



(51) International Patent Classification:

A61B 5/00 (2006.01) A61B 5/11 (2006.01)
A61B 5/024 (2006.01) A61B 5/113 (2006.01)
A61B 5/08 (2006.01)

(21) International Application Number:

PCT/IB2015/059341

(22) International Filing Date:

4 December 2015 (04.12.2015)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/090,534 11 December 2014 (11.12.2014) US

(71) Applicant: **KONINKLIJKE PHILIPS N.V.** [NL/NL];
High Tech Campus 5, 5656 AE Eindhoven (NL).

(72) Inventors: **LONG, Xi**; c/o High Tech Campus, Building 5,
5656 AE Eindhoven (NL). **HAAKMA, Reinder**; c/o High
Tech Campus, Building 5, 5656 AE Eindhoven (NL).
FONSECA, Pedro Miguel; c/o High Tech Campus,
Building 5, 5656 AE Eindhoven (NL). **AARTS, Ronaldus
Maria**; c/o High Tech Campus, Building 5, 5656 AE Eindh-
hoven (NL).

(74) Agents: **FREEKE, Arnold** et al.; High Tech Campus
Building 5, 5656 AE Eindhoven (NL).

(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM,

AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY,
BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM,
DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT,
HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR,
KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG,
MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM,
PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC,
SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN,
TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

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

Declarations under Rule 4.17:

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

Published:

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

(54) Title: SYSTEM AND METHOD FOR DETERMINING SPECTRAL BOUNDARIES FOR SLEEP STAGE CLASSIFICATION

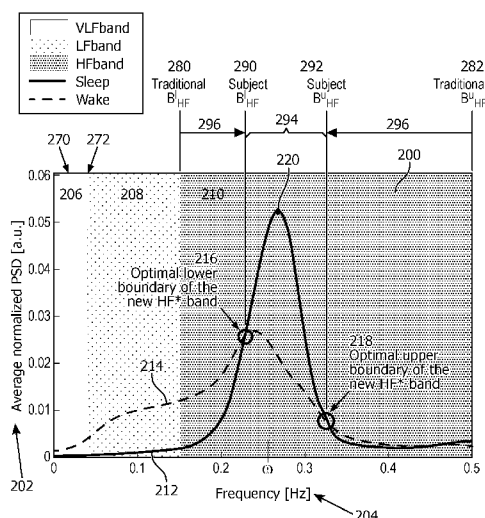


FIG. 2

(57) Abstract: The present disclosure pertains to a system (10) configured to determine spectral boundaries (216, 218) for sleep stage classification in a subject (12). The spectral boundaries may be customized and used for sleep stage classification in an individual subject. Spectral boundaries determined by the system that are customized for the subject may facilitate sleep stage classification with higher accuracy relative to classifications made based on static, fixed spectral boundaries that are not unique to the subject. In some implementations, the system comprises one or more of a sensor (16), a processor (20), electronic storage (22), a user interface (24), and/or other components.

SYSTEM AND METHOD FOR DETERMINING SPECTRAL BOUNDARIES FOR SLEEP STAGE CLASSIFICATION

BACKGROUND

1. Field

[01] The present disclosure pertains to a system and method for determining spectral boundaries for sleep stage classification.

2. Description of the Related Art

[02] Assessment of sleep quality based on monitoring sleep and wake phases during bedtime is known. Over-night polysomnography (PSG) recordings with manually scored hypnograms (done by sleep technicians) for analysis of sleep architecture and occurrence of specific sleep-related problems is known. The analysis is performed based on fixed spectral boundaries that are not individually adjusted for a particular subject.

SUMMARY

[03] Accordingly, one or more aspects of the present disclosure relate to a system configured to determine spectral boundaries for sleep stage classification in a subject. The system comprises one or more sensors, one or more physical computer processors, and/or other components. The one or more sensors are configured to generate output signals that convey information related to a respiratory wave amplitude metric for a sleep session of the subject. The one or more physical computer processors are configured by computer readable instructions to transform the information conveyed by the output signals in individual epochs of time into a frequency domain; determine individual frequencies of respiratory wave amplitude metric peaks within the individual epochs of time; determine an aggregated frequency of the respiratory wave amplitude metric peaks by aggregating the individual frequencies of the respiratory wave metric peaks within the individual epochs of time; determine the spectral boundaries for sleep

stage classification for the subject based on the aggregated frequency; and determine sleep stages of the subject during individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks using the determined spectral boundaries.

[04] Another aspect of the present disclosure relates to a method to determine spectral boundaries for sleep stage classification in a subject with a determination system. The determination system comprises one or more sensors, one or more physical computer processors, and/or other components. The method comprises generating, with the one or more sensors, output signals that convey information related to a respiratory wave amplitude metric for a sleep session of the subject; transforming, with the one or more physical computer processors, the information conveyed by the output signals in individual epochs of time into a frequency domain; determining, with the one or more physical computer processors, individual frequencies of respiratory wave amplitude metric peaks within the individual epochs of time; determining, with the one or more physical computer processors, an aggregated frequency of the respiratory wave amplitude metric peaks by aggregating the individual frequencies of the respiratory wave metric peaks within the individual epochs of time; determining, with the one or more physical computer processors, the spectral boundaries for sleep stage classification for the subject based on the aggregated frequency; and determining, with the one or more physical computer processors, sleep stages of the subject during individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks using the determined spectral boundaries.

[05] Still another aspect of the present disclosure relates to a system configured to determine spectral boundaries for sleep stage classification in a subject. The system comprises means for generating output signals that convey information related to a respiratory wave amplitude metric for a sleep session of the subject; means for transforming the information conveyed by the output signals in individual epochs of time into a frequency domain; means for determining individual frequencies of respiratory wave amplitude metric peaks within the individual epochs of time; means for determining

an aggregated frequency of the respiratory wave amplitude metric peaks by aggregating the individual frequencies of the respiratory wave metric peaks within the individual epochs of time; means for determining the spectral boundaries for sleep stage classification for the subject based on the aggregated frequency; and means for determining sleep stages of the subject during individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks using the determined spectral boundaries.

- [06] These and other objects, features, and characteristics of the present disclosure, as well as the methods of operation and functions of the related elements of structure and the combination of parts and economies of manufacture, will become more apparent upon consideration of the following description and the appended claims with reference to the accompanying drawings, all of which form a part of this specification, wherein like reference numerals designate corresponding parts in the various figures. It is to be expressly understood, however, that the drawings are for the purpose of illustration and description only and are not intended as a definition of the limits of the disclosure.

BRIEF DESCRIPTION OF THE DRAWINGS

- [07] FIG. 1 illustrates a system configured to determine spectral boundaries for sleep stage classification in a subject.
- [08] FIG. 2 illustrates a plot of average normalized power spectral density as a function of frequency.
- [09] FIG. 3 illustrates an example of spectral boundaries of a high frequency band determined for an individual subject using a linear regression model.
- [10] FIG. 4 illustrates a method to determine spectral boundaries for sleep stage classification in a subject with a determination system.

DETAILED DESCRIPTION OF EXEMPLARY EMBODIMENTS

- [11] As used herein, the singular form of “a”, “an”, and “the” include plural references unless the context clearly dictates otherwise. As used herein, the statement that two or more parts or components are “coupled” shall mean that the parts are joined or operate together either directly or indirectly, i.e., through one or more intermediate parts or components, so long as a link occurs. As used herein, “directly coupled” means that two elements are directly in contact with each other. As used herein, “fixedly coupled” or “fixed” means that two components are coupled so as to move as one while maintaining a constant orientation relative to each other.
- [12] As used herein, the word “unitary” means a component is created as a single piece or unit. That is, a component that includes pieces that are created separately and then coupled together as a unit is not a “unitary” component or body. As employed herein, the statement that two or more parts or components “engage” one another shall mean that the parts exert a force against one another either directly or through one or more intermediate parts or components. As employed herein, the term “number” shall mean one or an integer greater than one (i.e., a plurality).
- [13] Directional phrases used herein, such as, for example and without limitation, top, bottom, left, right, upper, lower, front, back, and derivatives thereof, relate to the orientation of the elements shown in the drawings and are not limiting upon the claims unless expressly recited therein.
- [14] FIG. 1 illustrates a system 10 configured to determine spectral boundaries for sleep stage classification in a subject 12. The spectral boundaries and/or other information related to respiratory activity in subject 12 may be used for sleep stage classification. This is because respiratory activity is associated with autonomic nervous activity (ANA) and breathing control which vary during sleep and wakefulness. The information related to respiratory activity that is used for sleep stage classification may be determined based on a spectral analysis (e.g., using the spectral boundaries) of respiratory effort by subject 12, for example. The spectral analysis may include determining and/or

otherwise analyzing spectral powers in different frequency bands (bounded by the determined spectral boundaries) including a very low frequency (VLF) band, a low frequency (LF) band, and a high frequency (HF) band, of respiratory signals from subject 12, and/or other analysis. Spectral boundaries determined by system 10 that are customized for subject 12 may facilitate sleep stage classification with higher accuracy relative to classifications made based on static, fixed spectral boundaries that are not unique to subject 12. In some implementations, system 10 comprises one or more of a sensor 16, a processor 20, electronic storage 22, a user interface 24, and/or other components.

- [15] Sensor 16 is configured to generate output signals conveying information related to respiratory activity in subject 12, cardiac activity in subject 12, movement of subject 12, and/or other information. The respiratory activity, cardiac activity, and/or movement of subject 12 may correspond to a respiratory effort of subject 12 and/or other characteristics of subject 12. The respiratory activity, cardiac activity, and/or movement of subject 12 may correspond to a sleep stage of subject 12 and/or other characteristics of subject 12. The sleep stage of subject 12 may be associated with rapid eye movement (REM) sleep, non-rapid eye movement (NREM) sleep, and/or other sleep. Sensor 16 may comprise one or more sensors that measure such parameters directly and/or indirectly. For example, one or more sensors 16 may generate an output based on a heart rate of subject 12 (e.g., sensor 16 may be a heart rate sensor located on the chest of subject 12, and/or be configured as a bracelet on a wrist of subject 12, and/or be located on another limb of subject 12), movement of subject 12 (e.g., sensor 16 may include a bracelet around the wrist and/or ankle of subject 12 with an accelerometer such that sleep may be analyzed using actigraphy signals), respiration of subject 12, and/or other characteristics of subject 12. In some embodiments, respiratory signals may be measured directly, for example with a chest belt and/or a nose cannula, and/or may be derived from other signals such as photoplethysmography (PPG) signals and/or heart rate, which can be easily be measured, for example, with a wrist-worn sensor device. Although sensor 16 is illustrated at a single location near subject 12, this is not intended to be limiting. Sensor 16 may include

sensors disposed in a plurality of locations, such as for example, within (or in communication with) user interface 24, coupled (in a removable manner) with clothing of subject 12, worn by subject 12 (e.g., as a headband, wristband, etc.), positioned to point at subject 12 while subject 12 sleeps (e.g., a camera that conveys output signals related to movement of subject 12), and/or in other locations.

[16] In some embodiments, sensor 16 is configured to generate output signals conveying information related to an amplitude and/or power of cardiac, respiratory, movement, and/or other (e.g., respiratory effort) signals from subject 12. The output signals may fluctuate with cardiac, respiratory, and/or movement signal wave amplitudes and/or powers as a function of time. In some embodiments, sensor 16 is configured to generate output signals that convey information related to a specific cardiac, respiratory, and/or movement wave amplitude metric for a sleep session of subject 12. This specific cardiac, respiratory, and/or movement wave amplitude metric may be and/or include a power spectral density and/or other metrics of the cardiac, respiratory, movement, and/or other (e.g., respiratory effort) signals from subject 12, for example.

[17] Processor 20 is configured to provide information processing capabilities in system 10. As such, processor 20 may comprise one or more of a digital processor, an analog processor, a digital circuit designed to process information, an analog circuit designed to process information, a state machine, and/or other mechanisms for electronically processing information. Although processor 20 is shown in FIG. 1 as a single entity, this is for illustrative purposes only. In some embodiments, processor 20 may comprise a plurality of processing units. These processing units may be physically located within the same device, or processor 20 may represent processing functionality of a plurality of devices operating in coordination.

[18] As shown in FIG. 1, processor 20 is configured to execute one or more computer program components. The one or more computer program components may comprise one or more of a respiratory activity component 30, a frequency component 32, a spectral boundary component 34, a sleep stage component 36, and/or other components. Processor 20 may be configured to execute components 30, 32, 34, and/or 36 by software;

hardware; firmware; some combination of software, hardware, and/or firmware; and/or other mechanisms for configuring processing capabilities on processor 20.

[19] It should be appreciated that although components 30, 32, 34, and 36 are illustrated in FIG. 1 as being co-located within a single processing unit, in embodiments in which processor 20 comprises multiple processing units, one or more of components 30, 32, 34, and/or 36 may be located remotely from the other components. The description of the functionality provided by the different components 30, 32, 34, and/or 36 described below is for illustrative purposes, and is not intended to be limiting, as any of components 30, 32, 34, and/or 36 may provide more or less functionality than is described. For example, one or more of components 30, 32, 34, and/or 36 may be eliminated, and some or all of its functionality may be provided by other components 30, 32, 34, and/or 36. As another example, processor 20 may be configured to execute one or more additional components that may perform some or all of the functionality attributed below to one of components 30, 32, 34, and/or 36.

[20] In some embodiments, respiratory activity component 30 is configured to facilitate low pass filtering (e.g., a 10th order Butterworth filter with a cut-off frequency of about 0.7Hz) and then normalization (e.g., by subtracting a median peak-trough amplitude estimated over an entire sleep session to remove a signal baseline) of the output signals (e.g., respiratory effort signals) from sensors 16. Then, respiratory activity component 30 is configured to transform the information conveyed by the filtered and/or normalized output signals into a frequency domain. Respiratory activity component 30 is configured to separate the output signals into signal segments that correspond to individual time epochs of information. The length of the individual time epochs may be determined by respiratory activity component 30, may be determined at manufacture, and/or may be determined by other methods. In some embodiments, an individual time epoch may be about 30 seconds long. In some embodiments, respiratory activity component 30 is configured to transform the output signals (or some derivation thereof) into a frequency domain epoch by epoch to create transformed signal segments. In some embodiments, creating the transformed signal segments may include performing a Fourier

Transform, a Fast Fourier Transform, or some other transform on segments of the output signal (or derivation thereof) that correspond to the individual epochs of time.

[21] In some embodiments, respiratory activity component 30 is configured to determine one or more respiratory (effort) wave amplitude metrics based on the transformed signal segments. Respiratory (effort, e.g., and/or cardiac and/or movement) wave amplitudes may be indicative of respiratory wave power. In some embodiments, respiratory activity component 30 is configured to determine a respiratory wave amplitude metric such as power spectral density and/or other metrics. Power spectral density describes how the strength of a respiratory wave signal is distributed in a frequency domain. Power spectral density describes power contributed to a wave, by a frequency, per unit frequency. Power spectral density describes a rate of variance in characteristics of a wave as a function of frequency. An integral of the power spectral density over a given frequency band gives an average power in a signal over that frequency band, for example. In some embodiments, respiratory activity component 30 is configured to determine a respiratory wave amplitude metric such as power spectral density, average power spectral density, average normalized power spectral density, and/or other metrics. Such determinations may be made for individual time epochs that correspond to the individual transformed signal segments (*e.g.*, on a per-segment basis).

[22] In some embodiments, within individual transformed (respiratory effort) signal segments, the logarithms of spectral powers within the VLF, the LF, and the HF bands may be determined by respiratory activity component 30 as well as a ratio of LF and HF band spectral powers, and/or other information. The power spectral density of the individual bands may be normalized by dividing the power spectral density of an individual band by the total spectral power in the LF and HF bands. In some embodiments, respiratory activity component 30 is configured to transform 30 second power spectral density epochs into the frequency domain, normalize the 30 second power spectral density epochs, average the normalized 30 second power spectral density epochs, and/or analyze the output signals in other ways.

[23] By way of a non-limiting example, FIG. 2 illustrates a plot 200 of average normalized power spectral density 202 as a function of frequency 204 (e.g., in the frequency domain) for an individual 30 second (for example) epoch (e.g., an individual transformed signal segment). FIG. 2 illustrates the VLF band 206, the LF band 208, and the HF band 210 at traditional fixed boundary positions (e.g., VLF is typically considered to be .01 - .05Hz, LF is typically considered to be .05 - .15 Hz, and HF is typically considered to be .15 - .5 Hz). FIG. 2 illustrates the normalized power spectral density of a sleep epoch 212 and a wake epoch 214. As described below, spectral boundaries determined for subject 12 correspond to locations 216, 218 where the average normalized power spectral density of wake epoch 214 and sleep epoch 212 cross over each other. The frequencies that correspond to locations 216 and 218 determined for subject 12 are not the same as the traditional fixed boundary frequencies of the HF band (thus determining sleep stages based on spectral boundaries that correspond to locations 216 and 218, instead of the traditional fixed boundary frequencies of the HF band, will produce more accurate sleep stage determinations for subject 12).

[24] Frequency component 32 (FIG. 1) is configured to determine individual frequencies of respiratory (effort) wave amplitude metric peaks (e.g., average normalized power spectral density peaks) within the individual epochs of time (e.g., within individual transformed signal segments). For example, in FIG. 2, frequency component 32 is configured to determine the frequency of peak 220 (e.g., about .26Hz) for the 30 second (for example) epoch of the transformed signal segment shown in FIG. 2. Frequency component 32 is configured to determine an aggregated frequency of the respiratory wave amplitude metric peaks by aggregating the individual frequencies of the respiratory wave amplitude metric peaks (e.g., peaks 220) within the individual epochs of time (e.g., multiple similar epochs/transformed signal segments to the one shown in FIG. 2). As used herein, an aggregated peak frequency refers to a value for frequency determined from consideration and/or combination of the peak frequency values of the respiratory wave amplitude metric of multiple transformed signal segments. For example, the aggregated peak frequency may be determined through one or more of averaging peak frequencies of the multiple

transformed signal elements, a summation of one or more characteristics of the multiple transformed signal elements, graphically overlaying multiple transformed signal elements and visually and/or graphically determining a peak frequency, and/or other methods. In some embodiments, determining the aggregated frequency of the respiratory wave amplitude metric peaks comprises averaging frequencies of average normalized power spectral density peaks (e.g., peak 220 in FIG. 2) from the individual epochs of time. In some embodiments, an average frequency of the power spectral density peaks from individual thirty second epochs of time during the sleep session is a mean respiratory frequency, ω , of subject 12 for a sleep session. In some embodiments, frequency component 32 (FIG. 1) is configured to determine power spectral density peaks (e.g., peaks 220) only within the traditional LF and HF bands (e.g., .01-.5Hz) because the respiration frequency of healthy people typically lies with this range.

[25] Spectral boundary component 34 (FIG. 1) is configured to determine spectral boundaries (e.g., that correspond to the frequencies of locations 216 and 218 in FIG. 2) in subject 12. The spectral boundaries may correspond to different levels of alertness (e.g., different sleep stages) in subject 12. For example, the spectral boundaries may define, describe, and/or be related to sleep stages such as light REM sleep, deep NREM sleep, and/or other sleep stages.

[26] The spectral boundaries (B) are determined based on the aggregated frequency (ω) determined by frequency component 32, and/or other information. In some embodiments, spectral boundary component 34 (FIG. 1) is configured to determine upper (B_{HF}^u) and lower (B_{HF}^l) spectral boundaries 290, 292 of the HF band (e.g., corresponding to frequencies of locations 218 and 216 in FIG. 2) of subject 12 (FIG. 1). The lower spectral boundary 290 of the HF band may be the same as the upper spectral boundary of the LF band (B_{LF}^u). Spectral boundary component 34 is configured such that the upper and lower boundaries 270, 272 of the VLF band remain fixed at .01 - .05 Hz (which means that the lower boundary of the LF band, B_{LF}^l , is also .05Hz). In some embodiments, the upper (B_{HF}^u) and lower (B_{HF}^l) spectral boundaries of the HF band are

determined for subject 12 based on the aggregated (respiratory) frequency ω of subject 12 using linear regression and regression coefficients according to the equation:

$$B = a \omega + b$$

where B is an individual boundary for a subject 12 (e.g., $B = B_{HF}^u$ or $B = B_{HF}^l$), ω is the aggregated (mean respiratory) frequency determined by frequency component 32, and a and b are regression coefficients (e.g., slope and intercept). Regression coefficients a and b may include upper regression coefficients a^u and b^u used when determining B_{HF}^u and lower regression coefficients a^l and b^l used when determining B_{HF}^l . By way of a non-limiting example, the equation $B_{HF}^u = a^u \omega + b^u$ may be used to determine the upper boundary of the HF band in subject 12 and the equation $B_{HF}^l = a^l \omega + b^l$ may be used to determine the lower boundary of the HF band in subject 12.

The regression coefficients a and b are determined by spectral boundary component 34 based on sleep information obtained from a population of users. The two regression coefficients a and b are determined using one or more methods such as the least square estimation (LSE) method, maximum likelihood estimation (MSE), and/or other methods, based on the equations:

$$a = \frac{n \sum_{i=1}^n \omega_i B_i - \left(\sum_{i=1}^n \omega_i \sum_{i=1}^n B_i \right)}{n \sum_{i=1}^n \omega_i^2 - \left(\sum_{i=1}^n \omega_i \right)^2}$$

$$b = \frac{1}{n} \left(\sum_{i=1}^n B_i - a \sum_{i=1}^n \omega_i \right)$$

where $B_i = \{B_1, B_2, \dots, B_i, \dots, B_n\}$ ($i = 1, 2, \dots, n$) is a set of previously determined boundaries from n different subjects in a population of subjects, and $\omega = \{\omega_1, \omega_2, \dots, \omega_i, \dots, \omega_n\}$ ($i = 1, 2, \dots, n$) are the corresponding previously determined mean respiratory

frequencies from n subjects (e.g., determined as described above for subject 12 for the individual subjects in the population). By way of a non-limiting example, upper regression coefficients a'' and b'' may be determined based on upper boundaries (e.g., B_{HF}^u) previously determined for the individual subjects in the population of subjects and lower regression coefficients a' and b' may be determined based on lower boundaries (e.g., B_{HF}^l) previously determined for the individual subjects in the population of subjects. In some embodiments, the previously determined information for the population of subjects may be programmed at manufacture of system 10, stored in electronic storage 22 and/or in other locations and obtained by spectral boundary component 34, entered and/or selected (e.g., by subject 12, a doctor, a caregiver, and/or other users) via user interface 24, and/or determined in other ways.

[27] Sleep stage component 36 (FIG. 1) is configured to determine sleep stages of subject 12. The sleep stages are determined for individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks (ω) using the determined spectral boundaries (B_{HF}^u and B_{HF}^l). In some embodiments, sleep stage component 34 is configured to determine sleep stages based on the output signals from sensors 16, the determined spectral boundaries, and/or other information. For example, FIG. 2 illustrates traditional upper and lower HF band spectral boundaries 280 and 282 and newly determined upper and lower HF band spectral boundaries 290 and 292 determined specifically for subject 12 (FIG. 1). The subject 12 HF band 294 is narrower 296 than the traditional HF band 210. Determining sleep stages based on spectral boundaries that correspond to locations upper and lower boundaries 290 and 292, instead of the traditional fixed upper and lower boundaries 280 and 282 will produce more accurate sleep stage determinations for subject 12.

[28] FIG. 3 illustrates an example of spectral boundaries of the HF band determined for an individual subject 12 using the linear regression model described above. FIG. 3 illustrates that boundaries for an individual subject 12 may be linearly estimated based on the aggregated (mean respiratory) frequency ω . FIG. 3 is a plot 300 of determined boundary frequency 302 versus mean respiratory frequency 304. FIG. 3

shows upper HF band boundary determinations 306 as a function of ω in subject 12 and a corresponding linear regression 308, as well as determined lower HF band boundary determinations 310 as a function of ω in subject 12 and a corresponding linear regression 312.

[29] Returning to FIG. 1, electronic storage 22 comprises electronic storage media that electronically stores information. The electronic storage media of electronic storage 22 may comprise one or both of system storage that is provided integrally (i.e., substantially non-removable) with system 10 and/or removable storage that is removably connectable to system 10 via, for example, a port (e.g., a USB port, a firewire port, etc.) or a drive (e.g., a disk drive, etc.). Electronic storage 22 may comprise one or more of optically readable storage media (e.g., optical disks, etc.), magnetically readable storage media (e.g., magnetic tape, magnetic hard drive, floppy drive, etc.), electrical charge-based storage media (e.g., EPROM, RAM, etc.), solid-state storage media (e.g., flash drive, etc.), and/or other electronically readable storage media. Electronic storage 22 may store software algorithms, information determined by processor 20, information received via user interface 24 and/or external computing systems, and/or other information that enables system 10 to function properly. Electronic storage 22 may be (in whole or in part) a separate component within system 10, or electronic storage 22 may be provided (in whole or in part) integrally with one or more other components of system 10 (e.g., processor 20).

[30] User interface 24 is configured to provide an interface between system 10 and subject 12, and/or other users through which subject 12 and/or other users may provide information to and receive information from system 10. This enables data, cues, results, and/or instructions and any other communicable items, collectively referred to as "information," to be communicated between a user (e.g., subject 12) and one or more of sensor 16, processor 20, and/or other components of system 10. For example, adjusted spectral boundaries may be displayed to a caregiver via user interface 24.

[31] Examples of interface devices suitable for inclusion in user interface 24 comprise a keypad, buttons, switches, a keyboard, knobs, levers, a display screen, a touch

screen, speakers, a microphone, an indicator light, an audible alarm, a printer, a tactile feedback device, and/or other interface devices. In some embodiments, user interface 24 comprises a plurality of separate interfaces. In some embodiments, user interface 24 comprises at least one interface that is provided integrally with processor 20 and/or other components of system 10.

[32] It is to be understood that other communication techniques, either hard-wired or wireless, are also contemplated by the present disclosure as user interface 24. For example, the present disclosure contemplates that user interface 24 may be integrated with a removable storage interface provided by electronic storage 22. In this example, information may be loaded into system 10 from removable storage (e.g., a smart card, a flash drive, a removable disk, etc.) that enables the user(s) to customize the implementation of system 10. Other exemplary input devices and techniques adapted for use with system 10 as user interface 24 comprise, but are not limited to, an RS-232 port, RF link, an IR link, modem (telephone, cable or other). In short, any technique for communicating information with system 10 is contemplated by the present disclosure as user interface 24.

[33] FIG. 4 illustrates a method 400 to determine spectral boundaries for sleep stage classification in a subject with a determination system. The determination system comprises one or more sensors, one or more physical computer processors, and/or other components. The one or more physical computer processors are configured to execute computer program components. The computer program components comprise a respiratory activity component, a frequency component, a spectral boundary component, a sleep stage component, and/or other components. The operations of method 400 presented below are intended to be illustrative. In some embodiments, method 400 may be accomplished with one or more additional operations not described, and/or without one or more of the operations discussed. Additionally, the order in which the operations of method 400 are illustrated in FIG. 4 and described below is not intended to be limiting.

[34] In some embodiments, method 400 may be implemented in one or more processing devices (e.g., a digital processor, an analog processor, a digital circuit

designed to process information, an analog circuit designed to process information, a state machine, and/or other mechanisms for electronically processing information). The one or more processing devices may include one or more devices executing some or all of the operations of method 400 in response to instructions stored electronically on an electronic storage medium. The one or more processing devices may include one or more devices configured through hardware, firmware, and/or software to be specifically designed for execution of one or more of the operations of method 400.

- [35] At an operation 402, output signals conveying information related to a respiratory wave amplitude metric are generated. In some embodiments, the respiratory wave amplitude metric is a power spectral density. In some embodiments, operation 402 is performed by one or more sensors the same as or similar to sensors 16 (shown in FIG. 1 and described herein).
- [36] At an operation 404, the information conveyed by the output signals in individual epochs of time is transformed into a frequency domain. In some embodiments, operation 404 is performed by a processor component the same as or similar to respiratory activity component 30 (shown in FIG. 1 and described herein).
- [37] At an operation 406, individual frequencies of respiratory wave amplitude metric peaks within the individual epochs of time are determined. In some embodiments, operation 406 is performed by a processor component the same as or similar to frequency component 32 (shown in FIG. 1 and described herein).
- [38] At an operation 408, an aggregated frequency of the respiratory wave amplitude metric peaks is determined by aggregating the individual frequencies of the respiratory wave metric peaks within the individual epochs of time. In some embodiments, determining the aggregated frequency of the respiratory wave amplitude metric peaks comprises averaging frequencies of power spectral density peaks from the individual epochs of time. In some embodiments, an average frequency of the power spectral density peaks from individual thirty second epochs of time during the sleep session is a mean respiratory frequency of the subject. In some embodiments, operation

408 is performed by a processor component the same as or similar to frequency component 32 (shown in FIG. 1 and described herein).

[39] At an operation 410, spectral boundaries are determined. The spectral boundaries are determined based on the aggregated frequency. In some embodiments, the spectral boundaries are determined based on the mean respiratory frequency using linear regression. In some embodiments, operation 410 is performed by a processor component the same as or similar to spectral boundary component 34 (shown in FIG. 1 and described herein).

[40] At an operation 412, sleep stages of the subject are determined. The sleep stages are determined during individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks using the determined spectral boundaries. In some embodiments, operation 412 is performed by a processor component the same as or similar to sleep stage component 36 (shown in FIG. 1 and described herein).

[41] In the claims, any reference signs placed between parentheses shall not be construed as limiting the claim. The word “comprising” or “including” does not exclude the presence of elements or steps other than those listed in a claim. In a device claim enumerating several means, several of these means may be embodied by one and the same item of hardware. The word “a” or “an” preceding an element does not exclude the presence of a plurality of such elements. In any device claim enumerating several means, several of these means may be embodied by one and the same item of hardware. The mere fact that certain elements are recited in mutually different dependent claims does not indicate that these elements cannot be used in combination.

[42] Although the description provided above provides detail for the purpose of illustration based on what is currently considered to be the most practical and preferred embodiments, it is to be understood that such detail is solely for that purpose and that the disclosure is not limited to the expressly disclosed embodiments, but, on the contrary, is intended to cover modifications and equivalent arrangements that are within the spirit and scope of the appended claims. For example, it is to be understood that the present

disclosure contemplates that, to the extent possible, one or more features of any embodiment can be combined with one or more features of any other embodiment.

What is claimed is:

1. A system (10) configured to determine spectral boundaries for sleep stage classification in a subject (12), the system comprising:
 - one or more sensors (16) configured to generate output signals that convey information related to a respiratory wave amplitude metric for a sleep session of the subject; and
 - one or more physical computer processors (20) configured by computer readable instructions to:
 - transform the information conveyed by the output signals in individual epochs of time into a frequency domain;
 - determine individual frequencies of respiratory wave amplitude metric peaks within the individual epochs of time;
 - determine an aggregated frequency of the respiratory wave amplitude metric peaks by aggregating the individual frequencies of the respiratory wave metric peaks within the individual epochs of time;
 - determine the spectral boundaries for sleep stage classification for the subject based on the aggregated frequency; and
 - determine sleep stages of the subject during individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks using the determined spectral boundaries.
2. The system of claim 1, wherein the one or more sensors and the one or more physical computer processors are configured such that the respiratory wave amplitude metric is a power spectral density.
3. The system of claim 2, wherein the one or more physical computer processors are configured such that determining the aggregated frequency of the respiratory wave amplitude metric peaks comprises averaging frequencies of power spectral density peaks from the individual epochs of time.

4. The system of claim 3, wherein the one or more physical computer processors are configured such that an average frequency of the power spectral density peaks from individual thirty second epochs of time during the sleep session is a mean respiratory frequency of the subject.

5. The system of claim 4, wherein the one or more physical computer processors are configured such that the spectral boundaries are determined based on the mean respiratory frequency using linear regression.

6. A method to determine spectral boundaries for sleep stage classification in a subject (12) with a determination system (10), the determination system comprising one or more sensors (16) and one or more physical computer processors (20), the method comprising:

generating, with the one or more sensors, output signals that convey information related to a respiratory wave amplitude metric for a sleep session of the subject;

transforming, with the one or more physical computer processors, the information conveyed by the output signals in individual epochs of time into a frequency domain;

determining, with the one or more physical computer processors, individual frequencies of respiratory wave amplitude metric peaks within the individual epochs of time;

determining, with the one or more physical computer processors, an aggregated frequency of the respiratory wave amplitude metric peaks by aggregating the individual frequencies of the respiratory wave metric peaks within the individual epochs of time;

determining, with the one or more physical computer processors, the spectral boundaries for sleep stage classification for the subject based on the aggregated frequency; and

determining, with the one or more physical computer processors, sleep stages of the subject during individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks using the determined spectral boundaries.

7. The method of claim 6, wherein the respiratory wave amplitude metric is a power spectral density.

8. The method of claim 7, wherein determining the aggregated frequency of the respiratory wave amplitude metric peaks comprises averaging frequencies of power spectral density peaks from the individual epochs of time.

9. The method of claim 8, wherein an average frequency of the power spectral density peaks from individual thirty second epochs of time during the sleep session is a mean respiratory frequency of the subject.

10. The method of claim 9, wherein the spectral boundaries are determined based on the mean respiratory frequency using linear regression.

11. A system (10) configured to determine spectral boundaries for sleep stage classification in a subject (12), the system comprising:

means (16) for generating output signals that convey information related to a respiratory wave amplitude metric for a sleep session of the subject;

means (20) for transforming the information conveyed by the output signals in individual epochs of time into a frequency domain;

means (20) for determining individual frequencies of respiratory wave amplitude metric peaks within the individual epochs of time;

means (20) for determining an aggregated frequency of the respiratory wave amplitude metric peaks by aggregating the individual frequencies of the respiratory wave metric peaks within the individual epochs of time;

means (20) for determining the spectral boundaries for sleep stage classification for the subject based on the aggregated frequency; and

means (20) for determining sleep stages of the subject during individual epochs of time in a subsequent sleep session as a function of the aggregated frequency of respiratory wave amplitude metric peaks using the determined spectral boundaries.

12. The system of claim 11, wherein the respiratory wave amplitude metric is a power spectral density.

13. The system of claim 12, wherein the means for determining the aggregated frequency are configured such that determining the aggregated frequency of the respiratory wave amplitude metric peaks comprises averaging frequencies of power spectral density peaks from the individual epochs of time.

14. The system of claim 13, wherein the means for determining the aggregated frequency are configured such that an average frequency of the power spectral density peaks from individual thirty second epochs of time during the sleep session is a mean respiratory frequency of the subject.

15. The system of claim 14, wherein the means for determining the spectral boundaries are configured such that the spectral boundaries are determined based on the mean respiratory frequency using linear regression.

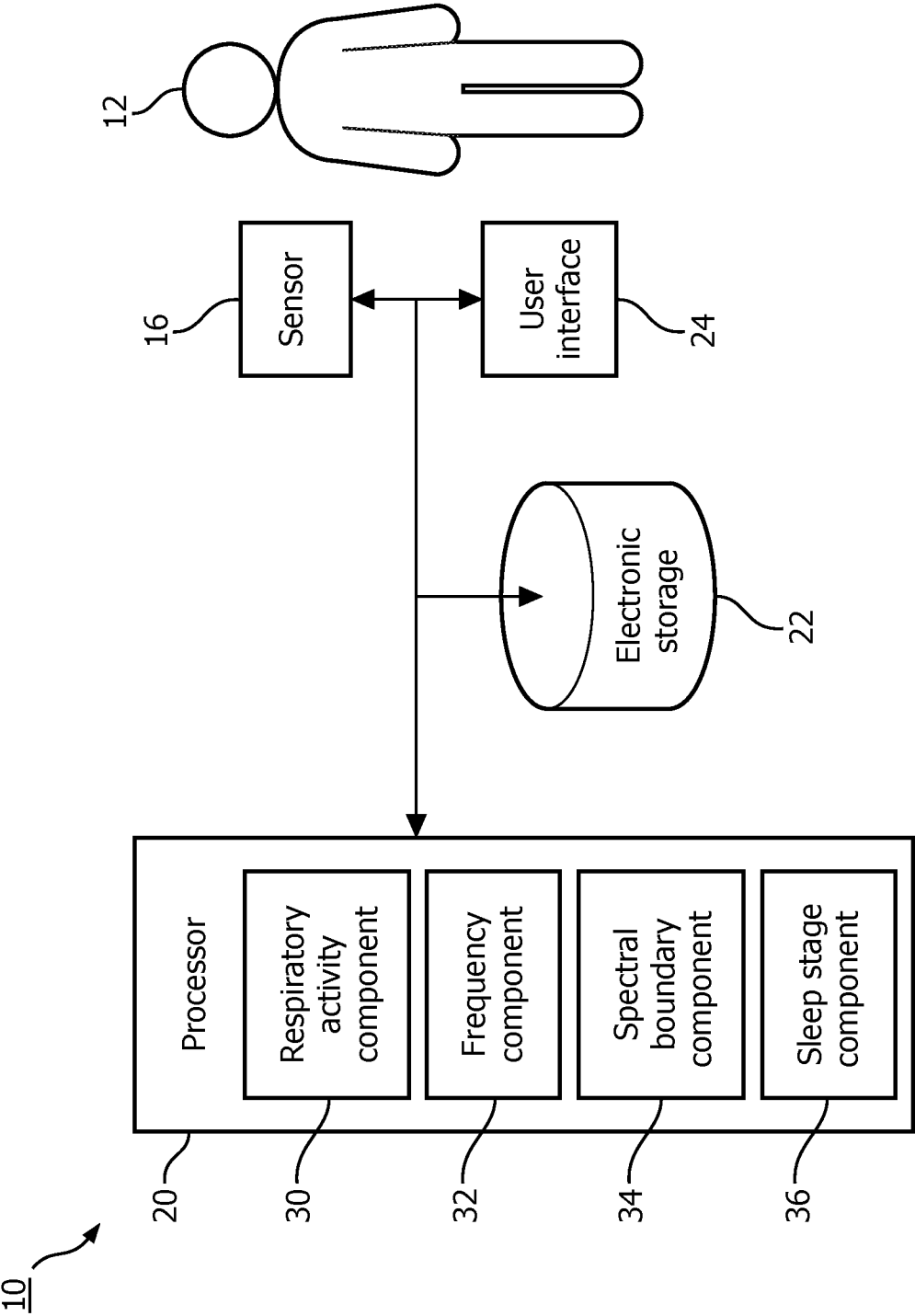


FIG. 1

2/4

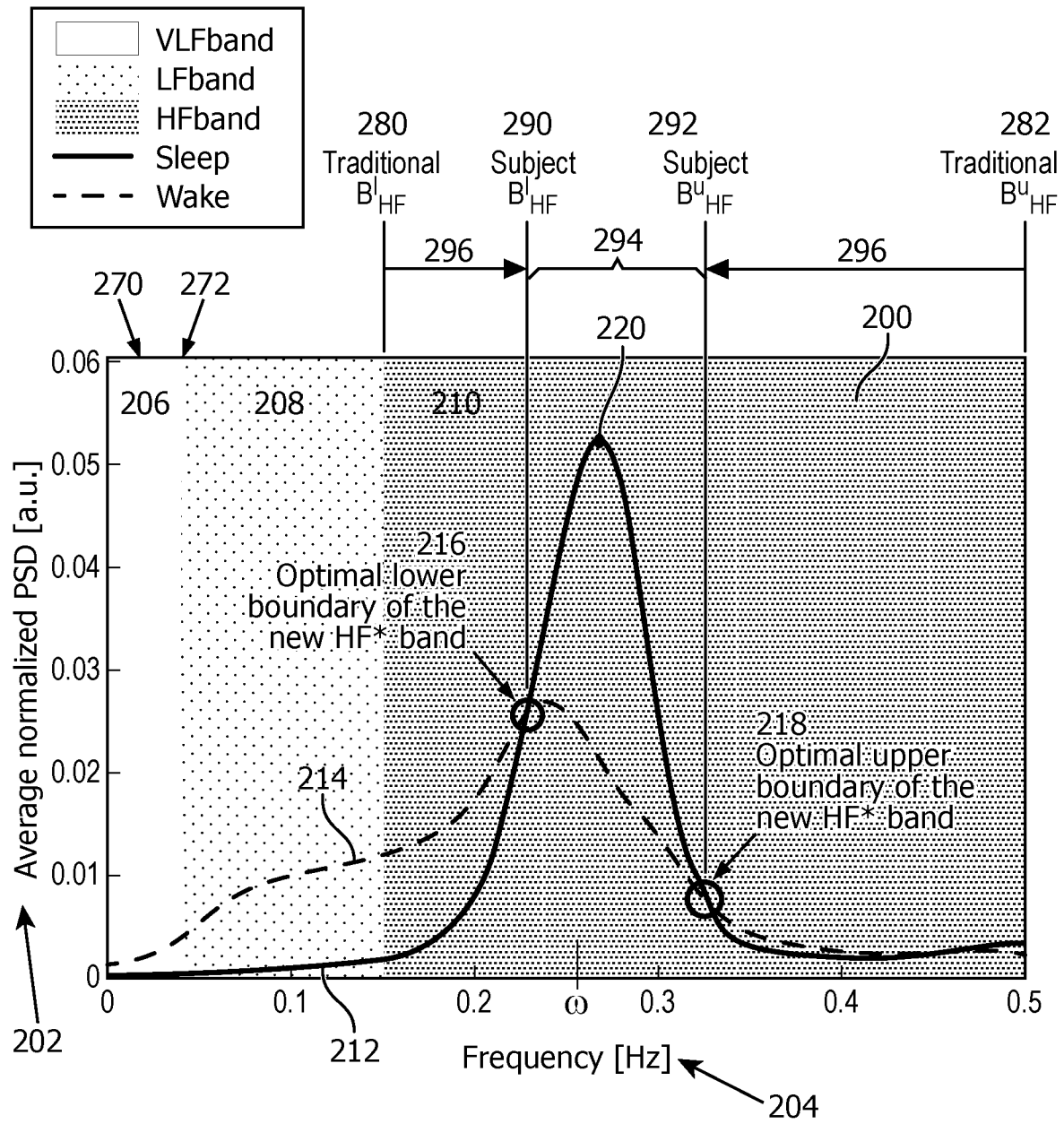


FIG. 2

3/4

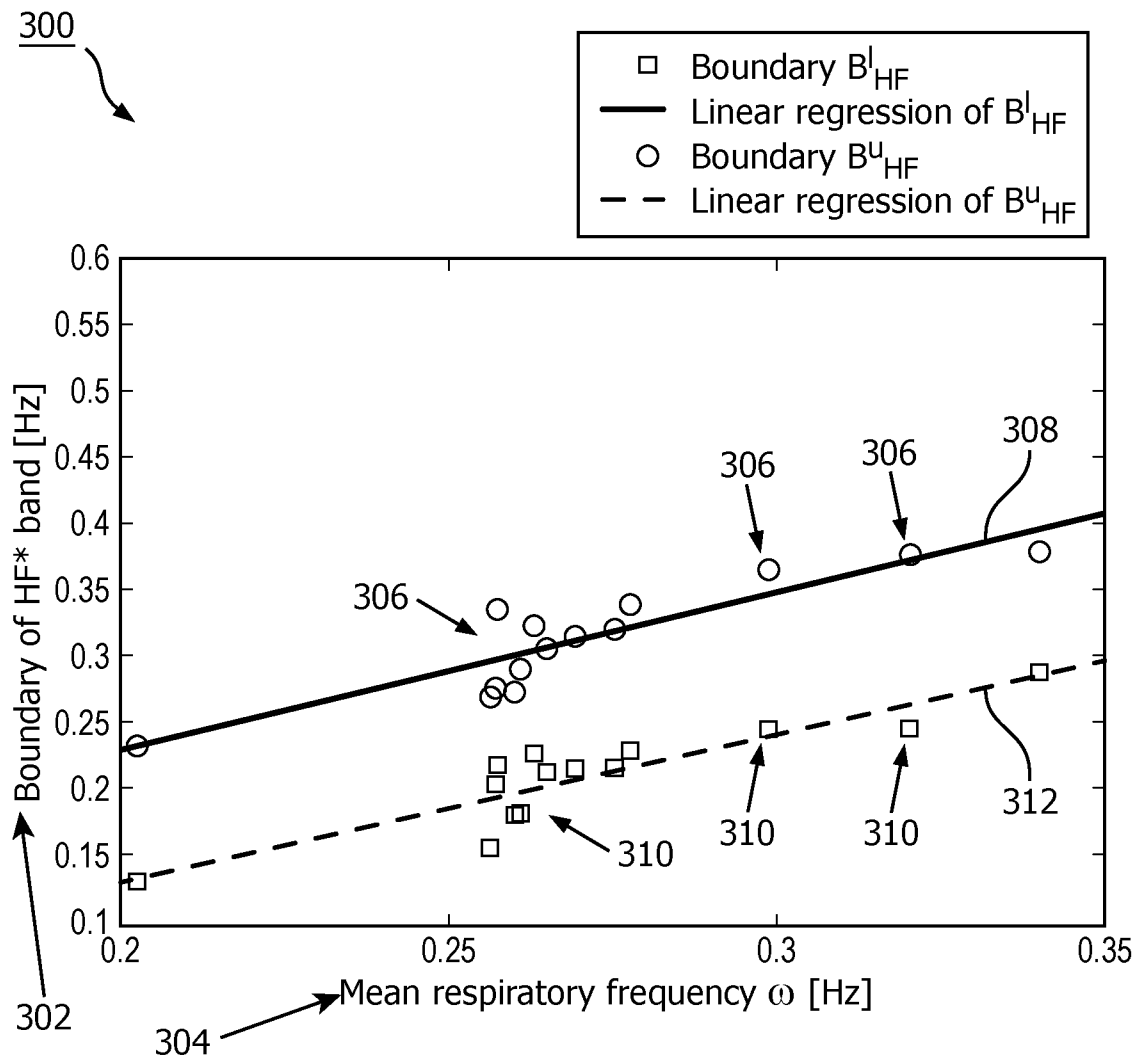


FIG. 3

4/4

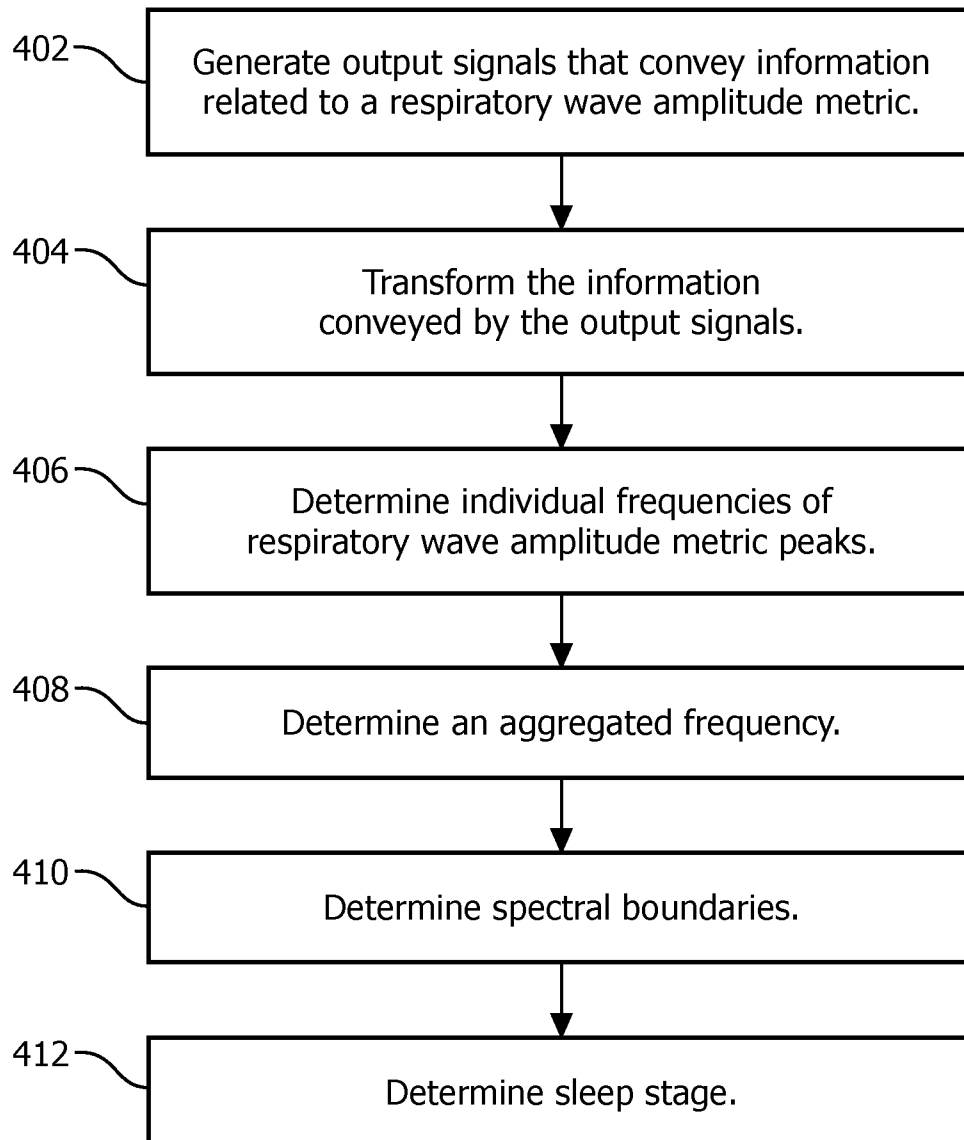
Method
400

FIG. 4

INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2015/059341

A. CLASSIFICATION OF SUBJECT MATTER INV. A61B5/00 ADD. A61B5/024 A61B5/08 A61B5/11 A61B5/113		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) A61B		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EP0-Internal, WPI Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 2 286 723 A1 (UNIV RAMOT [IL]) 23 February 2011 (2011-02-23) paragraphs [0220], [0333], [0334], [0346] - [0353]	1-15
X	----- XI LONG ET AL: "Spectral Boundary Adaptation on Heart Rate Variability for Sleep and Wake Classification", INTERNATIONAL JOURNAL OF ARTIFICIAL INTELLIGENCE TOOLS, vol. 23, no. 03, 28 May 2014 (2014-05-28), page 1460002, XP055250219, SG ISSN: 0218-2130, DOI: 10.1142/S0218213014600021 sections 2.1 to 2.6 ----- <div style="text-align: right;">-/-</div>	1-15
<div style="display: flex; justify-content: space-between;"> <input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex. </div>		
* Special categories of cited documents :		
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search <div style="text-align: center; font-size: 1.2em;">16 February 2016</div>		Date of mailing of the international search report <div style="text-align: center; font-size: 1.2em;">29/02/2016</div>
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer <div style="text-align: center; font-size: 1.2em;">Hemb, Björn</div>

INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2015/059341

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X,P	<p>LONG XI ET AL: "Improving sleep/wake detection via boundary adaptation for respiratory spectral features", 2015 37TH ANNUAL INTERNATIONAL CONFERENCE OF THE IEEE ENGINEERING IN MEDICINE AND BIOLOGY SOCIETY (EMBC), IEEE, 25 August 2015 (2015-08-25), pages 374-377, XP032810206, DOI: 10.1109/EMBC.2015.7318377 [retrieved on 2015-11-04] sections 2A to 2E figure 1</p> <p style="text-align: center;">-----</p>	1-15

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
PCT/IB2015/059341

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 2286723	A1	23-02-2011	AT 496577 T 15-02-2011
			AU 2003263571 A1 08-04-2004
			CA 2499547 A1 01-04-2004
			EP 1562472 A1 17-08-2005
			EP 2286723 A1 23-02-2011
			US 2006235315 A1 19-10-2006
			WO 2004026133 A2 01-04-2004
