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(54) **ELECTROSTIMULATION SYSTEM FOR THE TREATMENT OF SLEEP APNOEA**

ELEKTROSTIMULATIONSSYSTEM ZUR BEHANDLUNG VON SCHLAFAPNOE

SYSTÈME D'ÉLECTROSTIMULATION POUR LE TRAITEMENT DE L'APNÉE DU SOMMEIL

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Description

Field of the invention

[0001] The present invention relates to muscular stimulation, specifically stimulation using adjustable electrical signals in order to treat episodes of insufficient breathing produced during Obstructive Sleep Apnea (OSA) Syndrome.

Background of the invention

[0002] Various epidemiological studies have proven that sleep apnea is highly prevalent amongst sleep related conditions and affects between 4-6 % of males and 2-4% of females in the general middle-aged adult population.

[0003] The main problems related to sleep apnea include significant reduction in quality of life, arterial hypertension and the development of cardiovascular and cerebrovascular diseases.

[0004] Obstructive sleep apnea is therefore one of the most widespread and worrying sleep conditions.

[0005] The most extensive methods known to date for treating obstructive sleep apnea include:

US005245995A: A device named CPAP, consisting of a turbine that transmits a predetermined pressure via a nasal mask, adapted to the subjects face and fixed with a harness, with which the circuit is closed.

[0006] The device is effective in terms of continuously treating the illness, however presents various limitations:

- Nasal congestion and/or obstruction. Nasal congestion is the most common and treatment depends on the cause behind it. It usually arises as a result of edema and inflammation of the nasal mucus.
- Irritated skin. This occurs in the area that comes into contact with the CPAP mask.
- A dry throat. This is caused by water being lost from the soft palate, as a result of snoring and apnea and the air flow from the device itself. In order to be corrected, it is necessary to install a humidifier/heater connected to the CPAP.
- Noise. In addition to noise produced by the CPAP, which goes unnoticed after the first few weeks, patients and their companions complained of tonality changes between inhalation and exhalation produced in the mask.
- Cold. This is particularly significant in cold areas and houses without heating, where in winter, air from the CPAP may enter the VAS at 15 °C or lower (the temperature recommended for sleep being around 19 °C).

[0007] Surgical techniques: these involve generally aggressive surgical techniques, which are reserved in

many cases for when the CPAP does not work or for patients who reject it before looking into it further. It requires very precise prior anatomical exploration and must be carried out by experienced surgeons. The ill person must be informed of the chances of success exhaustively and may require more than one intervention.

[0008] Electric stimulation: The first research on electric stimulation treatments for obstructive sleep apnea was carried out in the 1980s, when Dr. Miki's team in Japan carried out various experiments on this usage.

[0009] Since then, several pieces of apparatus have been produced, which have attempted to apply principles related to electric stimulation in order to prevent the collapse of the upper airways by means of contracting the muscles involved. Some are internal, whilst others are external.

[0010] One device implanted into the subject internally is:

US006345202B2, which proposes a method for treating apnea using microelectrodes implanted into strategic places inside a patient. They are controlled wirelessly by an external transmitter, in order to stimulate the muscular and nerve tissue in a constructive way, which helps to open the airways blocked.

[0011] External Devices include:

US 2008/0021506A1, which uses electrodes fitted to the surface, to prevent snoring and sleep apnea. US 2006/0145878A1, which proposes a method and system for fighting apnea, wherein the subject is woken up.

WO 2006/054359, which proposes electrical stimulation by means of applying electrodes in the jaw portion of a patient, in order to prevent sleep apnea. It therefore applies electrical signals repeatedly during the night, without considering the state of obstruction in the upper airways.

[0012] One limitation common to all the above proposals is the lack of regulation. They do not facilitate automatic adjustment, in order to adapt to the morphological characteristics and the intensity level of apnea episodes of each patient.

[0013] The same goes for the CPAP device, wherein prior adjustment is required, thus increasing the costs of installing the apparatus for each patient considerably. It is currently substituted by an automatic graduation CPAP system known as the APAP (Roldán N et al. 2008).

[0014] Devices that detect sleep apnea by means of fragmenting and subsequent analyzing an acoustic signal generated by patient's breathing are known from WO02/13677, US2011/0288431 and US2011/0137195. Sleep apnea detection by means of counting inhalation and exhalation is disclosed by US2010048985. The apparatus described below compensates for the deficiencies identified in the state of the art and constitutes a

significant technological advancement, since it facilitates self-regulating detection and treatment, in order to adapt to a patient's morphology. It uses the minimal electric shock required in terms of both frequency and intensity in the muscles involved. Furthermore, it facilitates follow-up and control of the evolution of the condition.

Description of the invention

[0015] The invention is defined by claim 1 and is particularly suited for the continuous treatment of sleep and respiratory conditions, by means of self-regulating electrical stimuli, in order to adapt to a patient's type set.

[0016] The invention makes it possible to process and store information relative to apnea episodes and electric stimulation, in addition to making it possible to adjust the electrical stimuli in order to adapt to the evolution of the condition or problem. In turn, it makes it possible to follow-up over time in order to assess a patient's progress.

[0017] Muscular stimulation is carried out cutaneously on the patient's neck, without secondary side effects or alterations to sleep settings. It therefore acts on the muscles involved in the collapse of respiratory airways successfully.

[0018] The effect of stimulating muscles additionally serves to train and strengthen those responsible for keeping the upper airways open (the genioglossus).

[0019] The stimulation system is preferably built in such a way that it is portable and may be easily transported. Patients may therefore take it with them and use it both in their usual home environment and in other situations (for example during travel).

[0020] The system includes a neck piece and a central device. The neck piece in turn comprises an electronic support, responsible for collecting the signal from the upper airways and transmitting electrical impulses, in addition to an elastic band, designed in an anatomical format, which is molded to the patient's neck.

[0021] The device has an apnea detection module based on processing the acoustic signal produced by the upper airways. It is therefore possible to adjust detection criteria to the optimal breathing thresholds of each patient, within a continuous learning process. The signal produced in the respiratory airways is processed to calculate the strength thereof. In turn, a number of thresholds are defined, between which the strength must be situated. In the event of the strength being below the threshold linked to breathing, the duration thereof is thereby counted. If this duration exceeds another time threshold, it is concluded that there is a period of apnea and this result is communicated to the actuation module.

[0022] The actuation module controls the electric stimulator and facilitates the self-regulation of the intensity and frequency of electrical stimulus, according to the characteristics of the episode and patient. It is ensured that the stimulus is enough for each patient's layer of adipose tissue and for the severity of the collapse in the airways. A stimulus is therefore generated at a specific

intensity and frequency.

[0023] The device may store information relating to apnea episodes and the electrical stimuli in a memory (external, if necessary), thus creating a time sequence that facilitates the automatic assessment of the evolution of sleep apnea and modification of the electric stimulation program, in order to adjust the intensity and/or frequency of the electrical stimulus to be applied.

[0024] The device may include a communication system, by means of a cable or wireless, which makes it possible to connect to an external device. Said external device may in turn be in charge of other functions, for example, producing a report on the evolution of the apnea episodes.

Brief description of the drawings

[0025]

Figure 1. is a schematic block diagram of an electric stimulation system used to treat OSA, according to the present invention. The main components are shown.

Figure 2. (a) is an example of a neck piece (10) fitted around a patient's neck.

Figure 2. (b) represents the components of the neck piece (10) in detail.

Figure 2. (c) is a detailed view of the means used to fasten the neck piece (10).

Figure 3. shows how the elements of the neck piece (10) and central device (20) of the system are placed on a patient's body.

A neck piece (10) with free electrodes to be placed in the submental area of the muscles in order to facilitate electric stimulation and a microphone in contact with the neck area, in order to receive the sound of the air flow crossing the upper airways. Central device (20) with a module for processing acoustic signals, electric stimulation and the controller thereof, a battery and preferably an external memory. The sound signal is received from the microphone and processed and where necessary, the electrical signal is transmitted to the electrodes connected to the neck piece (10).

Figure 4. is a flow diagram of one embodiment.

Preferred embodiments

[0026] The design of the device makes it comfortable to wear for the whole night, without disturbing normal sleep cycles, with a microphone fitted in such a way that it comes into contact with the neck and two electrodes, which are placed in a submental position.

[0027] A small, central device (usually placed on the chest), with an apnea episode detection module based on analyzing the acoustic signal of the upper respiratory airways and an electric stimulation actuator, capable of producing a variable electrical signal, in terms of both

intensity and frequency. Furthermore, there is close communication between both modules, in such a way that controlled feedback is produced between the electrical signal applied to the muscles and the duration of the apnea episodes.

[0028] Below is a description of an exemplary embodiment of the invention, with reference to the figures.

[0029] According to Figure 1, the system for treating apnea comprises two modules: a neck piece (10) and a central device (20). It may also include a user interface (32) used to communicate with an external device (30) responsible for storing information of interest in a database (31) or for recovering data (history of apnea episodes) stored in the memory (26) of the central device (20) for subsequent follow-up and study of the evolution of the condition.

[0030] The neck piece (10) in turn comprises a microphone (11) adjusted to capture the sound of the upper airways. Furthermore, it preferably has two terminals (12) in order to connect the electrodes (18) that may be extracted and replaced in order to stimulate the muscles involved in the apnea.

[0031] The microphone (11) collects the acoustic signal of the upper respiratory airways generated by the patient in real time.

[0032] The electrodes (18) may be easily coupled to their adaptors (12) and are preferably flexible in order to adjust to various neck anatomies correctly.

[0033] The central device (20) incorporates a signal processor (23) for treating the acoustic signal coming from the microphone (11) and a method for detecting apnea episodes. The central device (20) receives the pre-filtered analogue signal, converts it into a digital signal and may furthermore include a filter (22) to be used on the digital signal, in order to reduce harmful noises.

[0034] The processing unit (23) recognizes an apnea episode when the strength of the acoustic signal falls below a certain strength threshold during a minimum period of time. It then decides to send a signal to the electric stimulator (25) controller. In turn, the controller (25) recovers the optimal frequency and intensity values previously adapted for the current episode, in such a way that it always administers the smallest shock necessary in order to stop an apnea episode. It then activates the electric stimulator (29), which in turn generates the electric shock to the electrodes (18). At the end of each stimulation, the processing unit (23) checks whether or not the apnea episode has been successfully corrected. Moreover, they are stored in a memory (26), which is external if necessary, providing a record of said episodes.

[0035] The patient may connect the apparatus to an external device (30) on a daily basis using a cable (15) (for example of the USB variety) and a specific user interface program (32) to download data from the external data base (31) and collect the data in order to create sleep apnea treatment follow-up reports, amongst other tasks.

[0036] As can be observed in Figure 2, the ergonomic

support (14) of the neck piece (10) may be made of a semi-rigid material, which may be shaped and adapted to the neck (for example, aluminum). This would contain the electronic portion. The inside may be covered with polyurethane foam. The microphone (11) is also incorporated into the internal portion of the support, preferably at one side and may be protected with a sponge against the air and reduce unwanted external noises.

[0037] The electronic support (10) may contain two terminals (12), wherein the replaceable electrodes (18) are connected.

[0038] The electronic support of the neck piece (10) may be coated with a textile material (17), which is preferably hypoallergenic, resistant to dust mites, porous, ergonomic and breathable, as well as being easy to wash. In the external portion of the electronic support (10), a Velcro coating (13) may form an easy fasten system.

[0039] In turn, the electronic support (10) may be shaped to the patient's neck by means of an elastic band with silicone straps (16) in order to provide a non-slip fit. The band is designed in an anatomical format that is shaped to the patient's neck. The electronic support (10) is joined to the band by means of a Velcro easy fasten system or pocket. In the external portion of the band, a Velcro coating (13) may serve as a closure.

[0040] Figure 3 shows the elements that form the piece of equipment for treating apnea, namely the neck piece (10) and the electric stimulator module (20) on the patient's neck.

[0041] The neck piece (10) leaves the electrodes (18) free in order to place them in the submental area for electric stimulation alongside a microphone (11) in contact with the area of the neck for receiving the sound of the air flow crossing the upper airways.

[0042] The central device (20) is formed by a digital signal processor (23), an electric stimulator (29), a controller (25) for the same, an external memory (26) if required and a battery.

[0043] The sound signal coming from the microphone (11) reaches the processor (23). The electronic impulses are transmitted from the central device (20) to the electrodes (18) placed in the patient's submental region.

[0044] Subsequently, Figure 4 describes the intelligence of the system in the apnea episode detection and treatment method using variable electronic impulses issued by the processor (23). Upon the device (20) starting up, the processor (23) recovers the optimal thresholds and settings, if necessary, from the external memory (26), updating them in a volatile memory. The first time the device (20) is used, said thresholds and settings are defined at a default value, the same being adjusted to their optimal operational value with each use, according to the conditions and characteristics of the patient and their illness.

[0045] The processing unit (23) receives the voice signal in real time, divides it into blocks of a pre-established size and calculates the strength of each block. In turn, blocks of said signal are stored so that when a minimum

value is reached, the self-correlation thereof is calculated, provided that said blocks do not contain apnea episodes. The self-correlation function provides information on the period in which the signal was issued, thus making it possible to update the threshold for low strength blocks (T_{LSB}), which shall serve as one of the criteria for detecting apnea.

[0046] When the strength of the current block exceeds the snoring threshold (T_{Snor}) this is because the patient is snoring. It therefore updates the T_{Snor} threshold, adjusting it to the patient's snoring characteristics.

[0047] In contrast, if the strength falls below the T_{Snor} threshold and in turn, above the breathing threshold (T_{Brea}), this is because the patient is breathing. It therefore updates the T_{Brea} threshold, adjusting it to the patient's breathing patterns.

[0048] In the opposite case, if the strength falls below the T_{Brea} , it indicates that an apnea episode has begun. In order to be sure of this, it starts to count the number of low strength blocks (C_{LSB}), which indicates the time in which the patient has gone without breathing.

[0049] When this time C_{LSB} exceeds the maximum threshold of low strength blocks (T_{LSB}), it recognizes an apnea episode and therefore sends a signal to the controller (25), in order to activate the electric stimulator (29) using the current frequency and intensity values of the processor.

[0050] Once the electric stimulator (29) has been activated, it starts to count the time (C_{AE}) in which it is active.

[0051] The processor sends an order to the controller to stop the electrical stimulator in one of the following two cases: if the active electrode counter C_{AE} exceeds the maximum activation time threshold (T_{MAT}) or if breathing or snoring is detected.

[0052] At the end of each stimulation, the processor (23) checks whether or not the apnea episode has been corrected successfully. This is confirmed when the electrode activation time is lower than the maximum activation time. The counter of successful cases ($C_{Successful}$) is also increased and the unsuccessful cases counter ($C_{Unsuccessful}$) is set to zero. If it failed, the unsuccessful cases counter ($C_{Unsuccessful}$) is thus increased and the successful cases counter ($C_{Successful}$) is set to zero.

[0053] Finally, the C_{LBP} and C_{AE} variables are restarted and the apnea event is stored in the events record found in a memory (26).

[0054] The signal processing unit (23) then decides to reduce the frequency and intensity values once the number of successful cases has exceeded the successful cases threshold ($T_{Successful}$). In the event of the opposite, it increases the frequency and intensity values once the number of unsuccessful cases has exceeded the unsuccessful cases threshold ($T_{Unsuccessful}$), always administering the smallest shock required to stop an apnea episode.

[0055] Before turning the device off, the processor (23) stores the breathing thresholds and electrical impulse settings adjusted to the patient in the memory (26), in

order to provide optimal operation the following night.

[0056] The processor (23) therefore adapts both the breathing thresholds (T_{Snor} and T_{Brea}), depending on the strength of the sound captured and the period of the signal (T_{LSB}), depending on the patient's breathing rate, i.e. the number of times the patient breathes or snores per minute. Said thresholds therefore vary depending on a patient's behavior, the precision of the detection of an apnea episode improving as they are adapted.

Abbreviations:

[0057]

T_{LSB}	low strength blocks threshold
T_{Snor}	snoring threshold
T_{Brea}	breathing threshold
T_{Frec}	electric stimulator frequency threshold
T_{Intens}	electric stimulator intensity threshold
N_{LSB}	number of low strength blocks
C_{AE}	active electrode counter
T_{MAT}	minimum activation time threshold
$C_{Successful}$	successful cases counter
F_{Opt}	optimal frequency values
I_{Opt}	optimal intensity values
$C_{Unsuccessful}$	unsuccessful cases counter

Numerical references

[0058]

10	Neck piece.
11	Microphone.
12	Electrode terminals.
13	Velcro.
14	Ergonomic support.
15	Cable.
16	Silicone strips.
17	Textile material.
18	Electrodes.
19	Pocket for storing electronic support 10.
20	Central device.
22	Filter.
23	Processing unit.
25	Controller.
26	Central device 20 memory.
29	Electric stimulator.
30	External device.
31	External database.
32	User interface.
40	Load thresholds: F_{Opt} , I_{Opt} , T_{Snor} , T_{Brea} , T_{Frec} , T_{Intens} , T_{LSB} , T_{MAT} , $T_{Successful}$, $T_{Unsuccessful}$.
41	Real time sample of the signal.
42	Divide the signal into blocks.
43	Update T_{LSB} .
44	Calculate the strength of the block.
45	Check strength is lower than T_{Snor} .
46	Update T_{Snor} .

47 Check strength is lower than T_{Brea} .
 48 Increase C_{LSB} .
 49 Check C_{LSB} is greater than T_{LSB} .
 50 Electric stimulator activation.
 51 Increase C_{AE} .
 52 Check C_{AE} is greater than T_{MAT} .
 53 Stop the electric stimulator 29.
 54 Check the electric stimulator 29 is active.
 55 Update T_{Brea} .
 56 Check C_{AE} is lower than T_{MAT} .
 57 Update F_{Opt} , I_{Opt} of the electric stimulator 29.
 58 Increase $C_{Successful}$.
 59 Restart $C_{Unsuccessful}$ counter = 0.
 60 Restart $C_{LSB} = 0$, $C_{AE} = 0$.
 61 Increase $C_{Unsuccessful}$ counter.
 62 Store event records.
 63 Check $C_{Successful}$ is greater than $T_{Successful}$.
 64 Decrease frequency using T_{Frec} .
 65 Decrease intensity using T_{Intens} .
 66 Check $C_{Unsuccessful}$ is greater than $T_{Unsuccessful}$.
 67 Increase frequency using T_{Frec} .
 68 Increase intensity using T_{Intens} .
 69 Restart the $C_{Successful}$ counter = 0.

Claims

1. An electric stimulation system for treating sleep apnea in a patient, **characterized in that** it comprises:
 - treatment means (22, 23) designed to treat the acoustic signal generated by the patient's breathing and divide it into a plurality of blocks, each block corresponds to a fragment of the acoustic signal within a time interval, said treatment means (22, 23) are furthermore designed to measure the strength of a plurality of blocks of the acoustic signal;
 - control means (23, 25, 26) designed to count, in the plurality of blocks measured, the number of low strength blocks C_{LSB} when the strength of the block is less than a strength threshold linked to breathing, T_{Brea} , said control means (23, 25, 26) furthermore being designed to determine the existence of an apnea episode when the number of low strength blocks C_{LSB} exceeds a threshold on the number of low strength blocks T_{LSB} , said control means (23, 25, 26) are also designed to selectively control, if there is an apnea episode, the activation of a number of electric stimulation means (12, 18, 25, 29), designed to apply an electric signal to the submental muscles linked to the respiratory airways.
2. The system according to claim 1, **characterized in that** the blocks are generated by the treatment means (22, 23), after having sampled and filtered the acoustic signal.
3. The system according to claim 1 or 2, **characterized in that** the low strength block threshold T_{LSB} is estimated by the control means (23, 25, 26) in coordination with the treatment means (22, 23), with said threshold being linked to the number of blocks present in the acoustic signal generated by the patient's breathing in a breathing interval.
4. The system according to any of the previous claims, **characterized in that** the electric stimulation means (18, 25, 29, 12) start a temporary counter that measures the activation time, C_{AE} , in such a way that if it exceeds a maximum activation limit T_{MAT} , the control means (23, 25, 26) deactivate the electric stimulation means (18, 25, 29, 12) independently of the existence of the apnea episode.
5. The system according to claim 4, **characterised in that** if the control means (23, 25, 26) still determine the existence of the apnea episode, they increase an unsuccessful episode count ($C_{Unsuccessful}$).
6. The system according to claim 4, **characterised in that** if the control means (23, 25, 26) still determine the non-existence of an apnea episode, they increase a successful episodes count, $C_{Successful}$.
7. The system according to any one of the claims 4 to 6, **characterised in that** the control means (23, 25, 26) adjust the value of the frequency and/or intensity of the electrical signal generated by the electric stimulation means (29, 12), in such a way that:
 - the intensity and/or frequency is decreased within a range depending on the value of the $C_{Successful}$ count and
 - the intensity and/or frequency is increased within a range depending on the value of the $C_{Unsuccessful}$ count.
8. The system according to any one of the previous claims, **characterised in that** the treatment means (22, 23) are configured to receive the signal from a microphone (11) designed to capture the acoustic signal generated by the patient during breathing.
9. The system according to any one of the previous claims, **characterised in that** it furthermore comprises storage means (26, 31), designed to store at least one of the following values:
 - T_{Brea} , T_{LSB} , C_{AE} , T_{MAT} , $C_{Unsuccessful}$, $C_{Successful}$, C_{LSB} , maximum and minimum frequency range for the electrical signal applied, maximum and minimum intensity range for the electrical signal applied.

Patentansprüche

1. Elektrostimulationssystem zur Behandlung von Schlafapnoe in einem Patienten, **dadurch gekennzeichnet, dass** es Folgendes umfasst:

- Bearbeitungsmittel (22, 23), welche dazu gestaltet sind, das durch die Atmung eines Patienten erzeugte Schallsignal zu bearbeiten und es in eine Vielzahl von Blöcken zu teilen, wobei jeder Block einem Fragment des Schallsignals innerhalb eines Zeitintervalls entspricht und die genannten Bearbeitungsmittel (22, 23) zusätzlich dazu gestaltet sind, die Stärke einer Vielzahl von Blöcken des Schallsignals zu messen;

- Steuermittel (23, 25, 26), welche dazu gestaltet sind, in der Vielzahl von gemessenen Blöcken, die Anzahl von Blöcken niedriger Stärke C_{LSB} zu zählen, wenn die Stärke des Blocks kleiner als eine mit der Atmung verbundene Schwelle, T_{Brea} , ist, wobei die genannten Steuermittel (23, 25, 26) zusätzlich dazu gestaltet sind, die Anwesenheit einer Apnoeepisode festzustellen, wenn die Anzahl von Blöcken niedriger Stärke C_{LSB} eine Schwelle der Anzahl von Blöcken niedriger Stärke T_{LSB} überschreitet,

wobei die genannten Steuermittel (23, 25, 26) auch dazu gestaltet sind, wenn eine Apnoeepisode vorliegt, die Aktivierung einer Anzahl von Elektrostimulationsmitteln (12, 18, 25, 29) selektiv zu steuern, welche dazu gestaltet sind, ein elektrisches Signal an die submentale Muskulatur, welche mit den Atemwegen verbunden ist, anzulegen.

2. System nach Anspruch 1, **dadurch gekennzeichnet, dass** die Blöcke durch die Bearbeitungsmittel (22, 23) erzeugt werden, nachdem das Schallsignal bemustert und filtriert worden ist.
3. System nach Anspruch 1 oder 2, **dadurch gekennzeichnet, dass** die Schwelle der Blöcke niedriger Stärke T_{LSB} durch die Steuermittel (23, 25, 26) in Koordination mit den Bearbeitungsmitteln (22, 23) geschätzt wird, wobei die genannte Schwelle mit der Anzahl von Blöcken, welche in dem durch die Atmung des Patienten in einem Atmungsintervall erzeugten Schallsignal vorliegen, verbunden ist.
4. System nach einem der vorhergehenden Ansprüche, **dadurch gekennzeichnet, dass** die Elektrostimulationssysteme (18, 25, 29, 12) einen temporären Zähler starten, welcher die Aktivierungszeit, C_{AE} , derart misst, dass wenn sie eine maximale Aktivierungsgrenze T_{MAT} überschreitet, die Steuermittel (23, 25, 26) die Elektrostimulationssysteme (18, 25, 29, 12) unabhängig von dem Vorliegen einer Ap-

noeepisode deaktivieren.

5. System nach Anspruch 4, **dadurch gekennzeichnet, dass** wenn die Steuermittel (23, 25, 26) noch das Vorliegen der Apnoeepisode feststellen, sie eine erfolglose Episoden-Zählung ($C_{Unsuccessful}$) erhöhen.
6. System nach Anspruch 4, **dadurch gekennzeichnet, dass** wenn die Steuermittel (23, 25, 26) noch das Nichtvorliegen einer Apnoeepisode feststellen, sie einer erfolgreichen Episoden-Zählung, $C_{Successful}$, erhöhen.
7. System nach einem der Ansprüche 4 bis 6, **dadurch gekennzeichnet, dass** die Steuermittel (23, 25, 26) den Wert der Frequenz und/oder Stärke des elektrischen, durch die Elektrostimulationssysteme (29, 12) erzeugten Signals derart einstellen, dass:
- die Stärke und/oder Frequenz innerhalb eines Bereichs in Abhängigkeit des Wertes der $C_{Successful}$ -Zählung verringert wird, und
 - die Stärke und/oder Frequenz innerhalb eines Bereichs in Abhängigkeit des Wertes der $C_{Unsuccessful}$ -Zählung, erhöht wird.
8. System nach einem der vorhergehenden Ansprüche, **dadurch gekennzeichnet, dass** die Bearbeitungsmittel (22, 23) dazu ausgebildet sind, das Signal aus einem Mikrophon (11), das dazu gestaltet ist, das während der Atmung durch den Patienten erzeugten Schallsignal zu erfassen, zu empfangen.
9. System nach einem der vorhergehenden Ansprüche, **dadurch gekennzeichnet, dass** es zusätzlich Speichermittel (26, 31) umfasst, welche dazu gestaltet sind, mindestens einen der folgenden Werte zu speichern:
- T_{Brea} , T_{LSB} , C_{AE} , T_{MAT} , $C_{Unsuccessful}$, $C_{Successful}$, C_{LSB} , maximalen und minimalen Frequenzbereich für das angelegte elektrische Signal, maximale und minimale Stärkebereich für das angelegte elektrische Signal.

Revendications

1. Système de stimulation électrique pour traiter l'apnée du sommeil chez un patient, **caractérisé en ce qu'il comprend :**
- des moyens de traitement (22, 23) désignés pour traiter le signal acoustique généré par la respiration du patient et le diviser en une pluralité de blocs, chaque bloc correspond à un fragment du signal acoustique dans un intervalle de temps,

lesdits moyens de traitement (22, 23) sont également désignés pour mesurer la force d'une pluralité de blocs du signal acoustique ;

- des moyens de contrôle (23, 25, 26) désignés pour compter, dans la pluralité de blocs mesurés, le nombre de blocs de force faible C_{LSB} lorsque la force du bloc est inférieure à un seuil de force relié à la respiration, T_{Brea} , lesdits moyens de contrôle (23, 25, 26) étant également désignés pour déterminer l'existence d'un épisode d'apnée lorsque le nombre de blocs de force faible C_{LSB} dépasse un seuil sur le nombre de blocs de force faible T_{LSB} .

lesdits moyens de contrôle (23, 25, 26) sont également désignés pour contrôler sélectivement, s'il y a un épisode d'apnée, l'activation d'un nombre de moyens de stimulation électrique (12, 18, 25, 29) désignés pour appliquer un signal électrique aux muscles sous-mentonniers reliés aux voies respiratoires.

2. Système selon la revendication 1, **caractérisé en ce que** les blocs sont générés par les moyens de traitement (22, 23), après avoir échantillonné et filtré le signal acoustique.
3. Système selon la revendication 1 ou 2, **caractérisé en ce que** le seuil de blocs de force faible T_{LSB} est estimé par les moyens de contrôle (23, 25, 26) en coordination avec les moyens de traitement (22, 23), ledit seuil étant relié au nombre de blocs présents dans le signal acoustique généré par la respiration du patient dans un intervalle de respiration.
4. Système selon l'une quelconque des revendications précédentes, **caractérisé en ce que** les moyens de stimulation électrique (18, 25, 29, 12) commencent un comptage temporaire qui mesure le temps d'activation, C_{AE} , de manière que s'il dépasse une limite d'activation maximale T_{MAT} , les moyens de contrôle (23, 25, 26) désactivent les moyens de stimulation électrique (18, 25, 29, 12) indépendamment de l'existence de l'épisode d'apnée.
5. Système selon la revendication 4, **caractérisé en ce que** si les moyens de contrôle (23, 25, 26) continuent à déterminer l'existence de l'épisode d'apnée, ils augmentent un comptage d'épisodes infructueux ($C_{Unsuccessful}$).
6. Système selon la revendication 4, **caractérisé en ce que** si les moyens de contrôle (23, 25, 26) continuent à déterminer la non-existence d'un épisode d'apnée, ils augmentent un comptage d'épisodes fructueux ($C_{Successful}$).
7. Système selon l'une quelconque des revendications

4 à 6, **caractérisé en ce que** les moyens de contrôle (23, 25, 26) ajustent la valeur de la fréquence et/ou l'intensité du signal électrique généré par les moyens de stimulation électrique (29, 12), de manière que :

- l'intensité et/ou la fréquence est réduite dans une plage dépendant de la valeur du comptage $C_{Successful}$ et
- l'intensité et/ou la fréquence est augmentée dans une plage dépendant de la valeur du comptage $C_{Unsuccessful}$.

8. Système selon l'une quelconque des revendications précédentes, **caractérisé en ce que** les moyens de traitement (22, 23) sont configurés pour recevoir le signal depuis un microphone (11) désigné pour saisir le signal acoustique généré par le patient durant sa respiration.
9. Système selon l'une quelconque des revendications précédentes, **caractérisé en ce qu'il** comprend en outre des moyens de stockage (26, 31), désignés pour stocker au moins un des valeurs suivantes :

T_{Brea} , T_{LSB} , C_{AE} , T_{MAT} , $C_{Unsuccessful}$, $C_{Successful}$, C_{LSB} , plage de fréquence maximale et minimale pour le signal électrique appliqué, plage d'intensité maximale et minimale pour le signal électrique appliqué.

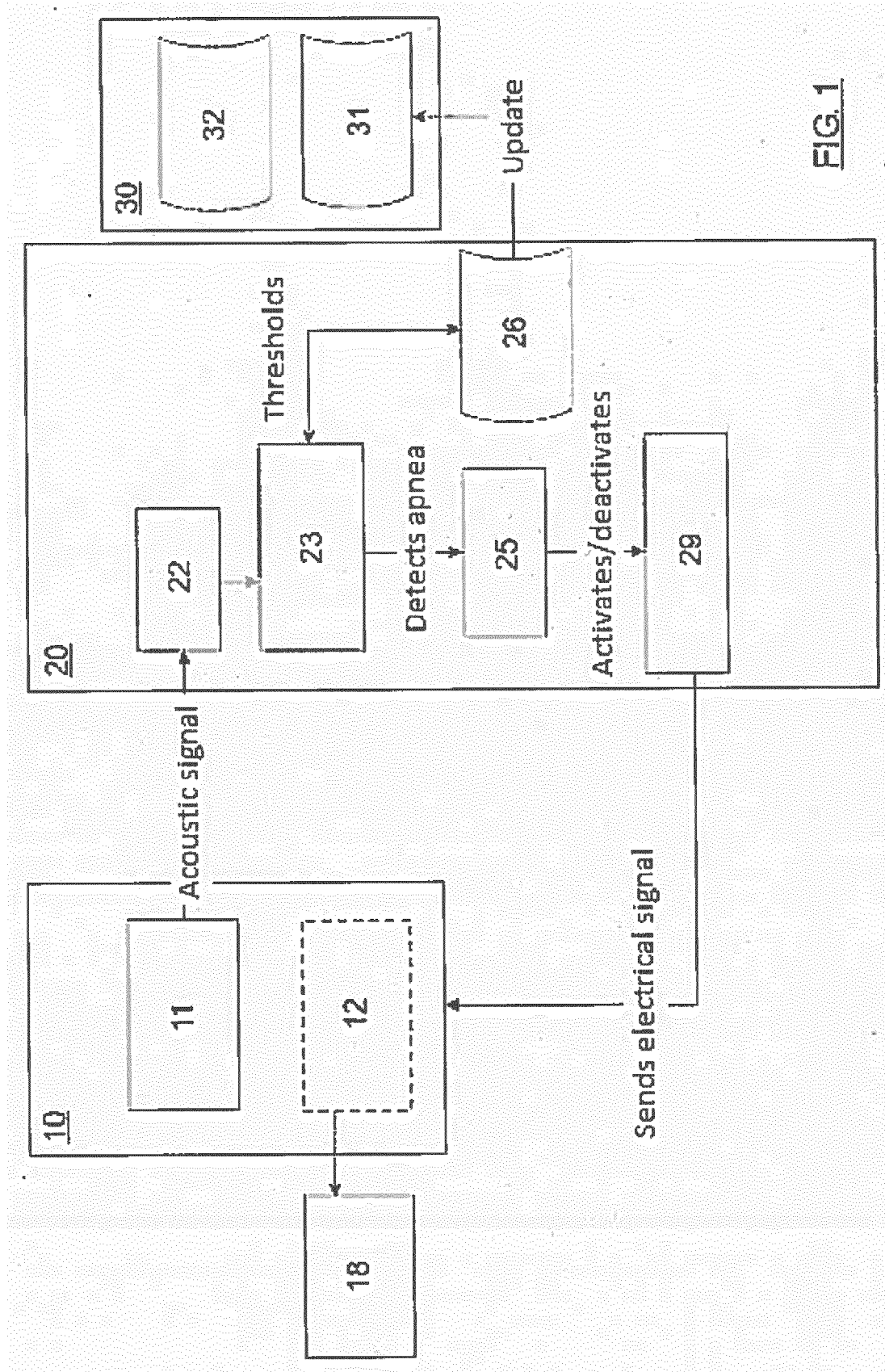


FIG. 1

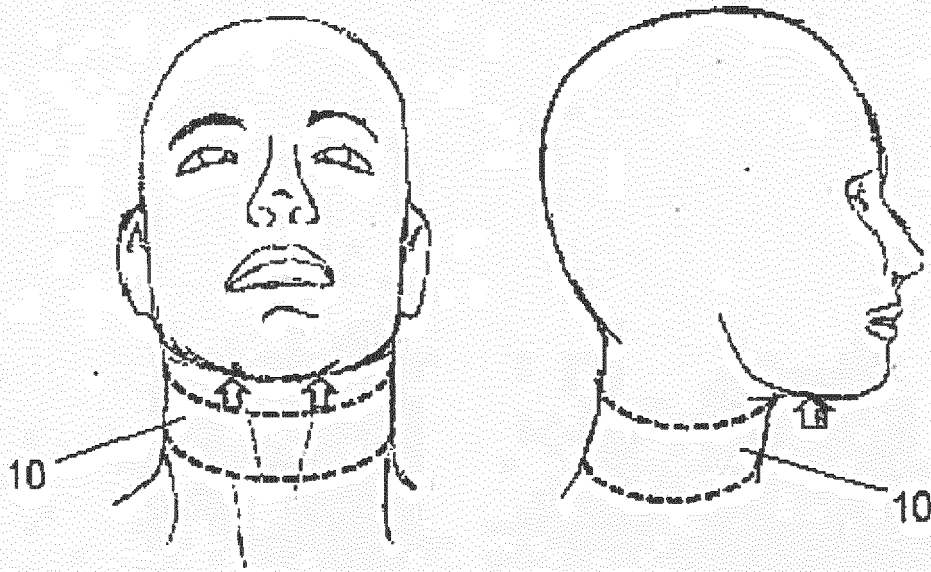


FIG. 2a

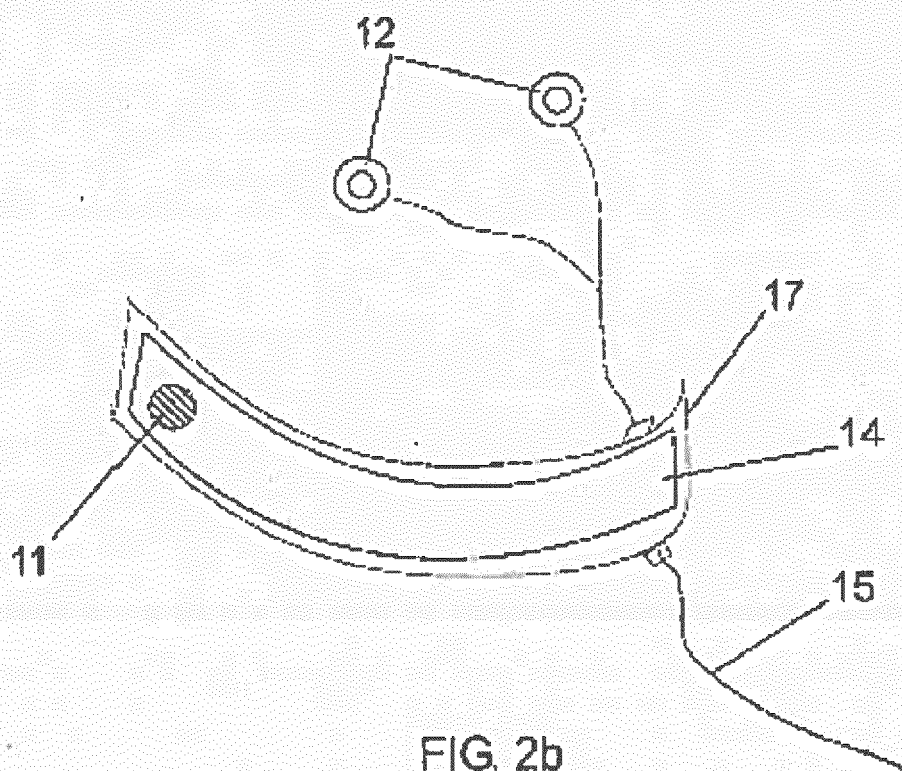


FIG. 2b

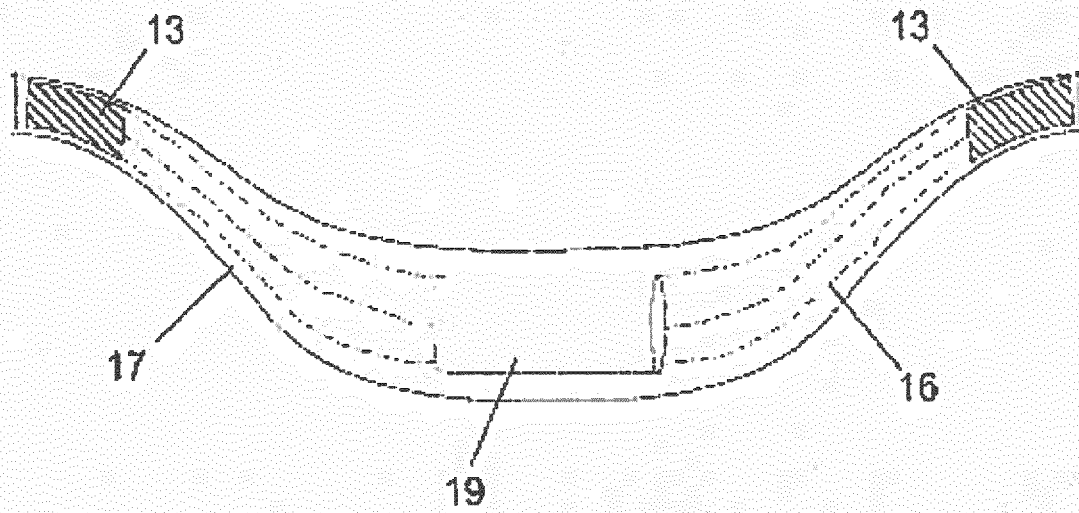


FIG. 2c

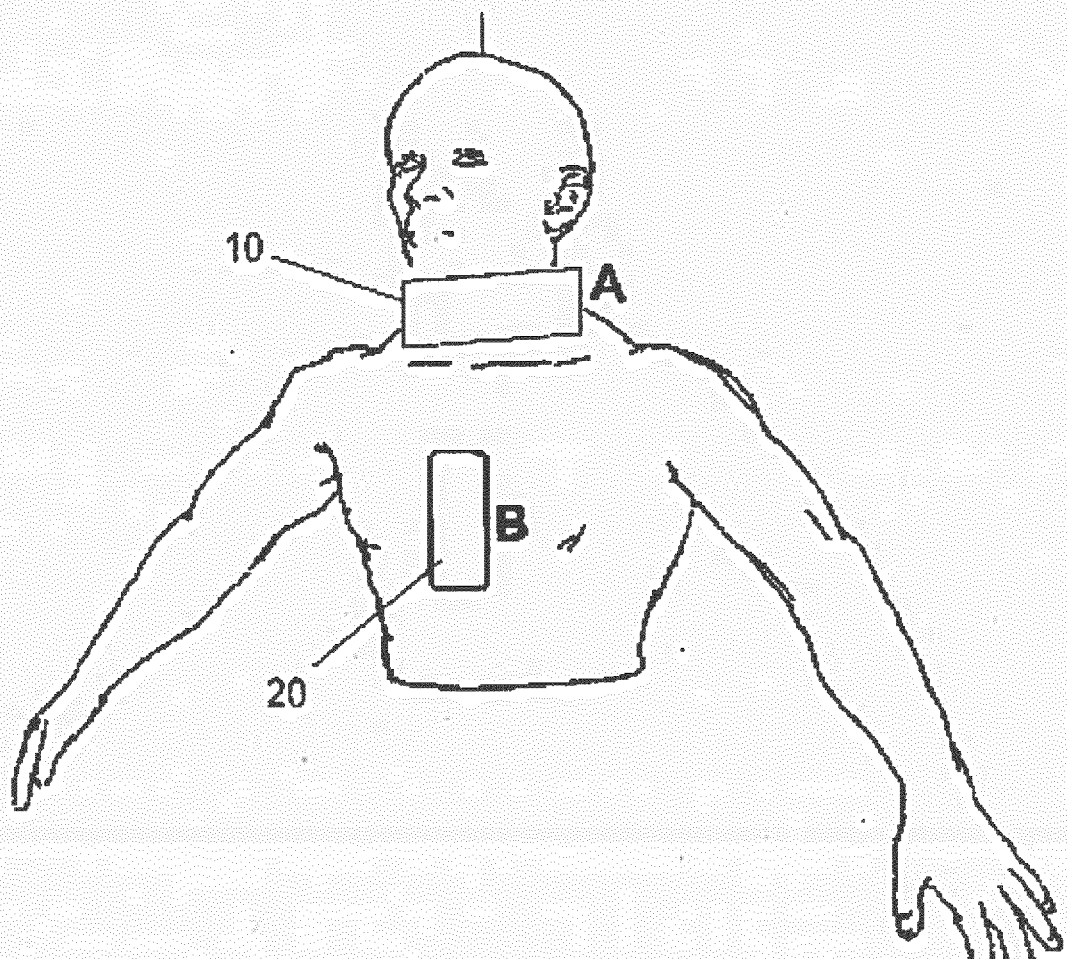


FIG. 3

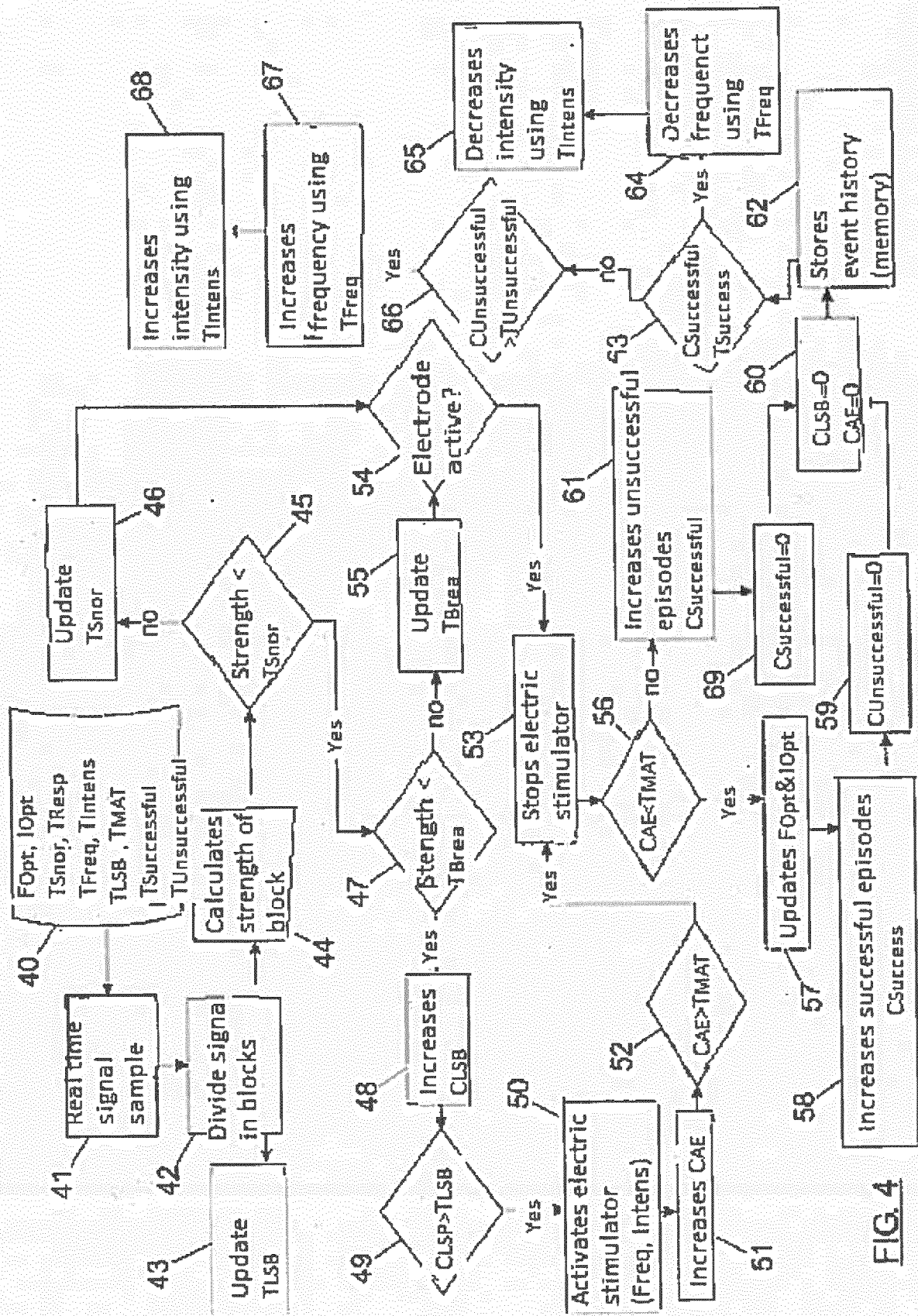


FIG. 4

REFERENCES CITED IN THE DESCRIPTION

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专利名称(译)	电刺激系统用于治疗睡眠呼吸暂停		
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其他公开文献	EP2810599A1 EP2810599A4		
外部链接	Espacenet		

摘要(译)

使用咽部 - 喉部肌肉上的外部致动器治疗阻塞性睡眠呼吸暂停 (OSA) 综合征的电刺激方法和系统。使用该致动器, 所涉及的肌肉接受电刺激, 目的是扩大肌肉开口并获得足够的气流以防止空气不足。通过分析上呼吸道的声音模式, 所述刺激仅在检测到呼吸暂停发作的情况下起作用。检测和治疗都是自我调节的, 以适应患者的形态和病情或问题的演变。

