer or automatic CPR device.

**[0100]** The user feedback system may also provide a rescuer or automatic CPR device with information concerning the compression phase quality and the decompression phase quality. Compression phase quality is the quality of compressions with respect to total compression depth, the duty cycle, the acceleration of compressions, smoothness of compressions, and other factors related to the compression phase. Decompression phase quality is the quality of compressions with respect to whether the rescuer returns to the actual initial position, the duty cycle, the acceleration of decompressions, the smoothness of decompressions, and other factors related to the decompression phase. The rescuer

or automatic CPR device may use this information to evaluate or prompt the kind and quality of compressions.

**[0101]** The user feedback system **62** may provide the rescuer or automatic CPR device with information concerning compression phase quality by combining information gained from the acceleration, velocity, and position waveforms. For example, the user feedback system can instruct the rescuer to increase compression force when the depth of compressions are less than recommended guidelines and to reduce compression force when the depth of compressions are greater than recommended guidelines. The user feedback system may also instruct the rescuer with regard to other compression phase parameters of a compression waveform. For example, the user feedback system can inform the rescuer or automatic CPR device if the time to achieve proper compression depth is too short or too long.

**[0102]** The user feedback system **62** may also provide the rescuer or automatic CPR device with information regarding decompression phase quality by combining information gained from the acceleration, velocity, and position waveforms. For example, the user feedback system can instruct the user or device on the proper position at which to rest after a decompression. Thus, the feedback system can instruct the user or device to allow the chest to fully relax if the rescuer or device is not allowing the chest to fully return to its initial starting position. Conversely, should it be medically indicated, the user feedback system can instruct the user or the device to return to a depth just below the initial chest position. In this case, the rescuer or device implements a "decompression hold" and maintains force on the chest even when the compression cycle reaches its minimum depth. In another case the feedback system can indicate different compression starting points at different times. Thus, the user feedback system can instruct the rescuer or device to apply incomplete decompression holds during compression cycles, but to allow the chest to return to its fully relaxed position during ventilation pauses. The user feedback system may also instruct the rescuer or device with regard to other decompression phase parameters, such as the decompression rate and the duty cycle of the decompression phase.

**[0103]** Taken together, the information gained from the compression phase quality and the decompression phase quality enable the user feedback system to prompt the rescuer on how to perform an optimum compression waveform and an optimum compression duty cycle. A rescuer performs a particular compression waveform by performing compressions at a pre-determined depth and rate, and by holding the chest at a pre-determined compression depth for a pre-determined time. A rescuer performs a particular duty cycle by compressing the chest for a pre-determined period and allowing the chest to relax for another pre-determined period.

**[0104]** Thus, the user feedback system can prompt the rescuer or automatic CPR device to perform at the appropriate compression rate, compression depth, compression velocity (the time required to compress or decompress the patient), compression acceleration, and compression hold time for each phase (compression and decompression) of the compression cycle. Accordingly, the compression waveform that the rescuer or device actually applies can conform to a complex compression waveform. Since research has shown that most patients benefit from more complex waveforms, patient survival is likely to increase if the rescuer or automatic CPR device uses a compression monitor with this user feedback system.

**[0105]** Similarly, the user feedback system **62** of Figure 5 can provide the rescuer or CPR device with feedback regarding the compression duty cycle. The duty cycle is the ratio of time under compression to the time under decompression for each compression cycle. (However, the duty cycle does not include time periods where no compressions are taking place, such as during ventilation.) If the duty cycle does not fall within pre-determined parameters, then the user feedback system may prompt the rescuer to adjust compression timing and compression rate in order to effect an optimal duty cycle.

**[0106]** The user feedback system described above comprises the last step in a particular solution to the problem of determining an accurate value for chest displacement from a raw acceleration signal. (Figure 5 is the flowchart for this solution). Many variations of that solution exist, as already described, though it is possible to view the problem from a general perspective and to create a general solution.

**[0107]** Figures 28 and 29 are block diagrams that represent the general problem to and the general solution for determining an accurate position from an acceleration measured during CPR. Figure 28 is a block diagram of how an actual chest compression acceleration is converted into a corrupted value for chest position. In broad terms, the actual acceleration **105**, signal noise **106**, external acceleration noise **107**, and some forms of drift **108** are combined by an unknown function **109** (which may be linear or non-linear and may include random or deterministic inputs). The unknown non-linear function is known as the system, which produces the corrupted acceleration **110** measured by the accelerometer. The corrupted acceleration is then integrated twice, which greatly compounds the problem introduced by the corruption in the acceleration. The increased error is referred to as integration error **111** (although it is assumed that the integration technique itself does not directly contribute errors into the position). Finally, additional sources of drift **112** can affect the final value for the corrupted position **113**.

**[0108]** Figure 29 is a block diagram of a general solution for converting a corrupted chest compression acceleration into an estimated actual depth of chest compressions. First, a reference sensor **119** may establish the actual starting point of a compression. Thus, the starting point of the acceleration will be known. (Although helpful, the reference sensor **119** is not necessary to the general solution). The actual acceleration **105** and the real or the estimated noise sources **120** (which comprise blocks 106 through 108 of Figure 28) are combined by the system **109** by an unknown function.



The result is the corrupted acceleration 110. The measured acceleration is then provided to a means for combining data 121 (which may comprise a linear or a non-linear function) and to a system identifier 122.

The system identifier comprises one or more functions (either linear or non-linear) that model the system. One or more noise references that can be correlated to the noise sources 120 may also be provided to the system identification function 122. For example, noise identified by a low frequency filter can be correlated to signal noise or a reference accelerometer can be correlated to the external acceleration noise.

The system identification function may also use various parameters of an automatic CPR device as noise source references, even if the reference itself does not produce noise in the acceleration. However, the noise source reference must somehow be correlated to a source of noise in the acceleration signal. For example, the accelerometer-based depth measurement reports a chest depth of 1.37 cm (0.5 inches).

However, a simultaneous current spike in the automatic CPR device informs the system that the CPR device is compressing the chest much harder than should be required to achieve a chest depth of 1.37 cm (0.5 inches). The discrepancy may be caused by external acceleration noise or by drift. Thus, the current spike may be correlated to a source of noise in the system.

This information may be used by the system identifier to help model the system. Likewise, voltage, shaft displacement, or optical or rotary encoders may be used as references by the system identifier to help model the system. (Again, the noise references are useful but not necessary).

**[0109]** The system identifier then combines or correlates the noise source references and the measured acceleration in order to produce the estimated noise 123 in the measured acceleration. The estimated noise 123 is then provided to the means for combining data 121. The means for combining data combines the estimated noise 123 and the measured acceleration 110 to produce an estimated actual acceleration 124. The estimated actual acceleration is then integrated 125 twice. Filters 126 may optionally be used during one or both integration steps to reduce the compounding effect of errors that may still linger in the estimated actual acceleration. The final result is an accurate and precise estimate of the actual position 127 of the accelerometer.

**[0110]** The system identification function 122 models the system and thus can be used to estimate the noise in the acceleration. (Once the noise is known it can be easily eliminated by combining the noisy acceleration with the measured acceleration.) In other words, system identification is the process of using the input and output data to model the function that combines the actual acceleration and the sources of noise in the acceleration. The system identification problem has a known or measured output and an input that may be known or unknown. The addition of known or measured input is beneficial to system identification, but not necessary. The system itself is an unknown arbitrary function that can be linear or non-linear, though some boundary conditions may be known.

**[0111]** A number of methods, both linear and non-linear, may be used to model the system. Each of these methods may comprise, alone or in combination, the system identification function in step 122. These methods may operate by taking many data samples per second, as opposed to operating only on the compression starting points or peak points. Nevertheless, these methods may also be performed on the compression starting points or the compression peaks. A partial list of these methods include: autoregression, autoregression with extra inputs, autoregressive moving average (which is one of the methods used in the techniques shown in Figures 5 and 6) autoregressive moving average with extra inputs, autoregressive integrated moving average, autoregressive integrated moving average with extra inputs, a Box-Jenkins model, an output error model, a hidden Markov model, a Fourier transform, a wavelet transform, wavelet de-noising, wavelet filtering, adaptive neural networks, recurrent neural networks, radial basis function nets, adaptive curve fitting (splines), Kalman filters, extended Kalman filters, adaptive Kalman filters, unscented Kalman filters, and kernel estimation. Algorithmic approaches that may be used to find the system identification function include maximum entropy, maximum likelihood, recursive least squares (or similar techniques), numerical methods, unconstrained global search or optimization, expectation minimization, and fast Fourier transforms.

**[0112]** In the case of recursive identification, the formula for general recursive identification may be expressed as:

$$(1) \quad X(t) = H(t, X(t-1), y(t), u(t))$$

and

$$(2) \quad \theta(t) = h(X(t)),$$

where  $X(t)$  is the state of the system at time  $t$ ;  $H$  is the state of the transfer function;  $X(t-1)$  is the past system state;  $Y(t)$  is the measured output;  $u(t)$  is the measured input;  $\theta(t)$  is the system, and  $h$  transforms the system state to the output. The system state can be converted to the system output by  $h(X(t))$ .

**[0113]** Since  $X(t)$  and  $\theta(t)$  are evaluated at each time point as  $u(t)$  and  $y(t)$  are collected, the total amount of previously

collected data has a much more profound effect on the system than do the most recently collected data.

**[0114]** Equations (1) and (2) may be simplified into equations (3) and (4):

$$(3) \quad X(t) = X(t-1) + \mu Q_x(X(t-1), y(t), u(t))$$

and

$$(4) \quad \theta(t) = \theta(t-1) + \gamma Q_\theta(X(t-1), y(t), u(t)),$$

where  $\mu$  and  $\gamma$  are small numbers that reflect the relative amount of information provided by the latest time step.  $Q$  is a function that relates inputs, outputs, and states. Equations (3) and (4) are more simple in the sense that it is more simple to compute the equations by recursive than equations (1) and (2).

**[0115]** A number of numerical algorithms may be used to solve equations (3) and (4). A partial list of numerical algorithms include recursive least squares, recursive (or recursion) instrumental variables, recursive prediction error methods, recursive pseudolinear regression, recursive Kalman filters (including time varying parameters), and recursive Kalman filters for time varying systems. These numerical techniques encompass many of the famous "named" techniques as special cases, including a Kalman filter, an extended Kalman filter, extended recursive least squares, and others.

Each algorithm has strengths and weaknesses, but all asymptotically approach a solution to equations (3) and (4).

**[0116]** The "named" special cases may be derived from general equations (3) and (4) when certain conditions or assumptions are made. Thus, the equations for each of the listed algorithms can be further specified. For example, when using a recursive least squares algorithm equation (4) may be expressed as:

$$(5) \quad \theta(t) = \theta(t-1) + L(t)[y(t) - \phi^T(t)\theta(t-1)]$$

where

$$(5a) \quad L(t) = \frac{P(t-1)\phi(t)}{\lambda(t) + \phi^T(t)P(t-1)\phi(t)}$$

and

$$(5b) \quad P(t) = \frac{P(t-1)}{\lambda(t)} - \frac{P(t-1)\phi(t)\phi^T(t)P(t-1)}{\lambda(t)[\lambda(t) + \phi^T(t)P(t-1)\phi(t)]}.$$

**[0117]** In equations 5 through 5(b)  $L(t)$ ,  $P(t)$ , and  $P(t-1)$  are terms used to simplify the equation,  $\phi(1)$  is a regression vector,  $\lambda(t)$  is a forgetting factor (described in more detail below), and  $\phi^T(t)$  is the transpose of the regression vector.

**[0118]** In addition, equation 4 can also be expressed for the cases of recursive instrumental variables, recursive prediction-error methods, recursive pseudolinear regression, a recursive Kalman filter for time-varying systems, and a recursive Kalman filter with parametric variation.

**[0119]** Once the system identification algorithm as been selected from the above set of algorithms, there are several additional parameters that may affect the quality of the model. These additional parameters include data weighting, choice of updating step, choice of updating gain, and model order selection. In the case of data weighting, when a system is time-varying the input-output data near the present time more accurately reflects the nature of the present system. Data recorded further back in time is more closely related to a past system state. To reflect this fact the data can be weighted to favor a more recent system state. Actual data weighting is accomplished by the "forgetting factor,"  $\lambda$ , in equations 5 through 5b. The selection of  $\lambda$  is made based on information about how fast the system changes state. A typical range for  $\lambda$  is between about 0.9800 and about 0.9999 (though  $\lambda$  may be 1.0000 if no "forgetting factor" is desired).

**[0120]** Another way of thinking about the effect of  $\lambda$  on the system identifier is to evaluate at what point a data sample is given a weight of about 36%. (36% is the value of the number  $e^{-1}$ , which is the value at which a data sample may be considered statistically insignificant). At this weight a data sample's statistical significance becomes relatively small. The sample number at which a data sample has a weight of about 36%, known as  $T_0$ , may be mathematically expressed as:

$$T_0 = \frac{1}{1 - \lambda}.$$

$T_0$  (and hence  $\lambda$ ) is selected with appropriate knowledge of the system state and can be used to tune system identification so that the estimated actual acceleration most closely approximates the actual acceleration if the actual acceleration were independently measured. Thus,  $T_0$  is pre-set before the compression monitor begins taking measurements.

**[0121]** The closer  $\lambda$  is to 1 the more samples are needed to reach the point where a given data sample is given a weight of about 36%. A smaller  $\lambda$  means that a given data sample is "forgotten" more quickly. For example, if  $\lambda$  is equal to 0.9800 then  $T_0 = 50$  samples, meaning that the 50<sup>th</sup> sample receives a weight of about 36%. However, if  $\lambda = 0.9999$  then  $T_0 = 10,000$ , meaning that sample number 10,000 receives a weight of 36%. In the upper limit, if  $\lambda = 1$  then  $T_0 = \infty$ , meaning that a data sample is never "forgotten" (it always receives a weight of 100%).

**[0122]** The sampling rate (how many times a second the acceleration is measured) affects the how  $\lambda$  changes the system identifier. If samples are taken 1000 times per second then data may be "forgotten" rapidly on the time scale of CPR compressions. For example, if the sample rate is 1000 times per second and  $T_0 = 1000$  then data from just 1 second in the past is given a weight of 36%. In practice, the sampling rate may vary from about 100 samples per second to about 2000 samples per second. A useful sample rate for signal processing acceleration measurements during CPR is about 500 samples per second. In other embodiments the sample rate may be faster, but every certain number of samples may be ignored. For example, samples may be taken at 1000 samples per second, but every other sample ignored. This process, known as decimation, has the same effect as a slower sample rate.

**[0123]** In the preceding discussion the forgetting factor was a fixed number; it did not change with time. However,  $\lambda$  can vary with time so that the system identifier may adapt to changing situations. For example,  $\lambda$  may vary during a ventilation pause and in one embodiment  $\lambda$  increases during a ventilation pause. The effect of increasing  $\lambda$  during ventilation pauses is to discard data points very quickly. Thus, the compression monitor will not report a change in compression depth during a ventilation pause.

**[0124]** In addition to adding a forgetting factor to the system identification function, the choice of updating step affects the quality of the model. (Although only some of the system identification techniques require an update; for example, a Kalman filter requires an updating step). The update step can be implemented using a variety of methods. Some system identifiers may be solved analytically, such as the Kalman filter, and the updating step may be solved analytically. Other system identifiers must be solved numerically. Three updating methods that may be used when a numerical solution is required are a Gauss-Newton update, a gradient update, and a Levenberg-Marquardt update. The Gauss-Newton update converges to an accurate fit of the actual solution, though it requires a large number of steps (and thus more computation time). The gradient update converges quickly but does not converge as accurately to the actual solution as the Gauss-Newton update. The methods may be combined. The gradient update is used first to converge the fit quickly and then the identifier switches to the Gauss-Newton update to achieve the final fit. This combined technique is known as a Levenberg-Marquardt update.

**[0125]** Mathematically the Gauss-Newton update may be expressed as:

$$R(t) = R(t-1) + \gamma(t) [\phi(t)\phi^T(t) - R(t-1)].$$

Mathematically the gradient update may be expressed as:

$$R(t) = I |\phi(t)|^2 = R(t-1) + \gamma(t) [I |\phi(t)|^2 - R(t-1)].$$

In both equations  $R(t)$  is the Hessian of the identification criterion,  $R(t-1)$  is the Hessian of the identification criterion in the previous time step,  $\gamma(t)$  is the updating gain (which is related to the forgetting factor), and  $\phi(t)$  is the regression vector.

**[0126]** The choice of updating gain is another step that is used in many recursive system identification functions. The choice of updating gain may be expressed mathematically as:

$$\gamma(t) = (1 + \lambda(t) / \gamma(t-1))^{-1}.$$

Thus, the updating gain is related to the forgetting factor.

**[0127]** With regard to model order selection, the recursive system identification techniques fit a system model to input-output data. The structure of that model must be determined before the recursive. The standard way of solving the model structure problem is to solve a wide range of model structures and then select which model fits the data best. A simple

measure of model fit, like the mean squared error, tends to over-estimate the model order and fit to the process or measurement noise. If a model is under-estimated, critical components of the system might be missed. Several measures of model fit, or metrics, are evaluated over a range of model orders. The model order is the number of terms used in the model. The smallest model order that minimizes the fit is the appropriate model.

**[0128]** Several techniques can be used to estimate smallest model order, including final prediction error (FPE), Akaike's information criterion (AIC), maximum description length (a variant of the AIC), and statistical hypothesis testing (such as the student's t-test). The final prediction error can be expressed mathematically as:

$$(11) \quad FPE = \frac{V(1+d/N)}{(1-d/N)},$$

where V is the quadratic loss function, d is the size of the model order, and N is the number of data points.

**[0129]** Akaike's information criterion may be expressed as:

$$(12) \quad AIC = \log[V(1+2(d/N))],$$

where V is the quadratic loss function. The quadratic loss function may be any quadratic function that relates the additional cost function of using additional terms.

**[0130]** The system identification techniques described above have been described in the context of solving the problem of estimating actual compression depth from a raw acceleration measurement. Although not explicitly in the wording of the claims, techniques that do not produce an estimated actual depth signal corresponding to the estimated actual depth of compressions by calculating an autoregressive moving average of the measured depth signal may also be used to process a

noisy ECG signal. Figure 30 is a block diagram illustrating the problem of ECG noise caused by CPR and other sources of noise. Stated differently, Figure 30 is a block diagram of how a theoretical actual ECG signal **135** and the noise sources are combined to produce the measured ECG (which contains the motion corrupting artifact). The ECG noise, comprising ECG noise due to compressions **136** and other sources of noise **137**, are combined with the actual ECG by an unknown linear or non-linear function known as the system **138**. The primary source of noise in the ECG is due to the motion of compressions, though other sources of noise exist and may be accounted for by the solution presented below. The system produces a corrupted ECG **139** which, if left unprocessed, cannot be used to accurately report the electrical activity of the patient's heart. In addition, the system combines the ECG noise and the actual ECG in a way that causes the ECG noise to overlap the actual ECG in the frequency domain. Thus, a simple bandpass filter is insufficient to accurately process the corrupted ECG. (A simple bandpass filter will eliminate important components of the actual ECG as well as eliminating the ECG noise).

**[0131]** Figure 31 is a block diagram of a general solution to the problem illustrated in Figure 30 and illustrates the process of converting a motion corrupted ECG signal into an estimated actual ECG signal. As with Figure 30, the system **138** combines the actual ECG **135** and the ECG noise **136** to produce the corrupted ECG **139** that is measured by an observer. Next, the measured ECG **139** and a reference corresponding to the ECG noise **136** are provided to a system identifier **140**. For example, since CPR induced motion is the largest cause of ECG noise, a signal corresponding to CPR induced motion may be provided to the system identifier. Specifically, a force transducer may be disposed on a compression monitor (or on the patient or rescuer) such that the force transducer measures force during a compression. A signal corresponding to the force is provided to the system identifier as a reference signal. Other signals corresponding to CPR induced motion may comprise various parameters of an automatic CPR device. For example, a signal correlating to the displacement of a drive shaft or other component can be correlated to the CPR motion, a signal corresponding to the change in current or voltage required to drive the device can be correlated to the CPR motion, or signals produced by optical or rotary encoders may be correlated to the CPR motion.

**[0132]** The system identifier models the system and then estimates the noise component of the measured ECG signal (the estimated noise **141**). The estimated ECG noise **141** and the measured ECG **139** are then provided to a means for combining signals **142**, which combines the ECG noise and the measured ECG to produce the estimated actual ECG **143**. Since the estimated actual ECG is produced during compressions, the signal processing method allows the ECG sensor to detect the heart's normal sinus rhythm even during compressions. Thus, there is no need to periodically pause compressions to check for the existence of a pulse. As a result, the overall quality of CPR increases and the patient is more likely to survive.

**[0133]** The system identifier **140** may comprise similar kinds of functions and methods as described in the context of the signal processing methods of Figure 29. For example, the recursive least squares method described in the context of Figure 29 may be used to identify the noise component of the measured ECG signal.

**[0134]** Figures 32 through 35 show the effect of using the method of Figure 31 to estimate a pig's actual ECG when the ECG is measured during chest compressions. For all four graphs each time marker **150** along time scale **151** corresponds to the same time marker in the other three graphs, thus making possible a direct comparison of one graph to each of the other graphs. However, the voltage scales **152** of Figures 32, 34, and 35 are slightly different from each other.

**[0135]** Figure 32 is a graph (millivolts versus milliseconds) of an actual pig's ECG signal that is corrupted by noise caused by chest compressions. Figure 32 represents the ECG measured during compressions without signal processing.

**[0136]** Figure 33 is a graph of force versus time for an actual CPR motion signal. The motion signal comprises a time varying force signal and corresponds to the force a CPR device places on the pig's chest while performing chest compressions. The force peaks **153** correspond to the maximum depth of compressions. In the case of a compression monitor, the motion signal could comprise a time varying force signal that corresponds to the force placed on the patient's chest while a rescuer or automatic CPR device performs chest compressions. In this case a force transducer disposed on a compression monitor (such as on the back of the compression monitor) measures the force of compressions and produces the force signal. The force signal is later correlated to the ECG noise. The force transducer or other force sensor may also be disposed under the patient's back and then operably connected to the compression monitor.

**[0137]** Figure 34 is a graph of voltage versus time for the estimated ECG noise signal caused by the chest compressions shown in Figure 33. A comparison of Figures 33 and 34 shows that the time varying pressure signal corresponds directly to incidence of ECG noise. In other words, the pressure peaks **153** caused by chest compressions correspond to the incidence of noise peaks **154**.

**[0138]** The system identifier **140** used to generate the estimated noise component of the noisy ECG comprises a recursive least squares method as described in the context of Figure 29. The autoregressive order was selected to be equal to 1. The moving average order was selected to be 10. The autoregressive order was selected to be 10. The derivative order was also selected to be 0. (The derivative order is a linear or non-linear term used in the system model; specifically it may be either a truncated positive derivative or a truncated negative derivative. The non-linear terms are extensions of the recursive least squares model fit algorithm). The forgetting factor,  $\lambda$ , was selected to be 1.0000.

**[0139]** Figure 35 is a graph of the pig's estimated actual ECG signal. The graph of Figure 35 is generated by subtracting the estimated noise signal of Figure 34 from the measured ECG signal of Figure 32. The estimated actual ECG signal corresponds closely to the pig's actual ECG signal.

**[0140]** The signal processing methods described in the context of noisy ECG signals (Figures 30 and 31) and noisy acceleration signals (Figures 28 and 29), as well as the techniques described in relation to the baseline limiter and the peak limiter, may also be used to estimate the actual value of the patient's transthoracic impedance (the chest's electrical resistance or impedance). The estimated actual value of the patient's transthoracic impedance may be used to determine the amount of voltage needed to shock the patient with a defibrillator.

**[0141]** As compressions are applied to the patient the measured value of the transthoracic impedance becomes noisy. The general signal processing solutions and the limiters already described may be used to identify, isolate, and eliminate the noise component of the measured transthoracic impedance.

Thus, the actual value of the transthoracic impedance may be estimated.

**[0142]** The estimated actual value of the transthoracic impedance may be provided to a means for defibrillating the patient. The means for defibrillating the patient uses the estimated actual value of the transthoracic impedance to determine the exact voltage necessary to apply an effective shock to the patient. Since the exact value of the voltage required is also known, the defibrillator can be used efficiently (thus preserving battery life and making the device safer).

**[0143]** Since both the estimated actual ECG and the estimated actual transthoracic impedance are known, an automated CPR device equipped with an AED (automated external defibrillator) may perform defibrillation shocks to a patient without stopping compressions. The device may determine when defibrillation is appropriate based on the estimated actual ECG and may apply the appropriate defibrillation voltage based on the estimated actual transthoracic impedance. Since compressions do not stop during defibrillation, the patient's blood flow does not stop (meaning the patient is more likely to survive).

**[0144]** The compression monitor using these signal processing techniques (for either chest depth measurement or ECG measurement) may be used with any means for compressing the chest of the patient. A means for compressing the chest may comprise manual CPR, electro-stimulation, a means for performing automatic CPR (including belts, straps, pistons, and plates that are driven by motors or manual levers), or other devices suitable for compressing the chest. Examples of automatic CPR devices may be found in our own patent Sherman et al., Modular CPR Assist Device, U.S. Patent 6,066,106 (May 23, 2000) and in our application CPR Assist Device with Pressure Bladder Feedback, U.S. Application 09/866,377 filed May 25, 2001. (Both devices use optical or rotary encoders disposed on a compression belt or a drive shaft or spool to measure the amount of belt pay out or compression depth). The accelerometer itself may be disposed in any location where the accelerometer experiences the downward or upward acceleration of compressions. For example, the accelerometer may be disposed within or otherwise disposed on the means for compressing the chest, such as a compression belt. (In the case of manual compressions, the compression monitor may be disposed beneath a rescuer's hands while the rescuer performs compressions, or the compression monitor may be otherwise disposed

on the rescuer's hands, wrist, or arms.)

**[0145]** If the compression monitor is provided with a means for sensing the tilt of the accelerometer, such as a three-axis accelerometer, three-axis load sensor, three-axis displacement measurement device, or the tilt sensor shown in our own U.S. Patent 6,390,996 to Halperin et al., then the user feedback system can prompt the rescuer with respect to compression efficiency. For most patients, compressions are most efficient when compressions are performed perpendicular to the sternum (straight down in most cases). The tilt sensor measures the angle at which compressions are performed and the compression monitor prompts the rescuer to adjust the angle if the angle falls outside a particular range.

**[0146]** The compression monitor and signal processor may also be operably connected to a defibrillator. While the rescuer or device is performing compressions the defibrillator or compression monitor tracks the patient's ECG. If the compression monitor's processor measures an ECG signal that indicates that the patient would benefit from a shock, then the rescuer would be instructed to apply a defibrillation shock or to allow an AED to administer a shock. A means for estimating the patient's actual ECG during compressions comprises our own method disclosed in U.S. Patent 6,390,996 to Halperin et al and the method disclosed in this application.

**[0147]** The compression monitor may also be operably connected to a means for performing ventilation. After the rescuer has performed an appropriate number of compressions, such as 15, the compression monitor will instruct the rescuer to pause compressions. The means for performing ventilation will then administer an appropriate number of ventilations. After ventilations the compression monitor may evaluate the patient's condition. If the patient still requires compressions, then the compression monitor will instruct the rescuer to resume compressions. The means for performing ventilation may comprise the rescuer, a bag or balloon, a positive pressure ventilator, an electro-ventilator such as those shown in our own patent Sherman et al., Chest Compression Device with Electro-Stimulation, U.S. Patent 6,213,960 (Apr. 10, 2001), or other means for performing ventilation.

**[0148]** The compression monitor may also be provided with a means for communication that allows the compression monitor to communicate with a remote network. The means for communication comprises a signal carrier and a compression monitor communicator. The signal carrier may comprise a telephone line, direct connection cables, a dedicated digital network, a cell phone network, a satellite communication array, radio or other electromagnetic waves, a LED, the internet, or other means for carrying a signal. The compression monitor communicator may comprise a radio or other electromagnetic wave transmitter and receiver, a LED, a modem, or other means for transmitting and receiving a signal carrier. The means for communication allows the compression monitor to upload or download information from a remote network. The compression monitor may also be provided with a global positioning satellite reader (GPS reader), speakers, keypads, telephones, modems, microphones, cameras, or visual displays to allow the user to receive and input information. Data may be exchanged by the means for communication using known communication standards, such as the bluetooth standard.

**[0149]** The remote network may provide other information to the compression monitor and may also receive information from the compression monitor. For example, the compression monitor may be updated with additional waveforms, patient treatment history, ventilation ratios, or other compression-related information of use in a subsequent or current emergency. In the case when a group of monitors are assigned to a single network, each compression monitor may be tracked separately by the remote network and provided with different information (if warranted). The remote network itself may comprise one or more computers, the internet, another CPR-related device, or a human operator capable of remotely programming the compression monitor or remotely prompting the rescuer.

**[0150]** In use, the compression monitor is provided at a location, such as a shopping mall or a public place, and is stored until needed. Upon activation, the compression monitor establishes communication between it and the remote network, which may be a computer located in a call center. Emergency responders, such as police, fire, and ambulance services, may be notified of the activation and directed to go to a location pinpointed by the GPS reader or to contact the person activating the compression monitor. The remote network computer and an operator trained to use the computer may provide voice assistance to the rescuer while monitoring real time streaming data from the compression monitor. The operator or computer may from time to time provide other information to the compression monitor or rescuer. For example, the compression monitor may be provided with data corresponding to what waveform the rescuer should be prompted to perform and the operator may verbally coach the rescuer. In turn, the compression monitor provides data, such as compression depth, rate, force, waveform, and duty cycle, to the operator and computer for medical analysis. Should the rescuer become fatigued then the operator can provide a substitute waveform that is easier for the rescuer to follow. Likewise, the operator can provide verbal encouragement to the rescuer.

**[0151]** Another use for the means for communication is to enable automatic or prompted maintenance of the compression monitor. Either periodically or continuously the compression monitor communicates with a remote network and transmits data such as battery life and status, number of uses, whether any parts need to be replaced, and other information concerning maintenance status. In response, the compression monitor either automatically performs maintenance on itself, such as a software upgrade, or it prompts a user to perform maintenance upon it. Another use for the means for communication is to enable the compression monitor to communicate with additional products used during an emergency response. Example of such a products include those products described above, a drug dispenser, or any

other product useful for responding to the emergency.

[0152] An example of combined product use begins with a rescuer beginning manual resuscitation with the compression monitor. Emergency medical personnel arrive and deploy an automated chest compression device equipped with an AED. The automated chest compression device is adapted to exchange information with the compression monitor. After the automated chest compression device is deployed, the compression monitor automatically communicates with the chest compression device and transfers relevant treatment history to it, such as time under compression, quality of compressions as compared to an ideal waveform, ECG history, and other relevant medical data. Based on the information provided by the compression monitor, the automated chest compression device may provide a defibrillation shock to the patient before beginning compressions. Conversely, the automated chest compression device may determine, based on the transferred data, that chest compressions must be continued before administering a shock. Another example of a combined product is a compression monitor and a separate signal processor. The signal processor may be provided as one or more physical chips (hardware) or may be provided as a computer program (software). In either case, the signal processor may be a separate unit or module and need not be built directly into the monitor. Accordingly, the signal processing units may be provided as stand-alone products. Thus, while the preferred embodiments of the devices have been described in reference to the environment in which they were developed, they are merely illustrative of the principles of the inventions. Other embodiments and configurations may be devised without departing from the scope of the appended claims.

## Claims

1. A device for measuring depth of chest compressions, said device comprising:

an accelerometer capable of producing an acceleration signal (110) corresponding to the acceleration of chest compressions;  
 a first filter (35) capable of producing a filtered acceleration signal from the acceleration signal;  
 a means (36) for integrating the filtered acceleration signal (34), capable of producing a velocity signal;  
 a second filter (37) capable of producing a filtered velocity signal from the velocity signal;  
 a means for integrating (38) the filtered velocity signal and capable of producing a measured depth signal corresponding to the measured depth of chest compressions;  
 a processor, operably connected to the means for integrating the filtered velocity signal (38), programmed to produce an estimated actual depth signal corresponding to the estimated actual depth of chest compressions by calculating an autoregressive moving average of the measured depth signal; and  
 a user feedback system (62) configured to:

receive the estimated actual depth signal;  
 provide, to a rescuer or automatic CPR device, information corresponding to the estimated actual depth of a current chest compression and information concerning compression phase quality and decompression phase quality, wherein the information concerning compression phase quality and decompression phase quality is provided by combining information gained from the acceleration, velocity and estimated actual depth signals.

2. The device of claim 1 further comprising a display (5) operably connected to the processor, and wherein the display (5) is capable of displaying the estimated actual depth signal.

3. The device of claim 1 wherein the user feedback system (5) is capable of providing feedback that indicates whether chest compressions fall within a predetermined range of values.

4. The device of claim 1 further comprising:

a means for measuring an ECG signal of a patient, said ECG signal having a noise component and an actual component;  
 a means for estimating the noise component of the ECG signal operably connected to the means for measuring the ECG signal and to the processor;  
 wherein the processor is further programmed to produce the estimated actual depth signal when the noise component of the ECG signal falls within a predetermined range.

5. The device of claim 4, further comprising a display (5) operably connected to the processor, and wherein the display

(5) is capable of displaying the estimated actual depth signal.

6. The device of claim 4, wherein the user feedback system (5, 62) is capable of providing feedback that indicates whether chest compressions fall within a predetermined range of values.
7. The device of claim 6, wherein the processor is further capable of measuring the rate of chest compressions and wherein the user feedback (5) system is further capable of displaying the rate of chest compressions.
8. The device of claim 6, wherein the processor is further capable of measuring the duty cycle of chest compressions and wherein the user feedback (5) system is further capable of displaying the duty cycle of chest compressions.
9. The device of claim 1 further comprising a means for performing ventilation capable of ventilating the patient during CPR.
10. The device of claim 9, wherein the means for ventilation comprises a means for performing electro-ventilation.
11. The device of claim 1, wherein information regarding the compression phase quality comprises at least one of duty cycle, compression acceleration, and smoothness of compressions.
12. The device of claim 1, wherein information regarding the decompression phase quality comprises at least one of whether a rescuer returns to an actual initial position, duty cycle, smoothness of decompressions, and decompression acceleration.
13. The device of claim 1, wherein the user feedback system comprises at least one of a speaker, a visual display, an LED, a vibrator, and a radio.
14. The device of any preceding claim, comprising a load, force or pressure sensor.

## Patentansprüche

1. Vorrichtung zum Messen einer Tiefe von Thoraxkompressionen, wobei die Vorrichtung Folgendes umfasst:

einen Beschleunigungsmesser, der geeignet ist, um ein Beschleunigungssignal (110) zu erzeugen, das der Beschleunigung von Thoraxkompressionen entspricht;  
einen ersten Filter (35), der geeignet ist, um ein gefiltertes Beschleunigungssignal aus dem Beschleunigungssignal zu erzeugen;  
ein Mittel (36) zum Integrieren des gefilterten Beschleunigungssignals (34), das geeignet ist, um ein Geschwindigkeitssignal zu erzeugen;  
einen zweiten Filter (37), der geeignet ist, um ein gefiltertes Geschwindigkeitssignal aus dem Geschwindigkeitssignal zu erzeugen;  
ein Mittel zum Integrieren (38) des gefilterten Geschwindigkeitssignals und das geeignet ist, um ein gemessenes Tiefensignal, das der gemessenen Tiefe von Thoraxkompressionen entspricht, zu erzeugen;  
einen Prozessor, der betriebsfähig mit dem Mittel zum Integrieren des gefilterten Geschwindigkeitssignals (38) verbunden ist, der durch Berechnen eines autoregressiven gleitenden Durchschnitts des gemessenen Tiefensignals zum Erzeugen eines geschätzten tatsächlichen Tiefensignals, das der geschätzten tatsächlichen Tiefe von Thoraxkompressionen entspricht, programmiert ist; und  
ein Benutzerfeedbacksystem (62), das zu Folgendem konfiguriert ist:

Empfangen des geschätzten tatsächlichen Tiefensignals;  
Bereitstellen von Informationen, einem Retter oder einer automatischen HLW-Vorrichtung, die der geschätzten tatsächlichen Tiefe einer aktuellen Thoraxkompression entsprechen, und von Informationen bezüglich einer Kompressionsphasenqualität und einer Dekompressionsphasenqualität, wobei die Informationen bezüglich der Kompressionsphasenqualität und der Dekompressionsphasenqualität durch Kombinieren der von den Beschleunigungs-, Geschwindigkeits- und geschätzten tatsächlichen Tiefensignalen gewonnenen Informationen bereitgestellt werden.

2. Vorrichtung nach Anspruch 1, ferner umfassend eine Anzeige (5), die betriebsfähig mit dem Prozessor verbunden



ist, und wobei die Anzeige (5) geeignet ist, um das geschätzte tatsächliche Tiefensignal anzuzeigen.

3. Vorrichtung nach Anspruch 1, wobei das Benutzerfeedbacksystem (5) geeignet ist, um ein Feedback bereitzustellen, das angibt, ob Thoraxkompressionen in einen vorbestimmten Wertebereich fallen.

4. Vorrichtung nach Anspruch 1, das ferner Folgendes umfasst:

ein Mittel zum Messen eines EKG-Signals eines Patienten, wobei das EKG-Signal eine Rauschkomponente und eine tatsächliche Komponente aufweist;

ein Mittel zum Schätzen der Rauschkomponente des EKG-Signals, das betriebsfähig mit dem Mittel zum Messen des EKG-Signals und mit dem Prozessor verbunden ist;

wobei der Prozessor ferner programmiert ist, um das geschätzte tatsächliche Tiefensignal zu erzeugen, wenn die Rauschkomponente des EKG-Signals in einen vorbestimmten Bereich fällt.

5. Vorrichtung nach Anspruch 4, ferner umfassend eine Anzeige (5), die betriebsfähig mit dem Prozessor verbunden ist, und wobei die Anzeige (5) geeignet ist, um das geschätzte tatsächliche Tiefensignal anzuzeigen.

6. Vorrichtung nach Anspruch 4, wobei das Benutzerfeedbacksystem (5, 62) geeignet ist, um ein Feedback bereitzustellen, das angibt, ob Thoraxkompressionen in einen vorbestimmten Wertebereich fallen.

7. Vorrichtung nach Anspruch 6, wobei der Prozessor ferner geeignet ist, um die Rate der Thoraxkompressionen zu messen, und wobei das Benutzerfeedbacksystem (5) ferner geeignet ist, um die Rate der Thoraxkompressionen anzuzeigen.

8. Vorrichtung nach Anspruch 6, wobei der Prozessor ferner geeignet ist, um einen Arbeitszyklus der Thoraxkompressionen zu messen, und wobei das Benutzerfeedbacksystem (5) ferner geeignet ist, um den Arbeitszyklus der Thoraxkompressionen anzuzeigen.

9. Vorrichtung nach Anspruch 1, ferner umfassend ein Mittel zum Durchführen einer Luftzufuhr, die geeignet ist, um dem Patienten während der HLW Luft zuzuführen.

10. Vorrichtung nach Anspruch 9, wobei das Mittel zum Luftzuführen ein Mittel zum Durchführen einer elektronischen Luftzufuhr umfasst.

11. Vorrichtung nach Anspruch 1, wobei die Informationen bezüglich der Kompressionsphasenqualität einen Arbeitszyklus, eine Kompressionsbeschleunigung und/oder eine Glätte von Kompressionen umfasst.

12. Vorrichtung nach Anspruch 1, wobei die Informationen bezüglich der Dekompressionsphasenqualität Folgendes umfasst: Ob ein Retter zu einer tatsächlichen Anfangsposition zurückkehrt, einen Arbeitszyklus, eine Glätte von Dekompressionen und/oder eine Dekompressionsbeschleunigung.

13. Vorrichtung nach Anspruch 1, wobei das Benutzerfeedbacksystem einen Lautsprecher, eine visuelle Anzeige, eine LED, einen Schwingungserzeuger und/oder ein Radio umfasst.

14. Vorrichtung nach einem der vorhergehenden Ansprüche, die einen Last-, Kraft- oder Drucksensor umfasst.

## Revendications

1. Dispositif de mesure de la profondeur des compressions thoraciques, ledit dispositif comprenant :

un accéléromètre capable de produire un signal d'accélération (110) correspondant à l'accélération des compressions thoraciques ;

un premier filtre (35) capable de produire un signal d'accélération filtré à partir du signal d'accélération ;

un moyen (36) permettant d'intégrer le signal d'accélération filtré (34), capable de produire un signal de vitesse ;

un second filtre (37) capable de produire un signal de vitesse filtré à partir du signal de vitesse ;

un moyen permettant d'intégrer (38) le signal de vitesse filtré et capable de produire un signal de profondeur mesurée correspondant à la profondeur mesurée des compressions thoraciques ;

un processeur, connecté de manière fonctionnelle aux moyens pour intégrer le signal de vitesse filtré (38), programmé pour produire un signal de profondeur réelle estimée correspondant à la profondeur réelle estimée des compressions thoraciques en calculant une moyenne mobile autorégressive du signal de profondeur mesurée ; et

un système de rétroaction d'utilisateur (62) configuré pour :

recevoir le signal de profondeur réelle estimée ;

fournir, à un secouriste ou à un dispositif de RCP automatique, des informations correspondant à la profondeur réelle estimée d'une compression thoracique actuelle et des informations sur la qualité de la phase de compression et la qualité de la phase de décompression, dans lequel les informations sur la qualité de la phase de compression et la qualité de la phase de décompression sont obtenues en combinant les informations obtenues à partir des signaux d'accélération, de vitesse et de profondeur réelle estimée.

2. Dispositif selon la revendication 1, comprenant en outre un affichage (5) connecté de manière fonctionnelle au processeur et dans lequel l'affichage (5) est capable d'afficher le signal de profondeur réelle estimée.

3. Dispositif selon la revendication 1, dans lequel le système de rétroaction d'utilisateur (5) est capable de fournir une rétroaction indiquant si les compressions thoraciques tombent dans une plage de valeurs prédéterminée.

4. Procédé selon la revendication 1, comprenant en outre :

un moyen permettant de mesurer un signal ECG d'un patient, ledit signal ECG étant doté d'une composante de bruit et d'une composante réelle ;

un moyen d'estimation de la composante de bruit du signal ECG connecté de manière fonctionnelle au moyen de mesure du signal ECG et au processeur ;

dans lequel le processeur est en outre programmé pour produire le signal de profondeur réelle estimée lorsque la composante de bruit du signal ECG tombe dans une plage prédéterminée.

5. Dispositif selon la revendication 4, comprenant en outre un afficheur (5) connecté de manière fonctionnelle au processeur, et dans lequel l'afficheur (5) est capable d'afficher le signal de profondeur réelle estimée.

6. Dispositif selon la revendication 4, dans lequel le système de rétroaction d'utilisateur (5, 62) est capable de fournir une rétroaction indiquant si les compressions thoraciques tombent dans une plage de valeurs prédéterminée.

7. Dispositif selon la revendication 6, dans lequel le processeur est en outre capable de mesurer le taux de compressions thoraciques, et dans lequel le système de rétroaction d'utilisateur (5) est en outre capable d'afficher le taux de compressions thoraciques.

8. Dispositif selon la revendication 6, dans lequel le processeur est en outre capable de mesurer le cycle de fonctionnement des compressions thoraciques, et dans lequel le système de rétroaction d'utilisateur (5) est en outre capable d'afficher le cycle de fonctionnement des compressions thoraciques.

9. Dispositif selon la revendication 1, comprenant en outre un moyen de ventilation permettant de ventiler le patient pendant la RCP.

10. Dispositif selon la revendication 9, dans lequel les moyens de ventilation comprennent des moyens permettant d'effectuer une électro-ventilation.

11. Dispositif selon la revendication 1, dans lequel les informations sur la qualité de la phase de compression comprennent au moins l'un des facteurs de cycle de fonctionnement, d'accélération de compression et de régularité des compressions.

12. Dispositif selon la revendication 1, dans lequel les informations sur la qualité de la phase de décompression comprennent au moins l'un des critères selon lequel un secouriste revient à une position initiale réelle, à un cycle de fonctionnement, à la régularité des décompressions et à l'accélération de la décompression.

13. Dispositif selon la revendication 1, dans lequel le système de rétroaction d'utilisateur comprend au moins un haut-parleur, un afficheur visuel, une LED, un vibreur et une radio.

**14.** Dispositif selon l'une des revendications précédentes, doté d'un capteur de charge, de force ou de pression.

5

10

15

20

25

30

35

40

45

50

55

Fig. 1

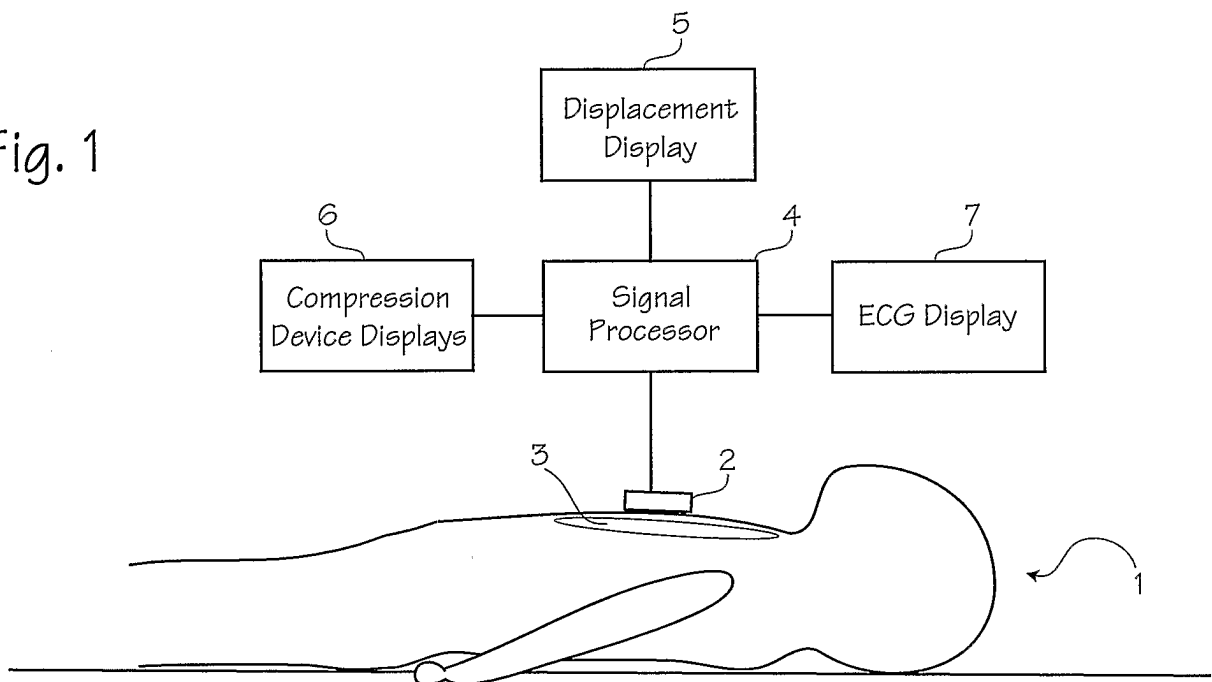


Fig. 2

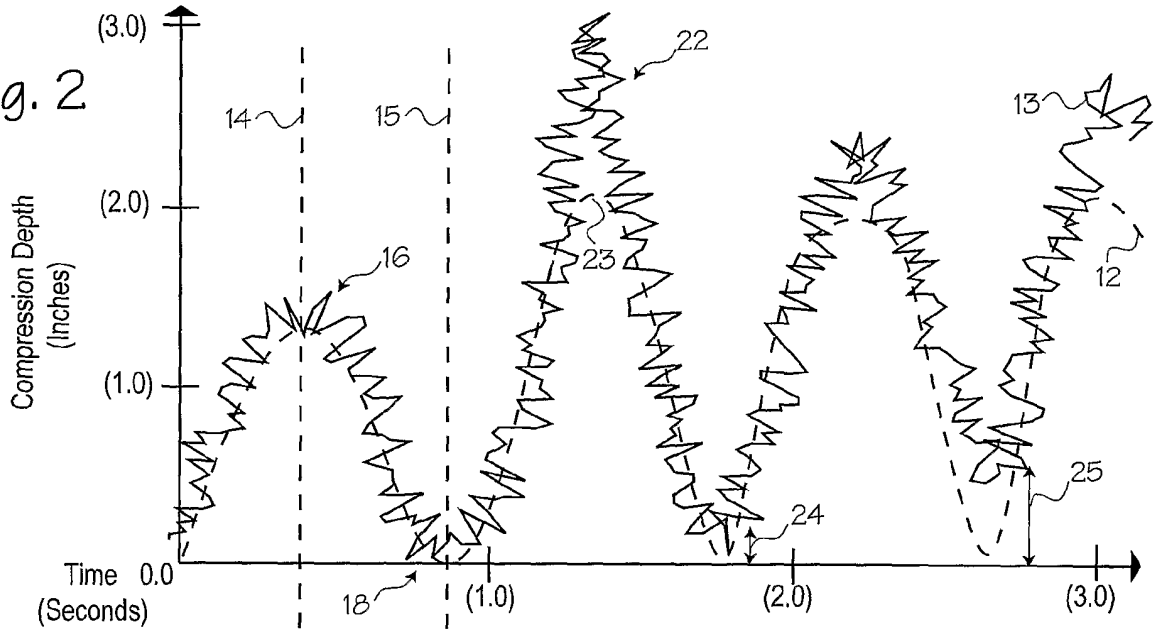


Fig. 3

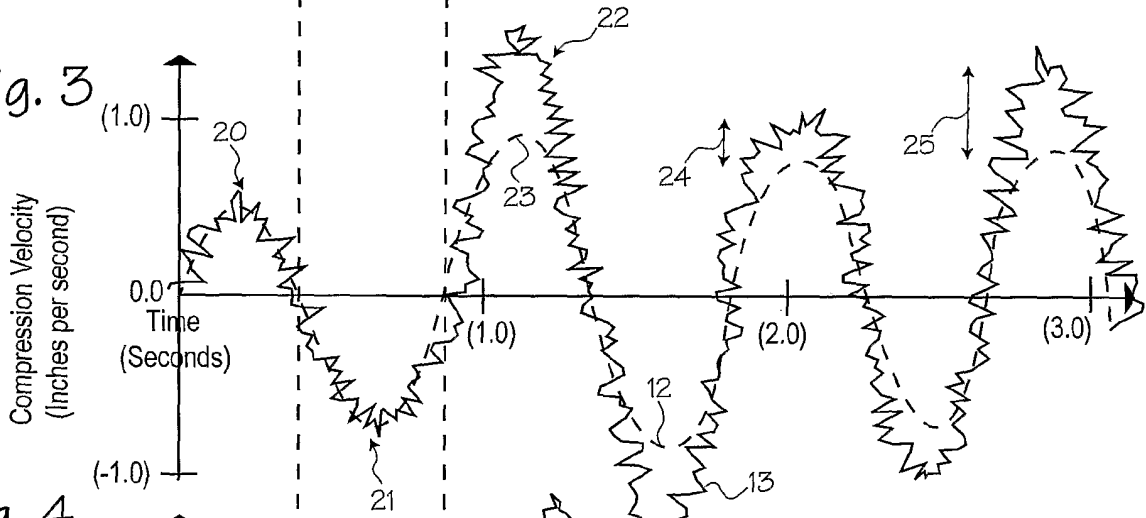


Fig. 4

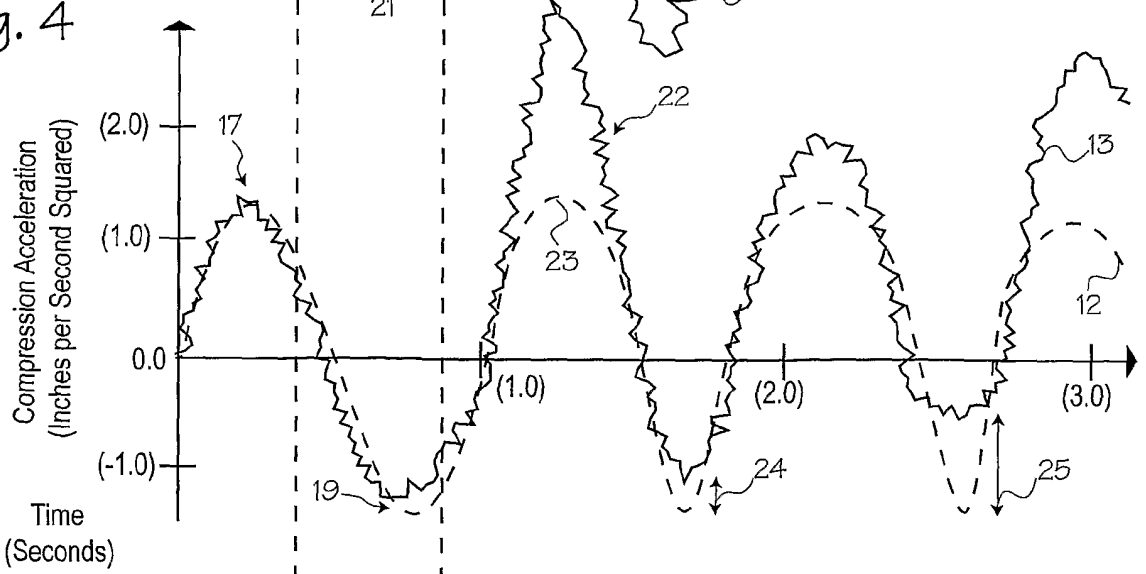


Fig. 5

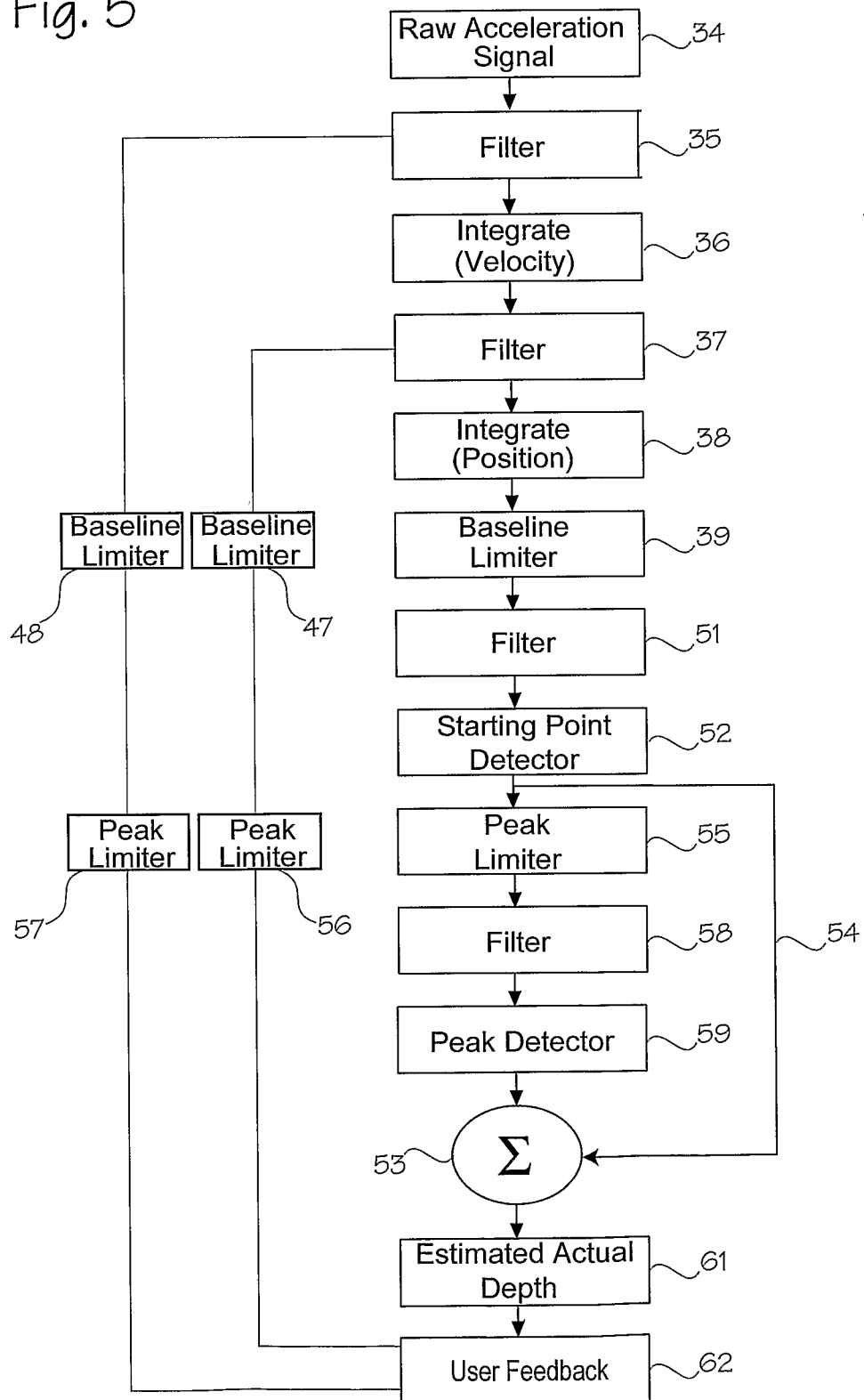


Fig. 6

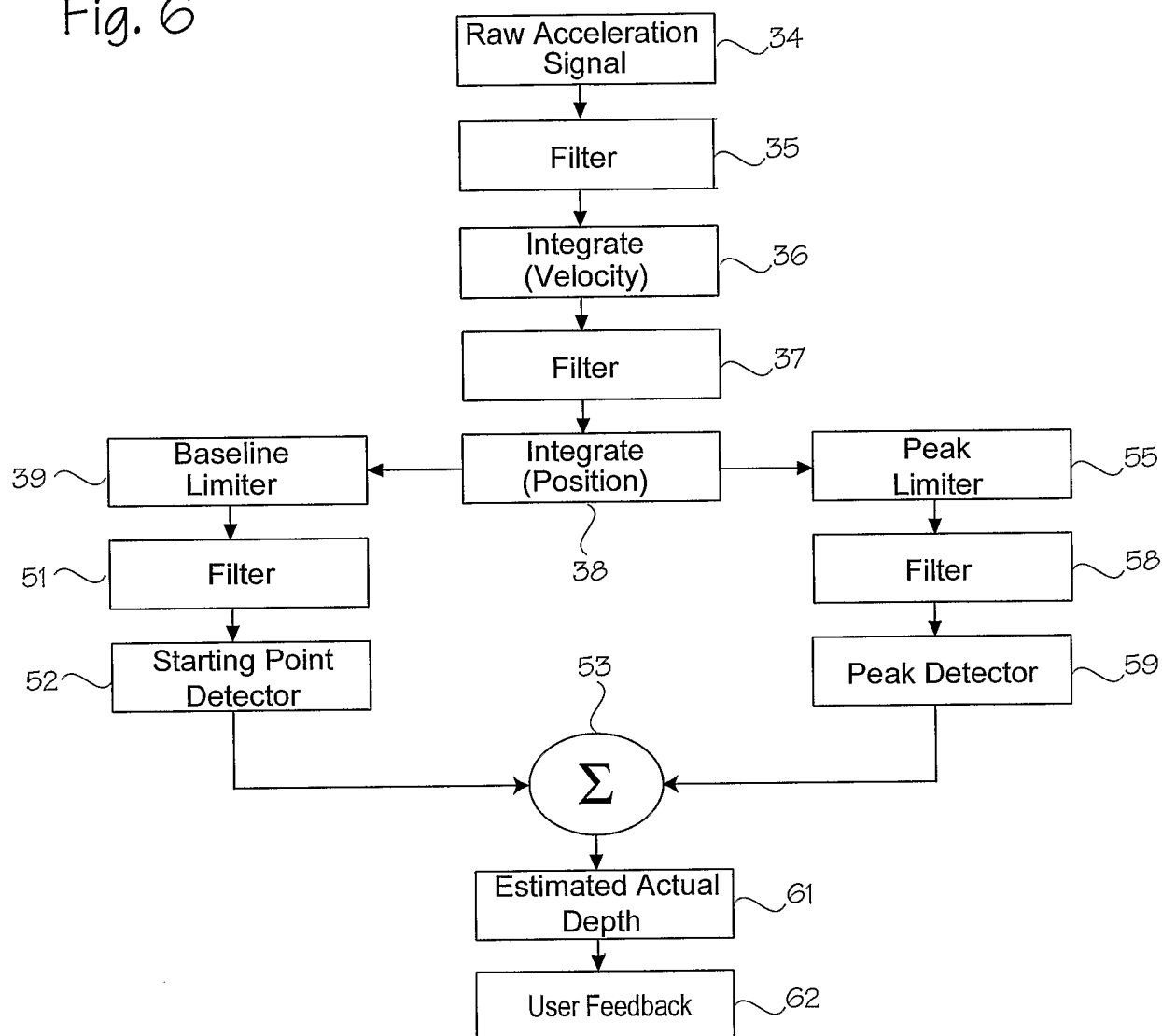


Fig. 7

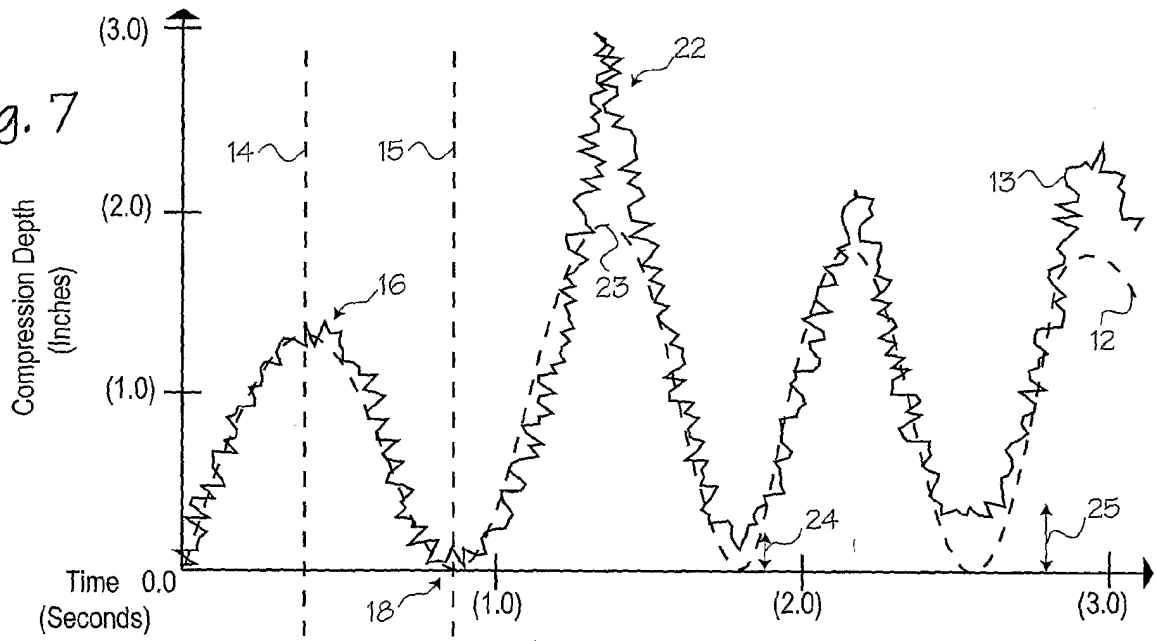


Fig. 8

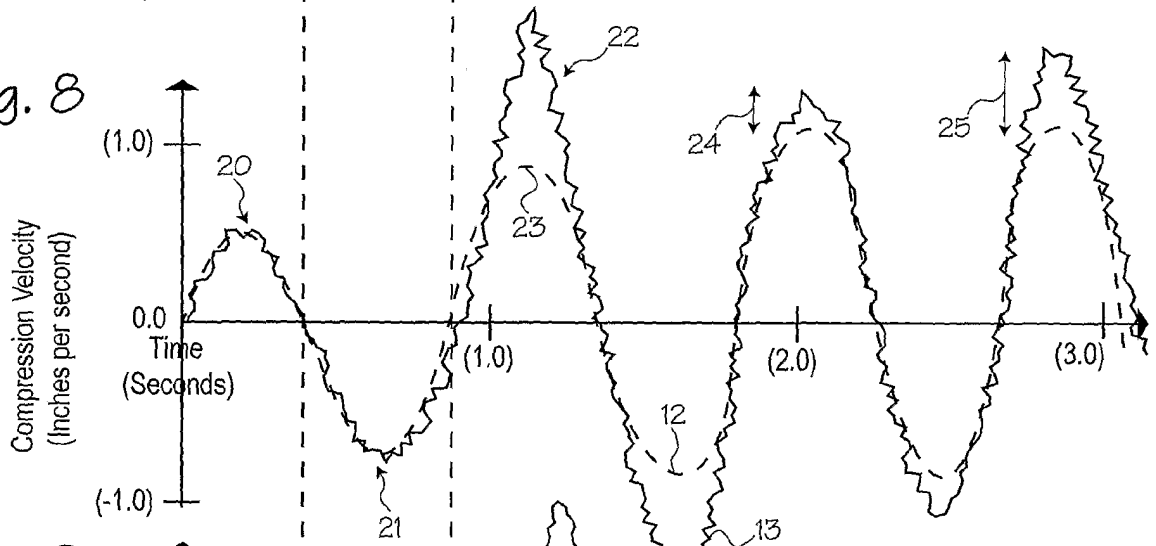


Fig. 9

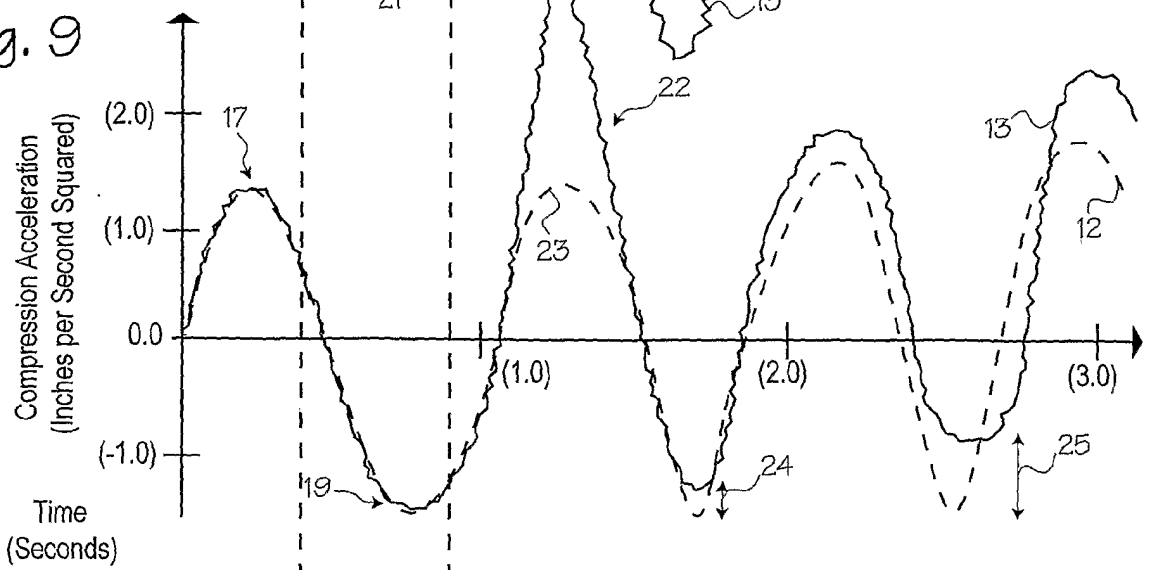




Fig. 10

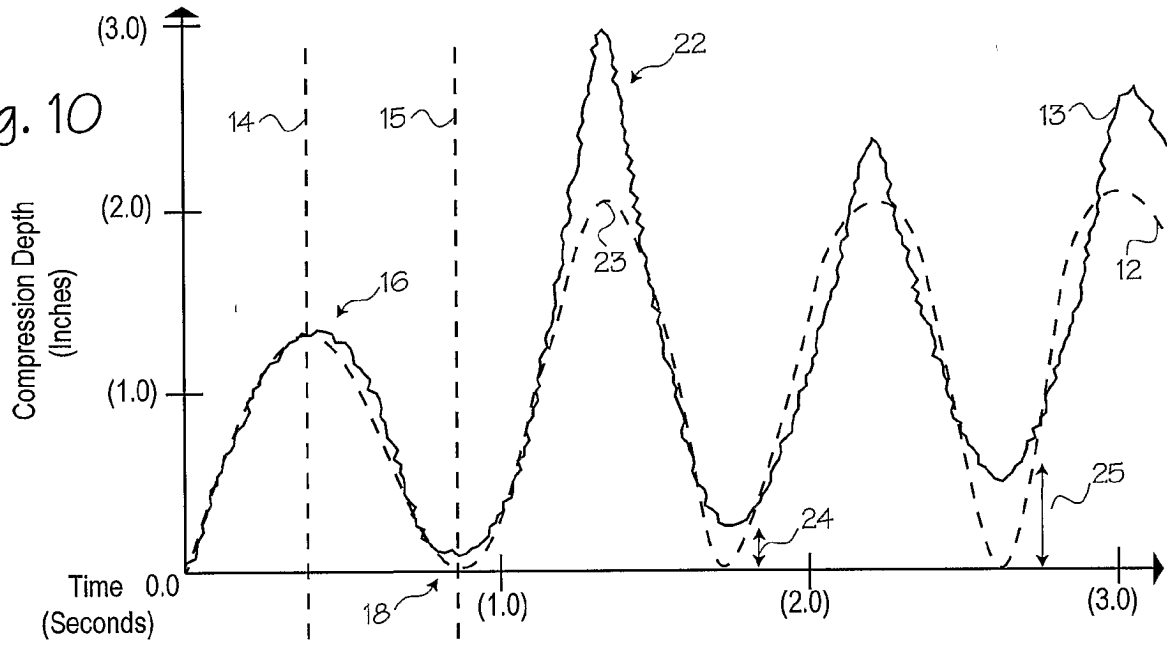


Fig. 11

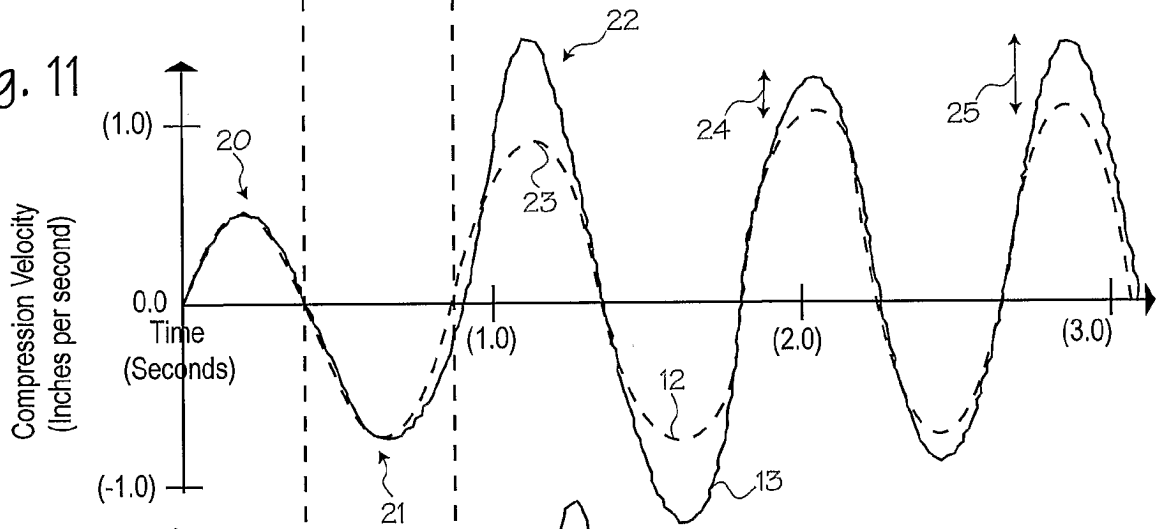


Fig. 12

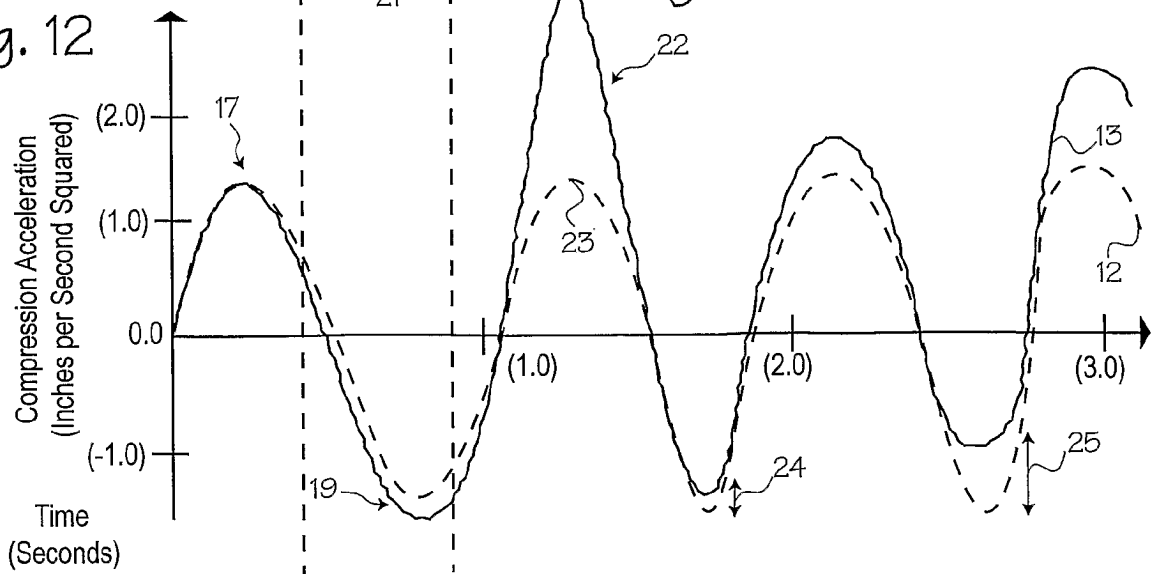


Fig. 13

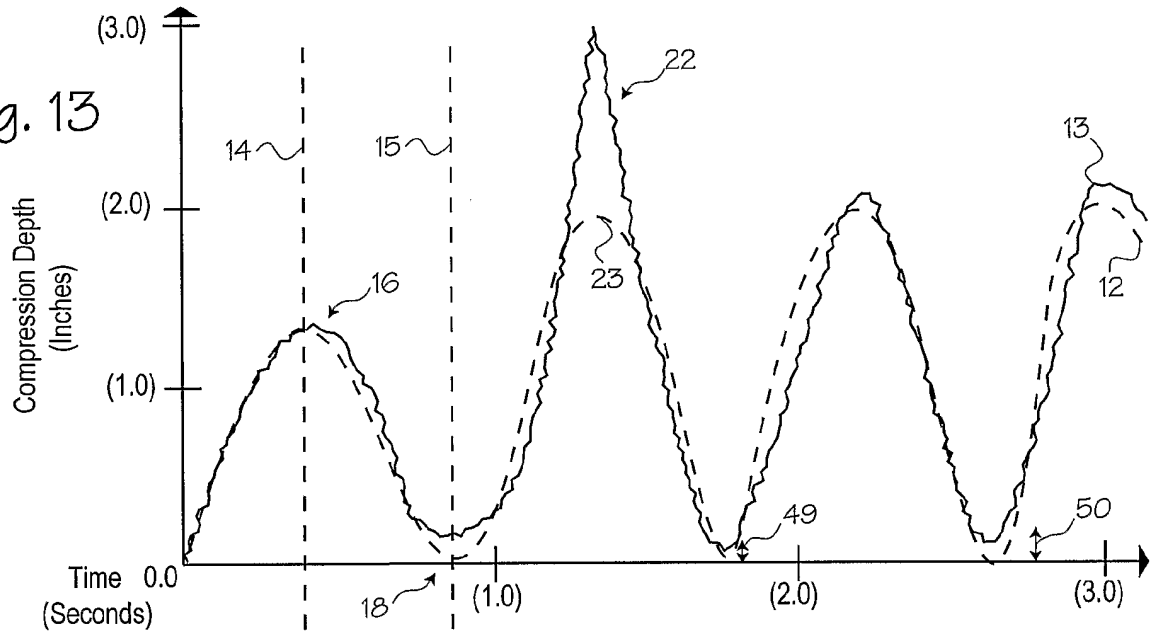


Fig. 14

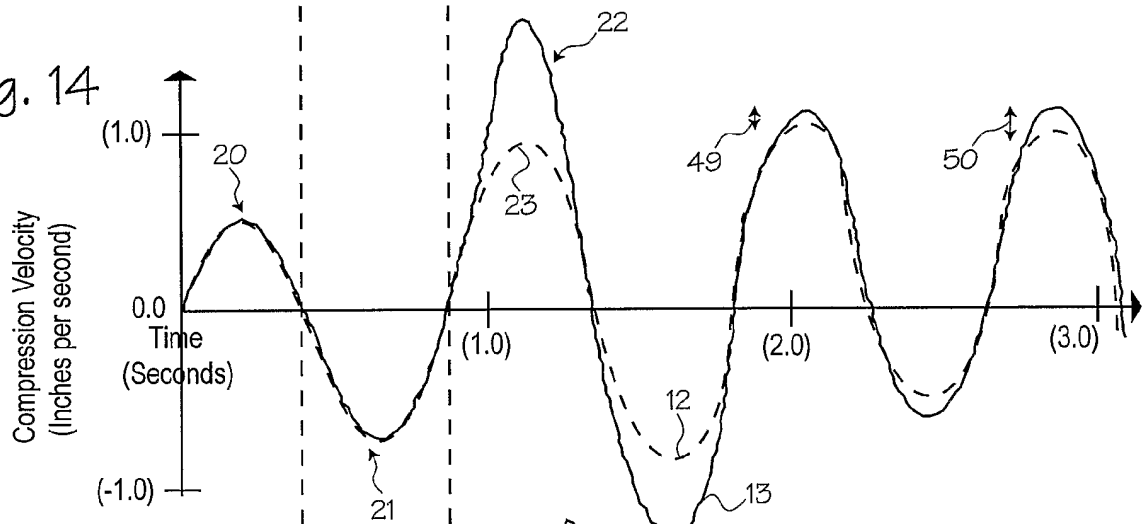


Fig. 15

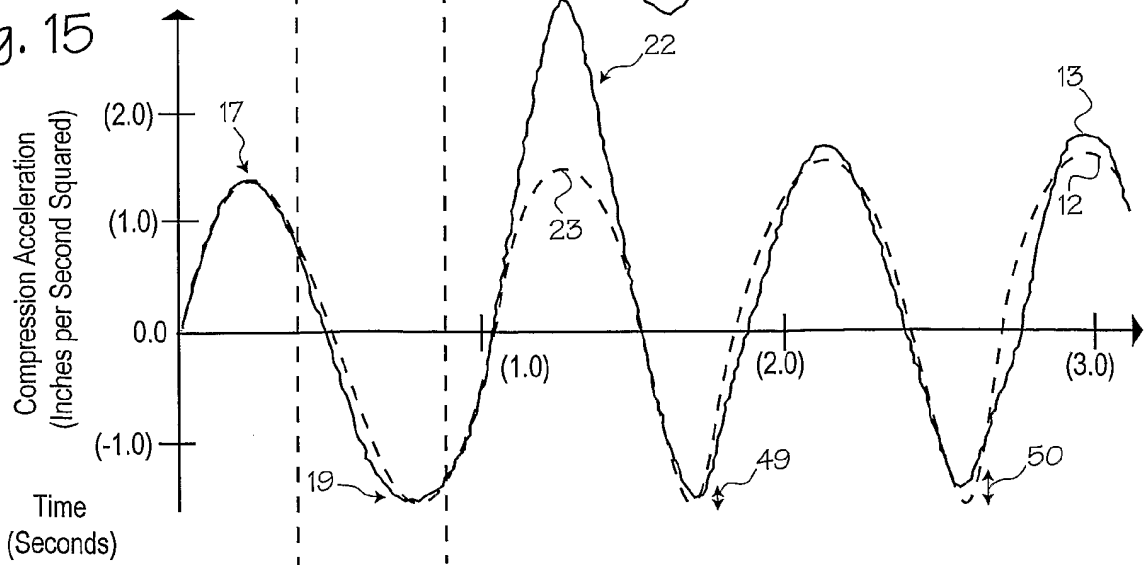


Fig. 16

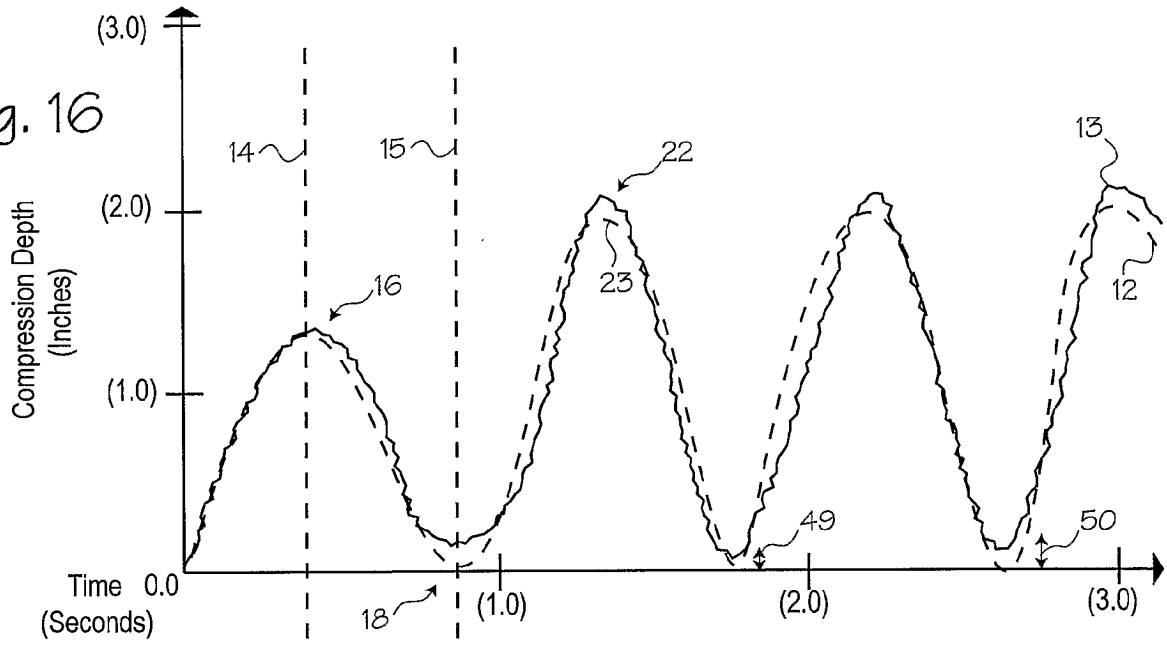


Fig. 17

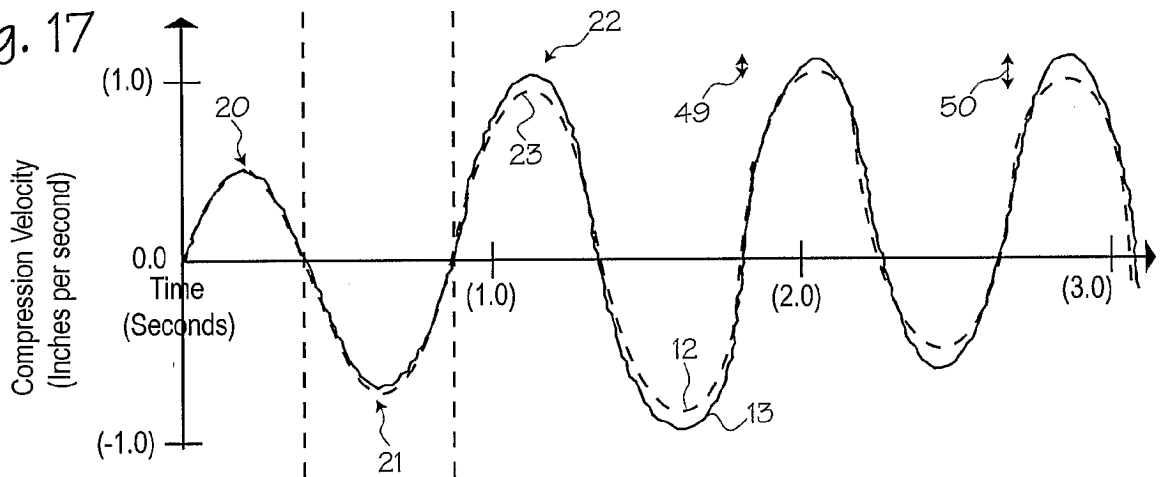


Fig. 18

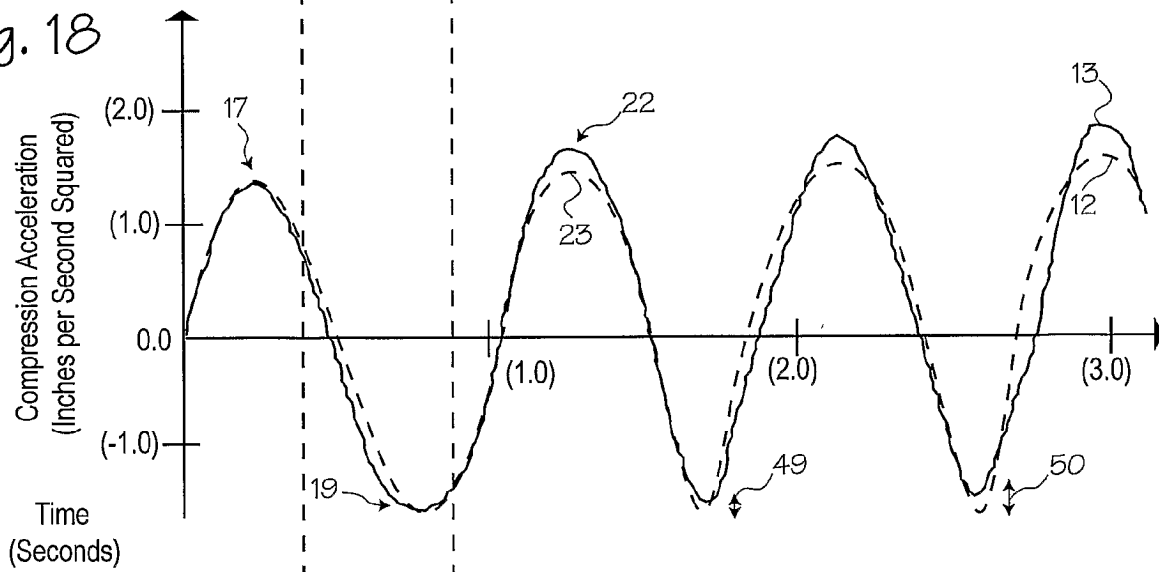
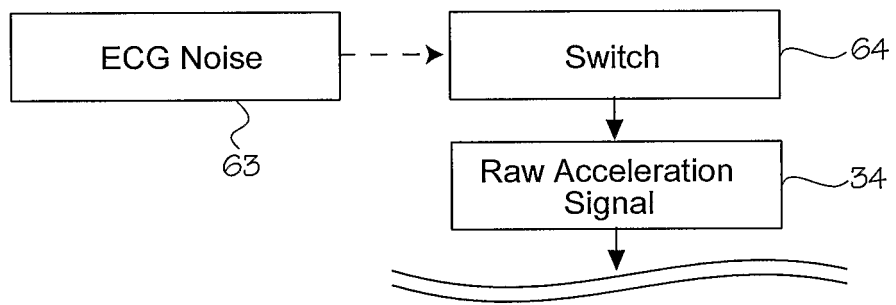


Fig. 19



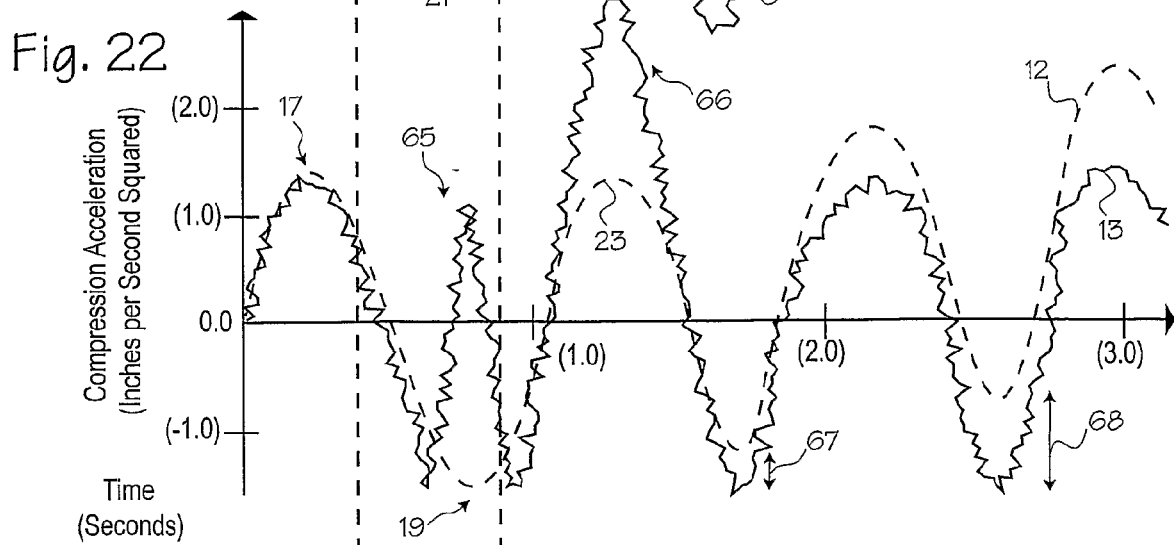
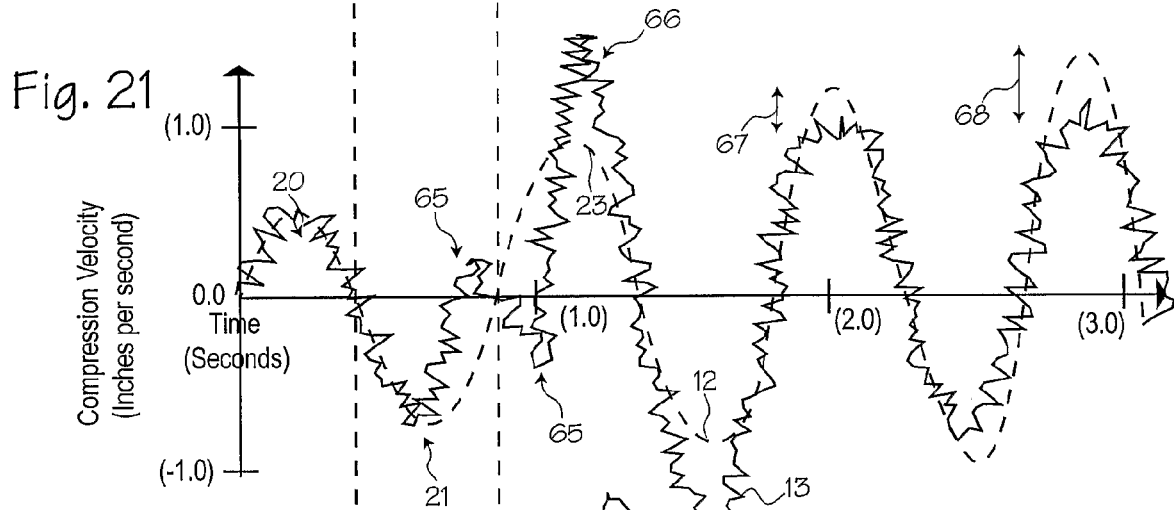
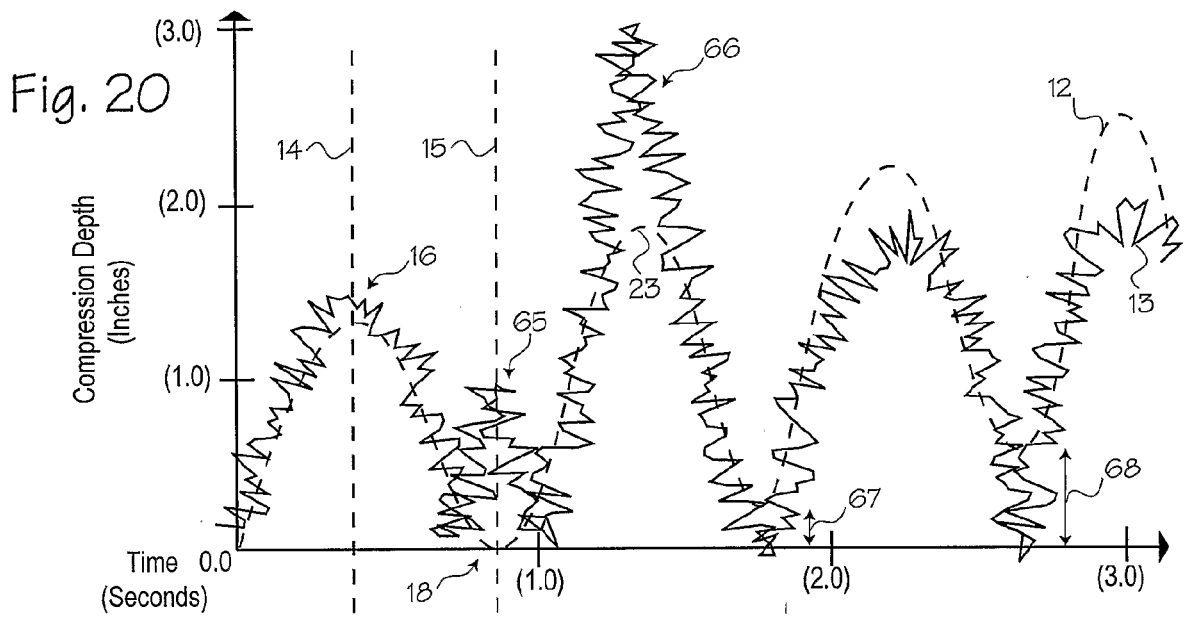


Fig. 23

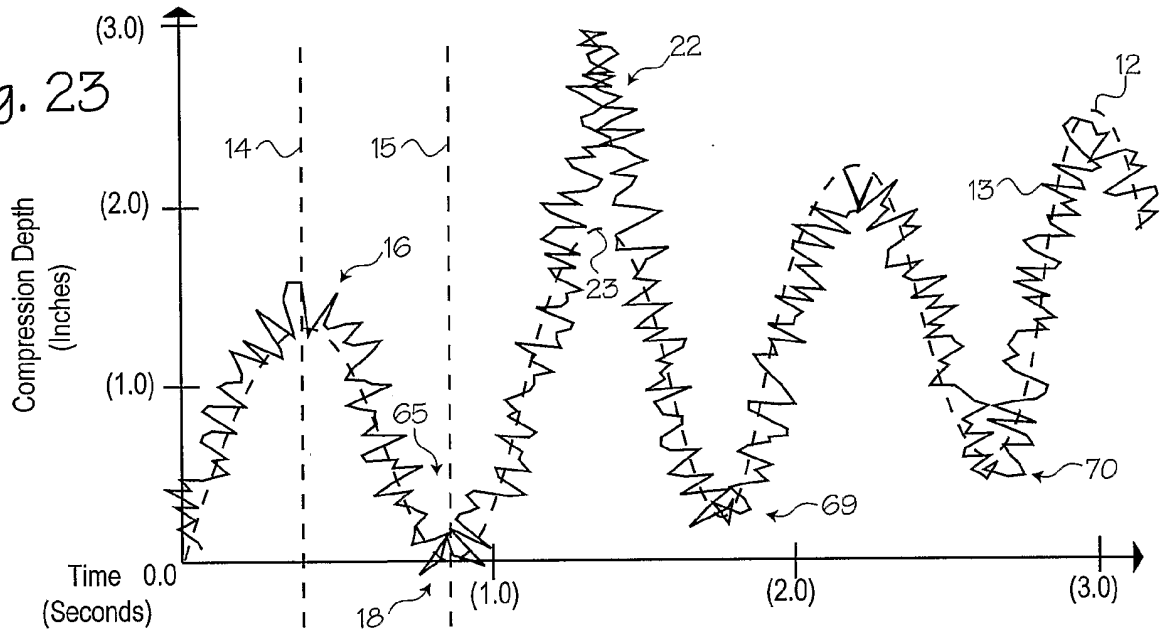


Fig. 24

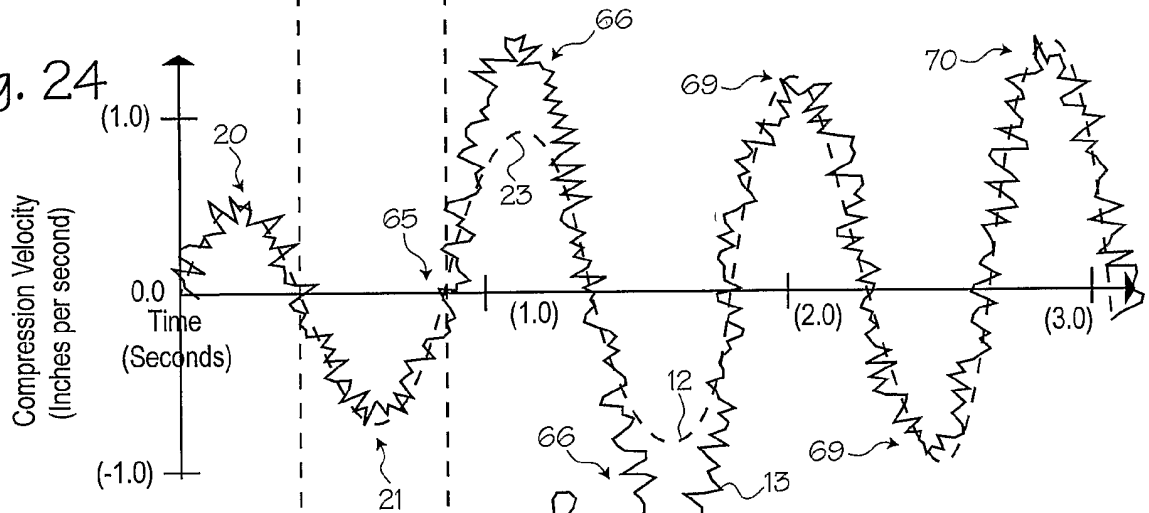


Fig. 25

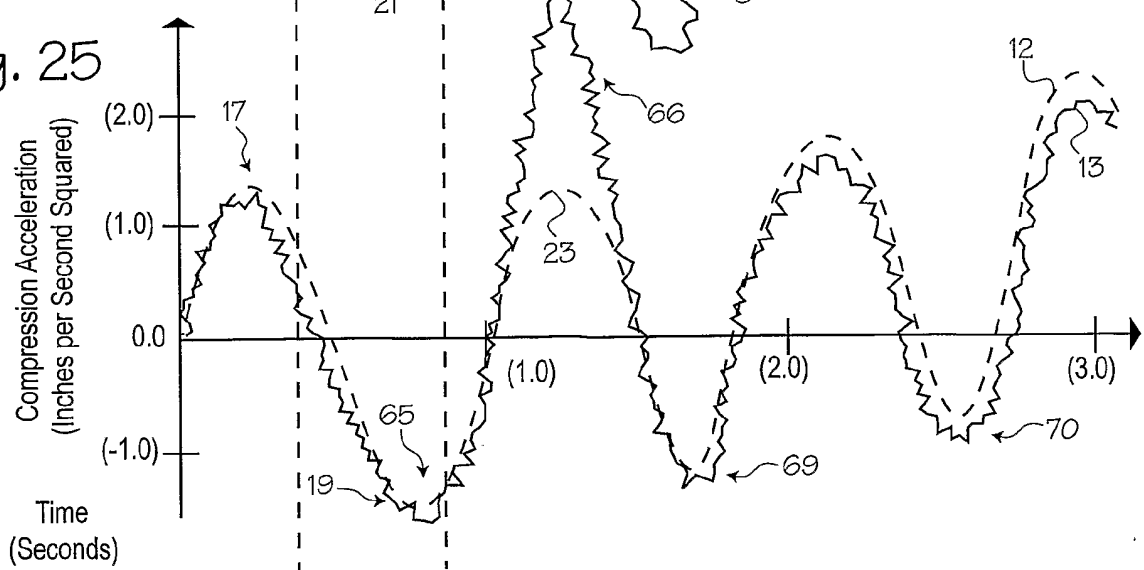


Fig. 26

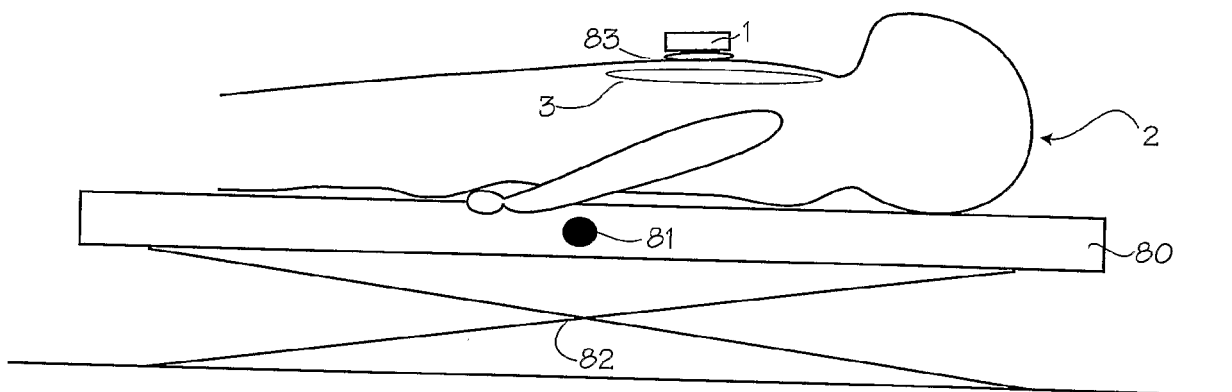


Fig. 27

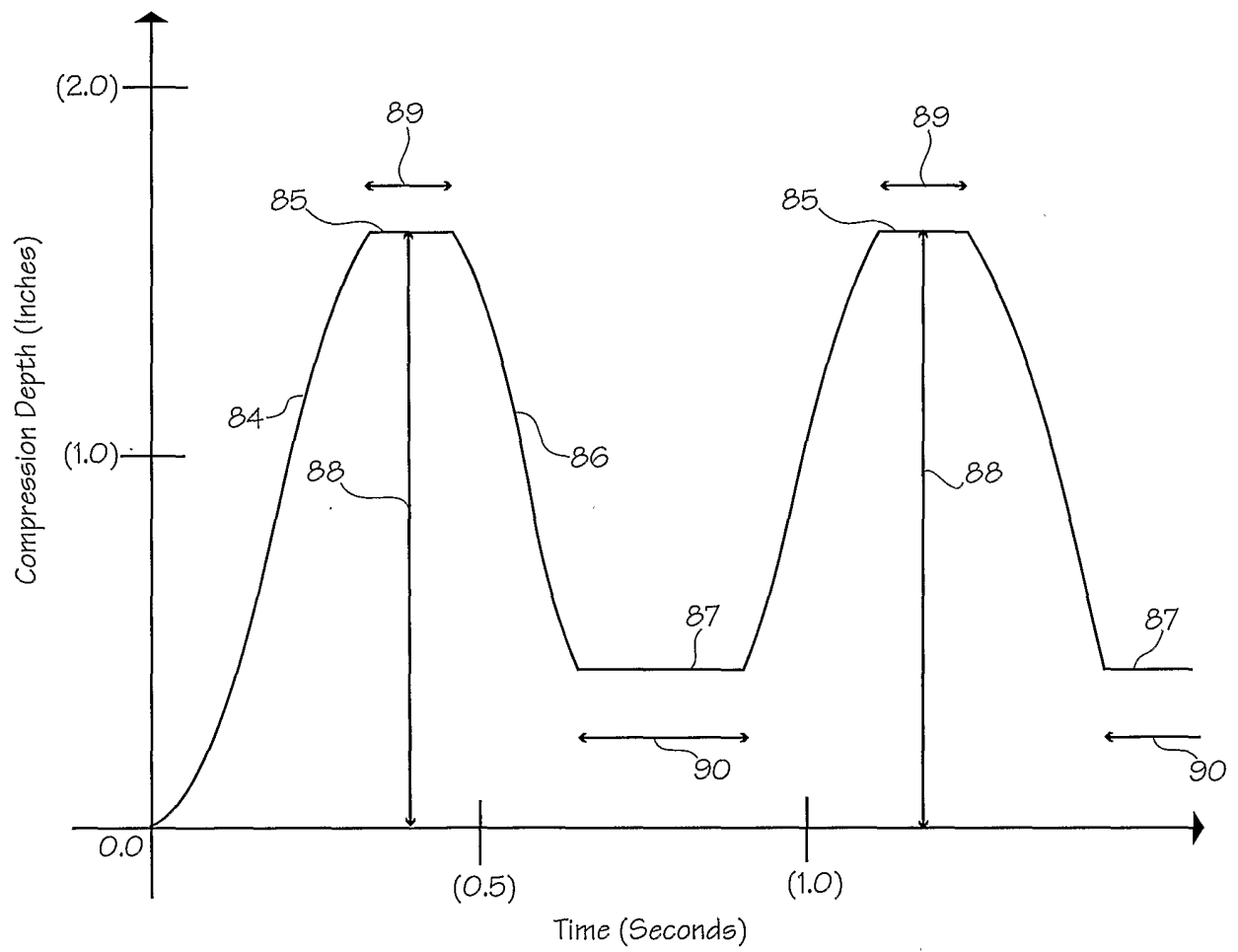




Fig. 28

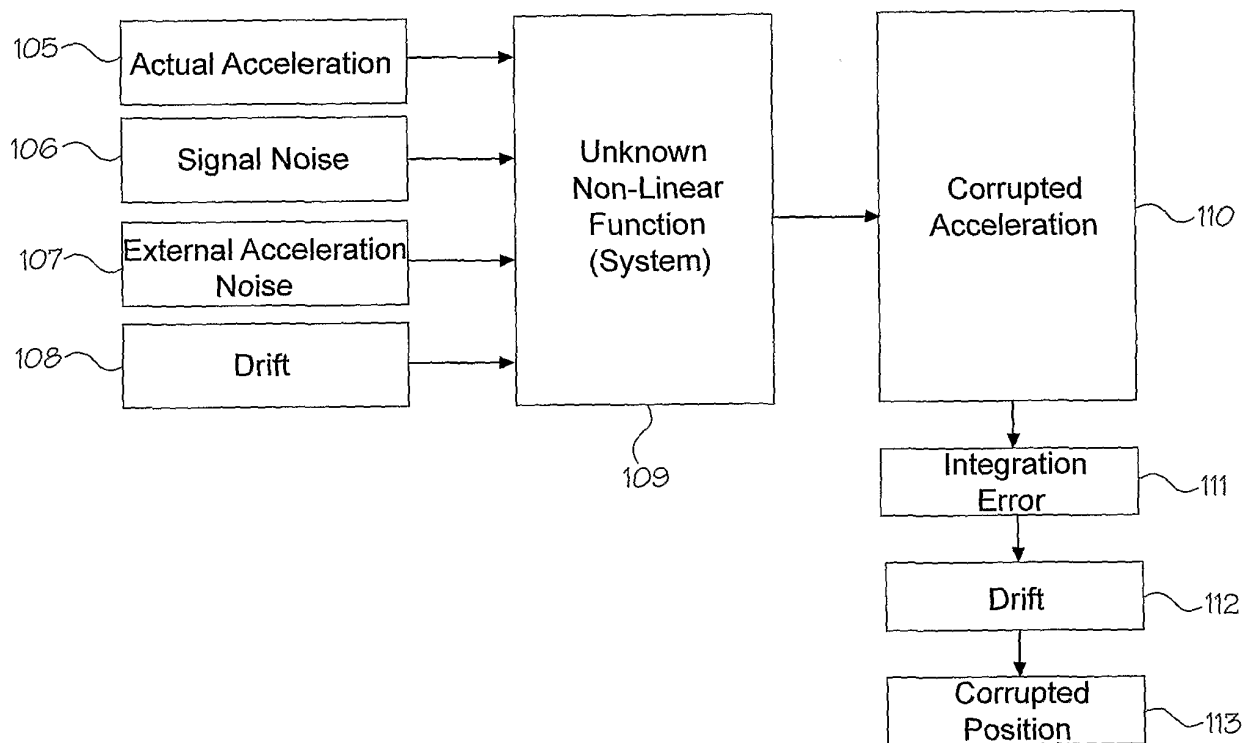


Fig. 29

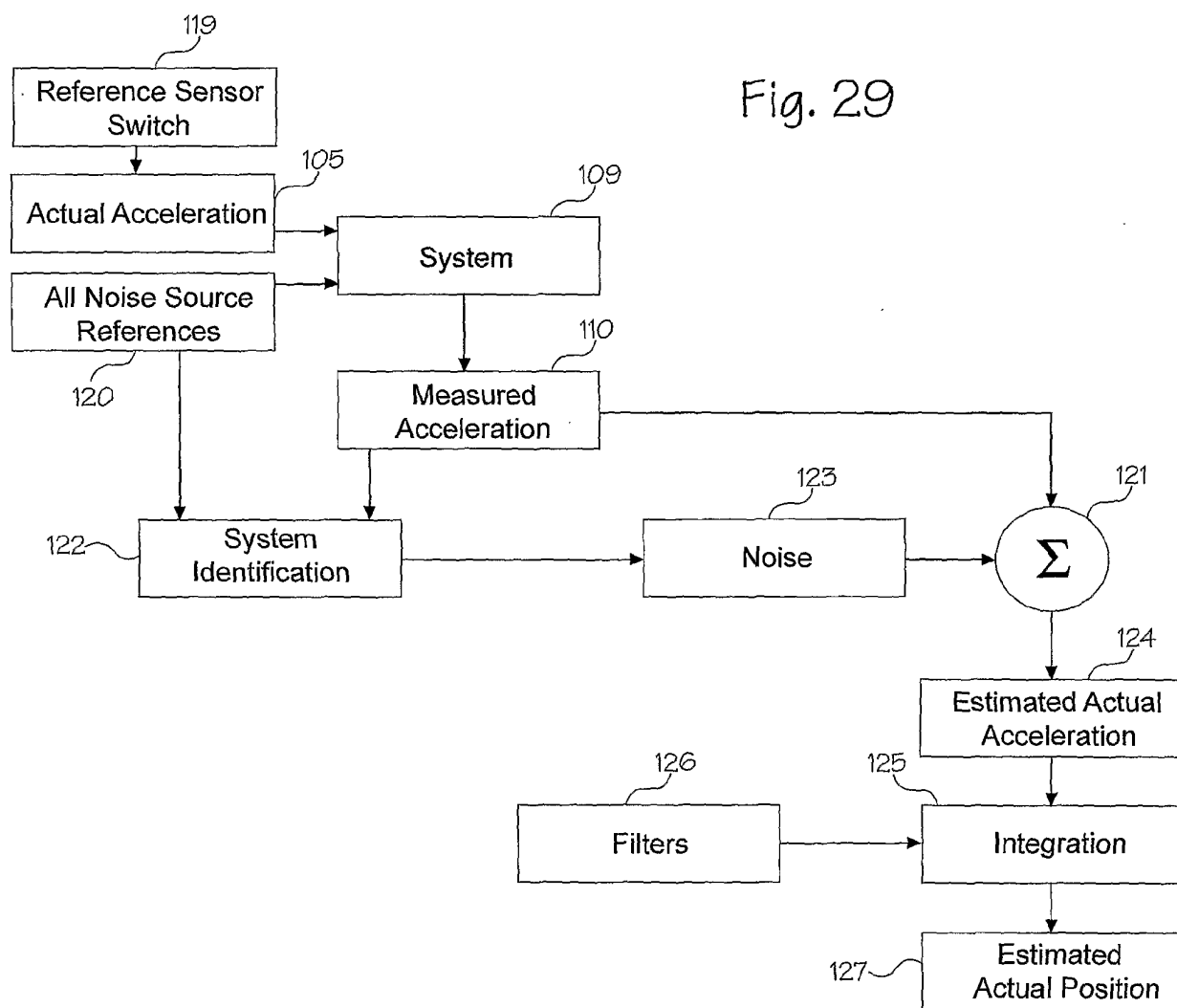


Fig. 30

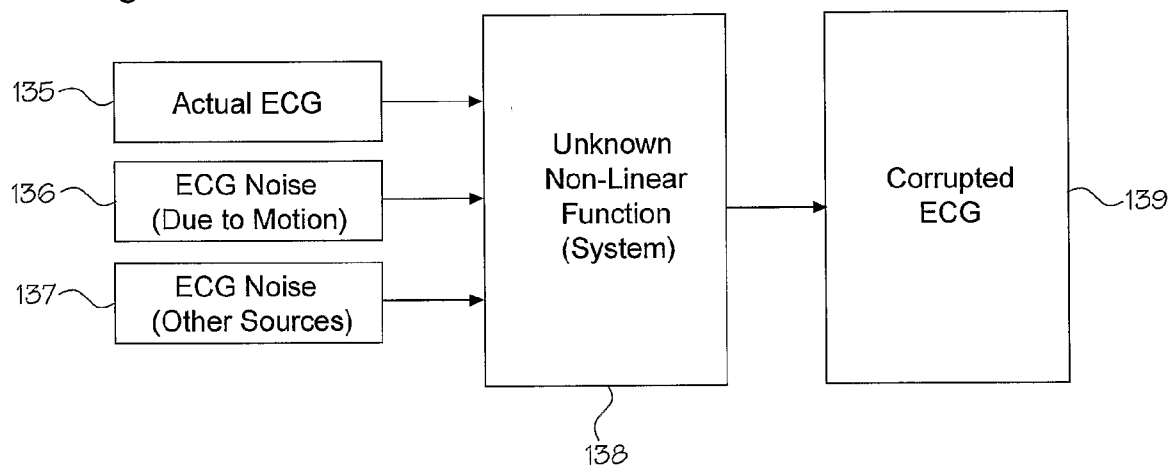
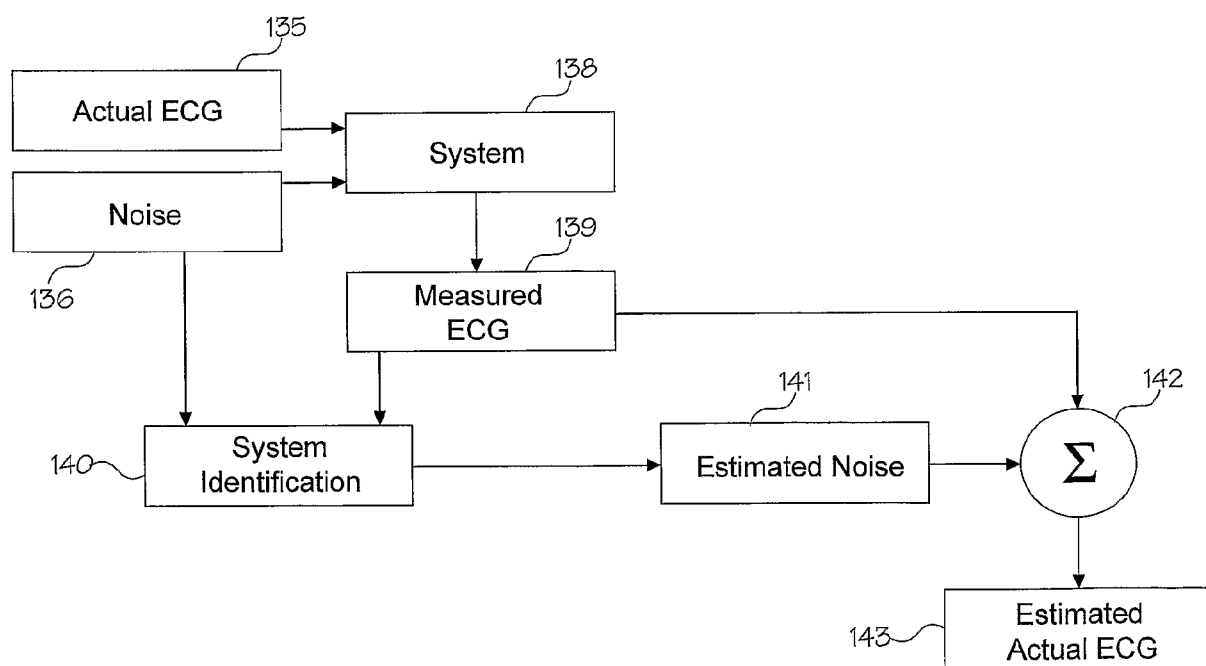
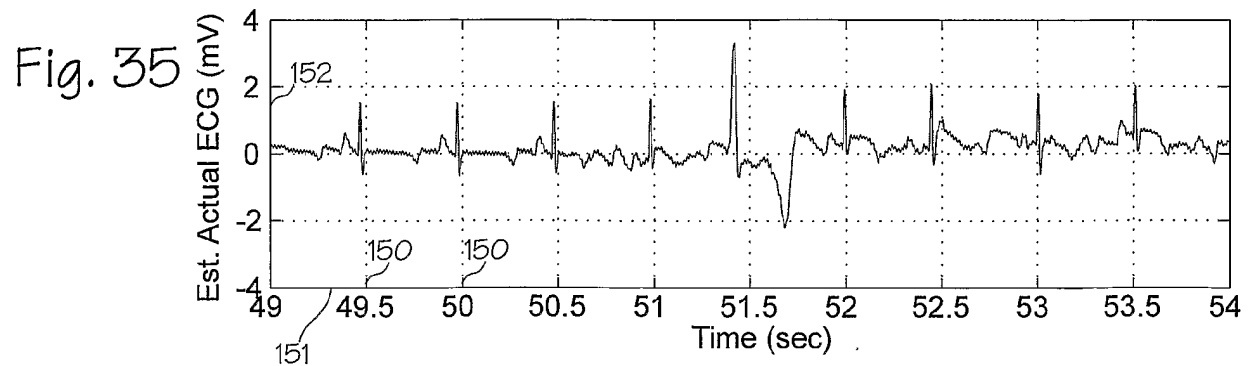
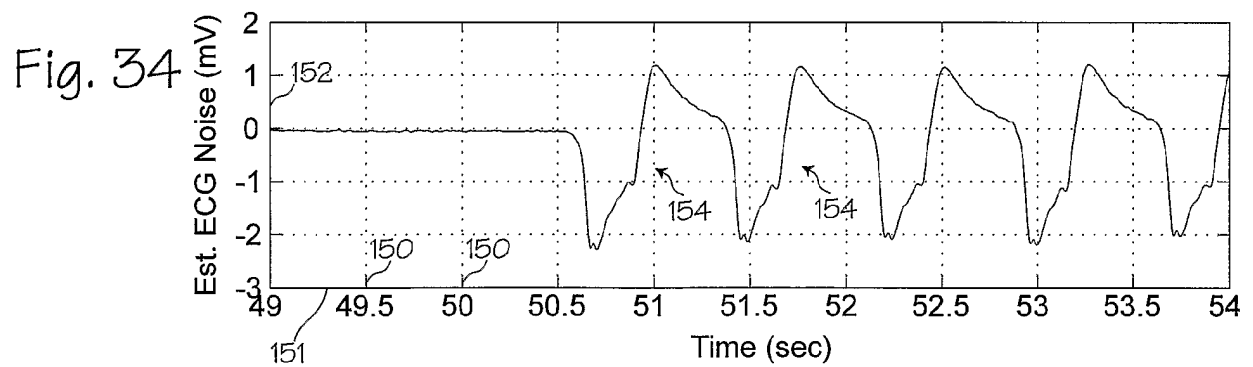
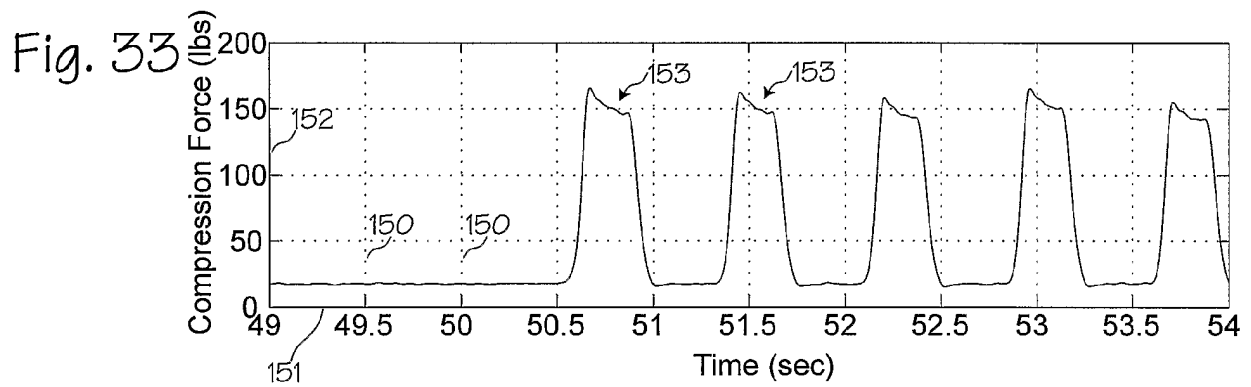
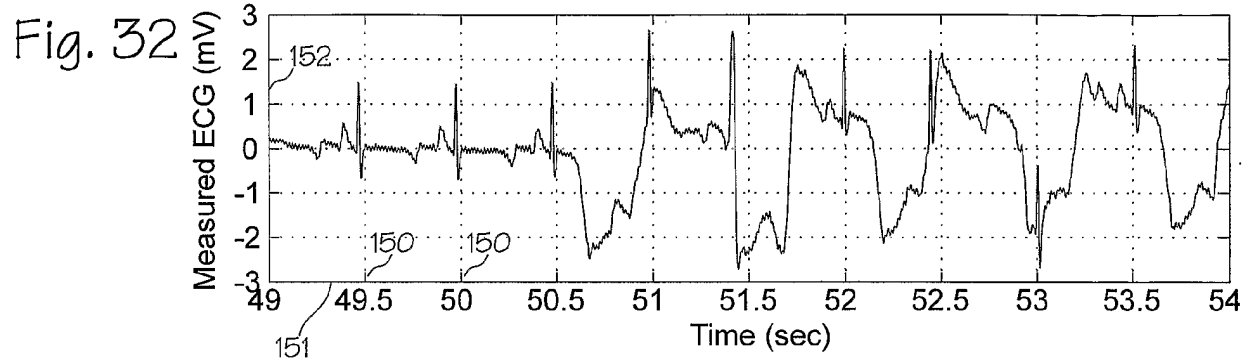


Fig. 31





## REFERENCES CITED IN THE DESCRIPTION

*This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.*

## Patent documents cited in the description

- US 5496257 A [0004]
- US 6125299 A [0004]
- US 6390996 B [0005] [0031] [0145] [0146]
- US 6390996 B1 [0005] [0018]
- US 20010047140 A [0007]
- US 6306107 B [0008]
- US 2002055694 A1 [0018]
- WO 0215836 A2 [0019]
- US 6171267 B1 [0022]
- US 6016445 A [0023]
- US 6263238 B1 [0024]
- US 6066106 A [0144]
- US 86637701 A [0144]
- US 6213960 B [0147]

## Non-patent literature cited in the description

- Guidelines 2000 for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care. *Circulation*, 2000, vol. 102 (I) [0002]
- **OCHOA et al.** The Effect of Rescuer Fatigue on the Quality of Chest Compressions. *Resuscitation*, vol. 37, 149-52 [0003]
- **HIGHTOWER et al.** Decay in Quality of Closed-Chest Compressions over Time. *Ann Emerg. Med*, September 1995, vol. 26 (3), 300-333 [0003]
- Citation Classics. *CC, CURRENT CONTENTS*, 30 October 1989, vol. 44, 22 [0020]
- **BOX, G E P ; JENKINS G M.** *Times series analysis: forecasting and control*, San Francisco, CA, Holden-Day, 1970, vol. 1976, 575 [0020]
- **SARAH MCCABE et al.** On the Use of Nonlinear Autoregressive Moving Average Models for Simulation and System Identification. *PROCEEDINGS OF THE AMERICAN CONTROL CONFERENCE*, 26 June 1991 [0021]
- *PROCEEDINGS OF THE AMERICAN CONTROL CONFERENCE*. IEEE, 26 June 1991, 1758-1763 [0021]

专利名称(译)	在心肺复苏期间确定按压深度的方法		
公开(公告)号	<a href="#">EP1558193A2</a>	公开(公告)日	2005-08-03
申请号	EP2003768544	申请日	2003-10-23
[标]申请(专利权)人(译)	REVIVANT		
申请(专利权)人(译)	REVIVANT CORPORATION		
当前申请(专利权)人(译)	REVIVANT CORPORATION		
[标]发明人	PALAZZOLO JAMES A BERGER RONALD D HALPERIN HENRY R SHERMAN DARREN R		
发明人	PALAZZOLO, JAMES, A. BERGER, RONALD D. HALPERIN, HENRY R. SHERMAN, DARREN, R.		
IPC分类号	A61H31/00 A61B5/00 A61B5/04 A61B5/044 A61B5/053 A61M16/00 A61N1/39 G06K9/00		
CPC分类号	A61B5/04012 A61B5/0535 A61B5/721 A61B5/7242 A61B5/725 A61H31/005 A61H31/006 A61H31/007 A61H31/008 A61H2201/5007 A61H2201/501 A61H2201/5012 A61H2201/5043 A61H2201/5048 A61H2201/5058 A61H2201/5084 A61H2201/5097 A61H2230/04 A61H2230/08 A61H2230/40 A61M16 /00 A61N1/39044 G06K9/0051 Y10S128/901 Y10S601/08 Y10S601/09 Y10S601/10 A61B5/044 A61B5 /053 A61M16/0078		
优先权	10/280220 2002-10-25 US		
其他公开文献	EP1558193A4 EP1558193B1		
外部链接	<a href="#">Espacenet</a>		

#### 摘要(译)

一种处理原始加速度信号的方法，该原始加速度信号由基于加速度计的压缩监测器测量，以产生准确且精确的胸部按压的估计实际深度。在积分期间过滤原始加速度信号，然后过去起始点的移动平均值估计实际当前起始点。然后以类似的方式确定估计的压缩实际峰值。从估计的实际峰值中减去估计的实际起始点，以计算估计的胸部按压的实际深度。另外，可以使用一个或多个参考传感器（例如ECG噪声传感器）来帮助建立压缩的起始点。可以单独使用参考传感器或者与其他信号处理技术组合使用参考传感器，以增强估计的实际压缩深度的准确度和精度。