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(54) **OBSTRUCTIVE SLEEP APNEA TREATMENT DEVICES**

VORRICHTUNGEN ZUR BEHANDLUNG OBSTRUKTIVER SCHLAFAPNOE

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Description

Field of the Invention

[0001] The inventions described herein relate to devices, systems and associated methods for treating sleep disordered breathing. More particularly, the inventions described herein relate to devices, systems and methods for treating obstructive sleep apnea.

Background of the Invention

[0002] Obstructive sleep apnea (OSA) is highly prevalent, affecting one in five adults in the United States. One in fifteen adults has moderate to severe OSA requiring treatment. Untreated OSA results in reduced quality of life measures and increased risk of disease including hypertension,

stroke, heart disease, etc.

[0003] Continuous positive airway pressure (CPAP) is a standard treatment for OSA. While CPAP is non-invasive and highly effective, it is not well tolerated by patients. Patient compliance for CPAP is often reported to be between 40% and 60%.

[0004] Surgical treatment options for OSA are available too. However, they tend to be highly invasive (result in structural changes), irreversible, and have poor and/or inconsistent efficacy. Even the more effective surgical procedures are undesirable because they usually require multiple invasive and irreversible operations, they may alter a patient's appearance (e.g., maxillo-mandibular advancement), and/or they may be socially stigmatic (e.g., tracheostomy). US 5 549 655 A discloses an apparatus for synchronized treatment of obstructive sleep apnea wherein stimulation is delayed when the patient is awake and stimulation is disabled if an arousal of the patient is indicated. During the delayed period the stimulation may not be completely suppressed but reduced to a level below the level of physiological response perceptible to the patient.

[0005] US Patent No. 4,830,008 to Meer proposes hypoglossal nerve stimulation as an alternative treatment for OSA. An example of an implanted hypoglossal nerve stimulator for OSA treatment is the Inspire™ technology developed by Medtronic, Inc. (Fridley, MN). The Inspire device is not FDA approved and is not for commercial sale. The Inspire device includes an implanted neurostimulator, an implanted nerve cuff electrode connected to the neurostimulator by a lead, and an implanted intrathoracic pressure sensor for respiratory feedback and timing of stimulation delivery. The Inspire device was shown to be efficacious (approximately 75% response rate as defined by a 50% or more reduction in RDI (Respiratory Disturbance Index) and a post RDI of ≤ 20) in an eight patient human clinical study, the results of which were published by Schwartz et al. and Eisele et al. How-

ever, both authors reported that only three of eight patients remained free from device malfunction, thus demonstrating the need for improvements.

Summary of the Invention

[0006] To address this and other unmet needs, the present invention provides, in exemplary non-limiting embodiments, devices, systems and methods for nerve stimulation for OSA therapy as described in the following detailed description. The invention is defined by the claims.

Brief Description of the Drawings

[0007] It is to be understood that both the foregoing summary and the following detailed description are exemplary. Together with the following detailed description, the drawings illustrate exemplary embodiments and serve to explain certain principles. In the drawings;

Figure 1 is a schematic illustration of a system according to an embodiment of the present invention, including internal (chronically implanted) and external components;

Figure 2 is a perspective view of a stimulation lead for use in the system shown in Figure 1, including a detailed view of the distal end of the stimulation lead; Figure 3A is a detailed perspective view of the cuff of the stimulation lead shown in Figure 2;

Figure 3B is a lateral cross-sectional view of the cuff shown in Figures 2 and 3A;

Figure 4A is a perspective view of a bifurcated respiration sensing lead which may be used in the system shown in Figure 1;

Figure 4A1 is a detailed perspective view of the proximal connector assembly of the respiration sensing lead shown in Figure 1;

Figure 4A2 is a detailed perspective view of the bifurcation section of the respiration sensing lead shown in Figure 1;

Figure 4A3 is a detailed perspective view of the contra-lateral distal body portion of the respiration sensing lead shown in Figure 1;

Figure 4A4 is a detailed perspective view of the ipsilateral distal body portion in the respiration sensing lead shown in Figure 1;

Figure 4B is a detailed perspective view of an alternative embodiment of the respiration sensing lead which may be used in the system shown in Figure 1;

Figure 4B1 is a detailed perspective view of the proximal connector assembly of the respiration sensing lead shown in Figure 4B;

Figure 4B2 is a detailed perspective view of the distal body portion of the respiration sensing lead shown in Figure 4B;

Figure 4C illustrates the implanted system shown in Figure 1 with the respiration lead shown in Figure 4A;

Figure 4D illustrates the implanted system shown in Figure 1 with the respiration lead shown in Figure 4B; Figure 4E illustrates an alternative embodiment of the respiration sensing lead which may be used in the system shown in Figure 1;

Figure 4F illustrates an alternative embodiment of the respiration sensing lead containing a loop-back region. This lead may be used in the system shown in Figure 1;

Figure 4G illustrates an alternative embodiment of the respiration sensing lead which may be used in the system shown in Figure 1;

Figure 4H illustrates an alternative embodiment of the respiration sensing lead which may be used in the system shown in Figure 1;

Figure 5A shows front, side and top views of an implantable neurostimulator for use in the system shown in Figure 1;

Figure 5B is a schematic block diagram of electronic circuitry for use in the implantable neurostimulator shown in Figure 5A;

Figures 6A, 6B, and 6C illustrate a bio-impedance signal, the corresponding physiological events, and stimulation delivery algorithms for use in the system shown in Figure 1;

Figures 6D, 6E, 6F, 6G, and 6H illustrate various stimulation pulse configurations for the implantable neurostimulator shown in Figure 1, as may be used for therapy or sleep titration, for example;

Figure 6I(traces 1-8) shows various stimulation modes for the implantable neurostimulator shown in Figure 1, as may be used for therapy or sleep titration, for example;

Figure 6J shows a stimulation regimen called core hours for the implantable neurostimulator shown in Figure 1, as may be used as a therapy mode;

Figure 7A is a schematic illustration of the programmer system for use in the system shown in Figure 1; Figures 7B and 7C are schematic block diagrams of electronic circuitry for use in the programmer system for shown in Figure 7A;

Figure 8A is a schematic illustration of the therapy controller for use in the system shown in Figure 1;

Figure 8B is a schematic block diagram of electronic circuitry for use in the therapy controller shown in Figure 8A;

Figure 9 is a top view of a magnet for use in the system shown in Figure 1;

Figure 10 is a schematic illustration of an interface of the system shown in Figure 1 and polysomnographic (PSG) equipment as may be used in a sleep study for therapy titration or therapy assessment, for example;

Figure 11A is an anatomical illustration showing the incision sites and tunneling paths that may be used for implanting the internal components shown in Figure 1;

Figure 11B is a perspective view of a disassembled

tunneling tool for use in tunneling the leads of the system shown in Figure 1;

Figure 11C is a detailed perspective view of the assembled tunneling tool shown in Figure 11B, but with the cap removed to expose the distal connector for attaching to the lead carrier disposed on the proximal end of a lead;

Figure 12A shows a flowchart of an idealized therapy process and the sub-processes that may be involved;

Figures 12B, 12C, and 12D show detailed flowcharts of idealized therapy sub-processes shown in Figure 12A;

Figures 13(traces 1-8) illustrate various stimulation output modes of the implantable neurostimulator shown in Figure 1;

Figure 14 illustrates an example of the effect of stimulation on airflow; and

Figure 15 illustrates a sleep wand, for wireless communication with the neurostimulator during sleep.

Detailed Description of Exemplary Embodiments

[0008] The following detailed description should be read with reference to the drawings in which similar elements in different drawings are numbered the same. The drawings, which are not necessarily to scale, depict illustrative embodiments and are not intended to limit the scope of the invention.

Overall System

[0009] Figure 1 schematically illustrates a hypoglossal nerve stimulation (HGNS) system 100 comprising internal components 1000 and external components 2000. The HGNS system 100 treats obstructive sleep apnea (OSA) by increasing neuromuscular activity to the genioglossus muscle via stimulation of the hypoglossal nerve (HGN) synchronous with inspiration to mitigate upper airway collapse during sleep. Stimulation is generated by an implantable neurostimulator (INS) 1100, synchronized with inspiration as measured by the respiration sensing lead (RSL) 1200 using bio-impedance, and delivered to the hypoglossal nerve by a stimulation lead (STL) 1300. A programmer system 2100 and a therapy controller 2500 are wirelessly linked to the INS 1100. The programmer system 2100 includes a computer 2300, a programmer interface 2400, and a programmer head 2200. The programmer system 2100 is used by the physician to control and program the INS 1100 during surgery and therapy titration, and the therapy controller 2500 is used by the patient to control limited aspects of therapy delivery.

[0010] The implanted components 1000 of the HGNS system 100 include the INS 1100, STL 1300, and RSL 1200. The INS is designed to accommodate one STL1300 and one RSL 1200. One STL 1300 may be used for unilateral implantation and unilateral hypoglos-

sal nerve stimulation. Similarly, one RSL 1200 may be used for respiration detection. Alternative embodiments of the RSL 1200 are described below and may be substituted. Therefore, for purposes of illustration not limitation, the INS 1100 is shown with STL 1300 and a bifurcated RSL 1200.

[0011] The implanted components 1000 may be surgically implanted with the patient under general anesthesia. The INS 1100 may be implanted in a subcutaneous pocket inferior to the clavicle over the pectoralis fascia. The distal end of the STL 1300 (cuff 1350) may be implanted on the hypoglossal nerve or a branch of the hypoglossal nerve in the submandibular region, and the proximal end of the STL 1300 may be tunneled under the skin to the INS 1100. The RSL 1200 may be tunneled under the skin from the INS 1100 to the rib cage and placed on the costal margin. The INS 1100 detects respiration via the RSL 1200 using bio-impedance.

Stimulation Lead (STL)

[0012] Figure 2 schematically illustrates the STL 1300 in more detail. The STL 1300 is designed to deliver the stimulation signal from the INS 1100 to the hypoglossal nerve and includes a proximal connector assembly 1310, a main tubular body 1330, and a distal cuff 1350. The main tubular body of the STL includes a sigmoid shaped section 1370 and a distal flexible transition section 1380 proximal of the cuff. The STL may have a nominal outside diameter of 0.062 inches to have minimal cosmetic impact, and a nominal overall length of 17.7 inches (45cm) (including cuff) to extend from the infraclavicular region (INS) to the submandibular region (hypoglossal nerve) and to accommodate anatomical variation.

[0013] The main tubular body 1330 of the STL 1300 is designed to withstand gross neck movement as well as mandibular movement and hypoglossal nerve movement caused by talking, chewing, swallowing, etc. To survive in this high fatigue environment, the main tubular body 1330 incorporates a highly compliant silicone jacket in the form of a sigmoid, and two conductors 1390 (one for cathode electrodes, one for anode electrodes) each comprising ETFE insulated MP35N multifilament cable disposed inside the jacket in the form of a bi-filar coil (not visible). This design provides high fatigue resistance and three-dimensional flexibility (bending and elongation).

[0014] The proximal connector assembly 1310 is designed to provide a reliable mechanical and electrical connection of the STL 1300 to the INS 1100. It has a number of strain relief elements that enable it to withstand handling during insertion and removal from the INS 1100, as well as adverse conditions encountered when implanted. The connector assembly 1310 includes two in-line stainless steel ring contacts (one for each conductor 1390) and two silicone ring seals. STL proximal connector contacts 1310 may have a nominal outside diameter of about 0.31 cm (0.122 inches). Set screws in the header of the INS 1100 bear down on the contacts, and together

with the ring seals, provide a sealed mechanical and electrical connection to the INS 1100. As an alternative, wound coil spring contacts may provide mechanical and electrical connections.

[0015] More detailed views of the cuff 1350 are shown in Figures 3A and 3B, wherein Figure 3A schematically illustrates the cuff 1350 in isometric view, and Figure 3B schematically illustrates the cuff 1350 in cross-sectional view. The cuff 1350 has a hinged oval-shaped silicone body (collectively 1352 and 1354) to define an oval lumen 1355 that provides secure and gentle retention around the hypoglossal nerve. The cuff 1350 may be designed to fit the nerve very closely to minimize tissue growth between the electrode and nerve. The cuff is designed to be self-sizing such that different nerve diameters may be accommodated safely. The self-sizing can safely adjust to larger sizes if swelling occurs. This reduces the likelihood of nerved damage caused by unsafe pressures. Thus, the cuff may be available in two sizes to accommodate nerves of different diameter: a small size to accommodate nerves having a diameter of up to about 2.5 - 3.0 mm, and a large size to accommodate nerves having a diameter of up to 3.0 - 4.0 mm. At 3.0 mm nerve diameter, either size cuff will fit the nerve with minimal open space for tissue in-growth. Using a large cuff on a 2.5mm nerve allows clearance between the nerve and electrode which promotes capsule formation between the cuff and nerve. This may cause an increase in capture threshold but will not affect safety. Conversely, a small cuff placed on a large nerve minimizes electrode coverage around the nerve and may fall off with swelling. The short side 1352 (e.g., 4.0 mm long) of the cuff body fits between nerve branches and connective tissue on the deep side of the nerve, thereby minimization nerve dissection. The long side 1354 (e.g., 10.0 mm long) of the cuff body rests on the superficial side of the nerve (where few branches exist) and is connected to the transition section 1380 of the main lead body 1330.

[0016] A silicone strap 1356 is connected to and extends from the short side 1352 of the cuff body. A silicone top plate comprising an integral base portion 1359 and loop 1358 is attached to and covers the exterior surface of the long side 1354 of the cuff body. The strap 1356 freely slides through the loop 1358, and wraps around the long side 1354 of the cuff body. The strap 1356 is removed from the loop 1358 for placement of the cuff 1350 around the nerve and reinserted into the loop 1358 to hold the cuff 1350 on the nerve. A mark may be disposed on the strap 1356 of the small size cuff to indicate that the cuff is too small and that a larger size cuff should be used if the mark does not pass through the loop 1358. The cuff body readily expands along a hinge line 1353 (defined at the junction of the short side 1352 to the long side 1354) as well as other portions of the cuff 1350 structure. Expansion of the cuff body accommodates nerves of different diameters and nerve swelling after implantation, while the strap 1356 remains in the loop 1358 to retain the cuff 1350 on the nerve. In the event of excess

nerve swelling (e.g., > 50% increase in nerve diameter) or traction from the lead 1300 (e.g., as may accidentally occur during implantation), the strap 1356 pulls out of the loop 1358 and releases the cuff 1350 from the nerve to minimize the potential for nerve damage.

[0017] The cuff body carries four platinum-iridium electrodes 1360 (e.g., 2.0 mm² exposed area each for small cuff, 3.0 mm² exposed area each for large cuff), with one cathode electrode 1360 on the short side 1352, another cathode electrode 1360 (not visible) diametrically opposed on the long side 1354, and two anode electrodes 1360 guarding the cathode electrode 1360 on the long side 1354. This guarded dual cathode arrangement provides a more uniform electrical field throughout the cross-section of the nerve while minimizing electrical field outside of the cuff. One conductor 1390 may be connected to the cathode electrode 1360 on the long side, to which the other cathode electrode 1360 on the short side is connected by a jumper wire. Similarly, the other conductor 1390 may be connected to the distal anode electrode 1360, to which the proximal anode electrode 1360 is connected by jumper wire. With this arrangement the cathode electrodes are commonly connected to one conductor 1390 and the anode electrodes are commonly connected to the other conductor 1390.

[0018] With the exception of the metal electrode contacts in the cuff, all external surfaces of the STL 1300 exposed to the body when implanted may comprise implantable grade polymers selected from the following: silicone, and fully cured silicone adhesive. The metal electrode contacts in the cuff may comprise implantable grade platinum-iridium and are secured to the silicone cuff body with silicone adhesive, for example.

Respiration Sensing Lead (RSL)

[0019] Figures 4A-4G schematically illustrates the respiration sensing lead (RSL) 1200 in more detail. The respiration sensing lead 1200 is designed to measure bio-impedance and includes a proximal portion with a proximal connector assembly 1210, a proximal tubular body 1220 ending in a bifurcation section 1280, and ipsi-lateral and contra-lateral distal portions extending from the bifurcation section. Each distal portion may include a tubular body 1220, a proximal sigmoid section 1230, a distal sigmoid section 1240, one or more current injection ring electrodes 1250, one or more voltage sensing ring electrodes 1260, anchor tabs 1270, and a suture hole 1290 in the most distal ring electrodes. Alternatively, the ring electrodes 1250 and 1260 may be dual-function, such that each electrode may function as either a current emitting electrode or voltage sensing electrode. The ipsi-lateral distal portion may contain three ring electrodes, the most distal being a current emitting electrode 1250 and containing a suture hole 1290, and the other two electrodes being voltage sensing electrodes 1260. The contra-lateral distal portion may contain two ring electrodes, the distal being a current emitting electrode 1250

and containing a suture hole 1290, and the proximal a voltage sensing electrodes. It may be advantageous to have the suture holes in the most distal ring electrodes since no wires pass through this point and because this provides a robust anchor point for the electrode to be sutured on the costal margin muscle fascia. The RSL 1200 may have a nominal outside diameter of about 0.072 inches to have minimal cosmetic impact. The RSL proximal connector contacts 1210 may have a nominal outside diameter of about 0.122 inches (same as the STL proximal connector contacts 1310). The distal ring electrodes (here, current emitting electrodes 1250), may also have a nominal outside diameter of 0.122 inches. This uniformity in diameters may be advantageous, allowing the same lead carrier 3100 to place both STL 1300 and RSL 1200 leads for tunneling.

[0020] The distance from the tip of the proximal connector 1210 to the bifurcation section 1280 may have an overall length of 8.9 inches (22.5 cm). The distance from the bifurcation section 1280 to the ipsi-lateral proximal anchor tab 1270 may be 9.6 inches (24.4 cm) unstretched and 12.2 inches (31 cm) stretched. The distance from the bifurcation section 1280 to the contra-lateral proximal anchor tab 1270 may be 13.5 inches (34.3 cm) unstretched and 16.1 inches (41 cm) stretched. The distance from the proximal anchor tab 1270 to the distal suture hole 1290 may be 2.8 inches (7 cm) unstretched and 3.1 inches (8 cm) stretched on both the contra-lateral and ipsi-lateral distal portions. The RSL 1200 may extend from the infraclavicular region (where the INS 1100 is implanted) to the ipsi-lateral and contra-lateral thorax where the RSL 1200 may be implanted to accommodate anatomical variation.

[0021] The bifurcated RSL 1200 design enables one RSL 1200 to sense bio-impedance on the contra-lateral and ipsi-lateral sides of the thorax. Two RSLs 1200, one on each side of the patient's chest, may also achieve this. The bifurcated design achieves this result while reducing the number of implanted components and reducing volume of the INS header 1110 since only one RSL port 1112 is required.

[0022] The main tubular lead body 1220 of the RSL 1200 is designed to withstand thoracic movement due to flexion, extension, rotation and breathing. To withstand this environment, the main tubular body 1220 may include a flexible silicone jacket formed such that each distal end has two sigmoid sections, 1230 and 1240, and conductors comprising small diameter ETFE insulated MP35NLT cables (not visible) disposed inside the jacket. An injection molded Y-fitting (yoke) connects the proximal portion of the RSL 1200 to the distal portions, creating the bifurcation section 1280. Conductors, here five, are continuously fed from the connector assembly through the proximal tubing body 1220 and proximal portion of the Y-fitting. Three of these conductors continue through the ipsi-lateral distal portion of the Y-fitting to the ipsi-lateral distal tubing body of the RSL. The other two conductors continue through the contra-lateral distal portion

of the Y-fitting and to the contra-lateral distal tubing body of the RSL. The tubing bodies may be adhesively bonded or molded to the Y-fitting. The number of conductors may equal the number of contacts in the INS header 1112, here five. Two of the conductors, one on each side, may connect proximally to current emitting header contacts, (e.g., RL and L1), and terminate distally in current emitting electrodes 1250. Three of the conductors may connect proximally to voltage sensing header contacts (e.g., R2, R3, and L3) and terminate distally in voltage sensing electrodes. As mentioned previously, dual-function electrodes may enable any electrode (ring electrode 1250 or 1260) to emit current or sense voltage. This switching may occur via components on the INS circuit board 1130. Alternatively, a bridge may be formed joining two contacts in the proximal connector assembly such that the corresponding electrode may function as either a current emitting electrode 1250 or voltage sensing electrode 1260. Dual-function electrodes enable more vectors in an implanted region without additional electrodes.

[0023] The proximal sigmoid section 1230 isolates movement of the INS 1100 from the electrodes 1250 and 1260, and accommodates anatomic variations in thoracic length. The distal sigmoid section 1240 allows adjustment in the distance between electrodes 1250 and 1260, and reduces strain applied between the anchor tabs 1270, which may be secured with sutures to the underlying fascia when implanted. The proximal sigmoid 1230 section may have 5 wavelengths with an outside peak-to-peak dimension of approximately 0.84 inches (2.1 cm) and an overall length of 7.0 inches (17.8cm). The distal sigmoid 1240 section may have half a wavelength with a center-to-center peak-to-peak dimension of approximately 0.43inches (2.1cm) and an overall length of 0.869 inches (2.2cm).

[0024] The two distal portions' electrodes 1250 and 1260 may comprise five electrodes total, and each may comprise MP35N rings having an exposed surface area. The distal electrode containing a suture hole 1290 may have an exposed surface area of 73.8 mm² including the suture hole 1290, and 66.4 mm² not including the suture hole 1290. The proximal electrode containing an anchor tab 1270 may have an exposed surface area of 30.5 mm² and the electrode without an anchor tab may have an exposed surface area of 32.0 mm². Tubular strain relief segments may be disposed on the lead body on either side of electrode 1250 or 1260. Where the strain relief segments are adjacent to each other, a gap may be provided there between the strain relief segments or the segments may abut one another to avoid a stress concentration point. Strain reliefs may also be disposed on each end of the electrodes 1250 or 1260 to avoid stress concentration points. The anchor tab 1270 may be disposed over an electrode leaving the proximal and distal extremities of the electrode exposed.

[0025] At any given time, the INS 1100 detects impedance along a vector, with each end of the vector defined by a current delivery electrode 1250 and a voltage sens-

ing electrode 1260. In each vector, a small excitation current is delivered between the two current emitting electrodes 1250, and the corresponding change in voltage is measured by the two voltage sensing electrodes 1260, The INS housing 1120 may also act as a current emitting and/or voltage sensing electrode, or contain smaller current emitting and/or voltage sensing electrodes on its surface. Changes in impedance are calculated by dividing the change in voltage by the excitation current, which correspond to movement of the diaphragm, lung, and other tissues to produce a signal indicative of respiratory activity.

[0026] The proximal connector assembly 1210 of the RSL 1200 is designed to provide a reliable mechanical and electrical connection of the RSL 1200 to the INS 1100. It has a number of strain relief elements that enable it to withstand handling during insertion and removal from the INS 1100, as well as adverse conditions encountered when implanted. The connector assembly 1210 may include five in-line stainless steel ring contacts (one for each conductor) and five silicone ring seals. Set screws in the header of the INS 1100 bear down on the contacts, and together with ring seals, provide a sealed mechanical and electrical connection to the INS 1100. Ring seals may be part of the RSL 1200 or the INS header 1110. With the exception of the distal electrodes, all external surfaces of the RSL 1200 exposed to the body when implanted may comprise implantable grade polymers selected from the following: silicone, and fully cured silicone adhesive. The distal electrodes may comprise implantable grade MP35N and are sealed to the lead body with silicone adhesive, for example.

[0027] A wide variety of respiration sensing lead designs may be employed to provide at least one bio-impedance vector (current injection electrode pair and voltage sensing electrode pair) from a point along the costal margin to a point along the opposite (trans-lateral) costal margin, to a point along the same side (ipsi-lateral) costal margin, or to a point in the infraclavicular region, as seen in Figure 4C. For example, an alternative embodiment of the RSL 1200 is shown in Figure 4B, wherein the bifurcation section 1280 and contra-lateral distal portion are eliminated. In this three electrode straight RSL 1200 embodiment, there is one current emitting ring electrode 1250 and two voltage sensing ring electrodes 1260. The lead body may contain three conductors. The connector assembly 1210 may include three in-line stainless steel ring contacts (one for each conductor) and three silicone ring seals. The RSL may have an overall length of 21.2 inches (53.9cm). The distance from the proximal tip of the proximal connector assembly 1210 to the first sigmoid may be 9.5 inches (24.1cm). The proximal sigmoid 1230 section may have 5 wavelengths with an outside peak-to-peak dimension of approximately 0.84 inches (2.1 cm) and an overall length of 7.0 inches (17.8cm). The distal sigmoid 1240 section may have a ½ wavelength with a center-to-center peak-to-peak dimension of approximately 0.43inches (2.1cm) and a length of 0.869 inches

(2.2cm). The RSL 1200 may be implanted ipsi-laterally on the ipsi-lateral costal margin, a less invasive surgery, while maintaining vectors from the ipsi-lateral costal margin to the infraclavicular region, see Figure 4D.

[0028] Further alternative embodiments are illustrated in Figures 4E, 4F, 4G, and 4H, wherein the RSL 1200 may contain four electrodes, one or more of which can function as either a current emitting electrode 1250 or voltage sensing electrode 1260, as described previously. Here, this is achieved by exposing the conductor of the bi-functional electrode to two contacts (one voltage sensing, one current emitting) in the INS header 1110, and selecting only one contact for sensing. Alternatively, this functionality may be built into the INS circuit board 1130. These embodiments of the RSL 1200 may be implanted ipsi-laterally (e.g. on the right costal margin), which is a less invasive surgery, while maintaining vectors from the ipsi-lateral costal margin to the infraclavicular region.

[0029] Figure 4E illustrates a straight four electrode RSL 1200. This is similar in design to the RSL 1200 of Figure 4B, wherein there are a first, second, third, and fourth electrodes, from proximal to distal.

[0030] Figure 4F illustrates a four electrode RSL 1200 with a loop back region 1255. This differs from the above embodiments in that there is an extra half sigmoid in the proximal sigmoid section 1230 after which the lead body enters the loop back region 1255. Here, the lead body runs in the medial direction, then loops back in the lateral direction. The loop back region 1255 may act as a strain relief and allow the medial anchor tab 1270 to be sutured at the intersection of the two tunneling paths (from INS incision to medial incision, and between two RSL incisions). This may allow the RSL 1200 to lie in an unbiased preferred configuration along the costal margin. Here again, the first and third most distal electrodes are current emitting, and the second, third, and fourth most distal electrodes are voltage sensing.

[0031] Figure 4G illustrates a four electrode RSL 1200 with a bifurcation section 1280 created by a T-fitting. An injection molded T-fitting connects the proximal portion of the RSL 1200 to the distal portions, creating the bifurcation section 1280. Conductors, here four, are continuously fed from the connector assembly through the proximal tubing body 1220 and proximal portion of T-fitting. Two of these conductors continue through the proximal distal portion of the T-fitting to the proximal distal tubing body of the RSL. The other two conductors continue through the medial distal portion of the T-fitting and to the medial distal tubing body of the RSL. The tubing bodies may be adhesively bonded or molded to the T-fitting. The anchor tab 1270 may be adhesively bonded to the bifurcation section 1280. Again, this may allow the RSL 1200 to lie in an unbiased preferred configuration along the costal margin. In addition, the T-fitting may act as a strain relief. Both medial and lateral distal electrodes may contain suture holes 1290.

[0032] Figure 4H illustrates an alternative embodiment of the RSL 1200, wherein the RSL 1200 has an overall

L-shape. In this embodiment, there may be four ring electrodes, numbered one through four from most proximal to furthest distal. The first and fourth electrodes may be current emitting electrodes 1250. The second and third electrodes may be voltage sensing electrodes 1260. The lead body may contain four conductors. The connector assembly 1210 may include four in-line stainless steel ring contacts (one for each conductor) and four silicone ring seals. The proximal portion of the RSL 1200 (including the proximal connector and proximal sigmoid) may have an overall length of 17.0 inches (43.2cm). The distance from the proximal tip of the proximal connector assembly 1210 to the first sigmoid may be 11.1 inches (28.1cm). The proximal sigmoid 1230 section may have 4.5 wavelengths, each wavelength 1.25 inches (3.2cm), and with an outside peak-to-peak dimension of approximately 0.84 inches (2.1 cm). The distal portion of the RSL 1200 (from the distal end of the proximal sigmoid 1230 to the distal suture hole 1290) may have an overall length of 4.9 inches (12.5cm). The length from the distal end of the proximal sigmoid 1230 to the proximal end of the distal sigmoid 1240 may be 2.2 inches (5.7cm). The length from the distal end of the distal sigmoid 1240 to the distal suture hole 1290 may be 1.8 inches (4.6cm). The distal sigmoid 1240 section may have a center-to-center peak-to-peak dimension of approximately 0.92inches (2.3cm). The RSL 1200 may be implanted ipsi-laterally on the ipsi-lateral costal margin, a less invasive surgery, while maintaining vectors from the ipsi-lateral costal margin to the infraclavicular region.

Implantable Neurostimulator (INS)

[0033] Figure 5A schematically illustrates the INS 1100 in more detail, including a front view, a top view and a side view. The INS 1100 is similar in certain aspects to commercially available implantable pulse generators and implantable neurostimulators, which may be obtained from suitable manufacturers such as CCC Medical Devices (Montevideo, Uruguay). The INS 1100 generally includes a header 1110 for connection of the STL 1300 and RSLs 1200, and a hermetically sealed housing 1120 for containing the associated electronics 1130, a battery 1140 (e.g., Greatbatch 9086), and an accelerometer 1150. The INS 1100 may contain an oxygen sensor (e.g., SaO_2 , SpO_2 , ion, etc.). Alternatively, the oxygen sensor may be incorporated in a lead with connection to the INS 1100.

[0034] The electronic circuitry 1130 contained in the INS 1100 enables telemetry communication with the programmer system 2100 and therapy controller 2500, detection of respiration via the RSL 1200, determination of the start time and duration of a stimulation signal, and delivery of a controlled electrical stimulation signal (pulse train) via the STL 1300. The INS 1100 also records therapy history data (device settings, status, measured data, device use, respiration data, stimulation delivery data, statistics based on measured signals, etc.).

[0035] The header 1110 may comprise epoxy that is hermetically sealed to the housing 1120. The housing 1120 may comprise titanium. As mentioned in the context of respiration sensing, the housing 1120 may be used as an electrode for bio-impedance respiration measurement. Similarly, electrodes 1360 may be used as an electrode for bio-impedance respiration measurement. For example, the housing 1120 may comprise a combination current emitting and voltage sensing electrode for respiration detection. Alternatively, separate electrodes may be included in the header of the device from which to sense or stimulate.

[0036] The header 1110 includes two ports: one RSL port 1112 (labeled "sense") for receiving the proximal connector of the RSL 1200 and one STL port 1114 (labeled "stim") for receiving the proximal connector of the STL 1300. The port configured to receive a STL 1300 includes two set screws (labeled "-" for cathode and "+" for anode) with associated set screw blocks and seals for mechanical and electrical connection to corresponding contacts on the proximal connector 1310 of the STL 1300. Similarly, the port that is configured to receive a RSL 1200 includes five set screws (two labeled R1 and L1 for current emitting electrodes and three labeled R2, R3, and L3, for voltage sensing electrodes) with associated set screw blocks and seals for mechanical and electrical connection to corresponding contacts on the proximal connector 1210 of the RSL 1200. Seals are located between electrical contacts as well as between the distal-most electrical contact and the remainder of the proximal connector assembly 1210. These seals electrically isolate each contact.

[0037] Alternatively, wound coil spring contacts may provide electrical connections between the INS header 1110 and the proximal connector assemblies 1210 and 1310. Typically, one electrical connection is still achieved with a set screw which also serves to hold the connector assembly in place. This embodiment provides a sealed mechanical and electrical connection of the RSL 1200 and STL 1300 to the INS 1100. An example of this technology is Bal Seal's Canted Coil™ Spring Technology.

[0038] The header 1110 further includes two suture holes 1116 for securing the INS 1100 to subcutaneous tissue such as muscle fascia using sutures when implanted in a subcutaneous pocket. As shown, component values and component configurations are given by way of example, not limitation.

[0039] The INS 1100 generates the stimulation output for delivery to the hypoglossal nerve by way of the STL 1300. For this purpose, the INS 1100 has a bipolar stimulation output channel corresponding to the STL port 1114, with the channel providing a pulse train of bi-phasic constant current pulses with a frequency range of 20 to 50Hz, a pulse width range of 30 to 215 μ s, an amplitude range of 0.4 to 5.0mA, and a stimulation duty cycle range of 41% - 69%, by way of example, not limitation.

[0040] The INS 1100 also generates the excitation signal and measures voltage by way of the RSL 1200 for

bio-impedance respiration detection. For this purpose, the INS 1100 also has two respiration sensing channels for simultaneous acquisition of bio-impedance sensing on different vectors. This may be achieved by sequential or alternating sampling of different vectors. The INS 1100 measures bio-impedance via the RSL port 1112, with each channel providing a small excitation current ("I") and measuring voltage ("V"). The excitation signal may comprise a 10Hz biphasic constant current pulse, with the positive and negative phases of each biphasic pulse having amplitude of 450 μ A, duration of 80 μ s, and charge of 36nC. Current ("I") may be fixed, allowing voltage ("V") to be a relative measure of impedance ("Z"), which corresponds to movement of the diaphragm, lung, and other structures to produce a signal indicative of respiratory activity.

[0041] With reference to Figure 5B, a block diagram of an example of the INS circuit 1130 is shown schematically. The INS circuit 1130 utilizes a microprocessor to control telemetry communications with the programmer system 2100, operating the sensing circuits to monitor respiration via the RSL 1200, controlling the delivery of output stimuli via the STL 1300, monitoring the accelerometer, magnetically sensitive reed switch and the real-time clock. The microprocessor contains built-in support circuits (RAM, flash memory, analog to digital (A/D) converter, timers, serial ports and digital IO) used to interface with the rest of the INS circuit 1130, including the accelerometer 1150. Two microprocessors communicating via a serial link may be used instead of one microprocessor, with the first microprocessor for telemetry communications, monitoring the accelerometer, magnetically sensitive reed switch and the real-time clock; and the second microprocessor for operating the sensing circuits and controlling the delivery of output stimuli. Alternatively, a single microprocessor could perform these functions.

[0042] The telemetry interface circuits consist of a tuned telemetry coil circuit and a telemetry driver/receiver circuit to allow pulse encoded communication between the external programmer system 2100 and the microprocessor. As an alternative to telemetry coils and an inductive link, RF antennae with associated circuitry may be used to establish a RF link to provide for arms-length telemetry. The reed switch provides a means for the INS 1100 to be controlled by using a magnet placed in close proximity thereto. The real-time clock provides the basic time base (768 Hz) for the INS circuit 1130 as well as a clock (year, day, hour, minute, second) which can be used to control the scheduled delivery of therapy. The clock is also used to time-stamp information about the operation of the system that is recorded on a nightly basis.

[0043] The respiratory sensing circuit is comprised of two main parts: the excitation current source (output) and the voltage sensing circuit (input). As will be described in more detail hereinafter, respiration is detected via the RSL 1200 using a 3 or 4-wire impedance measurement circuit. In a 4-wire measurement, an excitation current is

driven through a pair of electrodes, and the resulting voltage is measured on a separate pair of electrodes. The electrode switching circuits allow the INS 1100 to monitor one of several different vectors from the RSL electrodes 1250 and 1260. As mentioned previously, each physical electrode may function as a current emitting electrode 1250 or a voltage sensing electrode 1260, depending on the programmable vector configuration. In one embodiment of a 3-wire measurement, the INS housing 1120 may be used as both an excitation and sensing electrode. The excitation current circuit delivers biphasic pulses of low level (450uA) current to the selected electrode pair every 100ms during sensing. The voltage sensing amplifier circuit synchronously monitors the voltage produced by the excitation current on the selected electrode pair. The resulting output signal is proportional to the respiratory impedance (0.07Ω to 10Ω) and is applied to the A/D circuit in the microprocessor for digitization and analysis.

[0044] The stimulation output circuits deliver bursts of biphasic stimulation pulses to the STL 1300. These bursts may be synchronized to the sensed respiratory waveform to deliver stimulation and thus airway opening at the appropriate time. The stimulation output circuits include an electrode switching network, a current source circuit, and an output power supply. The electrode switching network also allows for a charge balancing cycle following each stimulation pulse during which the outputs are connected together with no applied output pulse. The timing and polarity of the pulse delivery is provided by control outputs of the microprocessor. The microprocessor selects the amplitude (e.g., 0.4mA to 5mA) of the output current from the current source circuit which is applied through the switching network. The output power supply converts battery voltage to a higher voltage (e.g., 5V to 14V) which is sufficient to provide the selected current into the load impedance of the STL 1300. The microprocessor measures the voltage output from the electrode switching network resulting from the delivered current and the load impedance. The microprocessor divides the output voltage by the output current resulting in a measure of the load impedance (400Ω to 2700Ω) which can be an indicator of integrity of the STL 1300.

[0045] The INS 1100 (or lead connected to the INS 1100) may contain an oxygen sensor to monitor oxygen levels, for example during a therapy session. This may be used to monitor efficacy as well to set stimulation settings during a therapy session. For example, the INS 1100 may be programmed to increase stimulation when oxygen de-saturations are detected at a programmable threshold rate and/or severity. In addition, the INS 1100 may turn stimulation on once de-saturations are detected, wherein thresholds of rate and severity are programmable. Desaturations may act to indicate the sleep state or wakefulness. In a similar manner, electroneurogram (ENG) may be used to monitor nerve activity, which may also be indicative of sleep state and/or wakefulness. The INS 1100 may use the indication of sleep state or wake-

fulness to change stimulation settings. For example, stimulation may be increased when the patient is in N3 or REM sleep. In addition, stimulation level may be decreased or turned off during stage N1, N2, or wakefulness.

[0046] The INS 1100 circuitry may contain a three-axis accelerometer 1150 that can be used to determine the patient's body position (supine, prone, upright, left, or right side) and/or detect motion events (wakefulness). These data may be used to change stimulation settings or inhibit output. The INS 1100 may be programmed to increase stimulation intensity when the patient is in specific body positions (e.g., supine, a more challenging position). The INS 1100 may segregate recorded therapy statistics (e.g., cycling detector events, oxygen desaturations) with respect to body position. For example, a patient's cycling detector may record very few events in the lateral position and many events in the supine position, indicative of the patient being treated in the lateral position.

[0047] With reference to Figure 6A, the bio-impedance respiration signal ("Z"), which is generated by dividing the change in measured voltage ("V") by the excitation current ("I"), tracks with diaphragm movement (DM) over time and therefore is a good measure of respiratory activity, and may be used to measure respiratory effort, respiratory rate, respiratory (tidal) volume, minute volume, etc. If the excitation current (I) is constant or assumed constant, then the bio-impedance (Z) is proportional to the measured voltage (V), and thus the voltage (V) may be used as a surrogate for bio-impedance (Z), thereby eliminating the division step. As used in this context, diaphragm movement includes movements and shape changes of the diaphragm and adjacent tissue that occur during normal breathing and during obstructed breathing. The bio-impedance waveform may be filtered to reduce noise and eliminate cardiac artifact, clarifying positive and negative peak occurrence. The signal may be filtered using a first order low pass filter. Alternatively, a higher order curve fit approach could be utilized to filter the signal. The (positive or negative) peak (P) of the impedance signal (Z) corresponds to the end of the inspiratory phase and the beginning of the expiratory phase. If the signal is normal (as shown), the positive peak is used; and if the signal is inverted, the negative peak is used. The beginning of the inspiratory phase occurs somewhere between the peaks and may not be readily discernable. Thus, the impedance signal provides a reliable fiducial (P) for end-inspiration and begin-expiration (also called expiratory onset), but may not provide a readily discernable fiducial for begin-inspiration (also called inspiratory onset). Therefore, algorithms described herein do not rely on begin-inspiration (or inspiratory onset) to determine the start of stimulation bursts as proposed in the prior art, but rather use a more readily discernable fiducial (P) corresponding to begin-expiration (or expiratory onset) in a predictive algorithm as described below. Other non-predictive (e.g., triggered) algorithms are de-

scribed elsewhere herein.

[0048] Gross body motion is often indicative of patient wakefulness and may change the bio-impedance signal (Z). A motion event may be detected, for example, by assessing variability in the bio-impedance peak-to-peak signal strength (P-P). Different thresholds of sensitivity may be utilized such that minor movements are not grouped with motion events. When a motion event is determined, stimulation may be turned off or turned down until motion stops or for a programmable duration of time. The frequency and duration of these motion events may be recorded in device history. The accelerometer 1150 could be utilized in a similar fashion to detect and record motion events

[0049] Waxing and waning of the bio-impedance signal (Z) is often indicative of apneas or hypopneas. Generally referred to as cycling, this pattern may be detected, for example, by assessing trends of increasing and decreasing average P-P amplitude values. Different thresholds of sensitivity may be utilized such that minor changes in P-P values are not declared cycling events. When cycling is detected, stimulation parameters may be initiated or changed (e.g., increased intensity, increased duty cycle, etc.) to improve therapy. The frequency and duration of these cyclic breathing patterns may be recorded in therapy history. These values may be used as an indicator of how well the patient is being treated, providing an estimate of AHI.

[0050] The INS 1100 may be programmed to change stimulation level between therapy sessions, days, or other programmable value. The stimulation level may be recorded alongside therapy session data, for example cycling rate (via the cycling detector), oxygen desaturation frequency and severity, stimulation time, variations in respiratory rate, variations in respiratory prediction, etc.

[0051] In people without OSA, inspiration is typically 25-50% of the respiratory cycle, with variations in respiration rate being common. Variations may cause actual inspiration to differ from predicted inspiration. The hypoglossal nerve usually activates approximately 300ms before inspiration and remains active for the entire inspiratory phase. To mimic this natural physiology, it is desirable to deliver stimulation to the hypoglossal nerve during the inspiratory phase plus a brief pre-inspiratory period of about 300ms. To maximize stimulation coverage of actual inspiration, it may be advantageous to account for this variability by centering stimulation on the predicted inspiration. As mentioned previously, there are reliable fiducials for the beginning of the expiratory phase (peak P) which may be used to deliver stimulation to cover the inspiratory phase plus brief pre and/or post-inspiratory periods.

[0052] Accordingly, an algorithm is used to predict respiratory period and determine the start of the stimulation burst. The predictive algorithm is contained in software and executed by a microprocessor resident in the INS circuitry 1130, thus enabling the INS 1100 to generate

stimulation synchronous with inspiration. One example of a prediction algorithm uses the respiratory period of previous breaths to predict the respiratory period of each subsequent breath. In this algorithm, a respiratory period is determined by calculating the time between peaks in the bio-impedance signal (Z). If the actual respiratory period is different from the predicted respiratory period, then the subsequent predicted respiratory period is resynchronized and updated to equal the actual period, up to a programmable value (e.g., 300ms). If the difference in respiratory period exceeds the programmable value, then the predicted respiratory period is incremented or decremented by this value.

[0053] One example of a predictive algorithm is illustrated in Figure 6B. In this example, the stimulation period is centered about a percentage (e.g., 75%) of the predictive respiratory period. The predictive algorithm uses historical peak data (i.e., begin-expiration data) to predict the time to the next peak, which is equivalent to the predicted respiratory period. The stimulation period is centered at 75%, for example, of the predicted respiratory time period. Thus, the time to start a stimulation burst is calculated by predicting the time to the next peak, adding 75% of that predicted time to the last peak, and subtracting 1/2 of the stimulation period (stimulation start time = time of last peak + 75% of predicted time to next peak - 1/2 stimulation period). A phase adjustment parameter (range: +/- 1000ms, for example) permits the stimulation period to be biased early or late.

[0054] A feature common to the predictive algorithms is illustrated in Figure 6C. This feature provides a sequence of predicted respiratory periods in case the respiration impedance signal ("Z") is temporarily lost (e.g., due to change in respiratory effort). Until a subsequent respiratory peak is detected, stimulation parameters which are based on the measured respiratory period (e.g., stimulation period) are unchanged. Thus, stimulation timing remains synchronous to the last detected peak.

[0055] The stimulation duty cycle may vary to meet efficacy and safety requirements. Generally, the stimulation duty cycle is used to determine the stimulation period as a percentage of the predicted respiratory period (stimulation period = duty cycle x predicted respiratory period). After a stimulation burst (pulse train) is started, stimulation continues until the end of the stimulation burst as set by the stimulation duty cycle, or until the next actual peak is detected, whichever occurs first. Alternatively, the feature of terminating a stimulation period when an actual peak is detected may be turned off.

[0056] The stimulation duty cycle may be fixed or adaptive. In the fixed mode, the stimulation duty cycle is set using to programmer system 2100 to a fixed percentage value. This fixed value may be increased when the respiratory signal is lost, increasing the likelihood of aligning with actual inspiration. In adaptive mode, the duty cycle is allowed to vary as a function of a characteristic of respiration. For example, the adaptive duty cycle may in-

crease when prediction is less accurate (higher variability in respiration rate) or when the respiratory signal is lost. Thus, in some instances, the stimulation duty cycle may run above normal (e.g., above 50% to 60%) to achieve a better likelihood of covering the inspiratory phase. Because above normal stimulation duty cycle may result in nerve and/or muscle fatigue if prolonged, it may be desirable to offset above-normal stimulation periods with below-normal stimulation periods to result in a net normal duty cycle. Thus, when the prediction is highly accurate (stable respiration rate), the stimulation duty cycle may be reduced.

[0057] The following stimulation duty cycle parameters are given by way of example, not limitation. In fixed mode, the maximum stimulation duty cycle may be set from 41% to 69% in 3% increments, and the default setting may be 50%. In adaptive mode, the stimulation duty cycle for a respiratory period may vary from 31% to 69% in 3% increments, and the maximum running average may be set to 53%. As mentioned above, the adaptive mode allows the duty cycle to decrease when respiratory period is stable and increase with respiratory period variability, for example, and the stimulation duty cycle may run in excess of 53% for a limited period of time, but those periods are proportionally offset by periods where the stimulation duty cycle runs less than 53% (e.g., according to an exponentially weighted moving average). For example, an adaptive duty cycle set to 69% would run at that level for no longer than 5 to 7 minutes before being offset by a lower stimulation duty cycle at 47% to result in a running average of 53%. This equation is approximate and may vary slightly depending on the averaging technique used. Other offset methods may be used as an alternative.

[0058] The stimulation duty cycle may be nominally 50%. A duty cycle limiter may be enabled such that it prevents the device from exceeding a programmable long term average stimulation duty cycle threshold (e.g., 53%). Long term average duty cycle may be calculated using a first order filter of duty cycle measured over a fixed time period (e.g., 6 seconds), with a programmable filter time constant (e.g., each iterative calculation is given a weighting of 1/32). If the long term average duty cycle reaches the programmable threshold, then stimulation duty cycle is decreased to a programmable value, (e.g., 44%) until the long term average drops below the nominal value (here, 50%), at which time the nominal duty cycle is restored. This safety mechanism may prevent nerve and muscle fatigue.

[0059] The INS 1100 may deliver stimulation as a train of pulses with constant pulse width and amplitude at a set frequency for a duration limited by duty cycle. This train of pulses may be described as a pulse train envelope and is illustrated in Figure 6D. The envelope describes a series of biphasic pulses delivered consecutively during a stimulation burst. When the stimulation level of the positive phase of each biphasic pulse is uniform, this level is the level of the stimulation burst. The INS 1100 may

also deliver stimulation in pulse train envelopes wherein the pulses are non-uniform (e.g., pulses may have different amplitudes).

[0060] The muscle(s) activated by the stimulation may not require the full stimulation intensity for the duration of the stimulation in order to maintain muscle contraction. Accordingly, the INS 1100 may be programmed to deliver a basic retention intensity pulse configuration, defined as a pulse train envelope wherein each pulse's intensity (e.g., amplitude) is less than or equal to the previous pulse's intensity, (e.g., a two second pulse wherein the first 1000ms is at 2mA and the subsequent 1000ms is at 1.7mA). This pulse configuration is illustrated in Figure 6E. This allows the muscle to activate to a level and then remain in that position with a less intense stimulation. This may be more comfortable and allow the patient to fall asleep more easily with the stimulation on, be less likely to cause arousal from sleep, and/or reduce the possibility of muscle/nerve fatigue. Alternatively, the pulse level (amplitude) could be decreased gradually during each burst (rather than abruptly) to reach the same final stimulation level.

[0061] A more gradual transition at the start of each burst may be more comfortable and be less likely to cause arousal from sleep, and/or reduce the possibility of muscle/nerve fatigue. Accordingly, the INS 1100 may be programmed to deliver a soft start pulse configuration, defined as a pulse train envelope wherein at the start of each burst, each pulse's intensity (e.g., amplitude) is greater than or equal to the previous pulse's intensity, (e.g., a two second pulse wherein the first 100ms is at 1.85mA, the second 100ms is at 1.95mA, the third 100ms is at 2.05mA and the remaining 1700ms is at 2.1mA). This pulse configuration is illustrated in Figure 6G. The pulse train envelope would thus have a stair-like appearance as stimulation increases to the full stimulation plateau.

[0062] In another embodiment, a pulse train envelope may employ a soft start to reach full stimulation and subsequently decrease intensity (amplitude) to a retention intensity for the remainder of the stimulation, (e.g., a two second pulse wherein the first 100ms is at 1.85mA, the second 100ms is at 1.95mA, the third 100ms is at 2.05mA, and the next 700ms is at 2.1mA, and the remaining 1000ms is at 1.8mA). This pulse configuration is illustrated in Figure 6F. This configuration may provide the benefits of both soft start and retention intensity, wherein the stimulation starts gradually to fully activate the muscle(s), then decreases to a level of less intense stimulation, with the muscle remaining in a contracted position. Figure 6H shows nested mode, a simplified version of the previously mentioned retention intensity, wherein there is one step up to the full amplitude, and then an equal step down to the retention intensity.

[0063] The INS provides two separate stimulation strengths (A & B) with independent parameters (amplitude, pulse width, frequency, duty cycle and phase adjust). Stimulation may be delivered in different therapy

modes, examples of which are shown in Figure 6I. Figure 6I(traces 1-8) illustrates some commonly used modes, all of which are inspiratory synchronous, meaning stimulation is automatically delivered according to an algorithm that predicts the inspiratory phase and initiates stimulation delivery at a desired time relative to inspiration, such as centered on the predicted inspiration. These modes may be used as standard therapy as well as to determine device settings during a PSG (e.g. sleep titration PSG). Additionally, these modes may be used to diagnose phenotypes of OSA or other diseases.

[0064] Figure 6I(trace 1) illustrates synchronous mode in which every stimulation has the same pulse configuration and amplitude, known as AAAA mode, the default therapy mode. The term AAAA mode means that four consecutive inspirations are covered by stimulation of level A, where A is 2.0mA, for example. Inspiration is shown in Figure 6I(trace 9) in the upward direction.

[0065] The inspiratory-synchronous ABAB mode, Figure 6I(trace 2), also delivers stimulation bursts synchronous with inspiration as determined by the device, therapy delivery algorithm settings, and sensed respiratory signal. This mode is similar to AAAA mode, except that the stimulation is delivered on four consecutive inspirations alternating between stimulation levels A and B and each burst where, for example, A is 2.0mA and B is 1.8mA.

[0066] Figure 6I(trace 3) illustrates a subset of ABAB mode known as A0A0 mode, wherein the B breath is not stimulated. A may be 2.0mA and B may be 0mA, for example.

[0067] Figure 6I(trace 4) illustrates A0B0 mode, wherein a first breath is stimulated at level "A," followed by an second breath that is unstimulated, followed by a third breath that is stimulated at level "B," followed by a fourth breath that is unstimulated, (e.g., A is 2.0mA and B is 1.8mA). This allows for simultaneous assessment of two different levels (A and B) when compared to adjacent non-stimulated breaths.

[0068] Figure 6I(trace 5) illustrates AABB mode wherein two breaths are stimulated at level "A" followed by two breaths stimulated level "B," (e.g. A is 2.0mA and B is 1.8mA). In this mode, every stimulated breath is adjacent to a stimulated breath at level "A" and a stimulated breath at level "B." The AABB mode may be used to test if there is a short-term residual cross-over effect when changing from one stimulation level to another stimulation level or from a stimulation level to no stimulation. For example, the airflow measured during the first "A" can be compared to the airflow measured during the second "A" in each sequence over many periods to determine if there is a measurable residual effect from the "B" level simulation.

[0069] Figure 6I(trace 6) illustrates xAB0 mode, wherein "x" number of breaths are stimulated at level "A," followed by a breath stimulated at level "B," followed by an unstimulated breath, (e.g. A is 2.0mA and B is 1.8mA). The illustration shows x equals 3 (3AB0), although x may be any number of breaths (e.g., 3, 5, 7).

[0070] Figure 6I(trace 7) illustrates xA0B mode, wherein "x" number of breaths are stimulated at level "A," followed by an unstimulated breath, followed by a breath stimulated at level "B," (e.g. A is 2.0mA and B is 1.8mA). The illustration shows x equals 3 (3A0B), although x may be any number of breaths (e.g., 3, 5, 7).

[0071] Figure 6I(trace 8) illustrates xAB mode, wherein "x" number of breaths are stimulated at level "A," followed by a breath stimulated at level "B," (e.g. A is 2.0mA and B is 1.8mA). The illustration shows x equals 4 (4AB), although x may be any number of breaths (e.g., 4, 6, 8),

[0072] Stimulation may also be delivered in two modes which are not inspiratory synchronous: manual stimulation and asynchronous (fixed) stimulation. Manual mode delivers stimulation at any frequency, pulse width, amplitude, pulse configuration, and/or duration (e.g., up to 12 seconds). In manual mode, stimulation is delivered by manually entering a command via the programmer system to initiate delivery of a stimulation burst or bursts. The stimulation continues until the burst duration expires or stimulation stop is commanded via the programmer system. Manually delivered stimulations may be delivered in any available pulse configuration.

[0073] Asynchronous mode (fixed mode) is when stimulation is delivered at regular programmable intervals (e.g., 2.5 seconds of stimulation, followed by 2.5 seconds off). The intervals may be set to a rate similar to a respiratory cycle, (e.g. 5 seconds). Alternatively, longer intervals (e.g. 8 seconds) may decrease the probability of missing two consecutive inspirations, and increase the probability of providing the patient with stimulation during an entire respiratory cycle. This may be used during daytime familiarization, ensuring that the patient receives stimulation in a regular fashion, as breathing patterns during wakefulness may be more irregular and difficult to predict than during sleep. In addition, this mode may be used to test the benefits of asynchronous stimulation compared to inspiratory synchronous stimulation. Asynchronous stimulation may be initiated by programming the device to fixed mode and starting a therapy session. Fixed mode stimulations may be in any available pulse train configuration.

[0074] Typically, stimulation is delivered during a therapy session having a start and a stop time. The patient or physician may start a therapy session using the therapy controller 2500 or programmer system 2100. Additionally, a therapy session may begin according to a programmable schedule. During a session, the start of stimulation may be delayed by a programmable delay, subject to patient preference. The patient or physician may stop a therapy session using therapy controller 2500 or programmer system 2100. Additionally, a therapy session may stop according to a programmable schedule or programmable maximum session duration.

[0075] A patient or physician may also pause a therapy session for a programmable time using the therapy controller 2500 or programmer system 2100. This pause function may be programmed to turn stimulation off or

reduce the stimulation intensity. The pause function may be programmed to smart pause, wherein the stimulation level is automatically reduced after a programmable number of pauses (e.g. after the second pause) in a programmable time period or session. Additionally, the smart pause may increase pause duration after a programmable number of pauses (e.g. the first pause is five minutes, the second pause is ten minutes). These pause functions, including smart pause, may allow a patient to reduce stimulation for brief periods following an arousal from sleep.

[0076] At the start of a therapy session or following an interruption in therapy such as a pause, stimulation level may increase incrementally from an initial stimulation level to an initial therapy level during a ramping period. The ramp may occur over a programmable number of stimulations, breaths, or time period. This ramp may also occur after a pause or a motion event. The ramping feature may be more comfortable, allowing the patient to fall asleep more easily with the stimulation on or reduce the likelihood of causing arousal from sleep.

[0077] A patient may be able to tolerate more intense stimulation as a therapy session progresses. This higher intensity stimulation may provide enhanced therapy efficacy. The INS 1100 may be programmed to change stimulation level (e.g., amplitude or pulse width) during a therapy session from an initial level to a second, possibly more efficacious level. This therapy stimulation configuration is illustrated in Figure 6J and is called core hours. This intensity change may occur after a programmable interval, for example after a fixed duration of time, number of breaths or stimulations (e.g., stimulation at 1.8mA for the first hour of a therapy session, after which stimulation is increased to 2.0mA). This feature and the related parameters may be programmed by a physician, for example based on patient feedback. This feature may allow a patient to fall asleep at a more tolerable level of stimulation, and then as the therapy session progresses, receive more appropriate therapeutic benefit.

[0078] Stimulation may be delivered during a therapy session, which may start and stop according to a programmable schedule or manual use of the therapy controller 2500. The therapy controller 2500 may also allow the patient to pause or adjust therapy settings. Summary history data from each session may be saved in the device memory. Data recorded may include: start, pause, and stop times of the therapy session, scheduled or manual starts/stops, motion detector outputs, cycling detector outputs, prediction algorithm outputs, respiration timing, signal stability outputs, accelerometer outputs, impedance data from STL 1300 and RSL 1200, number of breaths, number of stimulations in a session, average and median P-P sensing impedance ("Z") amplitude values, stimulation settings and changes in stimulation settings such as core hours, pulse configuration, and ramping. These summary data allow a physician or caretaker to understand how the patient is using the device, tolerating the stimulation, troubleshoot errors in program-

ming, and estimate the therapeutic effects of the neurostimulator. This feedback data may aid in determining if adjustments are needed to the patient's therapy (e.g. patient is ready for stimulation up-titration).

Programmer System

[0079] As shown schematically in Figure 7A, the programmer system 2100 includes a computer 2300, a programmer interface 2400, a programmer head 2200, and a sleep wand 2700. The programmer interface 2400 and programmer head 2200 are similar in certain aspects to commercially available programmers, which may be obtained from suitable manufacturers such as CCC Medical Devices (Montevideo, Uruguay). The programmer head 2200 is connected to the programmer interface 2400 via a flexible cable 2210, and the programmer interface 2400 is connected to the computer 2300 via a USB cable 2310. Cable 2210 may be coiled as shown or straight. As shown in Figure 15, the sleep wand 2700 may comprise a sleep wand head 2720, a flexible cable 2710, and an LED 2730. The sleep wand 2700 may connect to the programmer interface 2400 via a flexible cable 2710. The sleep wand head 2720 may be 3.2 inches in length, 2.1 inches in width, and .5 inches deep. The programmer system 2100 wirelessly communicates with the INS 1100 via a wireless telemetry link (e.g., 30 KHz) utilizing an antenna and associated circuitry in the programmer head 2200. The programmer may use long range telemetry such that the programmer head 2200 may rest beside the patient without interfering with sleep. The programmer interface 2400 provides analog to digital conversion and signal processing circuitry allowing the computer 2300 to control and program the INS 1100 via the programmer head 2200. The programmer head includes a power indication LED 2220, a signal strength LED array 2230 (signal strength to/from INS 1100), an interrogate button 2240 (to upload data from INS 1100), a program button 2250 (to download data/commands to the INS 1100) and a therapy-off button 2260 (to stop therapy/ stimulation output from the INS 1100). The computer 2300 may comprise a conventional laptop computer with software to facilitate adjustment of a variety of INS 1100 parameters, including, for example: stimulation parameters (stimulation pulse amplitude, stimulation pulse frequency, stimulation pulse width, stimulation duty cycle, etc.); respiration sensing algorithm parameters; stimulation trigger/synchronization algorithm parameters, therapy delivery schedule, and various test functions. The sleep wand 2700 functions like the programmer head 2200, but is reduced in size for patient comfort during sleep. There may be one LED 2730 to indicate signal presence. Frequency of LED light pulses may indicate signal strength. The sleep wand 2700 may exclude functional buttons (i.e. interrogate command, program command, and stop therapy command) found on the programmer head 2200.

[0080] With reference to Figure 7B, a block diagram of

example circuits 2420/2270 for the programmer interface 2400 and the programmer head 2200 are shown schematically. The programmer interface circuit 2420 is controlled by a microprocessor having a standard set of peripherals (RAM, flash, digital I/O, timers, serial ports, A/D converter, etc.). The microprocessor communicates with a standard personal computer (PC) 2300 through a Universal Serial Bus (USB) interface. Commands and data are passed from the computer 2300 to/from the microprocessor via the USB interface and cable 2310. The USB interface also provides DC power for the programmer interface circuit 2420 and the programmer head circuit 2270 via cable 2210. The microprocessor controls the cable interface leading to the programmer head circuit 2270 via cable 2210. The programmer head circuit 2270 contains telemetry driver and receiver electronics that interface to the telemetry coil. The telemetry coil is designed to inductively couple signals from the programmer head circuit 2270 to the coil in the INS circuit 1130 when the programmer head 2200 is placed over the INS 1100 with the coils aligned. As an alternative to telemetry coils and an inductive link, RF antennae with associated circuitry may be used to establish a RF link to provide for arms-length telemetry. The programmer head circuit 2270 also contains electronics that monitor the signal strength as received from the INS 1100. The outputs of the signal strength electronics drive display LED's for the user. Another LED indicates that power is available, for example, supplied by the computer 2300. The programmer interface microprocessor controls and receives analog input signals from an isolated sensor interface. The power and ground for the sensor interface are derived from the USB power input, but provide DC isolation for this circuitry to prevent leakage currents from flowing through any patient connections that may be present at the sensor inputs. The sensor inputs may be protected against external high voltages (i.e. defibrillation protection). The sensor input signals are amplified and filtered appropriately for the sensor type. The amplifier gain and filter characteristics may be controlled by microprocessor. The signals to/from the amplifier circuit are DC isolated to prevent leakage currents from flowing through any patient connections that may be present at the sensor inputs. The sensor signals are digitized by the microprocessor and are transmitted through the USB link to the PC along with the telemetered signals from the INS 1100 for recording and display.

[0081] With reference to Figure 7C, a block diagram of example circuit 2440 for the marker box 2430 is shown schematically. Generally, marker box 2430 and associated circuitry 2440 replace the D/A circuits and analog outputs 2410 of programmer interface circuit 2420 shown in Figure 7B providing for the alternative arrangement illustrated in Figure 10B. The marker box circuit 2440 is separately connected to a Universal Serial Bus (USB) port of the programmer computer 2300 via a USB cable. The USB interface also provides DC power for the marker box circuit 2440 via the USB cable. The power and

ground for the marker box circuit 2440 are derived from the USB power input, but provide DC isolation for this circuitry to prevent leakage currents from flowing through any equipment that may be connected to the patient. Analog marker output data signals are transmitted from the PC 2300 to control the digital to analog (D/A) converter outputs. These analog output signals may be connected to standard PSG recording equipment 2800. Signals from the INS 1100 (such as sensed respiration impedance and stimulation output timing and amplitude) can be represented by these outputs to allow simultaneous recording with other standard PSG signals (flow, belts, EMG/ECG, etc). The programmer 2300 may be enabled to automatically switch programmable settings at regular time intervals, allowing respiratory sensing vectors, stimulation levels, stimulation modes, or stimulation pulse configurations to be altered at specified intervals during sleep. Sampled values may be selected such that only a limited number of settings are sampled.

[0082] As mentioned previously, the INS 1100 records session summary data. The programmer computer 2300 may display these data using text and images to graphically display device settings, session data, and analyses of data. This data may be used to evaluate system performance and guide programming of settings. The patient's name or identifier may be stored in the INS 1100 and/or displayed on the programmer computer 2300. All text and symbols displayed by the programmer 2300 may be in a variety of selectable languages. The programmer 2300 may have the capability to connect to the internet. Through this connection files may be uploaded to a database to enable remote real time monitoring of device operation, recorded data and settings. The connection may also be used to update the programmer application software and or the (indirectly) the INS firmware.

[0083] The programmer 2300 may display and tag data to a variety of dates and times. These times may be programmed to take into account daylight savings time, local time, Greenwich Mean Time, or a free-running counter in the INS 1100.

[0084] The programmer 2300 may display the voltage (or other capacity measurements) of battery 1140. In addition, an elective replacement indicator (ERI) and end of life (EOL) indicator may be displayed on the programmer as the battery nears depletion. There may be several months from ERI to EOL or alternatively, two months from ERI to EOL, with an estimated one month of use after EOL.

Therapy Controller

[0085] As shown schematically in Figure 8A, the therapy controller 2500 may be used by the patient to control limited aspects of therapy delivery. The therapy controller 2500 is similar in certain aspects to commercially available patient controllers, which may be obtained from suitable manufacturers such as CCC Medical Devices (Montevideo, Uruguay). The therapy controller 2500 houses

a battery, an antenna, and associated circuitry (not visible) to control limited aspects of therapy delivery via a wireless telemetry link (e.g., 30 KHz) with the INS 1100. Therapy is normally operated in a manual mode but may also be set for automatic delivery according to a predefined schedule (set by physician using the programmer during titration). The therapy controller has a user interface including start button 2510 (to start therapy delivery), a stop button 2520 (to stop therapy delivery) and a pause button (to pause therapy delivery or reduce stimulation intensity to a programmable value), each with associated LED indicators 2540 which flash when the corresponding button is depressed and illuminate steadily when the command is received by the INS 1100. The buttons may be backlit when pressed for ease of use at night. The user interface also includes a schedule set LED 2550 that illuminates if a therapy delivery schedule has been programmed, and a contact physician LED 2560 that illuminates in the event of a low battery or a malfunction requiring a physician visit. In addition, this light may illuminate at ERI or EOL time points.

[0086] The therapy controller may have additional functionality (e.g., more buttons) which can be set to give the patient limited control over select therapy settings. These settings include, but are not limited to, stimulation intensity (e.g., amplitude), stimulation mode, pulse train configuration, core hours stimulation settings, ramp, programmable schedule, clock, and motion inhibit programmable values.

[0087] As mentioned previously, the INS 1100 contains data, including metrics from therapy sessions. The therapy controller 2500 may wirelessly communicate with the INS 1100 to download any data to the INS. This data may be stored in internal memory or removable memory such as a USB flash drive or smart card. This data may be uploaded (e.g. from the patient's home computer) for the physician to read. This may allow the physician to monitor device use, home efficacy, or the patient's acclimation.

[0088] An alternative embodiment of the user interface may include an LCD display or a touchscreen. This allows for multiple functions to be integrated into the therapy controller while keeping the interface simple. This may also allow for larger text.

[0089] With reference to Figure 8B, a block diagram of an example circuit for the therapy controller 2500 is shown schematically. The therapy controller circuit 2570 includes a battery powered microprocessor having a standard set of peripherals (RAM, flash, digital I/O, timers, serial ports, A/D converter, etc.). The microprocessor operates in a low power mode to conserve battery power. The microprocessor controls the telemetry driver and receiver electronics that interface with the telemetry coil. The telemetry coil is designed to inductively couple signals to the INS telemetry coil when aligned. The microprocessor monitors the membrane switches and reacts to switch closures by activating display LED's and initiating telemetry commands to the INS. As an alternative

to telemetry coils and an inductive link, RF antennae with associated circuitry may be used to establish a RF link to provide for arms-length telemetry. After communicating with the INS, status information can be displayed to the user. The microprocessor also controls a beeper which can provide audio feedback to the user when buttons are pressed and to indicate the success or failure of communications with the INS. The beeper may have a mute function or volume control.

Magnet

[0090] As schematically shown in Figure 9, an annular magnet 2600 may be provided to the patient to inhibit the INS 1100 in the event the therapy controller 2500 is not available or functioning. The magnet 2600 may comprise a permanent annular-shaped magnet made of ferrite strontium material coated with epoxy. The magnet 2600 may produce a strong field of 90 Gauss at 1.5 inches from the surface of the magnet along the centerline of the hole. The magnet 2600 may be used (or carried by) the patient in case of emergency. When temporarily (2 seconds or more) placed over the implanted INS 1100 on the skin or clothing, the magnet 2600 disables current and future therapy sessions. Although therapy sessions are disabled by the magnet 2600, all other functions of the INS 1100 may remain enabled including telemetry communication with the programmer system 2100 and therapy controller 2500. Therapy sessions may be re-enabled using the therapy controller 2500 by initiating a new therapy session. Alternatively, the therapy may be temporarily inhibited during placement of the magnet. If left in place for a specified time period (e.g. one minute), therapy may be deactivated.

Interface with PSG Equipment

[0091] The programmer interface 2400 may include an input/output (I/O) link 2410 to allow connection to polysomnographic (PSG) equipment 2800 as schematically shown in Figure 10A. Typical PSG equipment 2800 includes a computer 2810 connected to a plurality of sensors (e.g., airflow sensor 2820, respiratory effort belts 2830) via interface hardware 2840. The I/O link 2410 may be used in a number of different ways. For example, analog data signals from the PSG equipment 2800 may be downloaded to the computer 2300 of the programmer system 2100 to record and/or display PSG data (e.g. airflow) together with therapy data. Alternatively or in addition, digital data signals from the INS 1100 and/or the programmer system 2100 may be uploaded to the computer 2810 of the PSG equipment 2800 to record and/or display therapy data (e.g., stimulation amplitude, stimulation pulse width, and/or respiration data such as bio-impedance, vector, filter settings, prediction markers, or accelerometer data) together with PSG data. The circuitry corresponding to I/O link 2410 may be incorporated into the programmer interface 2400 as shown in Figure

10A, or may be incorporated into a separate marker box 2430 as shown in Figure 10B.

[0092] Synchronizing data from the sensors 2820/2830 of the PSG equipment 2800 with data from the INS 1100 via the programmer system 2100 may be beneficial to facilitate therapy titration and efficacy measurement. Although the programmer system 2100 and the PSG equipment 2800 may be directly connected by I/O link 2410, transmission delay in each system may result in asynchrony. Data synchronization between the systems may be addressed in a number of different ways. For example, if the delays in each system are relatively fixed and below an acceptable threshold (e.g., 0.5 to 1.0 second), no synchronization step need be taken. If the delays in each system are relatively fixed but above an acceptable threshold (e.g., above 0.5 to 1.0 second), data from the system with less delay may be offset (delayed) by a fixed time value to align with data from the system with more delay. As an alternative, a timing signal (e.g., from a clock signal generator separate from or integral with one of the systems) may be input into the PSG equipment 2800 and programmer system 2100 to allow time stamped data independently collected by each system to be merged and synchronized by post processing.

Treatment Overview

[0093] Figure 12A illustrates a treatment overview from implant 10100, to awake titration 10200, to daytime familiarization 10400, to sleep titration 10500, to nighttime familiarization 10700, to up-titration 10800, and finally to regular therapeutic use 11000 and therapy assessments 11100.

[0094] Beginning with a surgical implant 10100, figure 11A schematically illustrates the incision sites (solid thick lines) and tunneling paths (dotted lines) for implanting the INS 1100, STL 1300 and RSLs 1200. The implant procedure may be performed by a surgeon (e.g., otolaryngologist) in a 1 - 3 hour surgical procedure with the patient under general or local anesthesia, for example. In general, the implant procedure involves placing the cuff 1350 of the STL 1300 on the hypoglossal nerve via a submandibular dissection, and tunneling the lead body 1330 and sigmoid section 1370 of the STL 1300 subcutaneously down the neck to the INS 1100 in a subcutaneous pocket in the infraclavicular region. From the infraclavicular pocket, the RSL 1200 may be tunneled subcutaneously toward midline and then laterally along the costal margins.

[0095] After a healing period of a few weeks, an awake titration may be performed 10200 wherein the patient's tongue response to stimulation is observed over a range of comfortable stimulations, as illustrated in Figure 12B. In addition, a global system check may be performed to check the system integrity. The patient is then sent home for a period of daytime familiarization 10400 where the patient may turn on stimulation during wakefulness to introduce the sensation of stimulation. The patient sub-

sequently returns to the sleep lab for a sleep titration 10500 where a sleep technician, under the supervision of a certified sleep physician (e.g., pulmonologist), uses the programmer system 2100 to program the INS 1100 (e.g., set the therapy delivery schedule and titrate the stimulus to determine a range of efficacious settings during sleep). After the sleep titration, the patient may return home and begin the nighttime familiarization (acclimation) and therapy up-titration process, wherein stimulation may be increased over time to an efficacious range. For example, the patient may leave the sleep titration with stimulation programmed to turn on at 1.7mA and stimulation may be increased by 0.1mA at two week intervals up to a goal setting of 2.0mA.

[0096] Immediately after the titration visit, the patient may return home and begin using the device at the programmed stimulation level during nighttime familiarization 10700. A therapy delivery session may begin when the therapy controller 2500 is used to manually start, stop, and pause a therapy session. This may be beneficial when the patient has an irregular sleep schedule. At the beginning of a therapy delivery session, stimulus may be delayed for a period of time to allow the patient to fall asleep. The therapy delivery session may be programmed to not exceed a fixed number of hours (e.g., eight hours). In addition, a therapy delivery session may begin according to the pre-defined therapy delivery schedule, which may be set to coincide with when the patient normally goes to sleep. The therapy delivery session may end according to the pre-defined therapy delivery schedule, which may be set to coincide with when the patient normally wakes up, or with a manually stop command from the therapy controller. The patient may use the therapy controller 2500 to adjust limited aspects of therapy delivery as defined previously.

Tunneling System

[0097] Figure 11B and 11C schematically illustrate the tunneling system 3000 which may be used for tunneling the STL 1300 or RSL 1200. The tunneling system 3000 includes a relatively rigid tool 3010, a tubular sheath 3020, and a tip 3030, and lead carrier 3100.

[0098] The tool 3010 may be formed of stainless steel and include a handle 3016, a shaft 3012, and a distal connector 3018. The connector 3018 includes threads that mate with corresponding threads in the tip 3030. The connector 3018 may also include ring barbs that form an interference fit with the inside of the lead carrier 3100 for releasable connection thereto. The lead carrier 3100 may also form an interference fit with the RSL proximal connector 1210, the STL proximal connector 1310, or a distal ring electrode of the RSL 1250 or 1260.

[0099] The sheath 3020 is sized to be slid over the tool 3010 and secured in place via the tip 3030. The tip 3030 may include a radiopaque agent such as barium sulfate loaded at 18% by weight, for example.

[0100] The lead carrier 3100 may comprise a small pol-

ymeric tube with an inside diameter sized to form an interference fit with the distal connector 3018, the RSL proximal connector assembly 1210, an RSL distal electrode 1250 or 1260, or the STL proximal connector assembly 1310. During tunneling, the proximal end of the lead carrier 3100 may attach to the distal connector 3018 and the distal end of the lead carrier 3100 may attach to the RSL proximal connector assembly 1210, an RSL distal electrode 1250 or 1260, or the STL proximal connector assembly 1310.

[0101] The sheath 3020 may comprise a polymeric tube with two open ends, and the tip 3030 may comprise a polymeric tube with one threaded end and one closed end for blunt dissection. The proximal end of the tip 3030 includes internal threads to screw onto the connector 3018 and hold the sheath 3020 on the shaft 3012.

[0102] In the embodiment shown in Figures 11B and 11C, the tool 3010 may have a pre-bend length of 17.1 inches (43.4 cm) and a post-bend length of 16.9 inches (40.6 cm). The sheath 3020 may have an outside diameter of approximately 0.28 inches (0.711 cm), a pre-bend length of 12.4 inches (31.5 cm) and a post bend length of 12.25 inches (31.11 cm). The shaft 3012 may have a diameter of about 0.22 inches (0.559 cm), a pre-bend length of 12.375 inches (31.43 cm), and a post-bend length of 12.231 inches (31.07 cm), sufficient to fill the length of the sheath 3020. The handle 3016 may have a diameter of about 0.5 inches (1.27 cm) and a length of about 3.50 inches (8.89 cm). The tip 3030 may have an outside diameter tapering from approximately 0.13 inches (0.33 cm) and a length of about 1.0 inches (2.54 cm).

Surgical Implant Procedure

[0103] With continued reference to Figure 11A, the internal components 1000 may be implanted using the following surgical procedure 10100, which is given by way of example, not limitation. Unless specifically stated, the order of the steps may be altered as deemed appropriate. Although the INS 1100 may be surgically implanted on the right or left side, the right side is preferred to leave the left side available for implantation of cardiac devices that are traditionally implanted on the left side. The right side is also preferred for the RSL 1200 to provide a clean respiratory signal that is less susceptible to cardiac artifact than the left side.

[0104] Standard surgical instruments may be used for incisions, dissections, and formation of subcutaneous pockets. Commercially available nerve dissection instruments may be preferred for dissecting the hypoglossal nerve and placing the STL cuff 1350 on the nerve.

[0105] The patient is prepared for surgery using conventional practice including standard pre-operative care procedures, administration of antibiotics as appropriate, and administration of steroids as appropriate to reduce swelling around the nerve dissection. Because tongue movement must be observed during test stimulation, it is recommended that no long-acting muscle relaxants be

used during surgical preparation or during implant. General anesthesia is administered according to conventional practice and the patient is intubated using an endotracheal tube, taking care to position the endotracheal tube so that the tongue is free to protrude during test stimulation.

[0106] The neck is then extended to expose right submandibular region and a sterile field is created around the neck and thorax, taking care to avoid obstructing visualization of the oral cavity (a clear sterile drape over the mouth may be used). By way of a neck incision (A), the hypoglossal nerve is then exposed deep to the submandibular gland. Because the INS 1100 is preferably implanted on the right side to minimize cardiac artifact during respiratory sensing, this dissection is also preferably performed on the right side. A region of the hypoglossal nerve, preferably excluding the branch that innervates retrusive muscles (e.g. styloglossus or hyoglossus), is then identified and isolated. Confirmation of correct nerve location may be achieved by performing a test stimulation later in the procedure. The identified nerve branch is then circumferentially dissected to accommodate the cuff 1350. The short side 1352 of the cuff 1350 is designed to reside on the deep side of the nerve, and the long side 1354 of the cuff 1350 is designed to reside on the superficial side of the nerve.

[0107] The appropriate sized cuff 1350 is then selected based on the nerve diameter at the intended location for cuff placement. Nerve size may be assessed using reference size (e.g., forceps of known width), a caliper, or a flexible gauge that wraps around the nerve, for example. The cuff 1350 is then opened and placed around the nerve. The strap 1356 on the cuff 1350 may be used to facilitate placement of the cuff 1350 around the nerve. A curved forceps may be placed under the nerve to grasp the strap 1356 and gently pull the cuff 1350 onto the nerve. The strap 1356 is then placed through the loop (buckle) 1358 on the cuff 1350. The cuff 1350 may be available in two sizes (small and large), and the small cuff may have an indicator mark (not shown) on the strap 1356 that should be visible after insertion through the loop 1358. If a small cuff is selected and the indicator mark does not pass through the loop, the small cuff may be too small and should be replaced with a large cuff.

[0108] A strain relief loop (L) in the STL 1300 is then created by arranging approximately 6 cm of the STL sigmoid body 1370 in a C-shape inside a small subcutaneous pocket formed via the neck incision (A) by blunt dissection superficially along the lateral surface of the diaphragm muscle in a posterior direction.

[0109] The surgeon then verifies that the cuff 1350 is not pulling or twisting the nerve, and that there is contact between the inside of the cuff 1350 and the nerve.

[0110] A test-stimulation is then performed to confirm correct positioning of the cuff 1350 on the nerve. To conduct a test-stimulation, the proximal end of STL 1300 is plugged into the INS 1100 and the programmer system 2100 is used to initiate a test stimulation signal delivered

from the INS 1100 to the nerve via the STL 1300. The test stimulation is performed while observing, for example, tongue movement by direct visual observation, airway caliber by nasal endoscopy, fluoroscopy, cephalogram, etc. Correct placement of the cuff on the nerve may be confirmed by, for example, observing tongue protrusion, an increase in retro-glossal airway caliber, an increase in retro-palatal airway caliber, an increase in stiffness of the anterior and/or lateral walls of the retro-glossal airway with or without an increase in airway caliber, anterior movement without superior movement of the hyoid bone, among others. Incorrect placement of the cuff on the nerve is indicated, for example, when there is insufficient opening of the retro-palatal or retro-lingual space, when the tongue is observed to retract (posterior movement), a decrease in retro-glossal airway caliber, a decrease in retro-palatal airway caliber, superior movement and particularly unilateral superior movement of the hyoid bone, among others. If necessary, the cuff 1350 may be repositioned at a different location along the length of the nerve to obtain the desired effect. The capture threshold and impedance values are recorded and the STL 1300 is disconnected from the INS 1100. The surgeon may create a fascia wrap by suturing fascia around the cuff on the superficial side of the nerve.

[0111] A pocket for the INS 1100 is then created by making an incision (B) down to the pectoralis fascia approximately 2 finger breadths below the right clavicle. The INS 1100 is preferably implanted on the right side to minimize cardiac artifact during respiratory sensing. Blunt dissection inferior to the incision is used to create a pocket large enough to hold the INS 1100. The pocket should be inferior to the incision (B) such that the incision (B) does not reside over the INS 1100 when later placed in the pocket.

[0112] A tunnel is formed for the STL 1300 using the tunneling system 3000 (sheath 3020 and tip 3030 placed over tool 3010) to tunnel along a path (C) from the infraclavicular INS pocket to the neck incision (A). As shown in Figure 11C, the lead carrier 3100 is then placed on the most proximal electrical contact of the STL proximal connector 1310. The tip 3030 is removed from the sheath 3020 to expose the connector 3018 of the tool 3010 and attach to the lead carrier 3100. While holding the sheath 3020 in place, the tool 3010 is pulled proximally to pull back the STL 1300 through the sheath 3020, taking care not to pull out the C-shaped strain relief or disturb the cuff. If the C-shaped strain relief loop (L) is pulled out, it should be replaced into the small pocket. The tool 3010 is released from the lead carrier 3100 and the lead carrier 3100 is removed from the STL 1300. The sheath 3020 is then removed from the body leaving the STL 1300 in place. The neck incision (A) need not be closed at this time, but rather may be closed later in the procedure allowing confirmation that the C-shaped strain relief remains in the small pocket.

[0113] The following implant instructions refer to an INS 1100 implanted at the patient's right sub-clavicular

region. The right and left distal portions of the RSL 1200 are placed near the right and left costal margins, respectively, by making four small incisions (D and E) as shown. The lateral incisions (E) may be made approximately 80% (+/- 5%) of the distance from the midline to the mid-axillary line, and on the costal margin. The medial incisions (D) may be made such that the RSL 1200 is relaxed and all electrodes are on the costal margin. Using the tunneling system 3000 (sheath 3020, tip 3030 attached via connector 3018), a tunnel (G) is formed between the pocket (B) and the medial incision (D), such that the right distal portion of the RSL 1200 may be pulled through the tunnel (G) from the pocket (B) to the medial incision (D). A tunnel (F) is then formed between the medial incision (D) and lateral incision (E), such that the right distal portion of the RSL may be pulled through the tunnel (F) from (D) to (E). This is repeated for the left distal portion of the RSL 1200. Alternatively, if the embodiments of RSL 1200 shown in Figures 4F or 4G are used, a small pocket may be formed medial to (D) in order to accommodate the medial electrodes and/or loop back region 1255.

[0114] The previously described tunneling operations (F and G) may be performed as follows: The tunneling tool 3010 including the connector 3018 is inserted into the sheath 3020 and the tip 3030 is connected to the connector 3018, forming the tunneling system 3000. The tunneling system is placed in the origination incision site and pushed beneath the skin towards the destination incision site, forming a tunnel. After tunneling, the tip 3030 is removed from the connector 3018 of the tunneling tool 3010. If needed, the tool 3010 may be removed and reversed such that the connector 3018 is at the other end of the sheath 3020. With the tool 3010 inserted through the sheath 3020, the lead carrier 3100 is attached on its proximal end to the connector 3018 and on its distal end to the distal electrode 1250 or proximal connector 1210 of the RSL 1200. While holding the sheath 3020 in place, the tunneling tool handle 3016 is pulled and the attached lead carrier 3100 and RSL 1200 are pulled into the sheath 3020. This may be visualized through the semi-transparent tunneling tool. The sheath 3020 may then be slid towards the tunneling tool handle 3016, exposing the lead carrier. The lead carrier may then be disconnected from the connector 3018, leaving the RSL 1200 in place. This process may be used to tunnel from (D) to (B) and subsequently from (E) to (D). For tunnel (G), the RSL 1200 may be pulled completely through the sheath 3020 to expose and disconnect the lead carrier.

[0115] Each anchor tab 1270 and suture hole 1290 is secured to the underlying tissue by dissecting down to the adjacent muscle fascia and suturing each anchor tab 1270 or suture hole 1290 to the muscle fascia. Permanent sutures are recommended to avoid movement of the RSL 1200 and braided suture material is recommended for knot retention and to prevent corrosion through the silicone anchors.

[0116] The STL 1300 and RSL 1200 are then connected to the INS 1100. The RSL 1200 is plugged into the

RSL port 1112 and the STL 1200 is plugged into the STL port 1114. The set screws are tightened using a torque wrench.

[0117] A closed loop test may be performed to confirm proper operation by observation of tongue protrusion or airway opening in concert with inspiration. The INS 1100 and proximal portions of the leads 1200/1300 are then placed into the infraclavicular pocket, looping the excess lead length beneath or around the INS 1100. Care should be taken not to pull out the C-shaped strain relief loop (L) in the STL sigmoid lead body 1370 while manipulating the INS 1100 into place. The INS 1100 is then sutured to underlying fascia through both suture holes 1116 found in the header 1110 of the INS 1100. Permanent sutures are recommended to avoid movement of the INS both before tissue encapsulation and chronically, and braided suture material is recommended for knot retention. Another system test may be performed at this point. After confirming that the C-shaped strain relief loop (L) is present in small pocket at neck incision, the incisions may be irrigated (optionally with an antibiotic solution) and closed using conventional techniques. After a healing period of approximately one month, the patient may undergo a sleep study to confirm proper operation of the system and to titrate therapy.

Screening Methods

[0118] Prior to implant, patients may be screened to estimate the probability of a successful outcome.

[0119] The airway can be characterized during sleep by a Pcrit measurement in differing sleep stages and body postures, per the methods of Schwartz et al. A higher Pcrit value is indicative of a more collapsible airway which may be more difficult to treat with this therapy. A surrogate measure for Pcrit may be an auto-PAP device that adjusts airway pressure dynamically to eliminate flow limitation. For example, a patient may require 12cmH₂O of air pressure to maintain a patent airway in the supine position, yet only require 8cmH₂O of air pressure in the lateral position. The auto-PAP would automatically adjust for this.

[0120] The volume of air expired during a pressure drop may be measured. The pressure drop may occur during natural expiration during wakefulness. Alternatively, the patient may be asleep during the measurement. The timing of the pressure drop during expiration may occur at a certain point during expiration to ensure consistency. The duration of the pressure drop may be fixed. An example of this measure is $V_{NEP_0.5}$.

[0121] The airway may be visualized using an imaging modality such as, but not limited to, cephalogram, MRI, fMRI, CT, ultrasound, OCT, naso-endoscopy, photography, and video imaging. This imaging may be performed during sleep, under sedation, or during wakefulness. The patient may be asked to protrude the tongue, inhale/exhale at specific flow rates, or perform Muller's Maneuver. Tongue protrusion force may also be measured. Tongue

size may be observed and/or measured quantitatively or qualitatively (e.g., Modified Mallampati). BMI may also be a good predictor of patient response.

[0122] The following metrics (and others) may be measured and used in screening: size of tongue and soft palate, angle of the soft palate, redundancy of tissue, and length of soft palate.. Endoscopy is one method for obtaining these metrics. Additional size metrics include craniofacial structures, tonsil size, adenoids, pharyngeal fat pads.

[0123] Mechanical linkage or coupling between airway structures may also be assessed. For example, airway opening may be measured at different levels concurrent with other motions, (e.g. measuring opening of the airway at the retro-palatal space during voluntary tongue protrusion or anterior displacement of the tongue).

[0124] Nasal airway collapse may be measured using nasal peak inspiratory flow meters in different body positions. Additionally, acoustic rhinometry may provide another way to measure this.

[0125] Body Mass Index (BMI) may be a useful tool in screening. Additional metrics include % body fat, % visceral fat, neck circumference, % neck fat, and body fat distribution.

[0126] A patient's arousal threshold from sleep may be quantified by measuring intra-pleural arousal pressure. A nasal EPAP device may be used in screening. An EPAP device reduces airflow through the nares. This may increase airway patency during the expiratory phase of respiration. An example of EPAP is the ProVent device (Ventus Medical Inc., Belmont CA). Arousals and respiratory events may be assessed with and without the EPAP device. During therapy, the patient's tongue may protrude past the teeth. A dental examination (i.e. identify sharp teeth), patient's use of dentures, and tolerance to oral appliances may be used in screening.

[0127] The airway may be characterized using a dual air pump, and valve system, configured for connection to a mask on the patient. In this configuration, the two different pressures (e.g., difference of 1cm H₂O) are maintained by each pump which is connected to the valve. The valve may be attached via a tube to the mask such that the pressure at the mask is from only one of the pumps. The valve may then be automated to alternate between the two pressures at a programmable rate (e.g., 1 Hz). This allows the airway to fluctuate between pressures within breaths. An airway may be characterized by lowering the pressure to a level that brings flow limitation, and then observing what pressures remove this flow limitation.

Awake Titration

[0128] As described previously, the patient may undergo an awake titration 10200, an iterative process where the response to stimulation is documented over a range of comfortable stimulation levels (Figure 12B). These stimulations may be delivered manually (e.g., 2 second

commanded stimulation bursts) or synchronous with respiration. A range of amplitudes may be tested across multiple frequencies (range of 20 to 50Hz, nominal 40 Hz), and pulse widths (range of 30 to 215 μ s, nominal 90 μ s) 10220.

[0129] The awake titration may involve defining a wake-stimulation operating window, defined at its lower limit by a capture threshold and at its upper limit by a discomfort threshold. The capture threshold may be defined as the lowest stimulation level at which muscle contraction is visible, palpable, or perceptible (e.g., gross tongue movement or stiffening) is observed. The discomfort threshold may be defined as the lowest stimulation level at which the patient experiences an unacceptable sensation (e.g., discomfort, pain) while awake.

[0130] While determining this range, the patient may be in the supine position or alternatively, in a posture typical of sleep. In general, during the stimulation titration, it is preferable to begin with the lowest settings for pulse width (30 μ s) and amplitude (0.4mA) at a nominal frequency (40Hz). If stimulation produces pulsatile (vibrating) contractions, the frequency may be increased to 50Hz. The pulse width is incrementally increased to 60 μ s, then to nominal (90 μ s), keeping Amplitude at 0.4mA. With the pulse width set to 90 μ s, amplitude may be iterated according to the process described hereinafter. If maximum amplitude is reached and additional intensity is required, the pulse width may be increased while reducing amplitude to minimum (0.4mA). If maximum pulse width (215 μ s) is reached and additional intensity is required, frequency may be increased while reducing the pulse width to 90 μ s and the amplitude to minimum (0.4mA).

[0131] At each stimulation level, observations may be recorded such as: visible tongue motion, palpable genioglossus muscle contraction, perception of muscle movement, tongue protrusion, tongue retrusion, tongue depression, tongue flattening, tongue cupping, and tongue protrusion past the teeth 10230.

[0132] After awake titration 10200, the patient may be sent home at a stimulation level in this operating range, beginning the daytime familiarization period 10400. This may occur prior to the sleep titration night, such that the patient may acclimate to the sensation of stimulation. This may allow higher levels of stimulation to be assessed during the sleep titration without patient arousal. Patients have been observed to tolerate (i.e., not arouse) higher stimulation intensities while asleep compared to wakefulness, so the arousal threshold may be higher than the wake discomfort threshold.

Sleep Titration

[0133] As described previously, after implantation and a healing period of approximately one month, the patient may undergo a sleep (PSG) study to confirm proper operation of the system and to titrate therapy stimulation levels 10500. Titration may utilize the set-up illustrated

in Figure 10, wherein the programmer system 2100 interfaces with the PSG equipment 2800. Generally, it is preferable to use an oro-nasal mask in order to measure airflow. Alternatively, a nasal cannula may be used. The preferred configuration is a calibrated pneumotach with the oro-nasal mask to measure the patient's airflow 10510. Alternatively, an uncalibrated pneumotach may be used. A thermistor or a thermocouple may also be used to sense airflow. The thermistor or thermocouple may be calibrated or uncalibrated.

[0134] Figure 14 illustrates an example of a patient's airflow response to stimulation. As shown in Figure 14, the inspired airflow increases as stimulation delivered to the nerve increases. The airflow capture threshold is the stimulation level at which an increase in airflow is first observed. As stimulation continues to increase, muscle (i.e. genioglossus) activation and airflow also increase. Prior to full muscle activation, a stimulation level which first removes respiratory events is observed. At full muscle activation, increasing stimulation does not increase airflow, resulting in an airflow plateau. The patient may arouse due to stimulation at a level on the plateau. Data comprising this curve may be acquired during a sleep titration PSG for each patient.

[0135] The sleep titration process is illustrated schematically in Figure 12C. Titration generally involves establishing an effective range of stimulation settings 10540, where the lower end is defined by the lowest stimulation level where respiratory events (e.g., apneas, hypopneas, oxygen desaturations, etc.) 10550 begin to decrease or airflow begins to increase, and the range's upper end is defined by stimulation that arouses the patient. A goal setting at which the patient is effectively treated (i.e. respiratory events removed) may be estimated.

[0136] During a sleep titration, device settings may be programmed using the programmer system 2100. At the start of titration, most stimulation settings will be at their default values (40 Hz frequency, 90 μ s pulse width, 50% stimulation duty cycle, 0ms phase adjust, etc.). Titration typically consists of many series of "ramps" in different body positions and sleep stages, wherein each ramp is a series of intervals where the stimulation intensity is increased from the previous interval in a certain stimulation mode 10540, illustrated in Figure 13. For example, at 40Hz, 90 μ s, in A0A0 mode, stimulation level of the "A" breaths could be incremented every two minutes, from the nerve capture amplitude to the amplitude that causes arousal, as illustrated in Figure 13 trace 3. Ramps are typically performed in A0A0, ABAB, AABB, A0B0, or AAAA modes, as illustrated in Figure 13, traces 1, 2, 3, and 5. In addition, ramps may be performed using any of the previously discussed modes, also shown in Figure 13. At each interval, observations are made as to whether stimulation causes arousal, reduces respiratory events, or increases airflow (e.g., increased $V_{i,max}$).

[0137] Alternative stimulation modes (not illustrated) may be utilized during wakefulness or sleep, (e.g. during a sleep titration PSG). An alternative mode may be xAy0

mode, where stimulation is delivered for "x" breaths wherein "x" is a programmable number of breaths, followed by "y" breaths with no stimulation wherein "y" is a programmable number of breaths. For example, in mode 4A40, stimulation is delivered on four consecutive breaths followed by no stimulation on four consecutive breaths.

[0138] A0A0 mode may be useful in determining if a stimulated breath has more flow than an adjacent unstimulated breath. In the same manner, ABAB and AABB modes may be useful in determining if a stimulated "A" breath provides more flow compared to an adjacent breath with less stimulation, "B." Likewise, A0B0 mode may be useful in comparing flows during "A" stimulation, "B" stimulations, and unstimulated breaths. During a ABAB, AABB, and A0B0 ramps, the difference in stimulation between "A" and "B" stimulations may be constant. ABAB, AABB, and AAAA ramps may also be useful in determining absolute flow (e.g., tidal volume, minute volume, etc.) and determining what level of stimulation reduces or eliminates respiratory events, since stimulation is delivered every breath. These observations may be recorded for future reference (e.g. physician use). Each stimulation mode may be combined with any of the pulse configurations, such as nested stimulation, soft start, or retention intensity, as chosen by physician or technician.

[0139] When a stimulation level is estimated to be efficacious, it may be tested in AAAA mode for a fixed time (e.g., five minutes) after which stimulation may be turned off for a fixed time (e.g., five minutes). If stimulation noticeably reduced or eliminated respiratory events compared to no stimulation, the settings may be considered the goal setting.

[0140] Additional titration PSG studies may be performed, either as a separate sleep study or as a split night study. In order to compare multiple stimulation settings during sleep, a cross-over PSG study may be performed wherein the stimulation settings are changed at fixed intervals throughout the night, toggling through a select group of settings. For example, stimulation may alternate between two stimulation settings every five minutes. Afterwards, respiratory events may be determined for each interval such that an index (e.g., AHI, ODI, etc.) can be calculated for each stimulation setting. This may be useful to gauge whether an increase in stimulation would provide additional therapeutic benefits or to gauge whether a decrease in stimulation would not lessen any therapeutic benefits. In addition, the oxygen sensor of the INS 1100 may be used to measure oxygen de-saturations and calculate an ODI.

[0141] Another type of titration PSG may be a characterization night, per the methods of Schwartz et al, such that a Pcrit may be determined in REM/nREM sleep, both supine and lateral. Another type of sleep study is the home PSG study, which may be utilized to assess efficacy without the burden of an in-lab PSG. Many home PSG systems are available for use.

[0142] The patient may also undergo a vector-sweep-

ing sleep study, wherein the programming system 2100 and a respiratory signal (e.g., nasal cannula) are utilized, 10520. During this study, the secondary vector may be changed at regular intervals to cycle through a select group of vectors. These vectors are compared to the primary vector to determine an optimal sensing vector to deliver therapy. Selection may be based on maximum signal strength, consistent correlation to respiratory fiducials (e.g., offset of inspiration), and maximum signal stability/reliability across sleep stages, body positions, and disordered breathing events, for example. A stable signal has a minimum probability of signal inversion. A reliable signal has a minimum probability of signal loss, and may preferably have a minimum threshold of 0.1 to 0.5 Ohms peak-to-peak, for example. The optimal vector may be selected by incrementally scrolling through all or a preferred subset of possible vectors while sampling the respiration signal and comparing the signal against themselves or predefined thresholds. This scrolling technique may be performed manually (with inputs via the programmer system) or automatically (i.e., programmed). The sampling technique may also be performed manually (visual observation using programmer system) or automatically (i.e., programmed).

Post-Sleep-Titration: Nighttime familiarization and Up-Titration

[0143] After titration, the patient is typically sent home with the device active and stimulation set to a level at which she can fall asleep. This begins the post-sleep-titration process, as illustrated in Figure 12D. This process may include nighttime familiarization, up-titration of stimulation, and determining efficacy 10700, 10800, and 11100.

[0144] The stimulation settings at which the patient is sent home may initially be below the estimated goal settings. As the patient becomes acclimated to sleeping with the device on 10700, stimulation may be slowly incremented, over the course of days, towards the range where therapeutic effects were observed during the sleep titration and ultimately towards the estimated goal settings. This period is known as up-titration 10800. These increments may be performed by a physician or caretaker, or by the patient, if the physician allows the patient to have limited control of stimulation via the therapy controller 2500.

[0145] Several feedback parameters may aid in determining the appropriate time to up-titrate a patient 10810. These include the device's therapy history of frequency and duration of therapy sessions, patient feedback (e.g., more daytime energy, no tongue abrasions, stimulation not causing arousal or pain, etc.), patient's bed-partner feedback (e.g., reduced snoring, perceives patient to be less sleepy, etc.), and most notably, a PSG study. Taken together, these may show whether the patient needs more time to acclimate to stimulation, is ready to have stimulation increased 10840, is receiving therapeutic

benefits, or is fully treated. In addition, these feedback may provide data to adjust the patient's estimated goal settings 10820. For example, a patient's upper airway may, over time, undergo muscle remodeling such that less stimulation than originally estimated provides efficacious therapy. Alternatively, a patient may gain weight such that more stimulation than originally estimated may be needed to provide efficacious therapy.

[0146] If the stimulation is causing discomfort to the patient such that stimulation disrupts sleep or inhibits falling asleep with stimulation on, many options are available, 10830. Stimulation may be reduced and the patient may be given more time for nighttime familiarization. A different strategy that may increase the therapy provided to the patient is to utilize the device features of core hours, soft start, retention intensity, and nested stimulation. The core hours feature (Figure 6K) allows the patient to fall asleep at one level of stimulation and after a programmable time interval, have stimulation increased to a more therapeutic level. Patients are often able to tolerate higher stimulations when asleep compared to wakefulness and in addition, tolerate higher stimulations further into a therapy session than at the start of a session.

[0147] As mentioned previously, the INS 1100 may be programmed to change stimulation level between therapy sessions, days, or other programmable value, enabling comparisons between stimulation level and therapy session data. For example, the INS 1100 may be programmed to alternate between 1.8mA and 2.0mA, where the change occurs between therapy sessions. This may be used as a diagnostic mode to assess the incremental benefit of the higher stimulation level. A patient receiving daily therapy would thus receive therapy at 1.8mA on one day, 2.0mA the next, 1.8mA the next day, 2.0mA the next day, and so on. This may allow therapy session data to be compared with stimulation level. For example, a physician may compare the cycling rate (via the cycling detector) during therapy sessions at 1.8mA and 2.0mA over time to determine if there is an observable difference. If cycling is reduced at 2.0mA, the physician may increase stimulation and re-test. If there was no observable difference, the physician may decrease stimulation and re-test or simply set stimulation to the lower level. This may reduce the likelihood of delivering stimulation at a level that, compared to a lower level, provides no additional flow or no additional benefit. As mentioned previously, other therapy session data may be compared to stimulation level as well in a similar manner. Examples of other therapy session data are: oxygen desaturation frequency and severity, stimulation time, variations in respiratory rate, and variations in respiratory prediction.

[0148] Various pulse configurations may also help a patient acclimate to stimulation. The soft start stimulations (Figure 6G) may provide a smoother transition from unstimulated breaths to stimulated breaths, reducing the patient's perception of stimulation intensity. Retention intensity (Figures 6E and 6F) may also provide therapy with reduced patient perception of stimulation intensity,

since the full amplitude is only used for a part of the stimulation. Nested stimulations (Figure 6H) may be used in a similar manner.

[0149] The patient may use positive airway pressure (PAP) therapy (e.g., CPAP, bi-PAP, auto-PAP, etc.) in conjunction with the neurostimulator. This may allow the patient to receive therapeutic benefits in addition to what is provided by the stimulation. The pressure necessary to provide these benefits may decrease as stimulation is up-titrated, 10800. This progress towards lower pressures may be monitored using the auto-PAP technology which automatically adjusts pressure to the level necessary to remove flow limitation. In time, the patient may wish to stop PAP therapy altogether. In a similar manner, other therapies such as but not limited to positional therapy and mandibular advancement may be used in conjunction with the neurostimulator.

[0150] If the patient begins noticing tongue abrasions, a tooth guard or other tooth covering (e.g. dental wax) may be used such that the tongue does not scrape against the teeth when stimulation causes tongue protrusion. Tooth guards may be custom-made (e.g., by a dentist).

[0151] Additional strategies for acclimation may include another sleep titration night to examine different stimulation frequencies, pulse widths, and modes, as previously described. Different stimulation frequencies and pulse widths may capture different muscle groups in a more therapeutic manner. In additions, respiration vectors may need to be assessed during a vector sweeping titration.

[0152] Therapy efficacy may be measured using standard PSG techniques during and after familiarization and up-titration. Therapy efficacy may be evaluated by assessing indicia of sleep disordered breathing such as AHI, apnea index, hypopnea index, respiratory disturbance index, apnea-hypopnea index, ODI, FOSQ, ESS, BDI, PSQI, or other measures.

Alternative Embodiments

[0153] The stimulation may be delivered to the nerve by utilizing a variety of stimulation electrode configurations, in addition to the configurations previously mentioned.

[0154] Mono-polar stimulation may be delivered to the nerve wherein the cathode (or multiple cathodes) is an electrode (or multiple electrodes) in the STL nerve cuff 1350, and wherein the anode is the INS 1100. Similarly, far-field bi-polar stimulation may be delivered to the nerve wherein the cathode (or multiple cathodes) is an electrode (or multiple electrodes) in the nerve cuff, and wherein the anode is an RSL electrode 1250 or 1260.

[0155] Any combination of bi-polar and mono-polar stimulation may be utilized to deliver stimulation to the nerve. For example, mono-polar stimulation may be delivered between a cathode electrode in the STL nerve cuff 1350 and the INS 1100 anode. Simultaneously, far-

field bi-polar stimulation may be delivered between a different cathode electrode in the STL nerve cuff 1350 and an RSL electrode 1250 or 1260 anode.

[0156] The INS 1100 may be programmed to periodically change stimulation parameters throughout a therapy session to vary which muscle fibers are recruited at any given time. For example, stimulation may be delivered at an initial lower frequency (e.g. 30Hz) for 5 minutes, followed by stimulation delivered at a higher frequency (e.g. 50Hz) for 2 minutes. Each frequency may have a unique pulse width, pulse width, and/or amplitude. This sequence could be repeated throughout the night.

[0157] This may allow certain muscle fibers to rest during periods when other muscle fibers are active. This may reduce muscle fatigue. Another potential advantage is that stimulation at more than a single frequency and/or pulse width may be a more effective means of building muscle strength and endurance. Another potential advantage is that different stimulation settings (e.g. frequency, pulse width, and/or amplitude) may result in slightly different movement of the tongue. Varying stimulation settings may decrease the possibility of repetition-induced irritation, inflammation or injury.

[0158] Alternatively, it may be possible to effectively build muscle strength and endurance by deactivating a portion of the nerve fibers and activating a remaining subset of fibers. This may allow delivery of higher levels of stimulation to the remaining fibers. This may reduce subject discomfort which could have occurred if all the fibers had been activated at that same intensity. Additionally, this may increase airway patency or opening by selecting muscles whose activation results in tongue protrusion and deactivating tongue retrusion muscles.

[0159] Certain fibers in the nerve may be selectively deactivated by choosing the cathode of pulse delivery and sequence of pulse delivery such that the fibers are not recruited by subsequent or simultaneous pulses. For example, a nerve fiber that innervates a retrusor muscle may be deactivated by delivering a sub-threshold pulse from a nearby cathode. This may allow a subsequent or simultaneous pulse at a different cathode to activate a nerve protrusor muscle nerve fiber without recruiting the retrusor muscle.

[0160] In particular, the following embodiments are provided.

[0161] A device for stimulating a hypoglossal nerve of a person, comprising:

a neurostimulator for generating an electrical signal; and
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve,
wherein the electrical signal comprises a first stimulation signal generated when the person is awake followed by a second stimulation signal generated when the person is asleep, the first stimulation signal

having a first intensity that is within a range having a lower bound and an upper bound, the upper bound being a discomfort threshold, the second stimulation signal having a second intensity that is greater than the discomfort threshold.

[0162] A device for stimulating a hypoglossal nerve of a person, comprising:

a neurostimulator for generating an electrical signal; and
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve,
wherein the electrical signal comprises a first stimulation signal generated when the person is awake followed by a second stimulation signal generated when the person is asleep, the first stimulation signal having a first intensity that is within a wake-stimulation operating window defined by a lower limit corresponding to a capture threshold and an upper limit corresponding to a discomfort threshold, the second stimulation signal having a second intensity that is greater than the discomfort threshold.

[0163] A device for stimulating a hypoglossal nerve of a person, comprising:

a neurostimulator for generating an electrical signal; and
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve,
wherein the electrical signal comprises a first stimulation signal generated during an awake condition of the person followed by a second stimulation signal generated during a sleep condition of the person, the first stimulation signal having a first intensity that is less than a discomfort threshold, the second stimulation signal having a second intensity that is greater than the first intensity and that is between the discomfort threshold and an arousal threshold, the arousal threshold defined by a third intensity that is greater than the second intensity and at which the person awakes from the sleep condition.

[0164] A device for stimulating a hypoglossal nerve of a person, comprising:

a neurostimulator for generating an electrical signal; and
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve,
wherein the neurostimulator generates the electrical

signal in a first mode followed by a second mode, the first mode comprising a first stimulation signal having a first intensity that is within a range having a lower bound and an upper bound, the upper bound being a discomfort threshold, the second mode comprising a second stimulation signal having a second intensity that is greater than the discomfort threshold.

[0165] A device for stimulating a hypoglossal nerve of a person, comprising:

a neurostimulator for generating an electrical signal; and
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve,
wherein the neurostimulator generates the electrical signal in a first mode followed by a second mode, the first mode comprising a first stimulation signal having a first intensity that is within a wake-stimulation operating window defined by a lower limit corresponding to a capture threshold and an upper limit corresponding to a discomfort threshold, the second mode comprising a second stimulation signal having a second intensity that is greater than the discomfort threshold.

[0166] A device for stimulating a hypoglossal nerve of a person, comprising:

a neurostimulator for generating an electrical signal; and
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve,
wherein the neurostimulator generates the electrical signal in a first mode followed by a second mode, the first mode comprising a first stimulation signal having a first intensity that is less than a discomfort threshold, the second mode comprising a second stimulation signal having a second intensity that is greater than the first intensity and that is between the discomfort threshold and an arousal threshold, the arousal threshold defined by a third intensity that is greater than the second intensity.

[0167] In all of these embodiments one of the following features may be implemented

[0168] The electrical signal transitions from the first stimulation signal to the second stimulation signal when a fixed duration of time expires. The electrical signal transitions from the first stimulation signal to the second stimulation signal when a preset number of breaths are counted. The electrical signal transitions from the first stimulation signal to the second stimulation signal when the neurostimulator generates a preset number of electrical

signals. The electrical signal transitions from the first stimulation signal to the second stimulation signal when the neurostimulator generates a preset number of electrical signals. The electrical signal transitions from the first stimulation signal to the second stimulation signal when a sleep state is detected. The electrical signal transitions from the first stimulation signal to the second stimulation signal when an activity level is detected.

[0169] Moreover it is provided:

10 A device for stimulating a hypoglossal nerve, comprising:

a neurostimulator for generating an electrical signal and measuring a bio-impedance signal;
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve;
a respiration sensing lead having a proximal portion with a proximal end connected to the neurostimulator and a distal end terminating in a bifurcation with first and second distal portions extending in different directions therefrom, wherein the first and second distal portions are configured to extend over opposite sides of the thorax.

The first distal portion may be configured to extend over the ipsi-lateral side of the thorax, and the second distal portion may be configured to extend over the contra-lateral side of the thorax. The first distal portion may be configured to extend over the ipsi-lateral costal margin, and the second distal portion may be configured to extend over the contra-lateral costal margin. At least one of the first and second distal portions may include a sensing electrode, wherein the bio-impedance signal may be delivered via the sensing electrode to the neurostimulator. 5. At least one of the first and second distal portions may include a current injection electrode. The proximal portion and the distal portions of the respiration sensing lead may comprise tubes with electrical conductors extending therethrough. The bifurcation may include a yoke connecting the proximal portion with the distal portions of the respiration sensing lead.

[0170] A device for stimulating a hypoglossal nerve, comprising:

a neurostimulator for generating an electrical signal; a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve; and
wherein the electrical signal comprises a pulse train envelope of two or more pulses at two or more stimulation levels. Each pulse stimulation level may be greater than or equal to the previous pulse stimulation level in the pulse envelope. Each pulse stimulation level may be less than or equal to the previous pulse stimulation level in the pulse envelope. The pulse envelope may contain 2 series of pulses,

wherein the first series each pulse stimulation level is greater than or equal to the previous pulse stimulation and wherein the second series each pulse stimulation level is less than or equal to the previous stimulation. The stimulation level may be defined as stimulation amplitude.

[0171] A device for stimulating a hypoglossal nerve, comprising:

a neurostimulator for generating an electrical signal; a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve and configured for monitoring neural activity of the hypoglossal nerve; and wherein the device may change stimulation level based on the monitored neural activity.

[0172] A device for stimulating a hypoglossal nerve, comprising:

an oxygen sensor;
a neurostimulator for generating an electrical signal; a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve; and
wherein the device may change stimulation level based on oxygen sensor reading.

[0173] A device for stimulating a hypoglossal nerve, comprising:

an accelerometer;
a neurostimulator for generating an electrical signal; a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve; and
wherein the device may change stimulation level based on the accelerometer reading.

Claims

1. A device for stimulating a hypoglossal nerve of a person, comprising:

a neurostimulator for generating an electrical signal; and
a stimulation lead connected to the neurostimulator, the stimulation lead having a nerve cuff electrode configured for delivering the electrical signal to the hypoglossal nerve,
characterized in that
the electrical signal comprises a first stimulation signal generated when the person is awake fol-

lowed by a second stimulation signal generated when the person is asleep,
the first stimulation signal having a first intensity that is within a range having a lower bound and an upper bound, the upper bound being a wake discomfort threshold, within a wake-stimulation operating window obtained by awake titration to which the person was subjected defined by a lower limit corresponding to a capture threshold and an upper limit corresponding to a wake discomfort threshold, the second stimulation signal having a second intensity that is greater than the wake discomfort threshold.

2. The device of claim 1 wherein the electrical signal transitions from the first stimulation signal to the second stimulation signal when a fixed duration of time expires.

3. The device of claim 1 wherein the electrical signal transitions from the first stimulation signal to the second stimulation signal when a preset number of breaths are counted.

4. The device of claim 1 wherein the electrical signal transitions from the first stimulation signal to the second stimulation signal when the neurostimulator generates a preset number of electrical signals.

5. The device of claim 1 wherein the electrical signal transitions from the first stimulation signal to the second stimulation signal when the neurostimulator generates a preset number of electrical signals.

6. The device of claim 1 wherein the electrical signal transitions from the first stimulation signal to the second stimulation signal when a sleep state is detected.

7. The device of claim 1 wherein the electrical signal transitions from the first stimulation signal to the second stimulation signal when an activity level is detected.

8. The device according to one of the preceding claims wherein
the first stimulation signal is generated during an awake condition of the person followed by a second stimulation signal generated during a sleep condition of the person, the first stimulation signal has a first intensity that is less than a wake discomfort threshold, the second stimulation signal has a second intensity that is greater than the first intensity and that is between the wake discomfort threshold and an arousal threshold, the arousal threshold defined by a third intensity that is greater than the second intensity and at which the person awakes from the sleep condition.

9. The device according to one of the preceding claims wherein the neurostimulator is configured to generate the electrical signal in a first mode followed by a second mode, the first mode comprising the first stimulation signal having the first intensity and the second mode comprising the second stimulation signal having the second intensity.
10. The device according to one of the claims 1 to 8, wherein the neurostimulator is configured to generate the electrical signal in a first mode followed by a second mode, the first mode comprising the first stimulation signal having the first intensity that is less than the wake discomfort threshold, the second mode comprising the second stimulation signal having the second intensity that is greater than the first intensity and that is between the wake discomfort threshold and an arousal threshold, the arousal threshold defined by a third intensity that is greater than the second intensity.

Patentansprüche

1. Eine Vorrichtung zum Stimulieren eines hypoglossalen Nervs einer Person, umfassend:

einen Neurostimulator zum Erzeugen eines elektrischen Signals; und
 ein Stimulationskabel, das mit dem Neurostimulator verbunden ist, wobei das Stimulationskabel eine Nervmanschetten-Elektrode aufweist, die dazu ausgebildet ist, das elektrische Signal an den hypoglossalen Nerv zu liefern,
dadurch gekennzeichnet, dass
 das elektrische Signal ein erstes Stimulationssignal umfasst, das erzeugt wird, wenn die Person wach ist, dem ein zweites Stimulationssignal folgt, das erzeugt wird, wenn die Person schläft,
 wobei das erste Stimulationssignal eine erste Intensität aufweist, die in einem Bereich mit einer unteren Grenze und einer oberen Grenze liegt, wobei die obere Grenze eine Unannehmlichkeitsschwelle im Wachzustand ist, innerhalb eines Stimulationsbetriebsfensters im Wachzustand, das durch eine Dosierung im Wachzustand erhalten wird, der die Person ausgesetzt wurde und die durch eine untere Schranke entsprechend einer Erfassungsschwelle und eine obere Schranke entsprechend einer Unannehmlichkeitsschwelle im Wachzustand definiert ist, und wobei das zweite Stimulationssignal eine zweite Intensität aufweist, die größer als die Unannehmlichkeitsschwelle im Wachzustand ist.

2. Die Vorrichtung von Anspruch 1, in der das elektri-

sche Signal von dem ersten Stimulationssignal zu dem zweiten Stimulationssignal übergeht, wenn eine vorbestimmte Zeitdauer vergangen ist.

3. Die Vorrichtung von Anspruch 1, in der das elektrische Signal von dem ersten Stimulationssignal zu dem zweiten Stimulationssignal übergeht, wenn eine vorbestimmte Anzahl an Atmungsvorgängen gezählt wird.
4. Die Vorrichtung von Anspruch 1, in der das elektrische Signal von dem ersten Stimulationssignal zu dem zweiten Stimulationssignal übergeht, wenn der Neurostimulator eine vorbestimmte Anzahl an elektrischen Signalen erzeugt.
5. Die Vorrichtung von Anspruch 1, in der das elektrische Signal von dem ersten Stimulationssignal zu dem zweiten Stimulationssignal übergeht, wenn der Neurostimulator eine vorbestimmte Anzahl an elektrischen Signalen erzeugt.
6. Die Vorrichtung von Anspruch 1, in der das elektrische Signal von dem ersten Stimulationssignal zu dem zweiten Stimulationssignal übergeht, wenn ein Schlafzustand detektiert wird.
7. Die Vorrichtung von Anspruch 1, in der das elektrische Signal von dem ersten Stimulationssignal zu dem zweiten Stimulationssignal übergeht, wenn ein Aktivitätsniveau detektiert wird.
8. Die Vorrichtung gemäß einem der vorhergehenden Ansprüche, in der
 das erste Stimulationssignal während eines Wachzustands der Person erzeugt wird und von dem zweiten Stimulationssignal gefolgt wird, das während eines Schlafzustands der Person erzeugt wird, wobei das erste Stimulationssignal eine erste Intensität aufweist, die niedriger als eine Unannehmlichkeitsschwelle im Wachzustand ist und wobei das zweite Stimulationssignal eine zweite Intensität aufweist, die größer als die erste Intensität ist und die zwischen der Unannehmlichkeitsschwelle im Wachzustand und einer Aufweckschwelle liegt, wobei die Aufweckschwelle durch eine dritte Intensität definiert ist, die größer als die zweite Intensität ist und bei welcher die Person aus dem Schlafzustand aufwacht.
9. Die Vorrichtung gemäß einem der vorhergehenden Ansprüche, in der der Neurostimulator dazu ausgebildet ist, das elektrische Signal in einem ersten Modus gefolgt von einem zweiten Modus zu erzeugen, wobei der erste Modus das erste Stimulationssignal mit der ersten Intensität und der zweite Modus das zweite Stimulationssignal mit der zweiten Intensität umfasst.

10. Die Vorrichtung gemäß einem der Ansprüche 1 bis 8, in der der Neurostimulator dazu ausgebildet ist, das elektrische Signal in einem ersten Modus gefolgt von einem zweiten Modus zu erzeugen, wobei der erste Modus das erste Stimulationssignal mit der ersten Intensität, die niedriger als die Unannehmlichkeitsschwelle im Wachzustand ist, und der zweite Modus das zweite Stimulationssignal mit der zweiten Intensität, die größer als die erste Intensität ist und die zwischen der Unannehmlichkeitsschwelle im Wachzustand und einer Aufweckschwelle liegt, umfasst, wobei die Aufweckschwelle durch eine dritte Intensität definiert ist, die größer als die zweite Intensität ist.

Revendications

1. Dispositif permettant de stimuler un nerf hypoglosse d'une personne, comprenant:

un neurostimulateur pour générer un signal électrique; et

un conducteur de stimulation connecté au neurostimulateur, le conducteur de stimulation ayant une électrode nerveuse à coiffe configurée pour délivrer le signal électrique au nerf hypoglosse,

caractérisé en ce que

le signal électrique comprend un premier signal de stimulation généré lorsque la personne est éveillée, suivi d'un deuxième signal de stimulation généré lorsque la personne est endormie, le premier signal de stimulation ayant une première intensité comprise dans une plage ayant une limite inférieure et une limite supérieure, la borne supérieure étant un seuil d'inconfort d'éveil, dans une fenêtre de stimulation d'éveil obtenue par titrage d'éveil auquel la personne a été soumise, définie par une limite inférieure correspondant à un seuil de capture et une limite supérieure correspondant à un seuil d'inconfort d'éveil, le deuxième signal de stimulation ayant une deuxième intensité supérieure au seuil d'inconfort d'éveil.

2. Dispositif selon la revendication 1, dans lequel le signal électrique passe du premier signal de stimulation au deuxième signal de stimulation lorsqu'une durée de temps fixe arrive à son terme.
3. Dispositif selon la revendication 1, dans lequel le signal électrique passe du premier signal de stimulation au deuxième signal de stimulation au bout d'un nombre prédéfini de respirations.
4. Dispositif selon la revendication 1, dans lequel le signal électrique passe du premier signal de stimulation

au deuxième signal de stimulation lorsque le neurostimulateur génère un nombre prédéfini de signaux électriques.

5. Dispositif selon la revendication 1, dans lequel le signal électrique passe du premier signal de stimulation au deuxième signal de stimulation lorsque le neurostimulateur génère un nombre prédéfini de signaux électriques.

6. Dispositif selon la revendication 1, dans lequel le signal électrique passe du premier signal de stimulation au deuxième signal de stimulation lorsque l'on détecte un état de sommeil.

7. Dispositif selon la revendication 1, dans lequel le signal électrique passe du premier signal de stimulation au deuxième signal de stimulation lorsque l'on détecte un niveau d'activité.

8. Dispositif selon l'une des revendications précédentes, dans lequel, le premier signal de stimulation est généré pendant un état éveillé de la personne suivi d'un deuxième signal de stimulation généré pendant une condition de sommeil de la personne, le premier signal de stimulation a une première intensité qui est inférieure à un seuil d'inconfort d'éveil, le deuxième signal de stimulation présente une deuxième intensité supérieure à la première intensité et est comprise entre le seuil d'inconfort d'éveil et un seuil de réveil, le seuil de réveil étant défini par une troisième intensité supérieure à la deuxième intensité et à laquelle la personne se réveille de la condition de sommeil.

9. Dispositif selon l'une des revendications précédentes, dans lequel le neurostimulateur est configuré pour générer le signal électrique dans un premier mode suivi d'un deuxième mode, le premier mode comprenant le premier signal de stimulation ayant la première intensité et le deuxième mode comprenant le deuxième signal de stimulation ayant la deuxième intensité.

10. Dispositif selon l'une des revendications 1 à 8, dans lequel le neurostimulateur est configuré pour générer le signal électrique dans un premier mode suivi d'un deuxième mode, le premier mode comprenant le premier signal de stimulation ayant la première intensité qui est inférieure au seuil d'inconfort d'éveil, le deuxième mode comprenant le deuxième signal de stimulation ayant la deuxième intensité qui est supérieure à la première intensité et qui est comprise entre le seuil d'inconfort d'éveil et le seuil de réveil, le seuil de réveil étant défini par une troisième intensité qui est supérieure à la deuxième intensité.

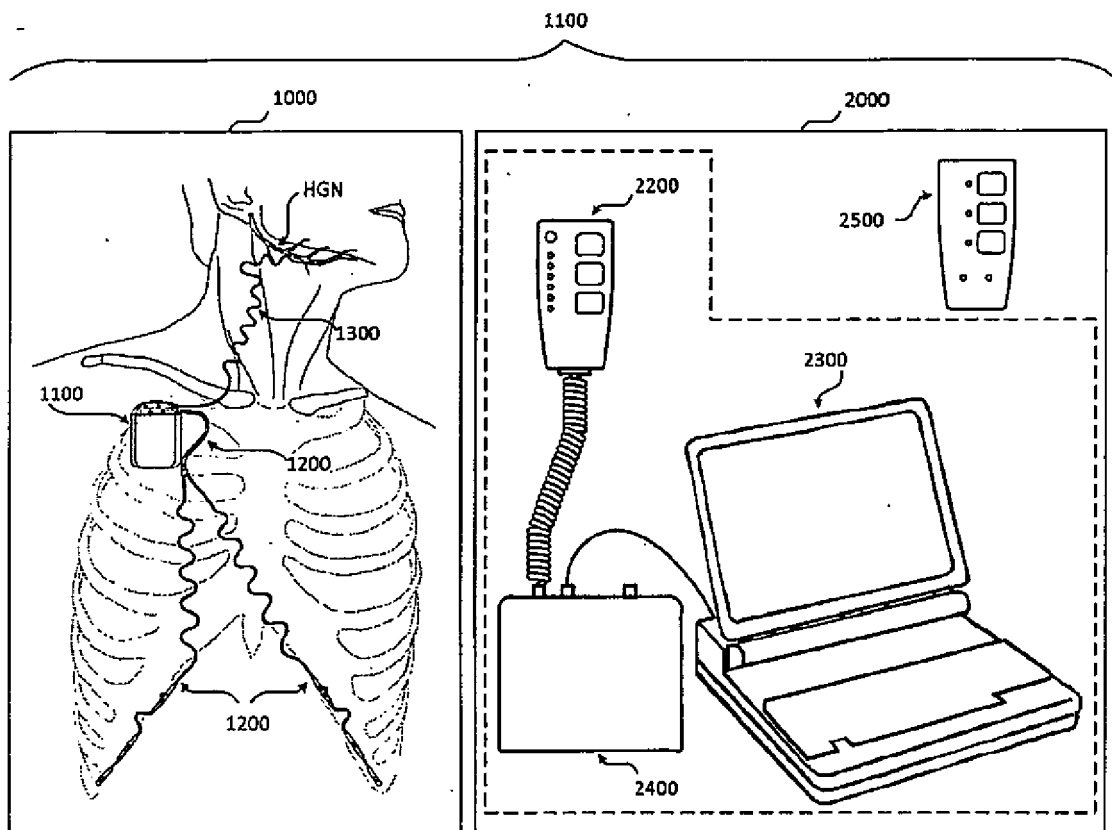
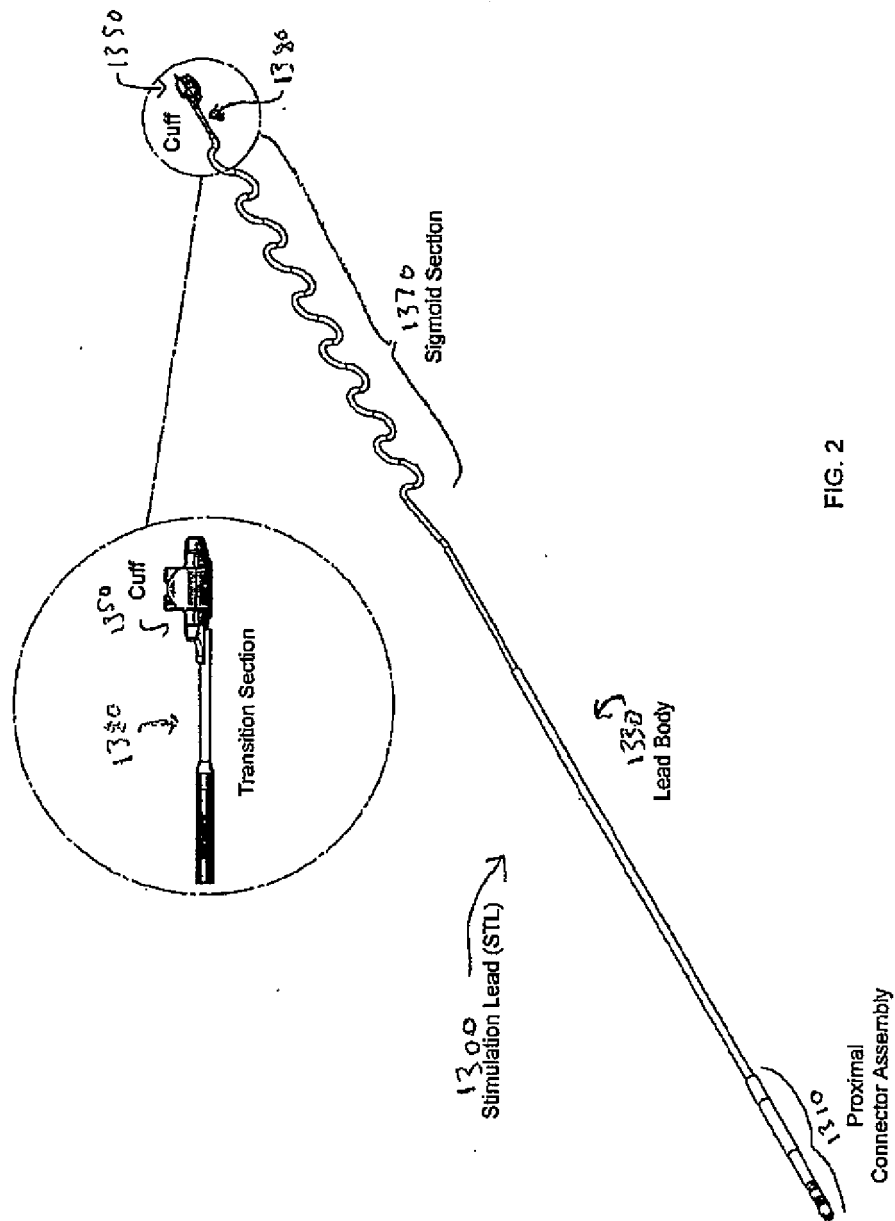


FIG. 1



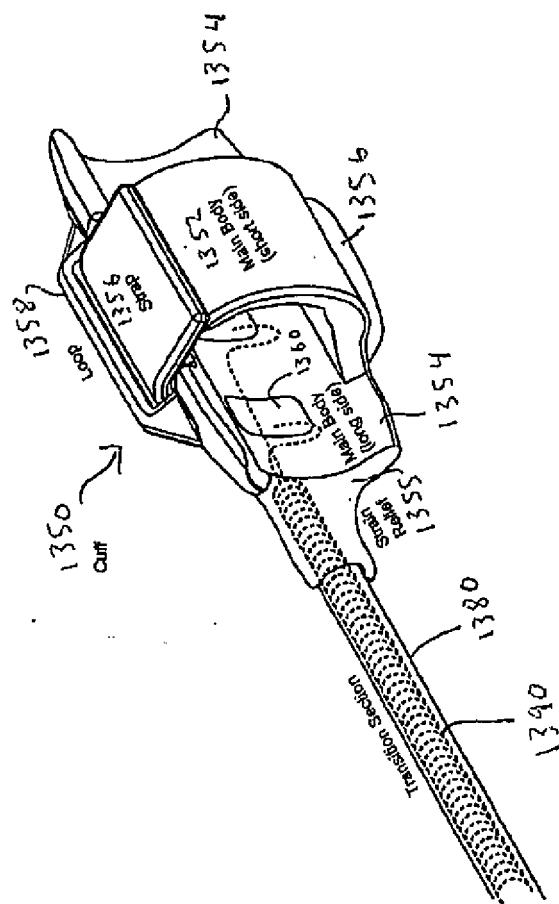


FIG. 3A

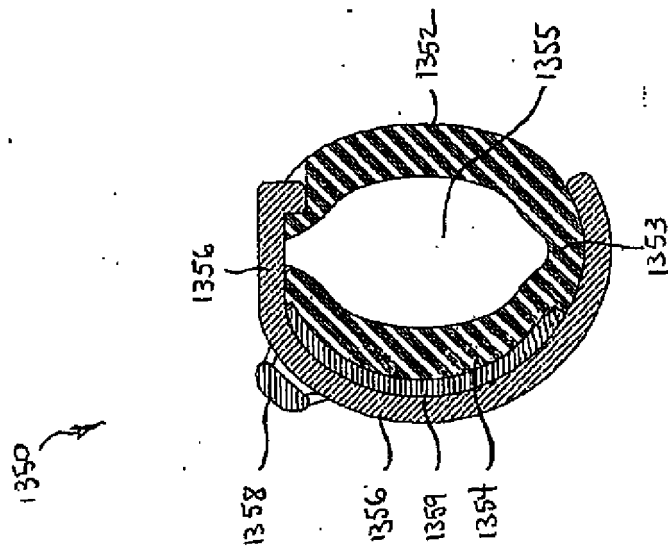


FIG. 3B

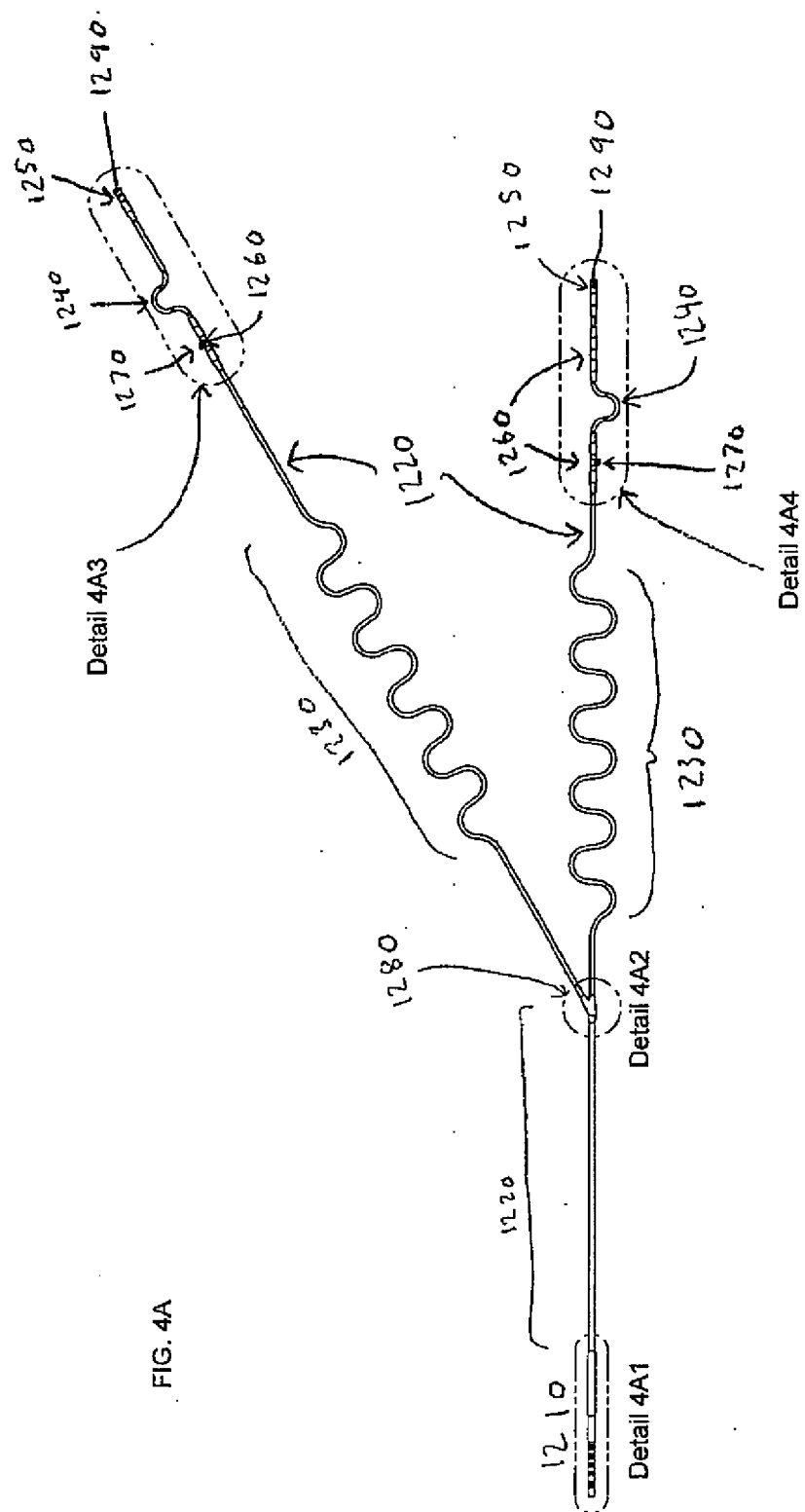
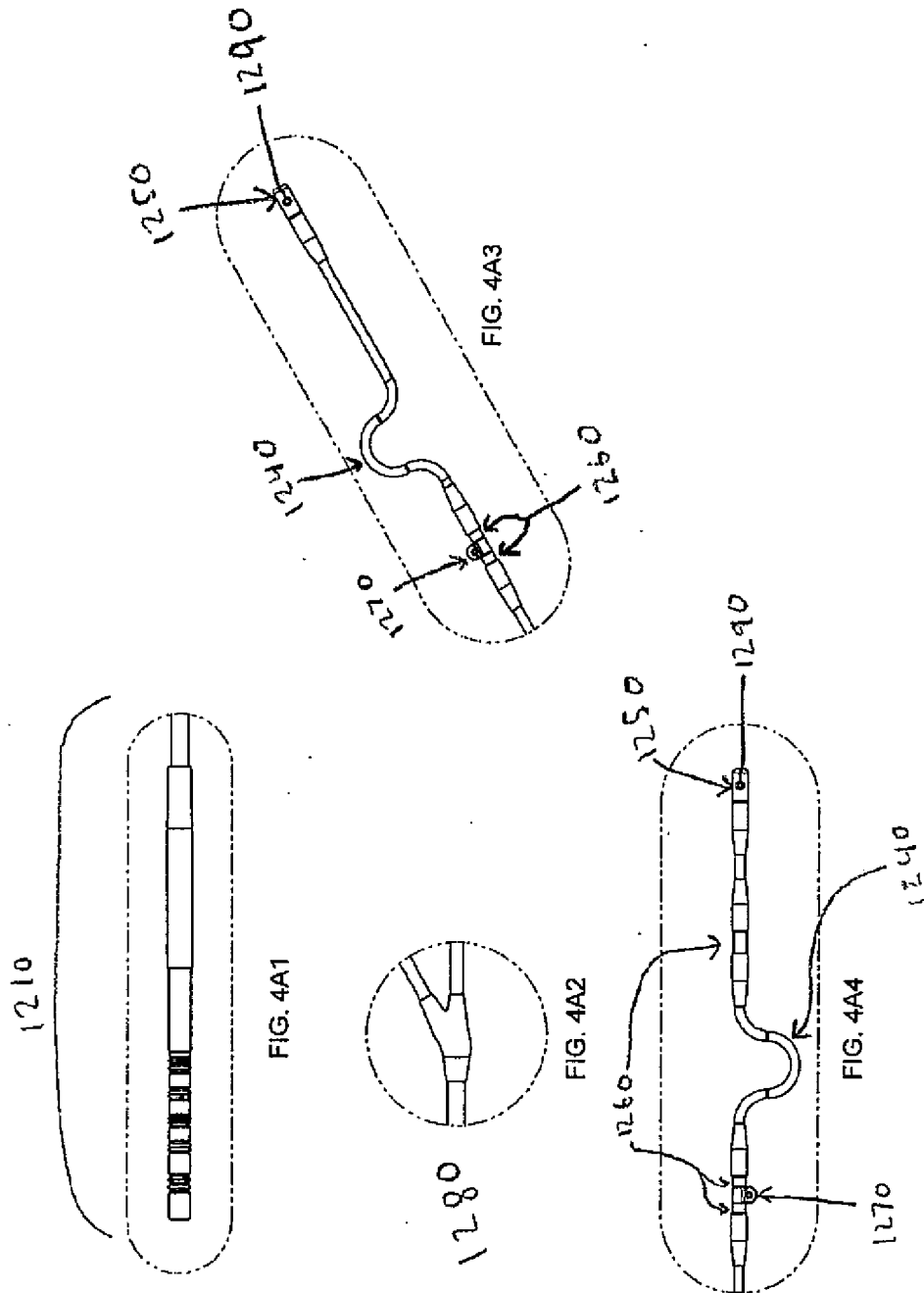
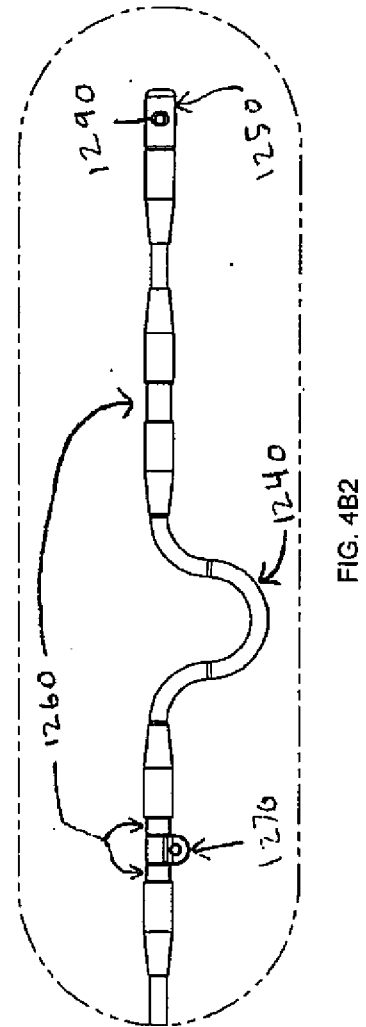
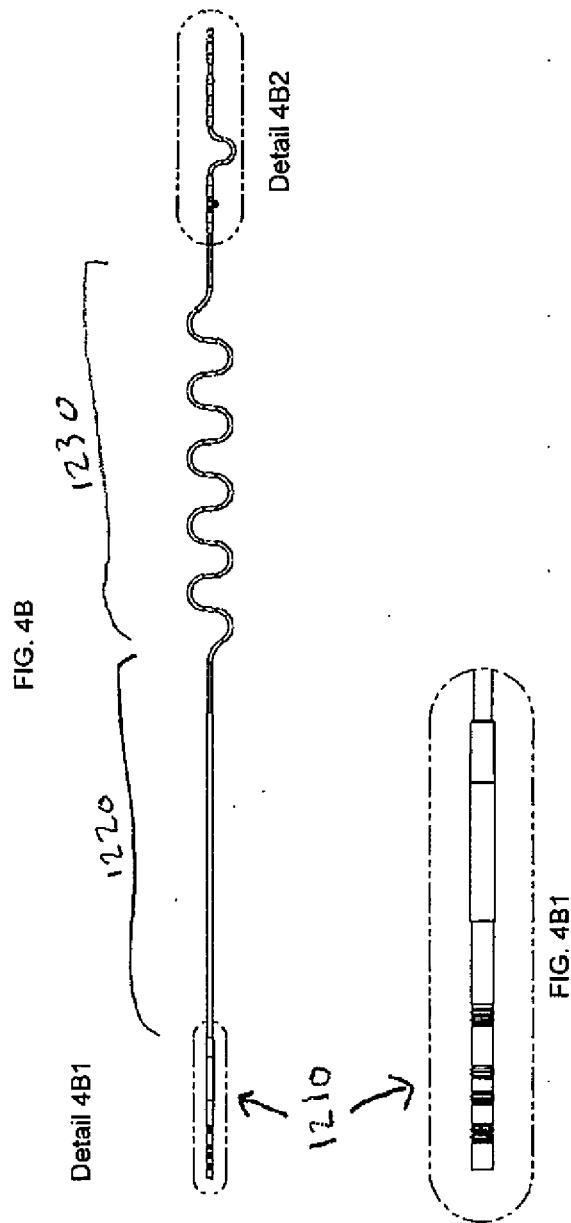


FIG. 4A





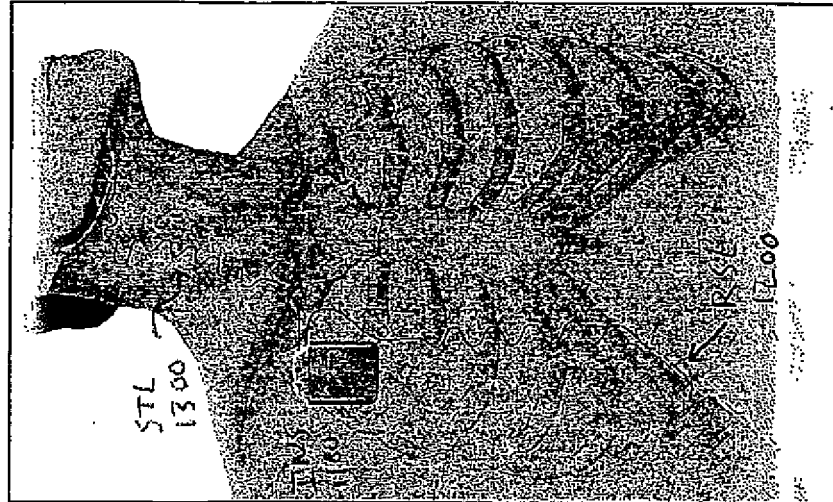


FIG. 4D

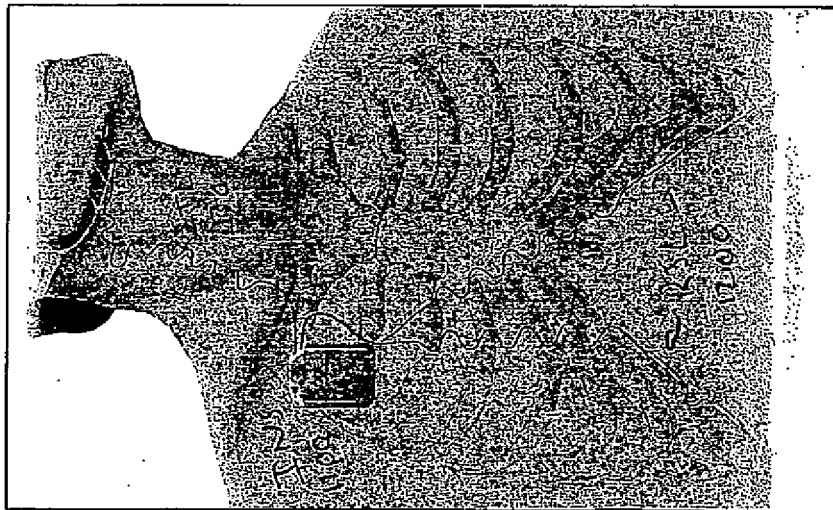
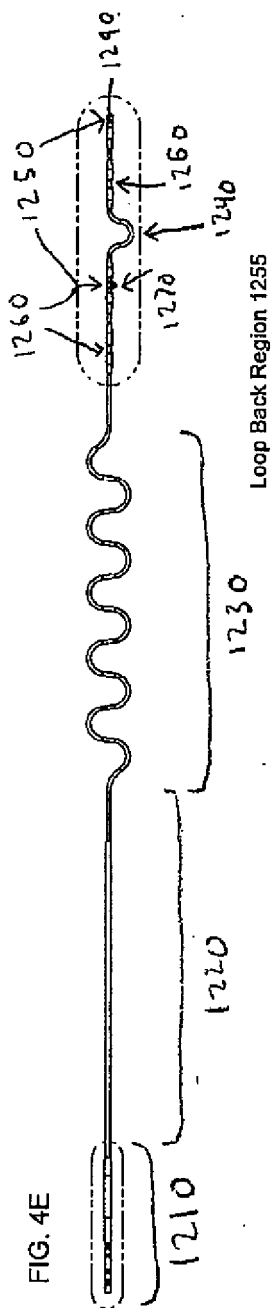


FIG. 4C



Loop Back Region 1255

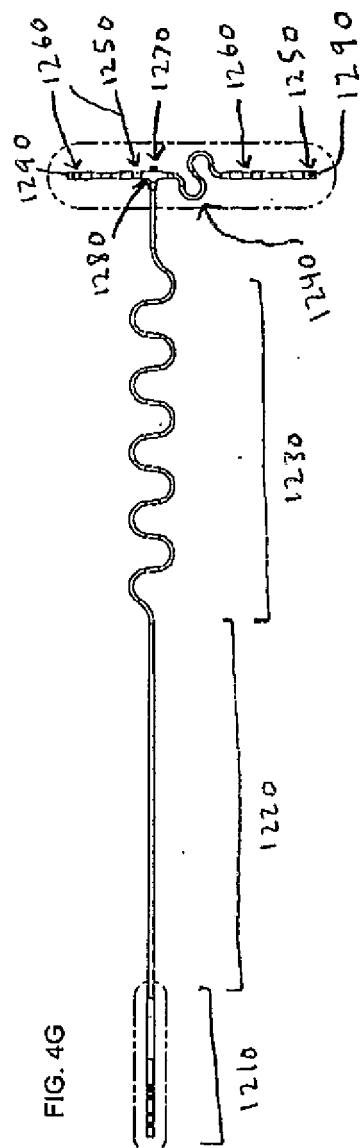
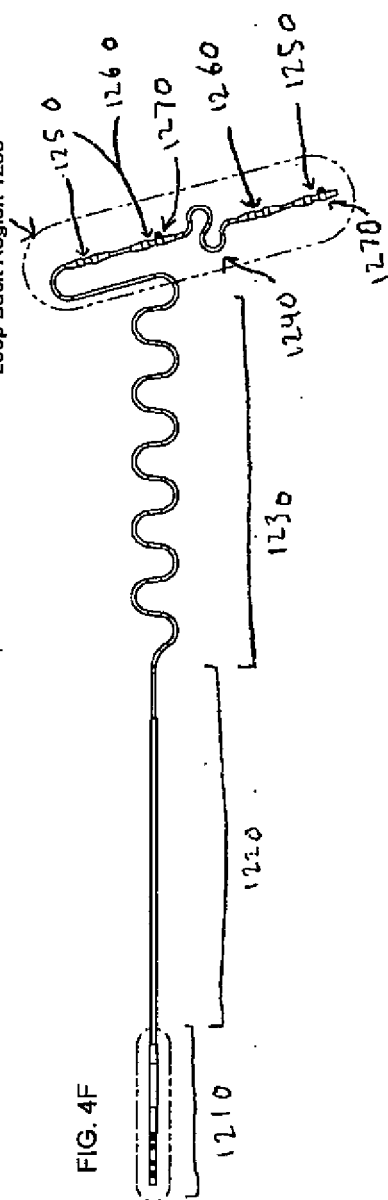
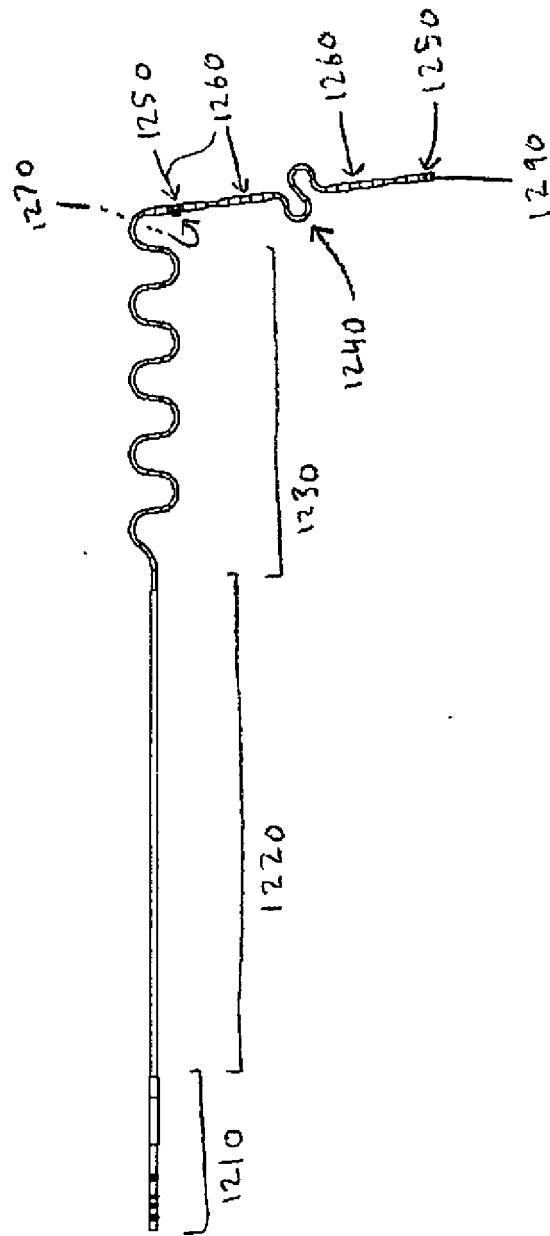


FIG. 4H



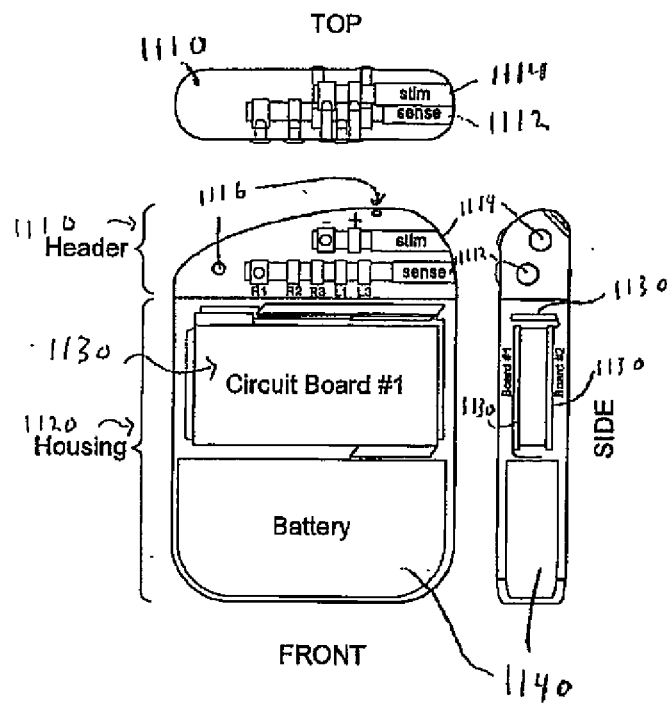


FIG. 5A

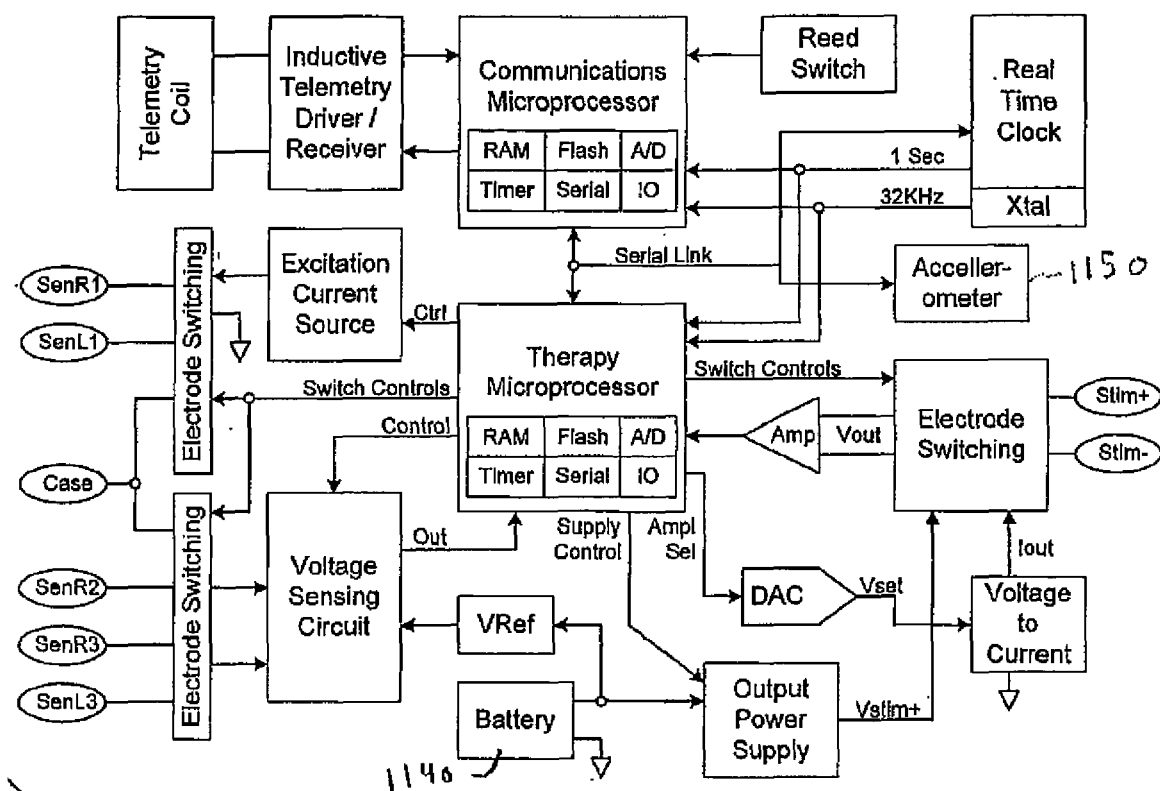


FIG. 5B

FIG. 6A

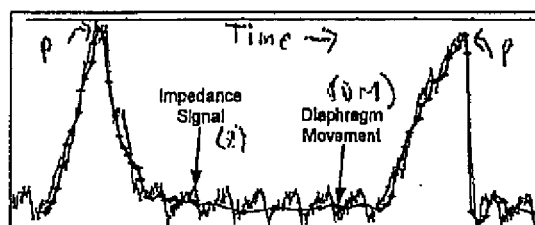


FIG. 6B

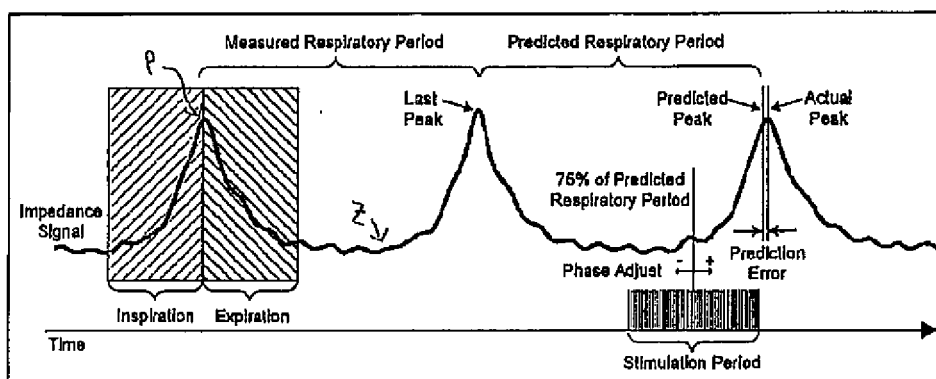


FIG. 6C

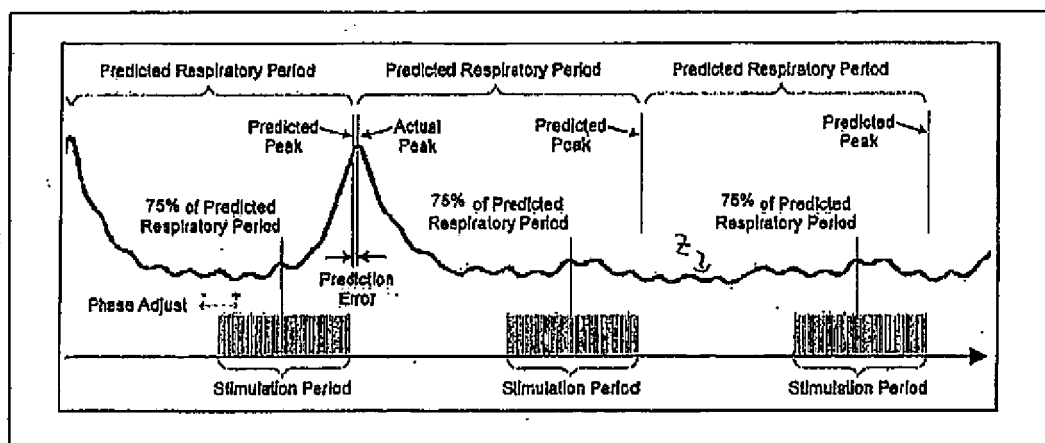
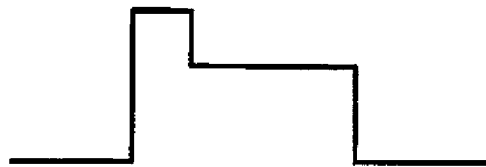


FIG. 6D



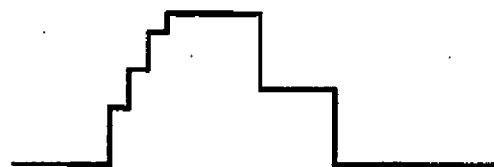
Default Stimulation

FIG. 6E



Basic Retention
Intensity

FIG. 6F



Retention Intensity

FIG. 6G



Soft Start

FIG. 6H



Nested Stimulation

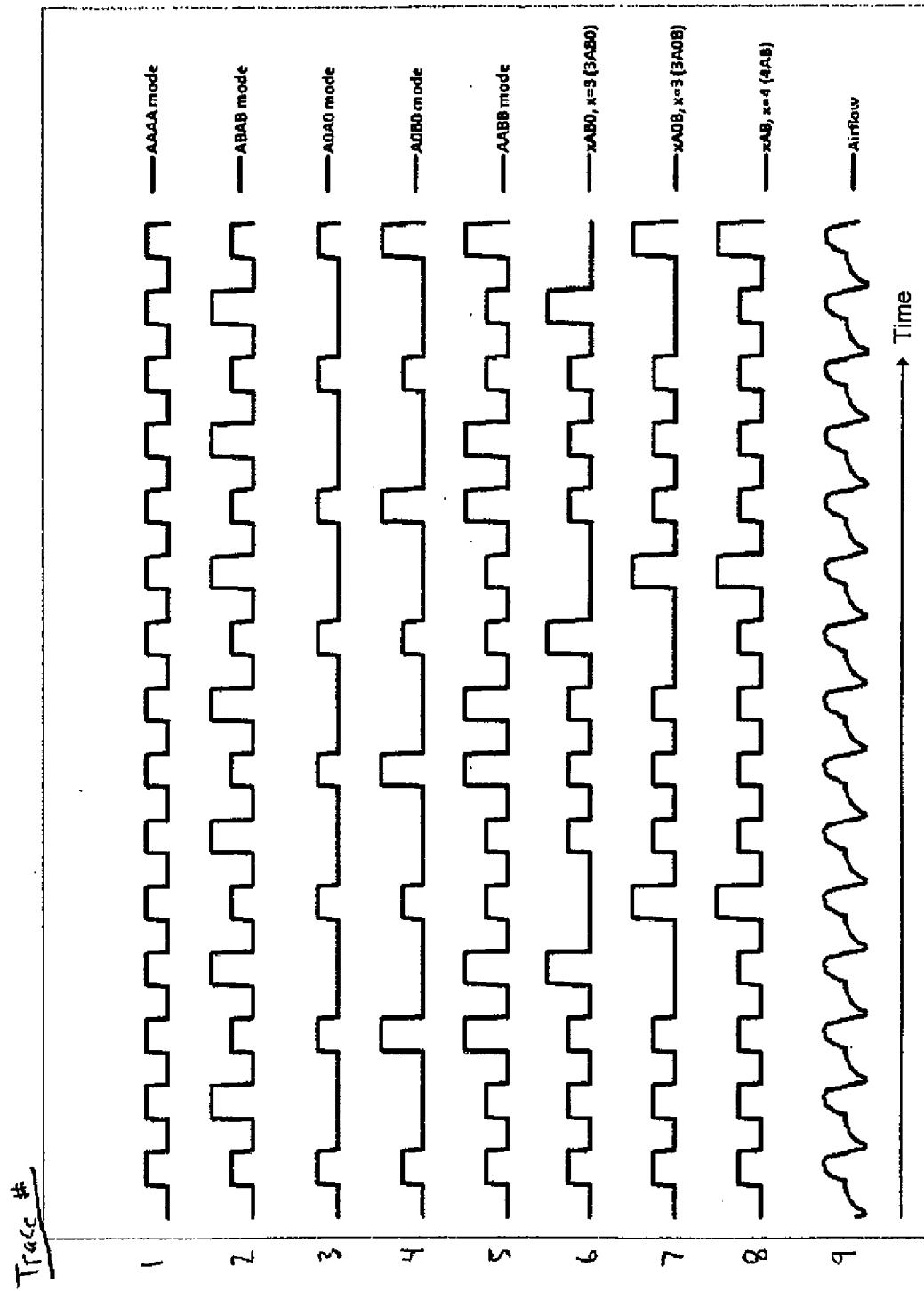


FIG. 6I

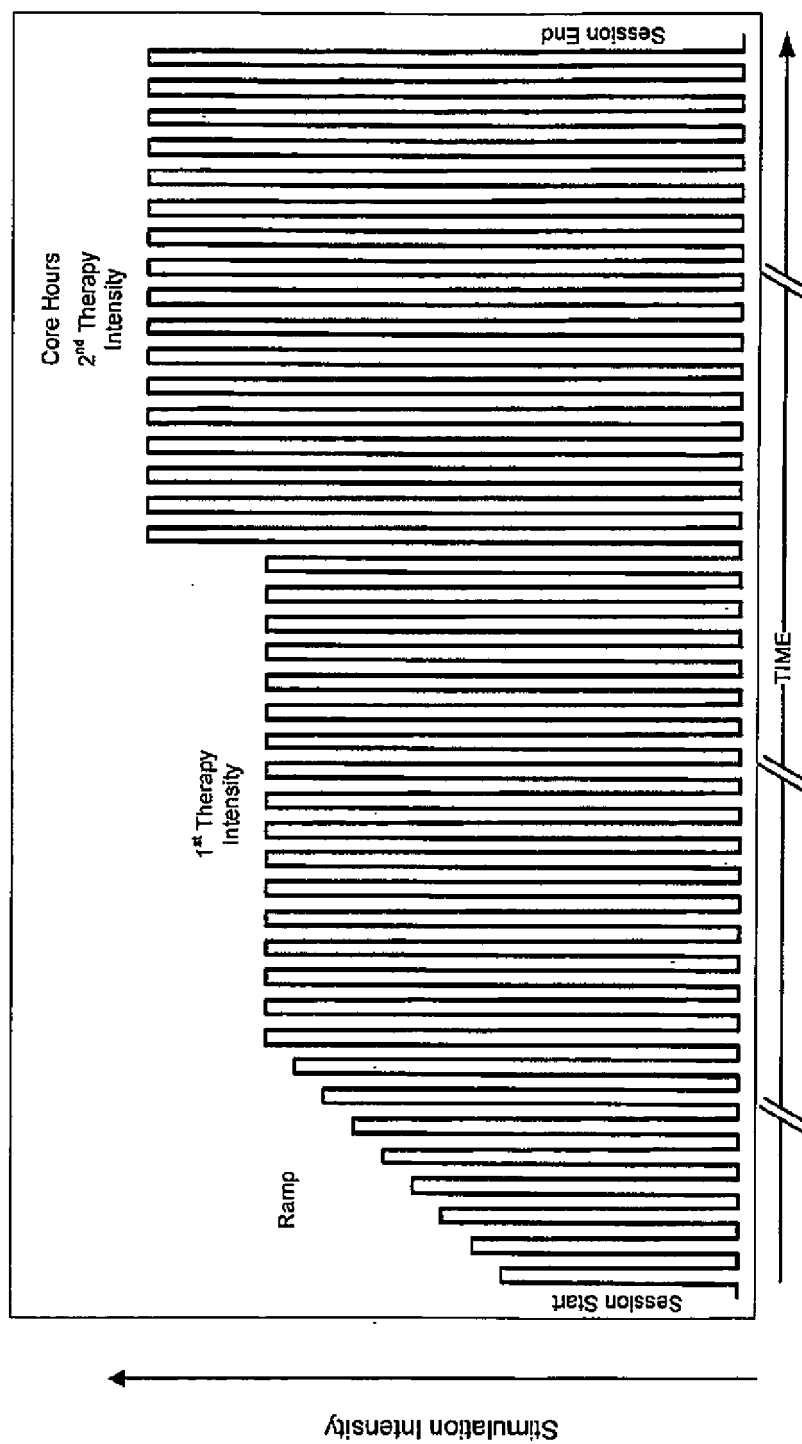


FIG. 6J

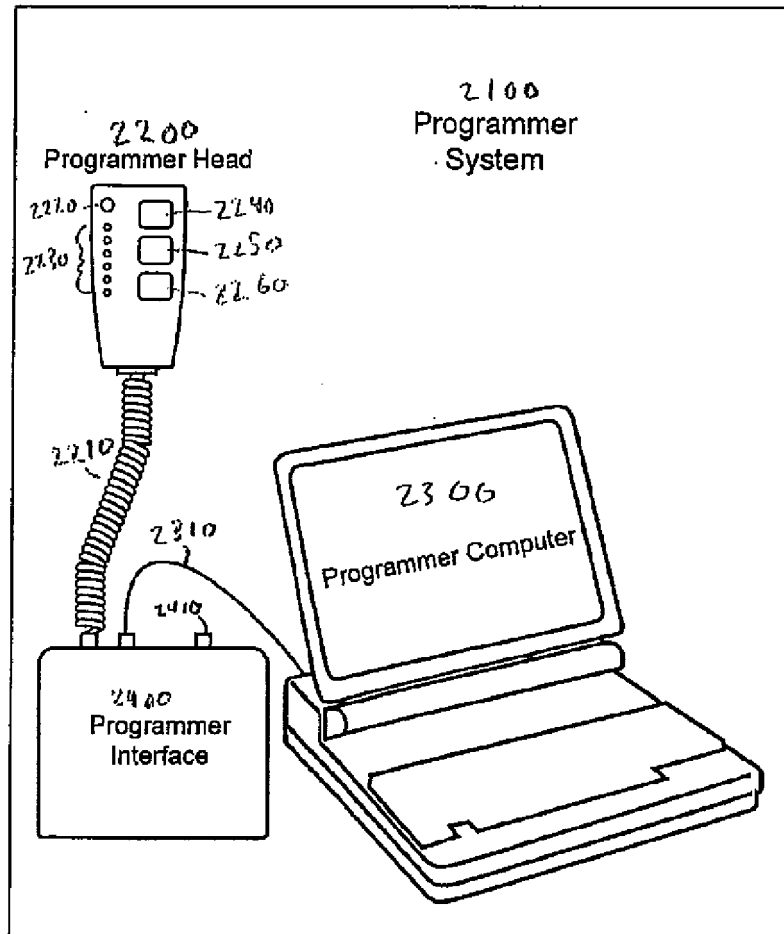


FIG. 7A

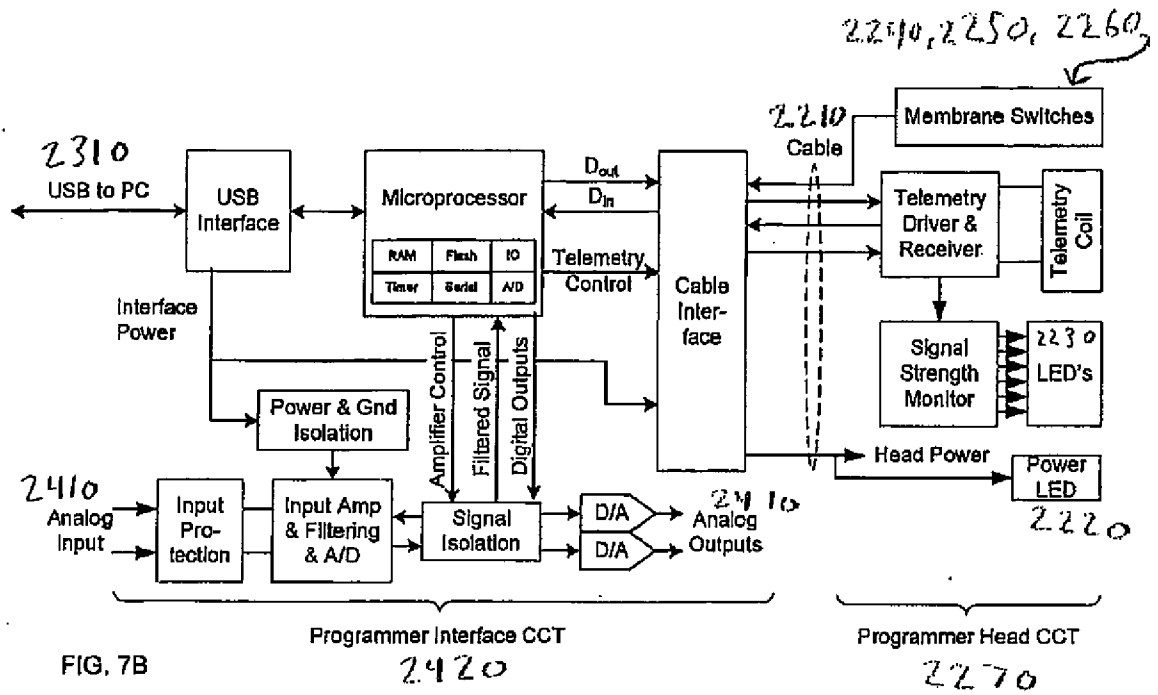


FIG. 7B

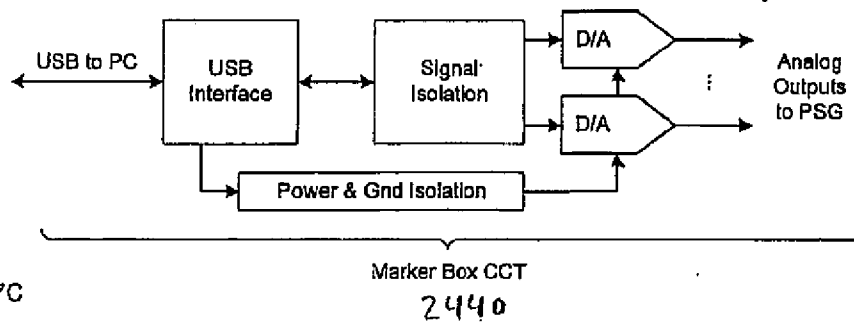


FIG. 7C

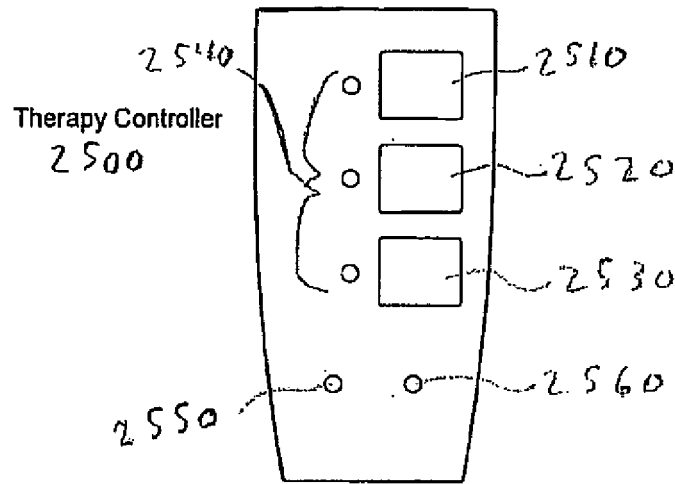


FIG. 8A

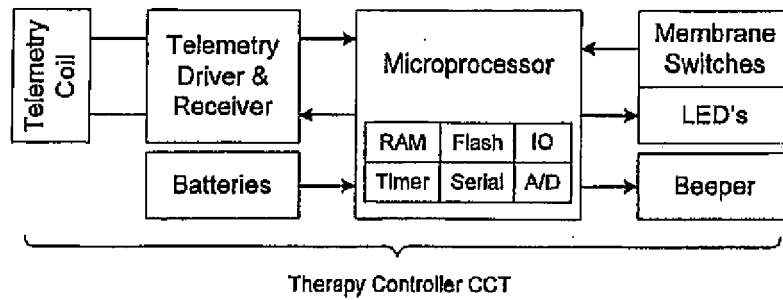


FIG. 8B

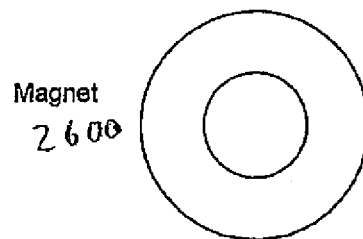


FIG. 9

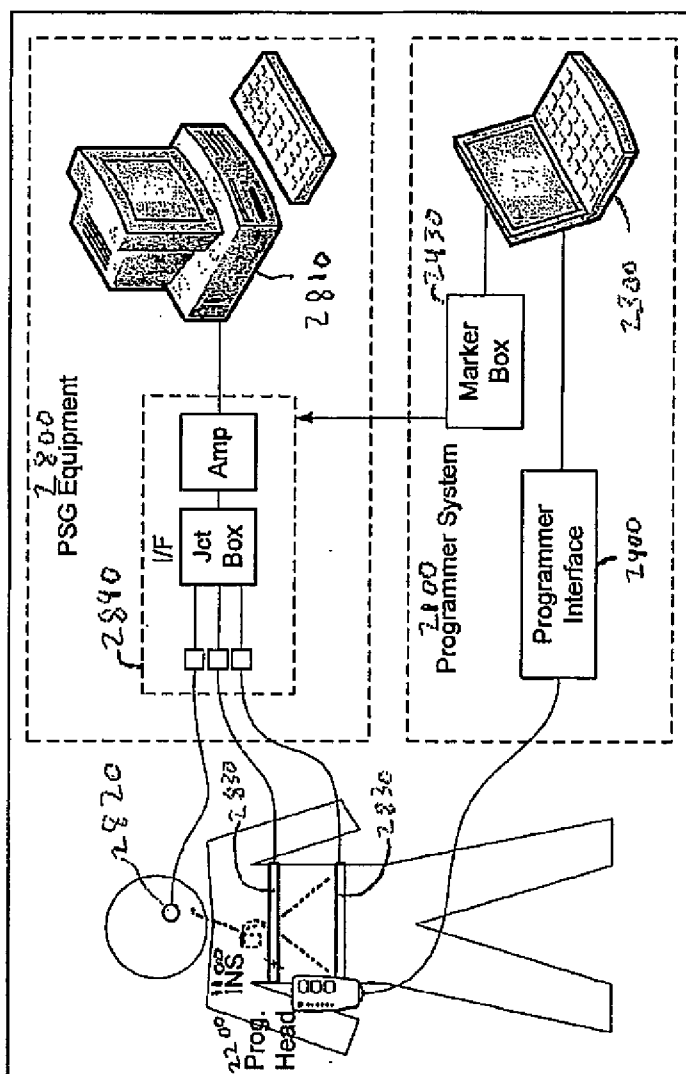


FIG. 10

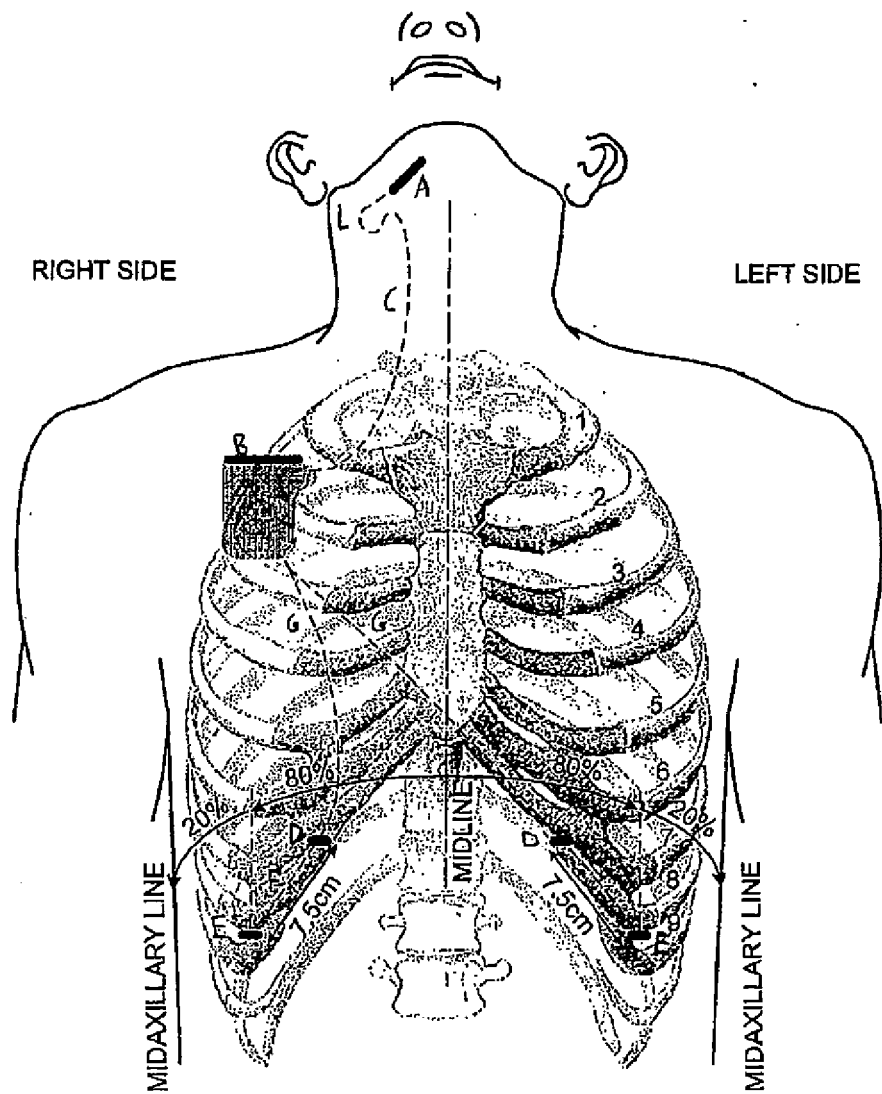


FIG. 11A

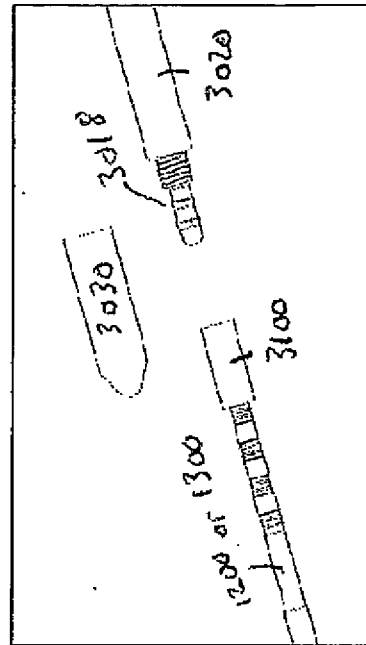
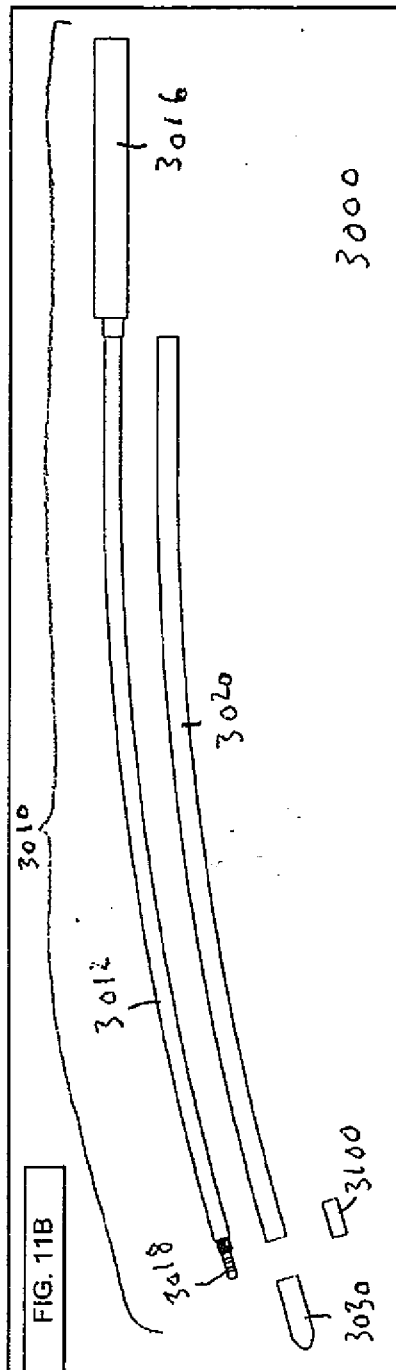


FIG. 11C

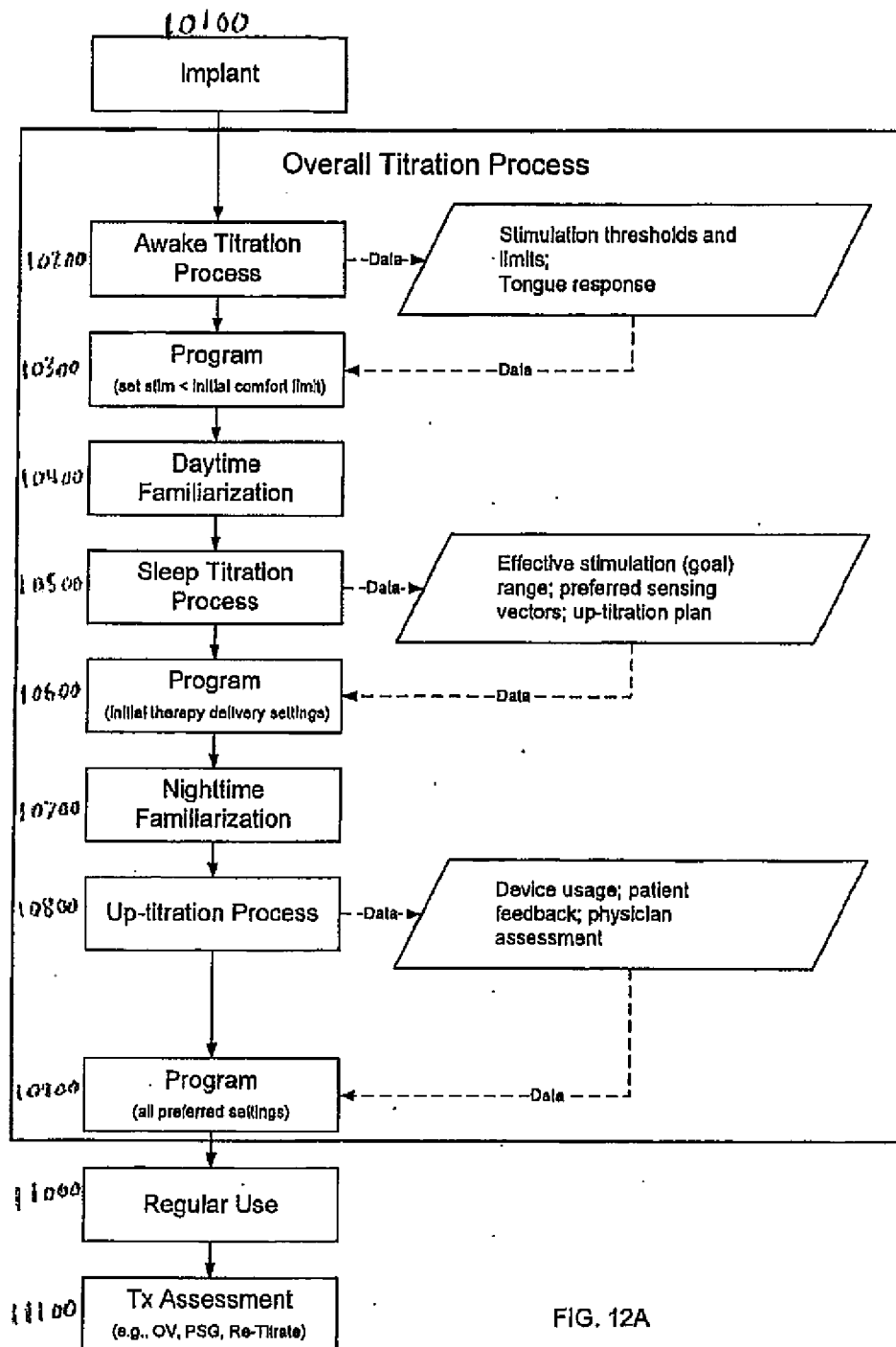


FIG. 12A

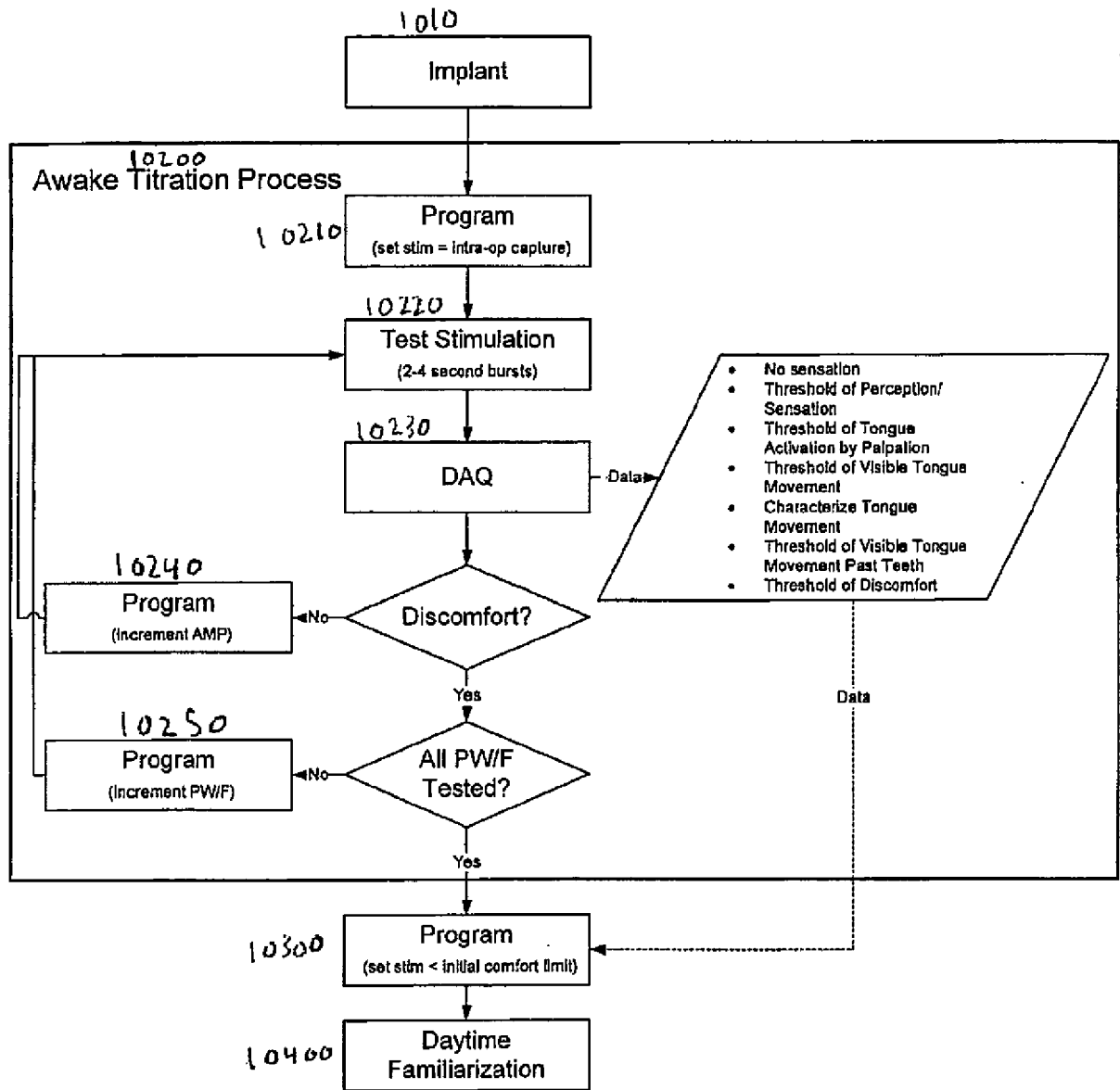


FIG. 12B

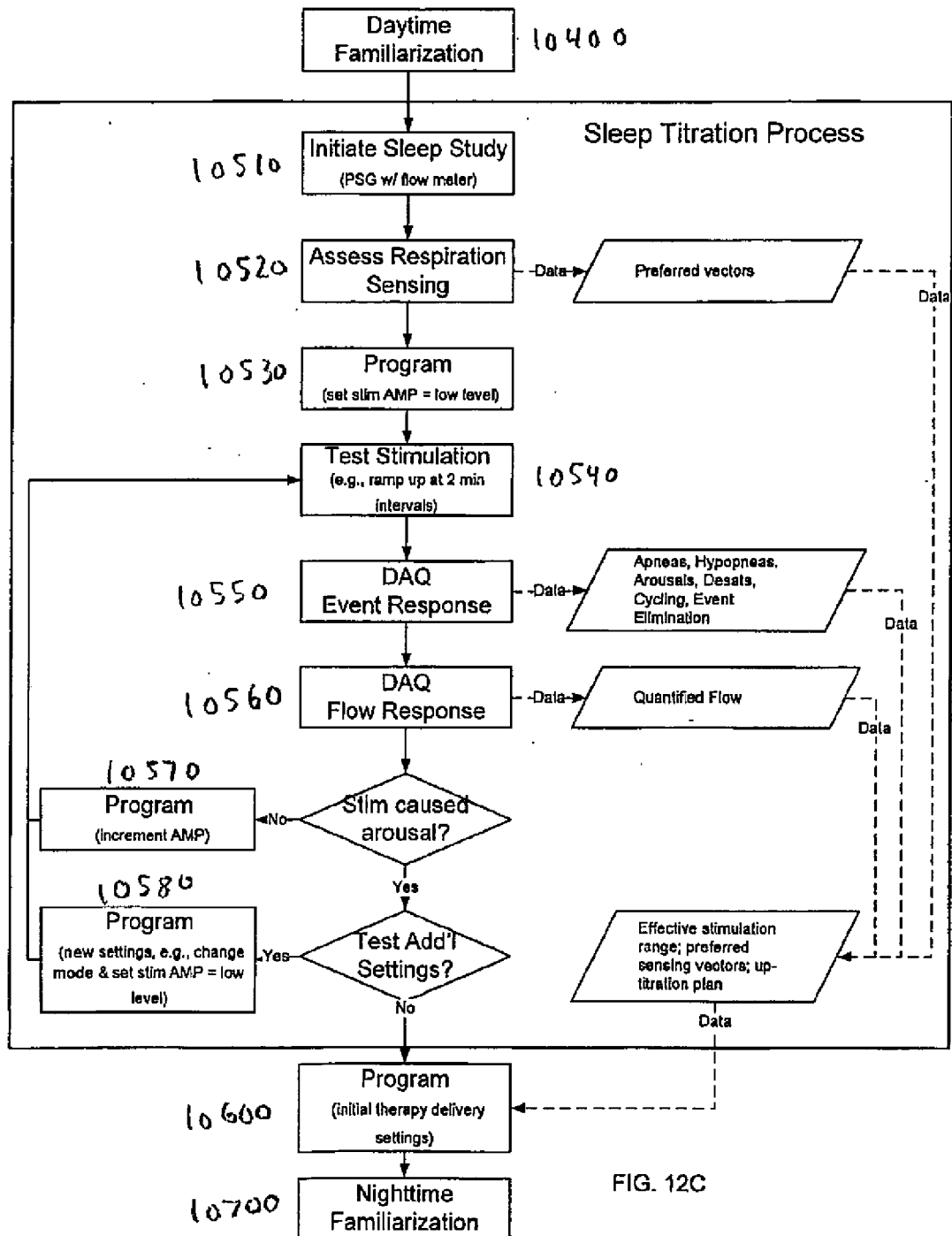


FIG. 12C

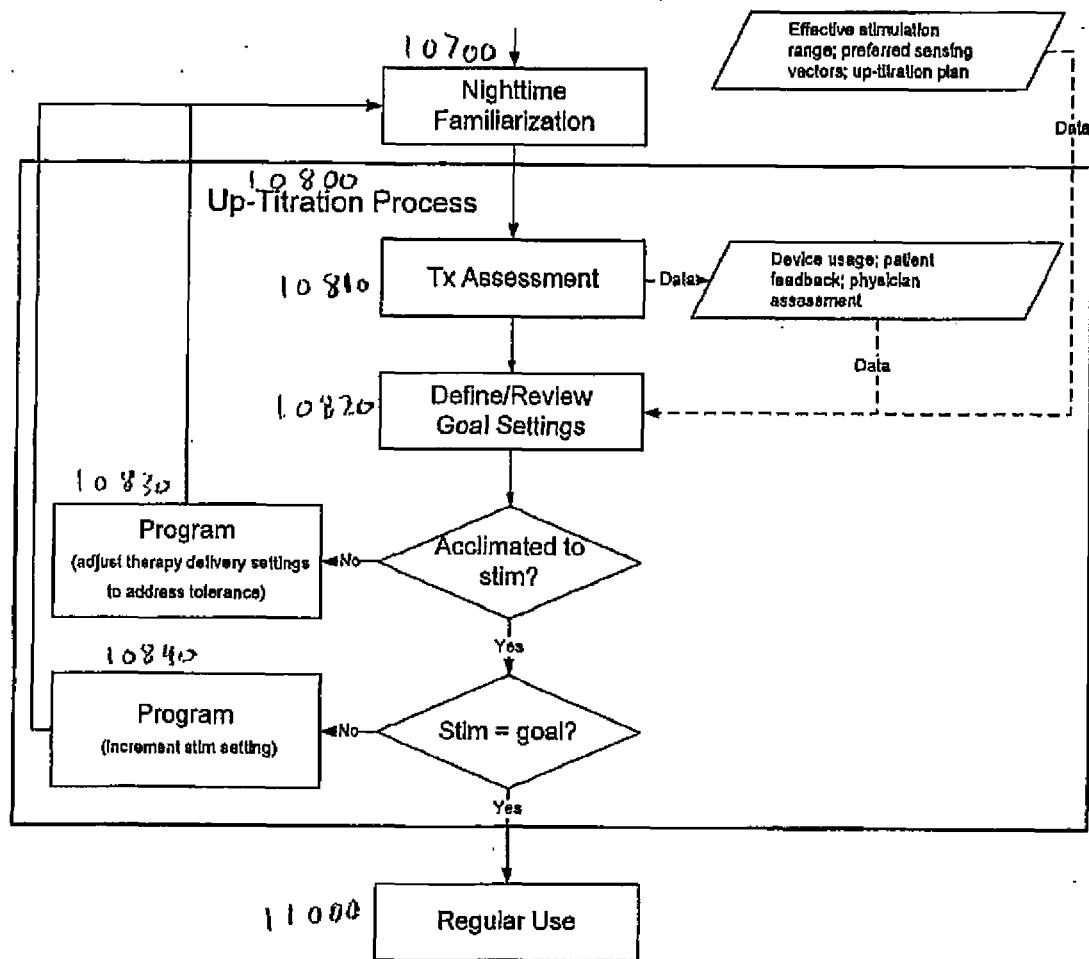


FIG. 12D

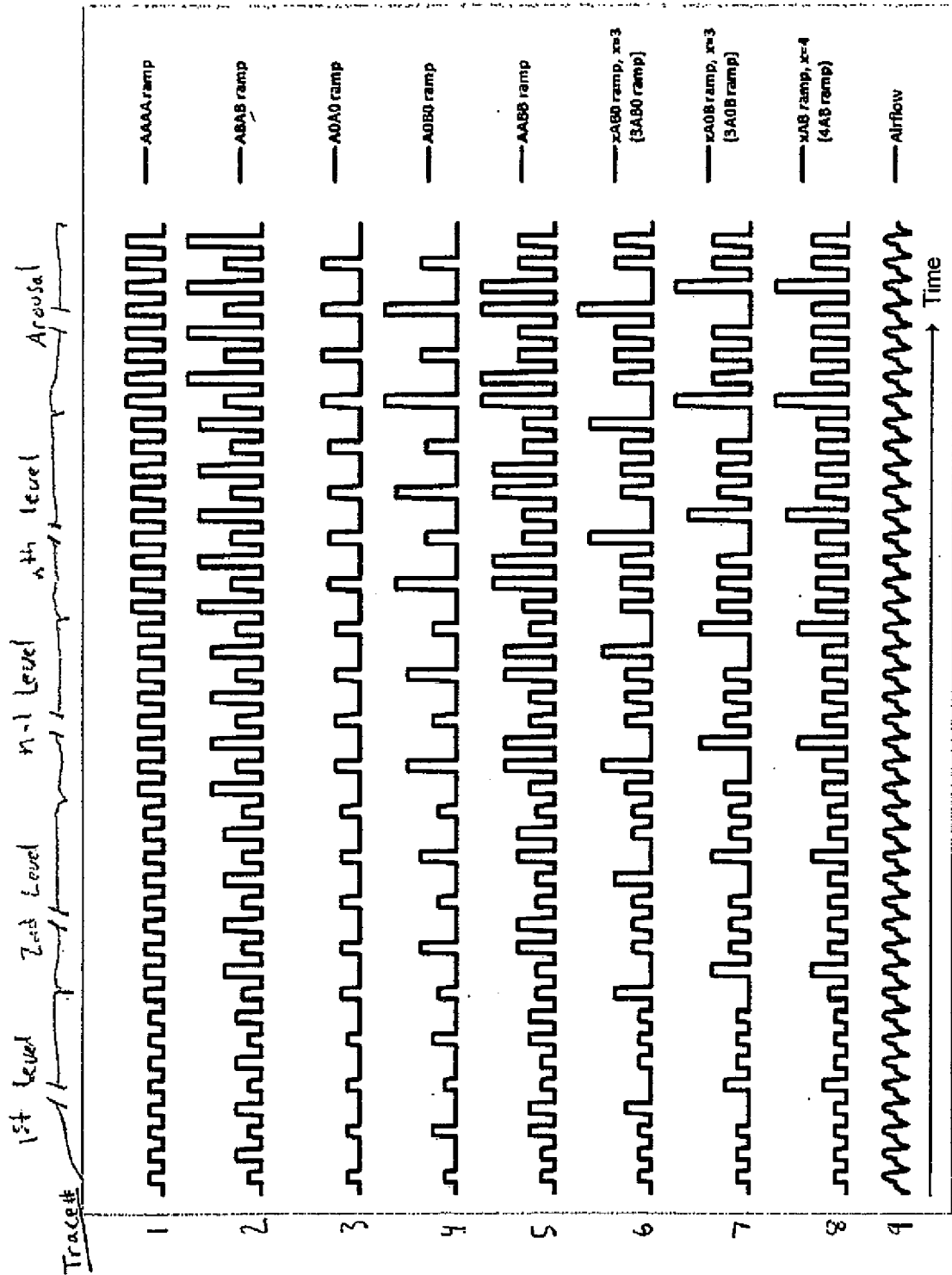


FIG. 13

Example of Airflow Response to Stimulation

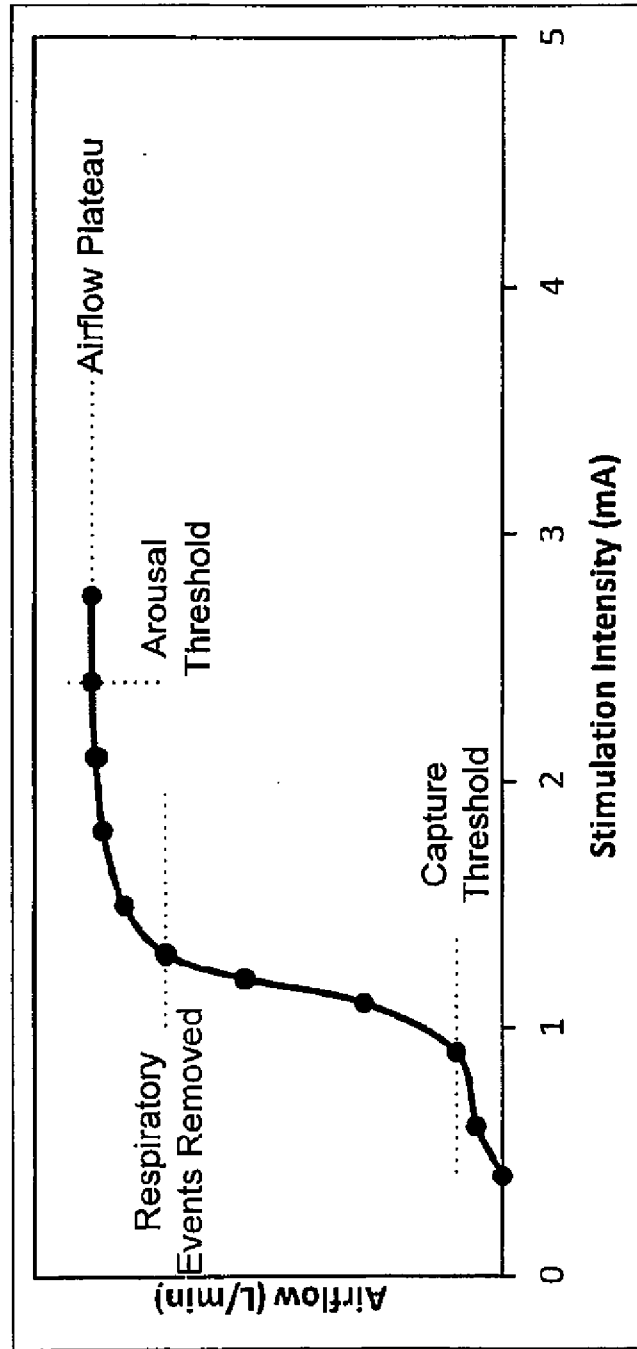
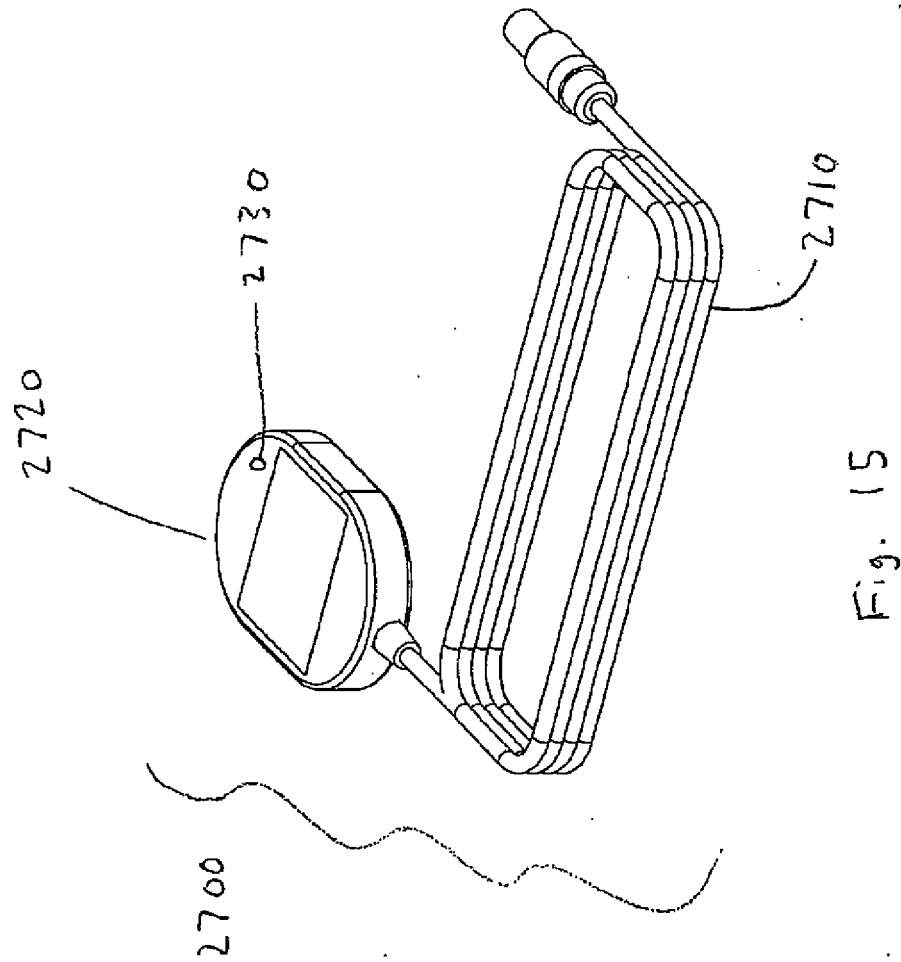


Fig. 14



REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

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- US 4830008 A, Meer [0005]

专利名称(译)	阻塞性睡眠呼吸暂停治疗装置		
公开(公告)号	EP3061493B1	公开(公告)日	2018-07-04
申请号	EP2016162666	申请日	2012-01-25
申请(专利权)人(译)	Cyberonics公司 , INC.		
当前申请(专利权)人(译)	Cyberonics公司 , INC.		
[标]发明人	BOLEA STEPHEN L HOEGH THOMAS B KUHNLEY BRIAN D SUILMANN DALE G PERSSON BRUCE J BECK JOHN P HAUSCHILD SIDNEY F TEFAYESUS WONDIMENEH SKUBITZ JASON J BOSSHARD MARK R PARRISH DANIEL A ATKINSON ROBERT E		
发明人	BOLEA, STEPHEN, L. HOEGH, THOMAS, B. KUHNLEY, BRIAN, D. SUILMANN, DALE, G. PERSSON, BRUCE, J. BECK, JOHN P. HAUSCHILD, SIDNEY, F. TEFAYESUS, WONDIMENEH SKUBITZ, JASON, J. BOSSHARD, MARK, R. PARRISH, DANIEL, A. ATKINSON, ROBERT, E.		
IPC分类号	A61N1/36 A61N1/05 A61B5/08 A61B5/00 A61N1/372		
CPC分类号	A61B5/0809 A61B5/0826 A61B5/4818 A61N1/0556 A61N1/3601 A61N1/3611 A61N1/36139 A61N1/3615 A61N1/37235 A61N1/36132		
优先权	61/437573 2011-01-28 US 13/106460 2011-05-12 US		
其他公开文献	EP3061493A1		
外部链接	Espacenet		

摘要(译)

用于治疗阻塞性睡眠呼吸暂停的神经刺激的装置，系统和方法。该系统适于基于来自呼吸感测引线的输入，在呼吸周期的适当阶段通过刺激导线将植入的神经刺激发送电信号到患者的神经。外部组件适于与神经刺激器进行无线通信。神经刺激器适于基于输入提供治疗刺激。

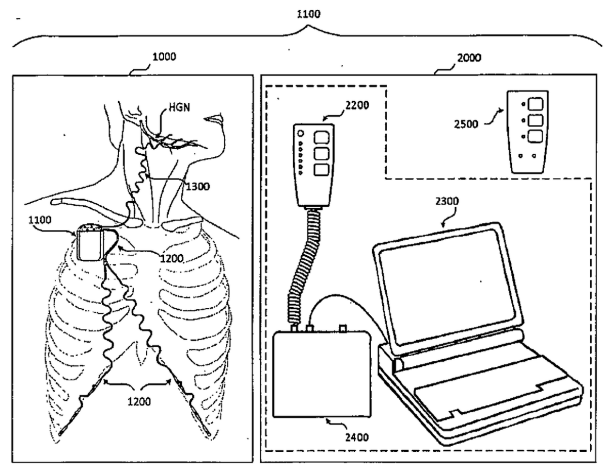


FIG. 1