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(54) **Title:** SYSTEMS AND METHODS FOR APPLYING ELECTRICAL STIMULATION FOR OPTIMIZING SPINAL CORD STIMULATION

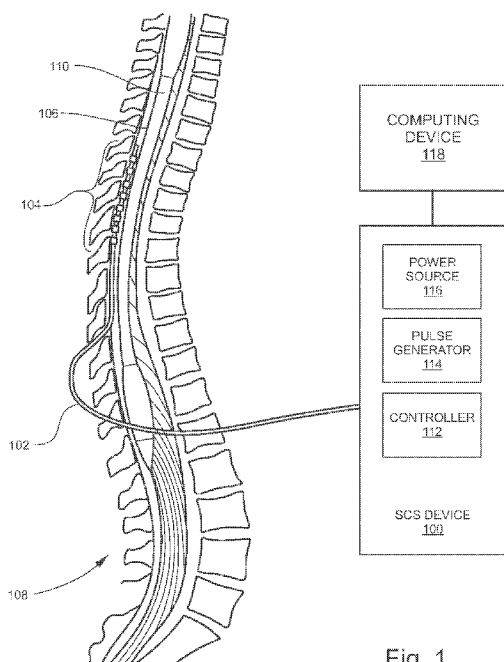


Fig. 1

(57) **Abstract:** Systems and methods for applying electrical stimulation to different sub-populations of targeted neurological tissue for optimizing spinal cord stimulation are disclosed. According to an aspect, a method includes applying a first pattern of electrical stimulation to a first sub-population of targeted neurological tissue of a subject. The method also includes applying a second pattern of electrical stimulation to a second sub-population of targeted neurological tissue of the subject, the second pattern of electrical stimulation being applied at a different frequency than the first pattern of electrical stimulation.

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SYSTEMS AND METHODS FOR APPLYING ELECTRICAL STIMULATION FOR OPTIMIZING SPINAL CORD STIMULATION

CROSS REFERENCE TO RELATED APPLICATION

[0001] This application claims the benefit of U.S. Provisional Patent Application Number 61/779,632, filed March 13, 2013 and titled SYSTEMS AND METHODS FOR OPTIMIZING SPINAL CORD STIMULATION, the disclosure of which is incorporated herein by reference in its entirety.

TECHNICAL FIELD

[0002] The presently disclosed subject matter relates to spinal cord stimulation, and more specifically, to applying electrical stimulation for optimizing spinal cord stimulation (SCS).

BACKGROUND

[0003] SCS has emerged as a therapy for chronic pain when kinetic (*e.g.*, physical rehabilitation), pharmaceutical, and surgical therapies have not been effective. However, between 1974 and 1991, according to studies the clinical success of SCS has been highly variable, with a mean of 54.2% and a standard deviation of 20%, and subsequent studies have shown very little improvement. Efforts to improve the clinical efficacy of SCS have focused on the development of more spatially selective electrodes, while only minimal attention has been paid to the temporal patterning of SCS or the effects of SCS on the activity of neurons in the dorsal horn pain processing circuit. Although there have been advances in SCS, there is a continuing need for improved techniques and systems for optimizing SCS.

BRIEF SUMMARY

[0004] Disclosed herein are systems and methods for applying electrical stimulation to different sub-populations of targeted neurological tissue for optimizing spinal cord stimulation. According to an aspect, a method includes applying a first pattern of electrical

stimulation to a first sub-population of targeted neurological tissue of a subject. The method also includes applying a second pattern of electrical stimulation to a second sub-population of targeted neurological tissue of the subject, the second pattern of electrical stimulation being applied at a different frequency than the first pattern of electrical stimulation. Further, the method includes controlling the first and second patterns of electrical stimulation for optimizing suppression of activity of wide-dynamic range (WDR) neurons to improve the efficacy of stimulation and / or reducing the average stimulation frequency to improve the efficiency of stimulation.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

- [0005]** The foregoing aspects and other features of the present subject matter are explained in the following description, taken in connection with the accompanying drawings, wherein:
- [0006]** FIG. 1 is an anatomic view of a system for stimulating targeted neurological tissue of a human subject in accordance with embodiments of the present disclosure;
- [0007]** FIG. 2 is a flow chart of an example method for SCS in accordance with embodiments of the present disclosure;
- [0008]** FIG. 3 are graphs showing that delivering SCS at different timings through different fiber populations can result in greater efficacy in response to a 1 Hz peripheral input;
- [0009]** FIG. 4 is a schematic of an example computational model for model-based design and evaluation of optimal temporal patterns of SCS;
- [0010]** FIGs. 5A and 5B are graphs showing example patterns of activity in peripheral primary afferent fibers;
- [0011]** FIGs. 6A and 6B are graphs showing 1-second long examples of non-harmonic and harmonic multi-frequency SCS, respectively;
- [0012]** FIG. 7 is a timeline of each experimental run in accordance with embodiments of the present disclosure;
- [0013]** FIG. 8 are Raster plots depicting example activity of a WDR neuron during the period of time in which multi-frequency SCS may be delivered;
- [0014]** FIG. 9 are Raster plots depicting comparisons of SCS efficacy and efficiency between multi-frequency SCS and conventional SCS at fixed frequency in response to

a 1 Hz conditioning input;

[0015] FIG. 10 are Raster plots depicting comparisons of SCS efficacy (WDR neuronal output) and efficiency (average stimulation frequency) between multi-frequency SCS and conventional SCS at a fixed frequency in response to a neuropathic input;

[0016] FIGs. 11A and 11B are Raster plots depicting comparisons of SCS efficacy (WDR neuronal output) and efficiency (average stimulation frequency) between several combinations of harmonic multi-frequency SCS and conventional SCS at a fixed frequency in response to a neuropathic input; and

[0017] FIG. 12 is an illustration of a regular, constant frequency stimulation train wherein the interpulse intervals are constant in time and examples of non-regular temporal patterns of stimulation wherein the interpulse intervals vary in time.

DETAILED DESCRIPTION

[0018] For the purposes of promoting an understanding of the principles of the present disclosure, reference will now be made to various embodiments and specific language will be used to describe the same. It will nevertheless be understood that no limitation of the scope of the disclosure is thereby intended, such alteration and further modifications of the disclosure as illustrated herein, being contemplated as would normally occur to one skilled in the art to which the disclosure relates.

[0019] Articles “a” and “an” are used herein to refer to one or to more than one (i.e. at least one) of the grammatical object of the article. By way of example, “an element” means at least one element and can include more than one element.

[0020] As used herein, the term “subject” and “patient” are used interchangeably herein and refer to both human and non-human animals. The term “non-human animals” of the disclosure includes all vertebrates, *e.g.*, mammals and non-mammals, such as non-human primates, sheep, dog, cat, horse, cow, chickens, amphibians, reptiles, and the like. In examples provided herein, the subject is a human patient in need of spinal cord stimulation.

[0021] As used herein, the term “neurological disorder” refers to any pathological condition relating to the brain and/or nervous system. Examples include, but are not limited to, pain, which includes chronic and acute neuropathic pain, migraine, trauma, and the like. As used herein, the term “pain” refers to the basic bodily sensation induced by a noxious stimulus,

received by naked nerve endings, characterized by physical discomfort (*e.g.*, pricking, throbbing, aching, etc.) and typically leading to an evasive action by the individual. As used herein, the term pain also includes chronic and acute neuropathic pain. The term “chronic pain” and “chronic neuropathic pain” are used interchangeably refer to a complex, chronic pain state that is usually accompanied by tissue injury wherein the nerve fibers themselves may be damaged, dysfunctional, or injured. These damaged nerve fibers send incorrect signals to other pain centers. The impact of nerve fiber injury includes a change in nerve function both at the site of injury and areas around the injury. Chronic neuropathic pain often seems to have no obvious cause, however, some common causes may include, but are not limited to, alcoholism, amputation, back, leg and hip problems, chemotherapy, diabetes, facial nerve problems, HIV infection or AIDS, multiple sclerosis, shingles, spine injury, and the like. For example, neuropathic pain may include phantom limb syndrome, which occurs when an arm or leg has been removed because of illness or injury, but the brain still gets pain messages from the nerves that originally carried impulses from the missing limb.

[0022] As referred to herein, the term “administering” refers to the delivery of an electrical impulse/signal/frequency to a subject to thereby cause stimulation to a nerve, nerve fiber, or group of nerve fibers. For example, electrical impulse/signal/frequency may be applied by use of one or more electrodes in electrical communication with a targeted neurological tissue region, such as sub-populations of dorsal column nerve fibers for example.

[0023] Unless otherwise defined, all technical terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this disclosure belongs.

[0024] In accordance with embodiments of the present disclosure, systems and methods of optimizing SCS are disclosed. A system for delivering SCS to a subject can include a pulse generator. The pulse generator may be configured to generate electrical signals for delivery to targeted neurological tissue of the subject. The system may also include one or more SCS electrodes in electrical communication with an output of the pulse generator. The contact(s) may be placed in contact with the targeted neurological tissue. A controller of the system may control the pulse generator to produce predetermined patterns of electrical stimulation to the targeted neurological tissue. The patterns may be controlled based on prior simulations that optimized suppression of activity of model wide-dynamic range (WDR) neurons to improve the efficacy of

treatment. The pattern may be controlled to reduce the average stimulation frequency to improve the efficiency of treatment.

[0025] FIG. 1 illustrates an anatomic view of a system for stimulating targeted neurological tissue of a human subject in accordance with embodiments of the present disclosure. The subject may be suffering from a neurological disorder, such as chronic pain. Referring to FIG. 1, the system includes an SCS device **100**, an electrical cord **102** and an electrode array generally designated **104**. The system is shown as being implanted in the subject. Particularly, the electrode array **104** is operatively positioned in the epidural space **106** of a vertebral column **108** of the subject. The electrode array **104** is positioned at the site of nerves that are the target of stimulation, *e.g.*, along the spinal cord **110**. Alternatively, the electrode array **104** may be suitably positioned in any other location for desired electrical stimulation of targeted neurological tissue. The cord **102** may include multiple lines or fibers such that different or the same electrical signals can be provided to contacts of the electrode array **104**. The SCS device **100** may be suitably implanted within the subject such as, but not limited to, implantation within the abdomen or buttocks. The electrical cord **102** may operatively connect an output of the SCS device **100** to the electrode array **104**.

[0026] The SCS device **100** may include a controller **112** and a pulse generator **114**. The controller **112** may include hardware, software, firmware, or combinations thereof for implementing functionality described herein. For example, the controller **112** may be implemented by one or more processors and memory. The controller **112** may be operatively connected to the pulse generator **114** for controlling the pulse generator **114** to generate electrical signals for applying patterns of electrical stimulation to targeted neurological tissue. The output signals may be received by the electrical cord **102** and carried to the electrode array **104** for electrical stimulation at targeted neurological tissue. The SCS device **100** may include a power source **116**, such as a battery, for supplying power to the controller **112** and the pulse generator **114**.

[0027] The system may also include an external computing device **118** that is not implanted within the subject. The computing device may communicate with the SCS device **100** via any suitable communication link (*e.g.*, a wired, wireless, or optical communication link). The communication link may also facilitate battery recharge. A clinician may interact with a user interface of the computing device for programming the output of the implanted pulse generator **114**, including the electrodes that are active, the stimulation pulse amplitude, the stimulation

pulse duration, the stimulation pattern (including pulse repetition frequency), and the like applied via each electrode contact to each sub-population.

[0028] Further, in accordance with embodiments of the present disclosure, the computing device **118** may determine one or more non-regular temporal patterns that results in predetermined WDR neuronal output and stimulation activity. The computing device **118** may communicate information for administering the temporal patterns to the SCS device **100**, which may then apply the non-regular temporal pattern(s) of electrical stimulation to targeted neurological tissue of the subject.

[0029] A patient may also interact with the user interface of the computing device **118**. In this embodiment, the patient may interact with the user interface for selecting among a set of pre-programmed stimulation parameter sets. These sets may have been programmed or otherwise set by the clinician and stored in the controller **112**.

[0030] FIG. 2 illustrates a flow chart of an example method for SCS in accordance with embodiments of the present disclosure. The example method is described as being implemented by the system and configuration shown in FIG. 1, although it should be understood that the method may alternatively be implemented by any other suitable system in any other suitable configuration.

[0031] Referring to FIG. 2, method includes applying **200** a first pattern of electrical stimulation to a first sub-population of targeted neurological tissue of a subject. For example, the controller **112** may be configured to control the pulse generator **114** to generate electrical signals that produce a predefined pattern of electrical stimulation to a particular sub-population of dorsal column nerve fibers. One or more contacts of the electrode array **104** may be placed in electrical communication and in position to apply the electrical stimulation to the sub-population of dorsal column nerve fibers. The pattern of electrical stimulation may include regular temporal patterns of stimulation (*i.e.*, constant interpulse intervals) or non-regular temporal patterns of stimulation (*i.e.*, interpulse intervals that vary in time).

[0032] The method of FIG. 2 includes applying **202** a second pattern of electrical stimulation to a second sub-population of targeted neurological tissue of the subject. It should be noted that steps **200** and **202** may occur simultaneously or one after the other. The second pattern of electrical stimulation may be applied at a different frequency than the first pattern of electrical stimulation. For example, the controller **112** may be configured to control the pulse generator **114** to generate electrical signals that produce a predefined pattern of electrical

stimulation to another sub-population of dorsal column nerve fibers. Another one or more contacts of the electrode array **104** may be placed in electrical communication and in position to apply the electrical stimulation to the other sub-population of dorsal column nerve fibers. The pattern of electrical stimulation may be applied at multiple different frequencies and at different timings. Further, for example, the patterns may be applied at different frequencies that are multiples of each other. The patterns may include regular temporal patterns of stimulation (*i.e.*, constant interpulse intervals) or non-regular temporal patterns of stimulation (*i.e.*, interpulse intervals that vary in time).

[0033] The method of FIG. 2 includes controlling **204** the first and second patterns of electrical stimulation for optimizing the effects of stimulation. For example, the patterns may optimize suppression of activity of WDR neurons and thereby achieve pain relief. For example, the controller **112** may control the pulse generator **114** to output electrical signals to the electrode array **104** for optimizing suppression of activity of WDR neurons. In an example, the controller **112** may implement an algorithm for optimization. In another example, the controller **112** may receive user input for control of the application of the patterns of electrical stimulation.

[0034] In accordance with embodiments, systems disclosed herein may provide multi-frequency, multi-fiber SCS for achieving suppression of nociceptive information from the spinal cord. Computational modeling work indicated that the activity of WDR neurons in the spinal cord that transmit nociceptive information (*i.e.*, pain signals) to the brain can be better suppressed by stimulation of sub-populations of dorsal column nerve fibers at different timings than by uniform stimulation at the same equivalent frequency. For example, FIG. 3 illustrates graphs showing that delivering SCS at different timings through different fiber populations can result in greater efficacy in response to a 1 Hz peripheral input. Referring to FIG. 3, SCS applied using the two population input set denoted in the right reduces the activity of the WDR neurons responsible for relaying nociceptive (pain) information to the brain to a greater extent and over a wider frequency range than application of SCS using the uniform input set denoted on the left across several simulated positions of the SCS electrode relative to the WDR neuron. The dotted line denotes the average frequency of the WDR neuron's activity when no SCS was applied. This finding indicates that delivering multiple frequencies of SCS to multiple sub-populations of dorsal fibers will yield more effective (reduction in WDR firing) or more efficient (fewer SCS pulses delivered, and thereby less power consumption) SCS than constant frequency stimulation.

[0035] FIG. 4 illustrates a schematic of an example computational model for model-based design and evaluation of patterns of SCS. Referring to FIG. 4, the computational model may include a network of biophysical neurons that are connected to represent a dorsal horn pain processing network. Inputs to the model include 30 A and 30 C primary afferent fibers that convey information from the periphery, and SCS may be delivered to the network via the A fibers to simulate dorsal column fiber activation. Multiple A/C fibers and excitatory interneurons may be used to account for the effects of temporal summation on neuronal activity as well as to add variability to the inputs. In addition, to simulate realistic signal propagation from a peripheral or dorsal column nerve fiber, propagation delays based on the conduction velocities of A and C fibers may be incorporated into all inputs for all simulations. In FIG. 4, the “IN” node represents inhibitory interneuron, the “EX” node represents excitatory interneuron, the “WDR” node represents WDR projection neurons. Synapses **400** denote excitatory connections. Synapse **402** denotes an inhibitory connection. SCS using the optimization algorithm may be delivered via the A-fiber input.

[0036] FIGs. 5A and 5B illustrate graphs showing example patterns of activity in peripheral primary afferent fibers. Referring to FIG. 5A, the graphs show representative uniform 1 Hz inputs. FIG. 5B shows randomized inputs representing a neuroma. A 5-second interval (x-axis) of each is shown for all fiber inputs (y-axis; split by A and C fibers). Each black dot on the graph represents a time point at which a spike is registered by a corresponding input to the model. In FIG. 5B, 30% of the A-fiber inputs exhibit bursting behavior. During multi-frequency SCS, the bursty inputs were split such that half of these inputs received one frequency of stimulation while the other half received the other frequency.

[0037] Computational experiments were conducted to demonstrate the utility of the present subject matter. For example, FIGs. 6A and 6B illustrate graphs showing 1-second long examples of non-harmonic (*i.e.*, a first stimulation frequency applied to subpopulation one and a second stimulation frequency applied to subpopulation two were not integer multiples of one another) and harmonic (*i.e.*, a first stimulation frequency applied to subpopulation one and a second stimulation frequency applied to subpopulation two were integer multiples of one another) multi-frequency SCS, respectively. Briefly, one second of simulation time was allowed to elapse to allow the model to initialize, and periphery sensory input including either a constant 1 Hz pulse train synchronized across all fibers or a random spike train based on a Poisson process whose characteristics match those taken from the firing behavior of a peripheral neuroma

(as shown in FIGs. 5A and 5B) was then delivered for 15 seconds. SCS using two frequencies – with half of the input A fibers receiving one frequency (a first subpopulation receiving a first stimulation frequency) and the other half receiving the other frequency (a second subpopulation receiving a second stimulation frequency) – was then delivered for the remaining 5 seconds while the output of the WDR neuron was recorded; these frequencies may be harmonic or non-harmonic (see FIG. 7 for example). In harmonic multi-frequency SCS, the higher frequency of stimulation (40 Hz) was set to be an integer multiple of the lower frequency of stimulation (10 Hz). In non-harmonic multi-frequency SCS, the lower frequency was drawn from a uniform random distribution ranging from 40 Hz to 50 Hz and checked to ensure that the higher frequency was not an integer multiple of the lower frequency. The output of the WDR neuron as well as the number of pulses used during stimulation were compared with the corresponding metrics resulting from the first lower frequency SCS delivered to all A fibers, the average of the two applied frequencies delivered to all A fibers, and the second higher frequency delivered to all A fibers. In FIGs. 6A and 6B, each black dot on the graphs represent a time point at which an SCS spike is fed into an A-fiber unit to the computational model during a time period shown in FIG. 7, which illustrates a timeline of each experimental run in accordance with embodiments of the present disclosure. In both the non-harmonic and harmonic cases, half of the A-fibers receive the lower frequency while the other half of the A-fibers receive the higher frequency.

[0038] Referring to FIG. 7, SCS is delivered following a brief model initialization period and 15 seconds of conditioning stimulation using either constant 1 Hz or randomized inputs similar to those recorded from neuromas in live preparations. The output of the WDR neuron as well as the average frequency of SCS delivered – a measure of power consumption – are used to gauge respectively the efficacy and efficiency of multi-frequency SCS (*i.e.*, a first stimulation frequency applied to a first subpopulation of nerve fibers and a second stimulation frequency applied to second subpopulation of nerve fibers) versus conventional SCS (one stimulation frequency delivered to all nerve fibers).

[0039] In experimentation, it has been shown that the techniques and systems disclosed herein are effective at suppressing WDR neuron behavior and efficient with respect to pulses delivered (power consumption) versus high frequency stimulation through testing of the prototype algorithm using a computational model of pain. In the experiments, it was shown that the application of non-harmonic and harmonic multi-frequency SCS inhibits the activity of the WDR neuron compared to the case in which no SCS was applied. The application of 12 Hz / 42

Hz non-harmonic SCS reduced the activity of the WDR neuron by 92.7% in response to a 1 Hz input (41 Hz to 3 Hz) and by 88.0% (73 Hz to 8.8 Hz) in response to a neuropathic input. Application of 10 Hz / 50 Hz harmonic SCS reduced the activity of the WDR neuron by 91.5 % (41 Hz to 3.5 Hz) in response to a 1 Hz input and by 90.8% (73 Hz to 6.75 Hz) in response to a neuropathic input. For example, FIG. 8 illustrates raster plots depicting example activity of a WDR neuron, such as shown in FIG. 4, during the period of time in which multi-frequency SCS may be delivered. Referring to FIG. 8, each blank line on the graph represents a time point at which a spike is output by the WDR neuron. The top row depicts the activity of the WDR neuron in response to the 1 Hz input and the neuropathic input during no SCS. The middle row depicts the activity of the WDR neuron in response to these inputs during non-harmonic (12 Hz / 42 Hz) multi-frequency SCS. The bottom row depicts the activity of the WDR neuron in response to these inputs during harmonic frequency (10 Hz / 42 Hz) multi-frequency SCS.

[0040] Further, in experimentation, it was demonstrated that both non-harmonic SCS and harmonic SCS are more effective at suppressing WDR neuronal activity versus single frequency stimulation at low frequencies and more efficient at suppressing WDR neuronal activity versus single frequency stimulation at high frequencies during a 1 Hz peripheral input. For example, FIG. 9 illustrates raster plots depicting comparisons of SCS efficacy and efficiency between multi-frequency SCS and conventional SCS at fixed frequency in response to a 1 Hz conditioning input. Each black line represents a time point at which a spike is output by the WDR neuron. The output from a given pair of stimulation frequencies is compared to the output due to the stimulation at the lower frequency (12 Hz, 10 Hz – top) and higher frequency (42 Hz, 40 Hz – bottom). Both non-harmonic and harmonic SCS significantly reduce the activity of the WDR neuron versus constant frequency stimulation at the lower frequency (top of FIG. 9) in response to a 1 Hz input (92.7% vs. 17.1% - non-harmonic; 90.8% vs. 15.9% - harmonic). Although simulation using the higher frequency reduces the activity of the WDR neuron to a slightly greater degree, multi-frequency SCS is able to achieve comparable results (92.7% non-harmonic and 90.8% harmonic reduction vs. 97.0% and 96.3% reduction during respective single frequency stimulation) using an average frequency that is 15 Hz lower than the higher frequency of SCS (27 Hz vs. 42 Hz non-harmonic; 25 Hz vs. 40 Hz harmonic), corresponding to 35.7% (non-harmonic) and 37.5% (harmonic) less power used if stimulation frequency is taken as a direct measure of power consumption (bottom of FIG. 9).

[0041] The trends observed above hold when SCS was applied during a

neuropathic input to the computational model as well. Both non-harmonic and harmonic SCS significantly reduce the activity of the WDR neuron versus constant frequency stimulation at the lower (88.0% vs. 47.7% - non-harmonic; 90.8% vs. 38.0% - harmonic – see FIG. 10 at the top). In addition, multi-frequency SCS is able to achieve comparable results (88.0% non-harmonic and 90.8% harmonic reduction vs. 88.7% and 90.8% reduction during respective single frequency stimulation) using an average frequency that is 15 Hz lower than the higher frequency of SCS (35.7% (non-harmonic) and 37.5% (harmonic) less power – see the bottom of FIG. 10). FIG. 10 illustrates Raster plots depicting comparisons of SCS efficacy (WDR neuronal output) and efficiency (average stimulation frequency) between multi-frequency SCS and conventional SCS at a fixed frequency in response to a neuropathic input. Each black line on the graph represents a time point at which a spoke is output by the WDR neuron. The output from a given pair of stimulation frequencies is compared to the output due to stimulation at the lower frequency (12 Hz, 10 Hz – top) and higher frequency (42 Hz, 40 Hz – bottom).

[0042] Further, it was shown experimentally that harmonic multi-frequency stimulation is both more effective and more efficient at suppressing WDR neuronal activity during a neuropathic input (see FIGs. 11A and 11B, for example). To this end, 3 combinations of harmonic SCS were tested against the neuropathic input – 10/20 Hz, 10/30 Hz, and 10/40 Hz SCS – and compared efficacy (WDR neuronal output) against stimulation at the lower frequency (10 Hz) and at the average frequency (15 Hz, 25 Hz, 35 Hz) as well as efficiency against the higher frequency (20 Hz, 30 Hz, 40 Hz). In all cases, harmonic multi-frequency SCS suppressed WDR neuronal activity (*i.e.*, is more effective) to a greater extent than single frequency stimulation at the lower frequency (56.5% - 10/20 Hz; 82.9% - 10/30 Hz, 88.0% - 10/40 Hz versus 38.0% 10 Hz). In addition, multi-frequency SCS was also more efficient and in some cases more effective than stimulation at the higher frequency: stimulation using 10/20 Hz, 10/30 Hz, and 10/40 Hz reduced the WDR neuron's activity by 56.5%, 82.9%, and 88.0%, respectively, versus 61.0%, 60.3%, and 88% reduction by stimulation using 20 Hz, 30 Hz, and 40 Hz, but stimulation using 10/20 Hz, 10/30 Hz, and 10/40 Hz was 25.0%, 33.3%, and 37.5% more efficient than stimulation using 20 Hz, 30 Hz, and 40 Hz alone. Finally, stimulation using harmonic frequencies (*i.e.*, single frequency stimulation using equal power consumption): 10/20 Hz stimulation suppressed WDR activity by 82.9% versus 61.0% using 20 Hz constant stimulation; 10/40 Hz stimulation suppressed WDR activity by 88.0% versus 68.2% using 25 Hz constant stimulation.

[0043] FIGs. 11A and 11B show raster plots depicting comparisons of SCS efficacy (WDR neuronal output) and efficiency (average stimulation frequency) between several combinations of harmonic multi-frequency SCS and conventional SCS at a fixed frequency in response to a neuropathic input. Each black line on the graph represents a time point at which a spike is output by the WDR neuron. The output from a given pair of stimulation frequencies is compared to the output due to stimulation at the lower frequency (10 Hz – top), average frequency (15 Hz, 20 Hz, 25 Hz – middle), and higher frequency (20 Hz, 30 Hz, 40 Hz – bottom).

[0044] In accordance with embodiments, systems and methods of the present disclosure may be implemented as an algorithm within an SCS pulse generator device. An on-board controller may deliver multiple frequencies of SCS through different output channels to different contacts on the spinal cord stimulation electrode. By virtue of stimulation through multiple contacts, different populations of axons (*e.g.*, sub-populations of dorsal column nerve fibers) traversing the dorsal column may be activated at different frequencies, resulting in greater suppression of the neurons responsible for transmitting nociceptive information to the brain. Values of the stimulation frequencies and the electrodes through which these frequencies are delivered can be input by either a physician or a patient through a user interface. Alternatively, the device can be pre-programmed with specific combinations of frequencies to use. The applied frequencies can be multiples of each other (harmonic) or not (non-harmonic), and they may or may not be offset from each other at the start of stimulation. In addition, multi-frequency SCS may be limited to 2 frequencies, as many frequencies and axon populations as the stimulation technology will allow can be delivered to the patient. The algorithm may be toggled on and off (*e.g.*, between multi-frequency and single frequency SCS) by either the physician or patient, or it can be coupled to an internal feedback-driven algorithm for automatic control.

[0045] FIG. 12 illustrates a regular, constant frequency stimulation train wherein the interpulse intervals are constant in time and examples of non-regular temporal patterns of stimulation wherein the interpulse intervals vary in time.

[0046] The present subject matter may be a system, a method, and/or a computer program product implemented on an SCS device, a smartphone, tablet computer, or the like. The computer program product may include a computer readable storage medium (or media) having computer readable program instructions thereon for causing a processor to carry out aspects of the present subject matter.

[0047] The computer readable storage medium can be a tangible device that can retain and store instructions for use by an instruction execution device. The computer readable storage medium may be, for example, but is not limited to, an electronic storage device, a magnetic storage device, an optical storage device, an electromagnetic storage device, a semiconductor storage device, or any suitable combination of the foregoing. A non-exhaustive list of more specific examples of the computer readable storage medium includes the following: a portable computer diskette, a hard disk, a random access memory (RAM), a read-only memory (ROM), an erasable programmable read-only memory (EPROM or Flash memory), a static random access memory (SRAM), a portable compact disc read-only memory (CD-ROM), a digital versatile disk (DVD), a memory stick, a floppy disk, a mechanically encoded device such as punch-cards or raised structures in a groove having instructions recorded thereon, and any suitable combination of the foregoing. A computer readable storage medium, as used herein, is not to be construed as being transitory signals *per se*, such as radio waves or other freely propagating electromagnetic waves, electromagnetic waves propagating through a waveguide or other transmission media (e.g., light pulses passing through a fiber-optic cable), or electrical signals transmitted through a wire.

[0048] Computer readable program instructions described herein can be downloaded to respective computing/processing devices from a computer readable storage medium or to an external computer or external storage device via a network, for example, the Internet, a local area network, a wide area network and/or a wireless network. The network may comprise copper transmission cables, optical transmission fibers, wireless transmission, routers, firewalls, switches, gateway computers and/or edge servers. A network adapter card or network interface in each computing/processing device receives computer readable program instructions from the network and forwards the computer readable program instructions for storage in a computer readable storage medium within the respective computing/processing device.

[0049] Computer readable program instructions for carrying out operations of the present subject matter may be assembler instructions, instruction-set-architecture (ISA) instructions, machine instructions, machine dependent instructions, microcode, firmware instructions, state-setting data, or either source code or object code written in any combination of one or more programming languages, including an object oriented programming language such as Java, Smalltalk, C++ or the like, and conventional procedural programming languages, such as the "C" programming language or similar programming languages. The computer readable

program instructions may execute entirely on the user's computer, partly on the user's computer, as a stand-alone software package, partly on the user's computer and partly on a remote computer or entirely on the remote computer or server. In the latter scenario, the remote computer may be connected to the user's computer through any type of network, including a local area network (LAN) or a wide area network (WAN), or the connection may be made to an external computer (for example, through the Internet using an Internet Service Provider). In some embodiments, electronic circuitry including, for example, programmable logic circuitry, field-programmable gate arrays (FPGA), or programmable logic arrays (PLA) may execute the computer readable program instructions by utilizing state information of the computer readable program instructions to personalize the electronic circuitry, in order to perform aspects of the present subject matter.

[0050] Aspects of the present subject matter are described herein with reference to flow chart illustrations and/or block diagrams of methods, apparatus (systems), and computer program products according to embodiments of the subject matter. It will be understood that each block of the flow chart illustrations and/or block diagrams, and combinations of blocks in the flow chart illustrations and/or block diagrams, can be implemented by computer readable program instructions.

[0051] These computer readable program instructions may be provided to a processor of a general purpose computer, special purpose computer, or other programmable data processing apparatus to produce a machine, such that the instructions, which execute via the processor of the computer or other programmable data processing apparatus, create means for implementing the functions/acts specified in the flow chart and/or block diagram block or blocks. These computer readable program instructions may also be stored in a computer readable storage medium that can direct a computer, a programmable data processing apparatus, and/or other devices to function in a particular manner, such that the computer readable storage medium having instructions stored therein comprises an article of manufacture including instructions which implement aspects of the function/act specified in the flow chart and/or block diagram block or blocks.

[0052] The computer readable program instructions may also be loaded onto a computer, other programmable data processing apparatus, or other device to cause a series of operational steps to be performed on the computer, other programmable apparatus or other device to produce a computer implemented process, such that the instructions which execute on the computer, other programmable apparatus, or other device implement the functions/acts

specified in the flow chart and/or block diagram block or blocks.

[0053] The flow chart and block diagrams in the Figures illustrate the architecture, functionality, and operation of possible implementations of systems, methods, and computer program products according to various embodiments of the present subject matter. In this regard, each block in the flow chart or block diagrams may represent a module, segment, or portion of instructions, which comprises one or more executable instructions for implementing the specified logical function(s). In some alternative implementations, the functions noted in the block may occur out of the order noted in the figures. For example, two blocks shown in succession may, in fact, be executed substantially concurrently, or the blocks may sometimes be executed in the reverse order, depending upon the functionality involved. It will also be noted that each block of the block diagrams and/or flow chart illustration, and combinations of blocks in the block diagrams and/or flow chart illustration, can be implemented by special purpose hardware-based systems that perform the specified functions or acts or carry out combinations of special purpose hardware and computer instructions.

[0054] Any patents or publications mentioned in this specification are indicative of the levels of those skilled in the art to which the present subject matter pertains. These patents and publications are herein incorporated by reference to the same extent as if each individual publication was specifically and individually indicated to be incorporated by reference.

[0055] One skilled in the art will readily appreciate that the present subject matter is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. The present examples along with the methods described herein are presently representative of various embodiments, are exemplary, and are not intended as limitations on the scope of the present subject matter. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the present subject matter as defined by the scope of the claims.

CLAIMS

What is Claimed is:

1. A method comprising:
 - applying a first pattern of electrical stimulation to a first sub-population of targeted neurological tissue of a subject;
 - applying a second pattern of electrical stimulation to a second sub-population of targeted neurological tissue of the subject, the second pattern of electrical stimulation being applied at a different frequency than the first pattern of electrical stimulation; and
 - controlling the first and second patterns of electrical stimulation for optimizing the effects of stimulation.
2. The method of claim 1, wherein applying a first pattern of electrical stimulation comprises:
 - placing a first contact in electrical communication with the first sub-population of targeted neurological tissue; and
 - using the first contact for applying the first pattern of electrical stimulation to the first sub-population of targeted neurological tissue.
3. The method of claim 2, wherein applying a second pattern of electrical stimulation comprises:
 - placing a second contact in electrical communication with the second sub-population of targeted neurological tissue; and
 - using the second contact for applying the second pattern of electrical stimulation to the second sub-population of targeted neurological tissue.
4. The method of claim 1, wherein the first and second patterns of electrical stimulation are applied at different timings.
5. The method of claim 1, further comprising receiving user input for control of the application of the first and second patterns of electrical stimulation.
6. The method of claim 1, wherein the frequency of the first pattern of electrical stimulation is a multiple of the frequency of the second pattern of electrical stimulation.
7. The method of claim 1, further comprising:
 - providing an electrode comprising a first contact and a second contact;
 - placing the first contact and the second contact in electrical communication with the first sub-population and the second sub-population, respectively, of the targeted neurological tissue;

using the first contact for applying the first pattern of electrical stimulation to the first sub-population of targeted neurological tissue; and

using the second contact for applying the second pattern of electrical stimulation to the second sub-population of targeted neurological tissue.

8. The method of claim 1, wherein the targeted neurological tissue comprises dorsal column nerve fibers.

9. The method of claim 1, further comprising selecting the first and second patterns of electrical stimulation from among other patterns for one of maximizing suppression of WDR neuron firing and minimizing average stimulation frequency.

10. The method of claim 1, further comprising selecting the first and second patterns of electrical stimulation from among other patterns for maximizing suppression of pain and minimizing average stimulation frequency.

11. The method of claim 1, further comprising:

selecting the first and second patterns of electrical stimulation from among other patterns for one of maximizing efficacy and efficiency;

minimizing average stimulation frequency is a proxy for efficiency; and

suppressing wide-dynamic range neuron firing in a prior simulation is a proxy for efficacy.

12. The method of claim 1, further comprising selecting the first and second patterns of electrical stimulation from among other patterns for one of minimizing patient pain and device power consumption during clinical use.

13. The method of claim 1, further comprising selecting the first and second patterns of electrical stimulation from among other patterns for minimizing average stimulation frequency while maintaining suppression of pain.

14. The method of claim 1, further comprising selecting the first and second patterns of electrical stimulation from among other patterns for optimizing efficacy and efficiency;
suppression of activity of wide-dynamic range neurons in a prior simulation (efficacy)
minimizing average stimulation frequency in a prior simulation (efficiency)
minimizing patient pain and stimulation frequency in clinical practice.

15. A system comprising:

a pulse generator;

one or more electrodes; and

a controller configured to:

control the pulse generator to output a first electrical signal to the one or more electrodes for applying a first pattern of electrical stimulation to a first sub-population of targeted neurological tissue of a subject;

control the pulse generator to output a second electrical signal to the one or more electrodes for applying a second pattern of electrical stimulation to a second sub-population of targeted neurological tissue of the subject, the second pattern of electrical stimulation being applied at a different frequency than the first pattern of electrical stimulation; and

control the first and second patterns of electrical stimulation for optimizing the effects of stimulation.

16. The system of claim 15, wherein the one or more electrodes comprise a first contact in electrical communication with the first sub-population of targeted neurological tissue for application of the first pattern of electrical stimulation to the first sub-population of targeted neurological tissue.

17. The system of claim 16, wherein the one or more electrodes comprises a second contact in electrical communication with the second sub-population for application of the second pattern of electrical stimulation to the second sub-population of targeted neurological tissue.

18. The system of claim 15, wherein the controller is configured to control the pulse generator to apply the multiple different frequencies of the first pattern at different timings, and to apply the multiple different frequencies of the second pattern at different timings.

19. The system of claim 15, further comprising a user interface for receipt of user input for control of the application of the first and second patterns of electrical stimulation.

20. The system of claim 15, wherein the frequency of the first pattern of electrical stimulation is a multiple of the frequency of the second pattern of electrical stimulation.

21. The system of claim 15, wherein the one or more electrodes comprise a first contact and a second contact, wherein the first contact and the second contact are positioned in electrical communication with the first sub-population and the second sub-population, respectively, of the targeted neurological tissue, and

wherein the controller is configured to:

control the pulse generator to output an electrical signal to the first contact for applying the first pattern of electrical stimulation to the first sub-population of targeted

neurological tissue; and

control the pulse generator to output an electrical signal to the second contact for applying the second pattern of electrical stimulation to the second sub-population of targeted neurological tissue.

22. The system of claim 15, wherein the targeted neurological tissue comprises dorsal column nerve fibers.
23. The system of claim 15, wherein the controller is configured to select the first and second patterns of electrical stimulation from among other patterns for one of maximizing suppression of WDR neuron firing (efficacy) and minimizing average stimulation frequency.
24. The system of claim 15, wherein at least one processor and memory are configured to select the first and second patterns of electrical stimulation from among other patterns for maximizing suppression of pain and minimizing average stimulation frequency.
25. The system of claim 15, wherein at least one processor and memory are configured to:
 - select the first and second patterns of electrical stimulation from among other patterns for one of maximizing efficacy and efficiency;
 - minimize average stimulation frequency in a prior simulation is a proxy for efficiency;
 - and
 - suppress wide-dynamic range neuron firing in a prior simulation is a proxy for efficacy.
26. The system of claim 15, wherein at least one processor and memory are configured to select the first and second patterns of electrical stimulation from among other patterns for one of minimizing patient pain and device power consumption during clinical use.
27. The system of claim 15, wherein at least one processor and memory are configured to select the first and second patterns of electrical stimulation from among other patterns for minimizing average stimulation frequency while maintaining suppression of pain.
28. The system of claim 15, further comprising a computing device including a user interface for receipt of selection of the first and second patterns.
29. The system of claim 15, wherein at least one processor and memory are configured to select the first and second patterns of electrical stimulation from among other patterns for optimizing suppression of activity of wide-dynamic range neurons.

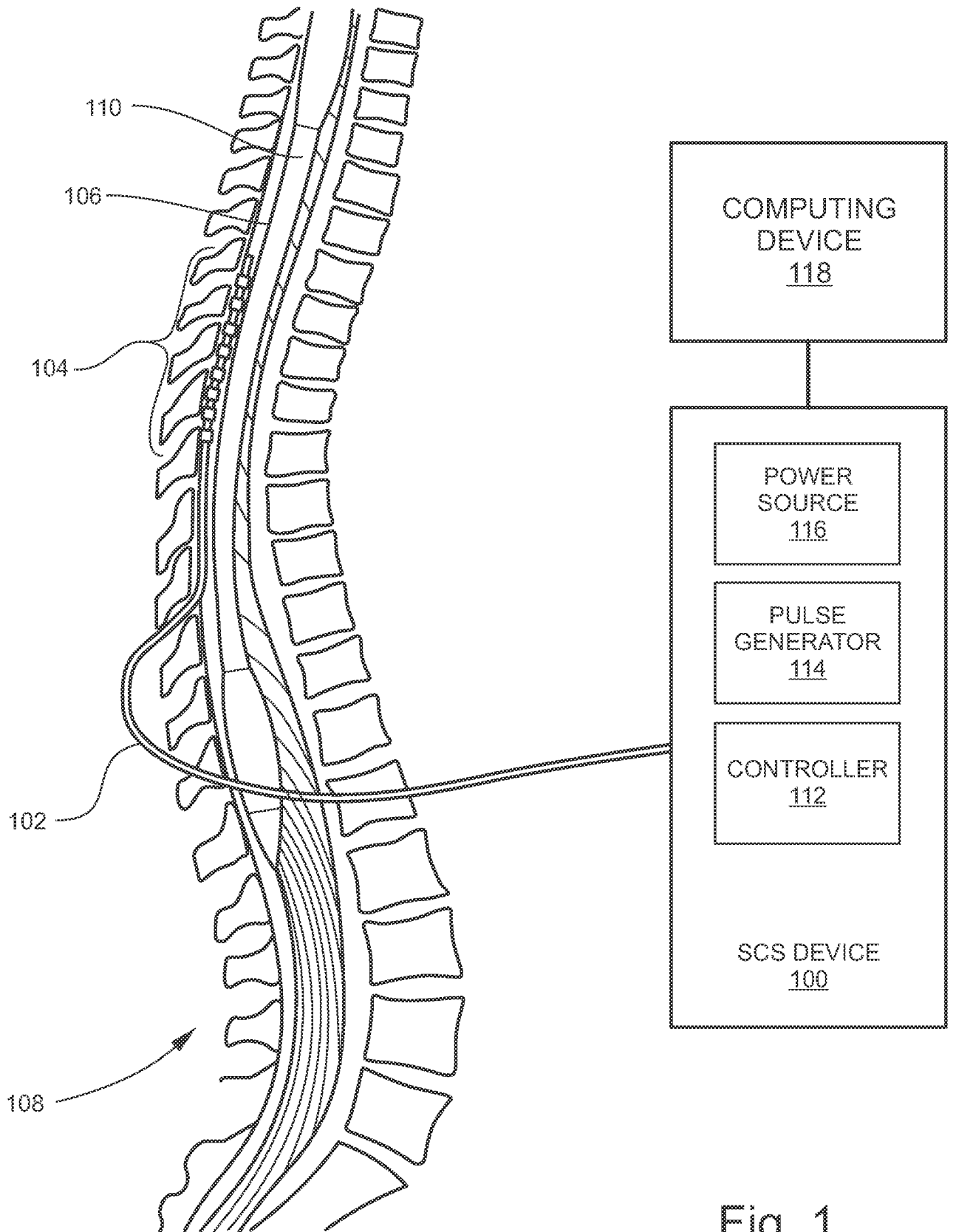


Fig. 1

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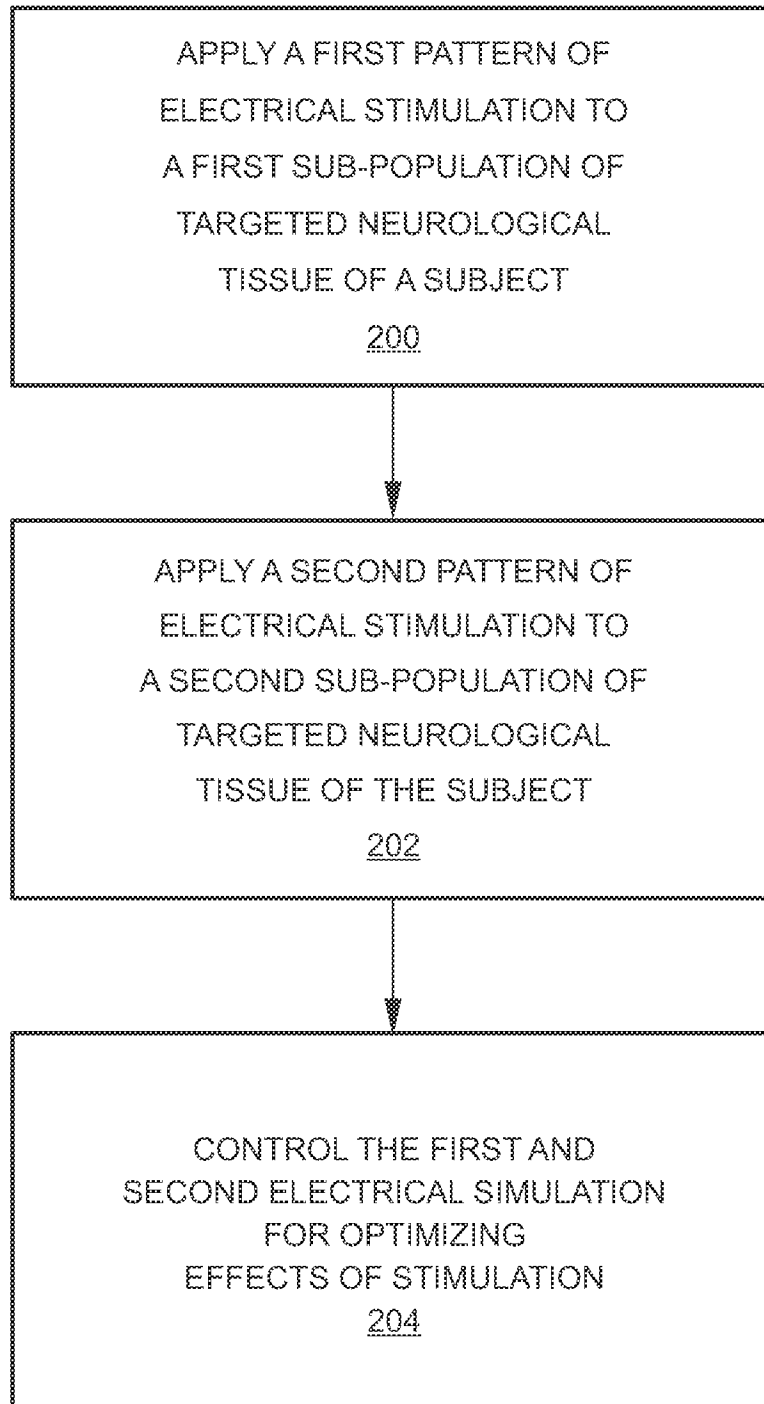
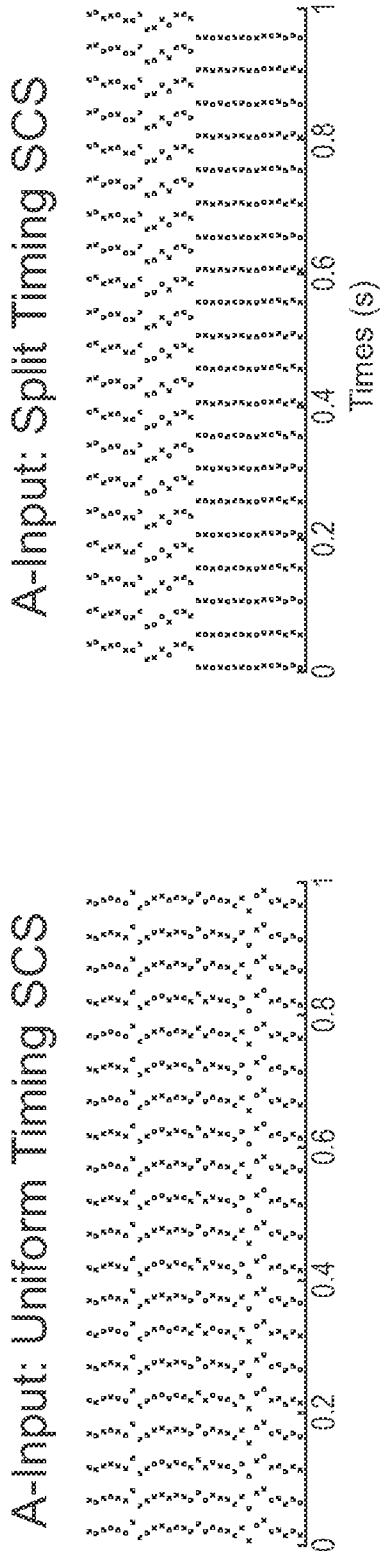
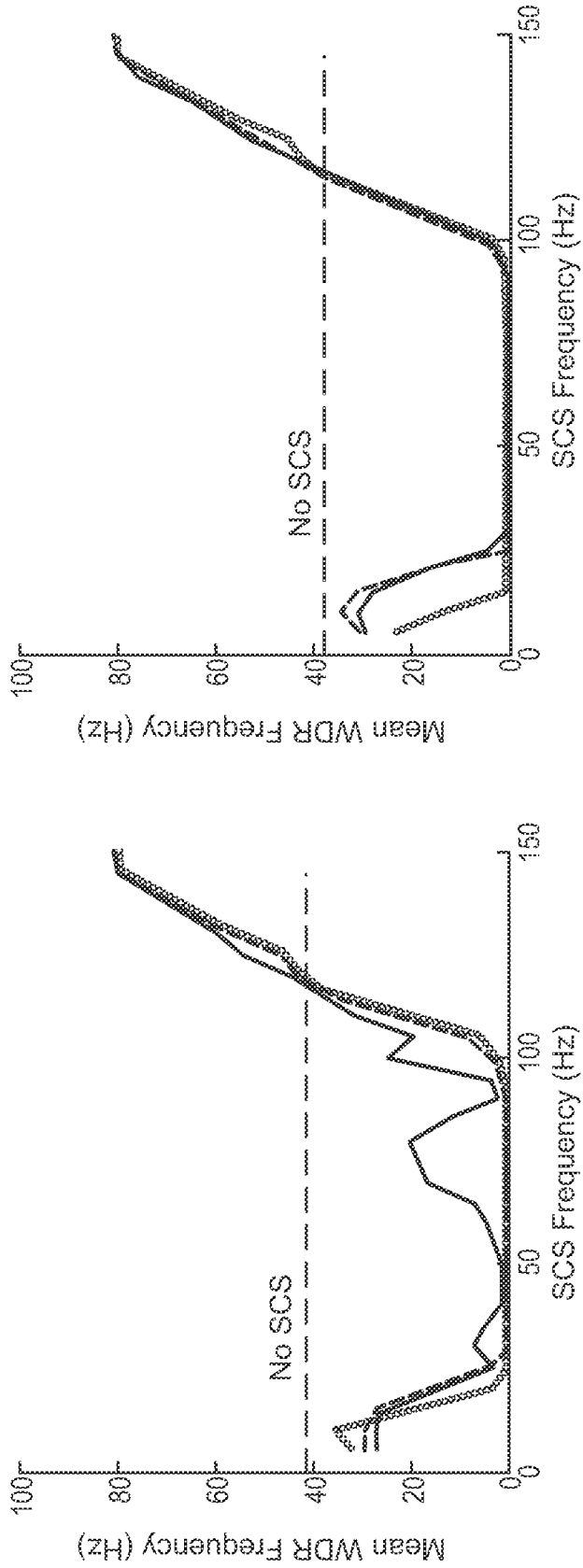


Fig. 2



WDR Neuron Output vs. SCS Frequency



Distance between SCS lead and WDR

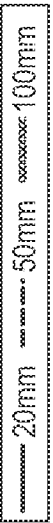
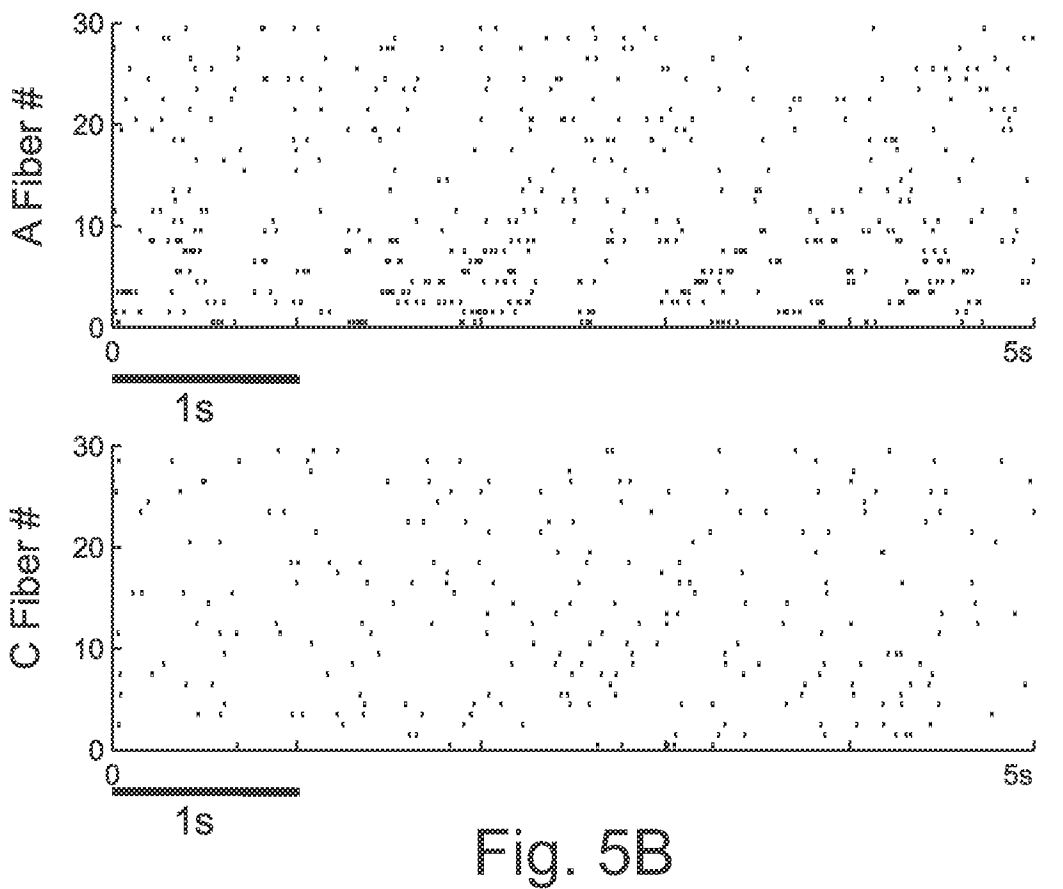
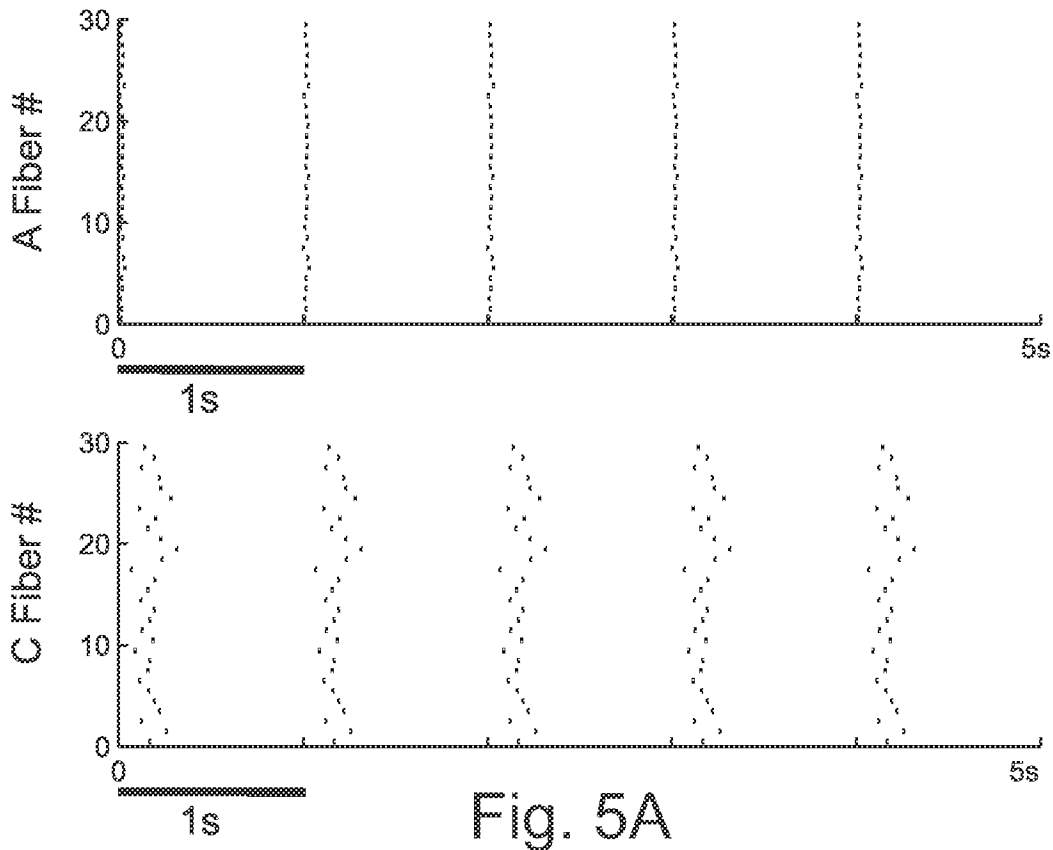
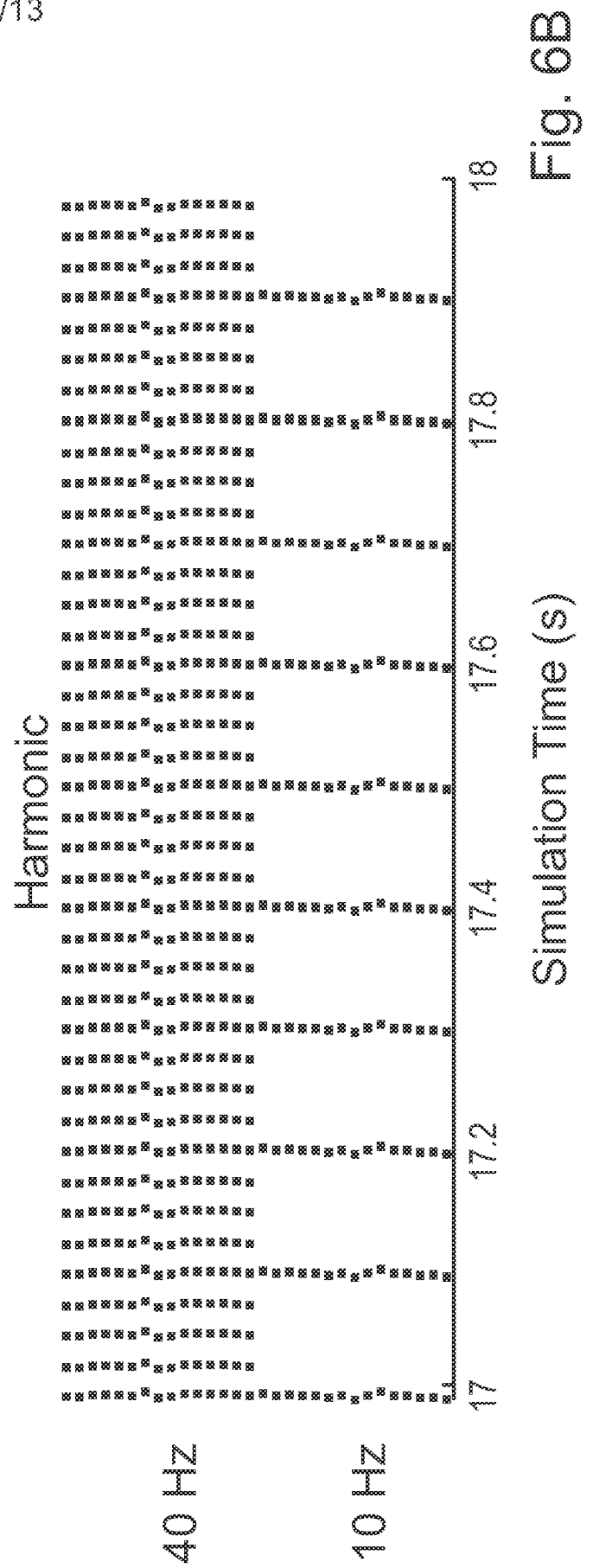
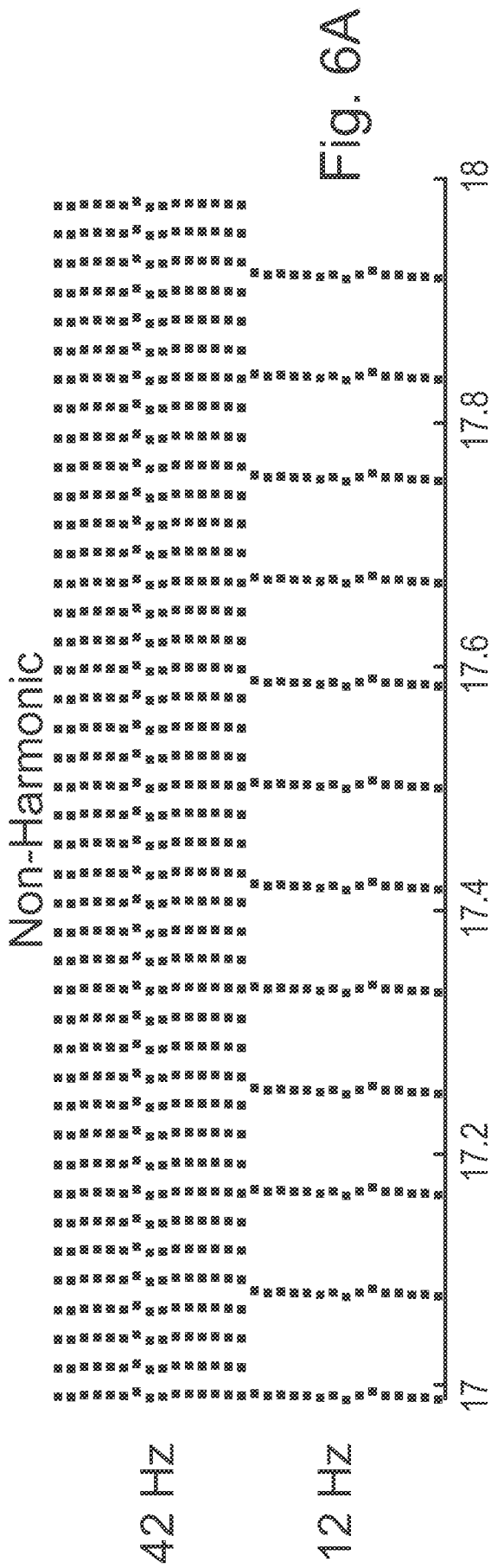
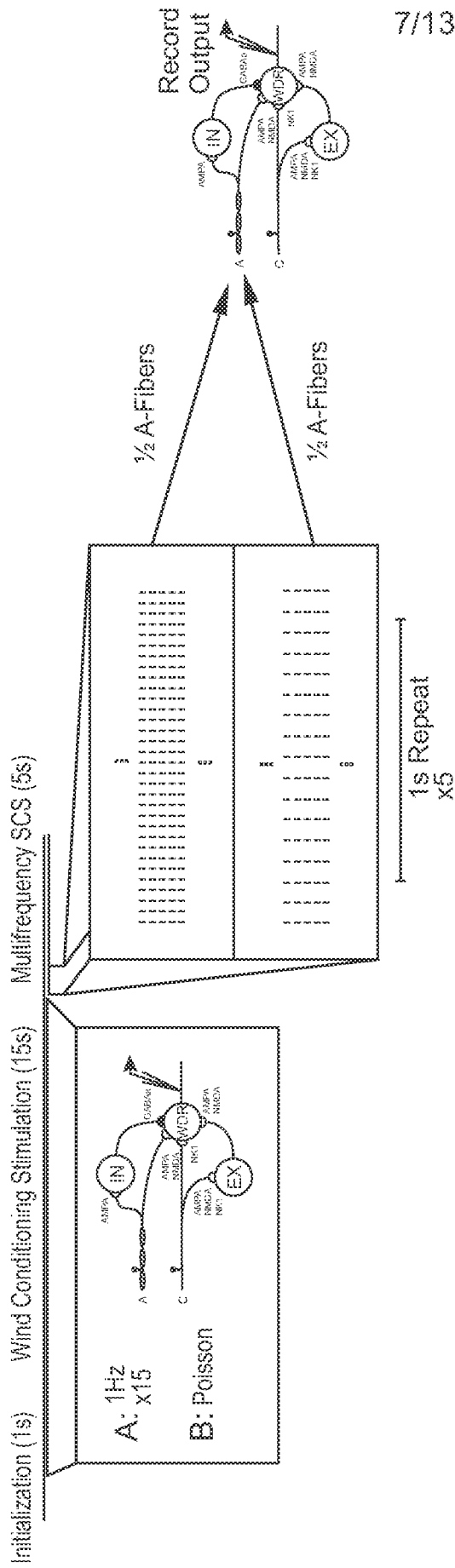


Fig. 3

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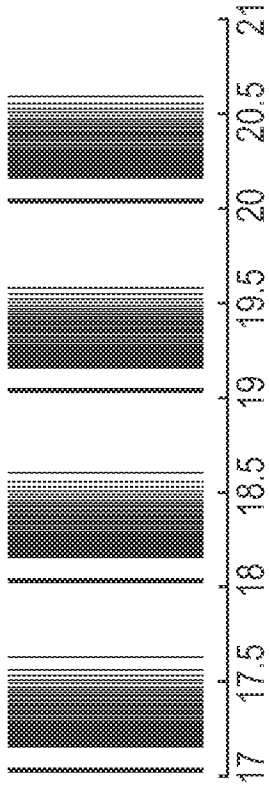


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Fig. 7

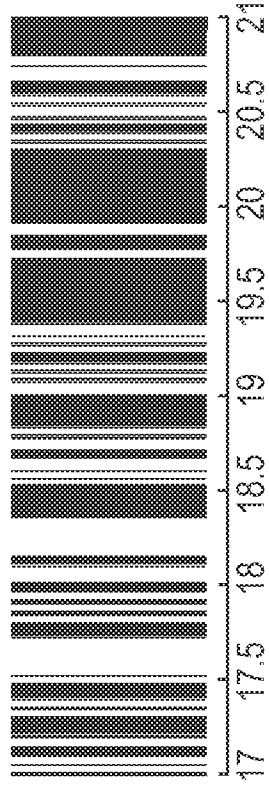
1 Hz Input

1 Hz: WDR Average Frequency 41Hz

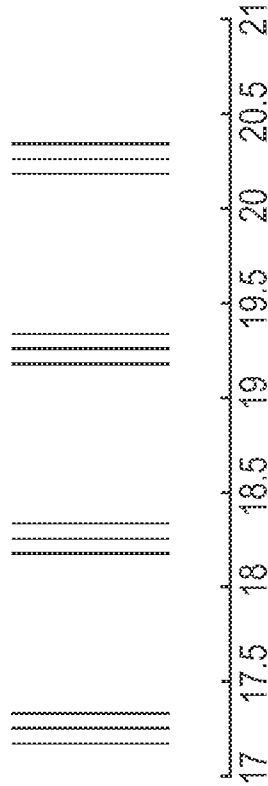


Neuropathic Input

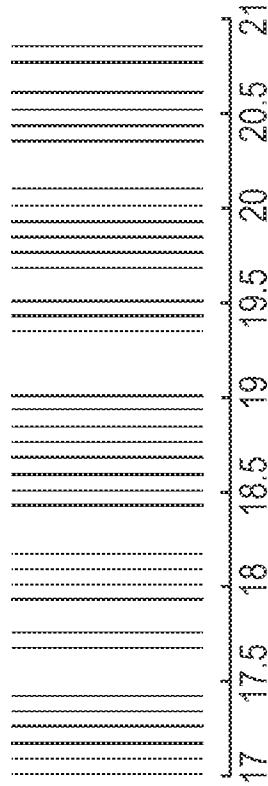
12/42 Hz: WDR Average Frequency 73Hz



12/42 Hz: WDR Average Frequency 3Hz

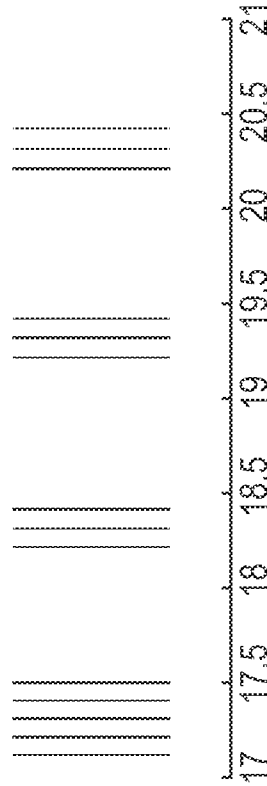


12/42 Hz: WDR Average Frequency 8.75Hz

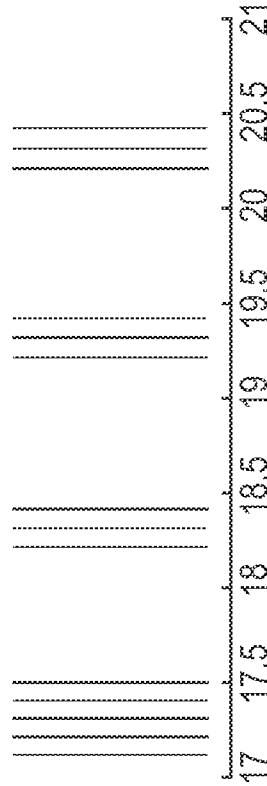


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10/40 Hz: WDR Average Frequency 3.5Hz



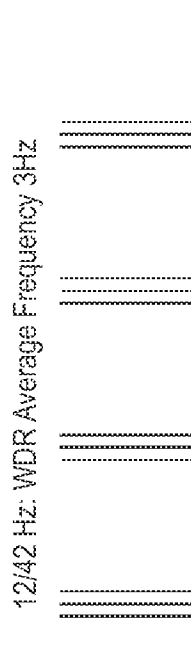
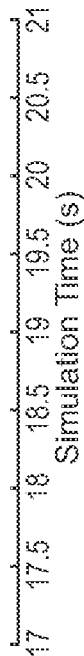
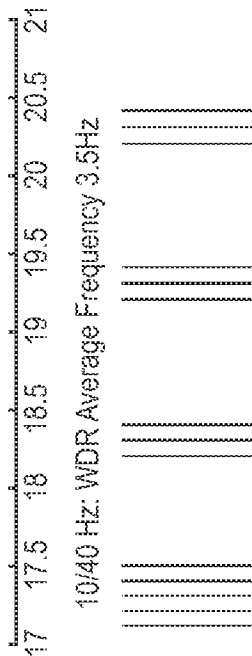
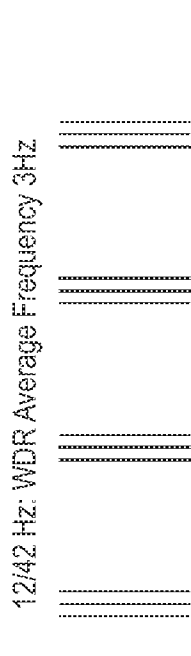
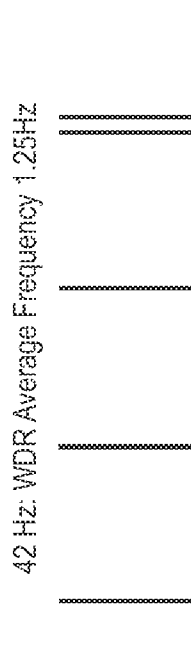
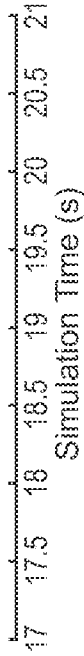
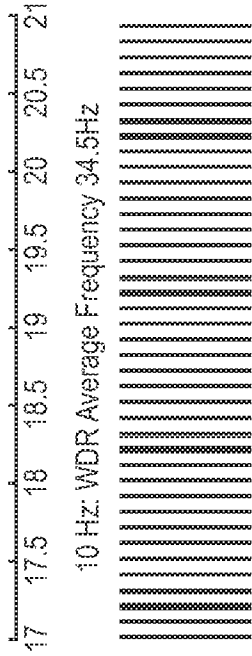
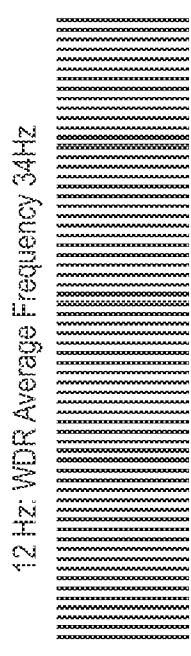
10/40 Hz: WDR Average Frequency 6.75Hz



Simulation Time (s)

Simulation Time (s)

Fig. 8



Efficacy

Efficiency

Fig. 9



Fig. 10

Efficiency

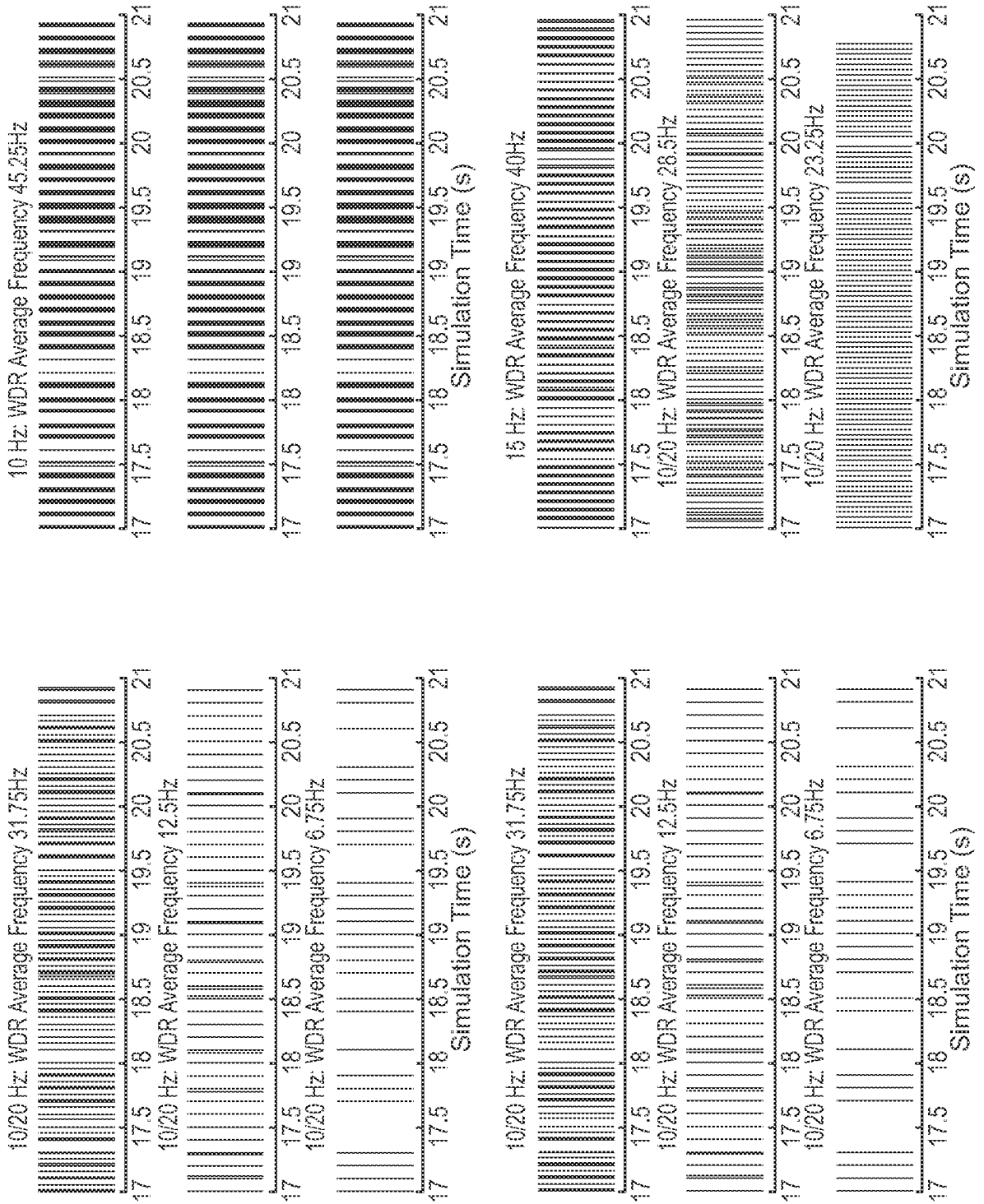


Fig. 11A

Efficiency

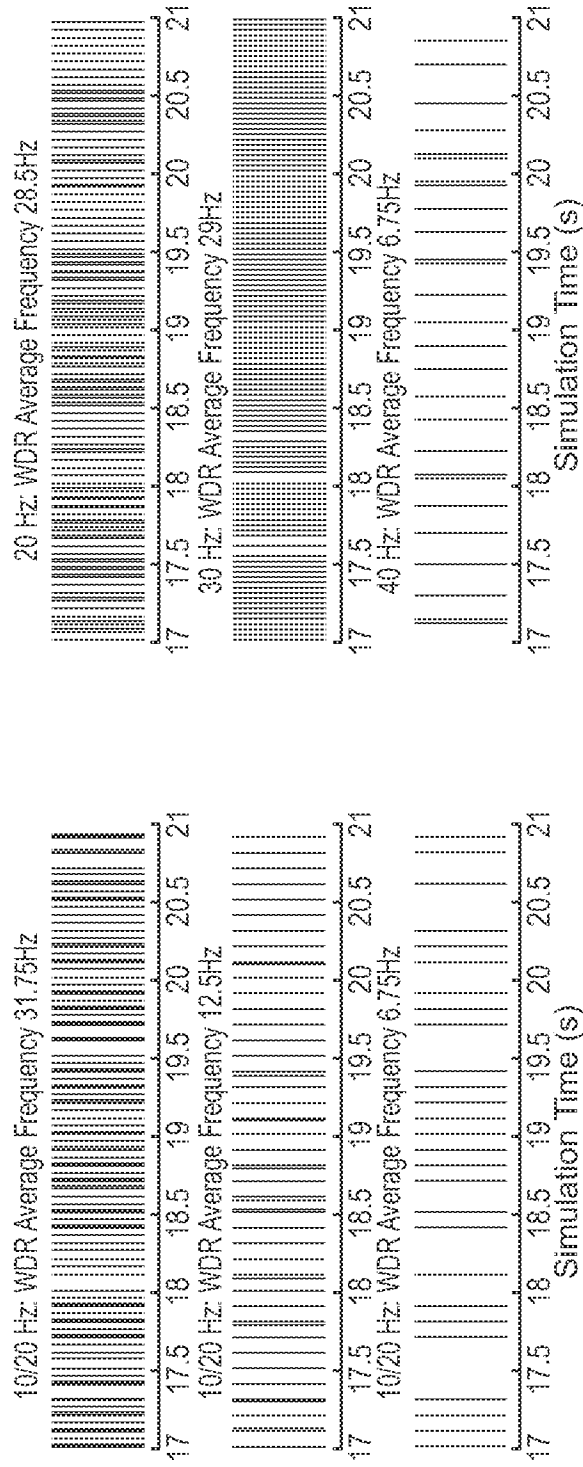


Fig. 11B

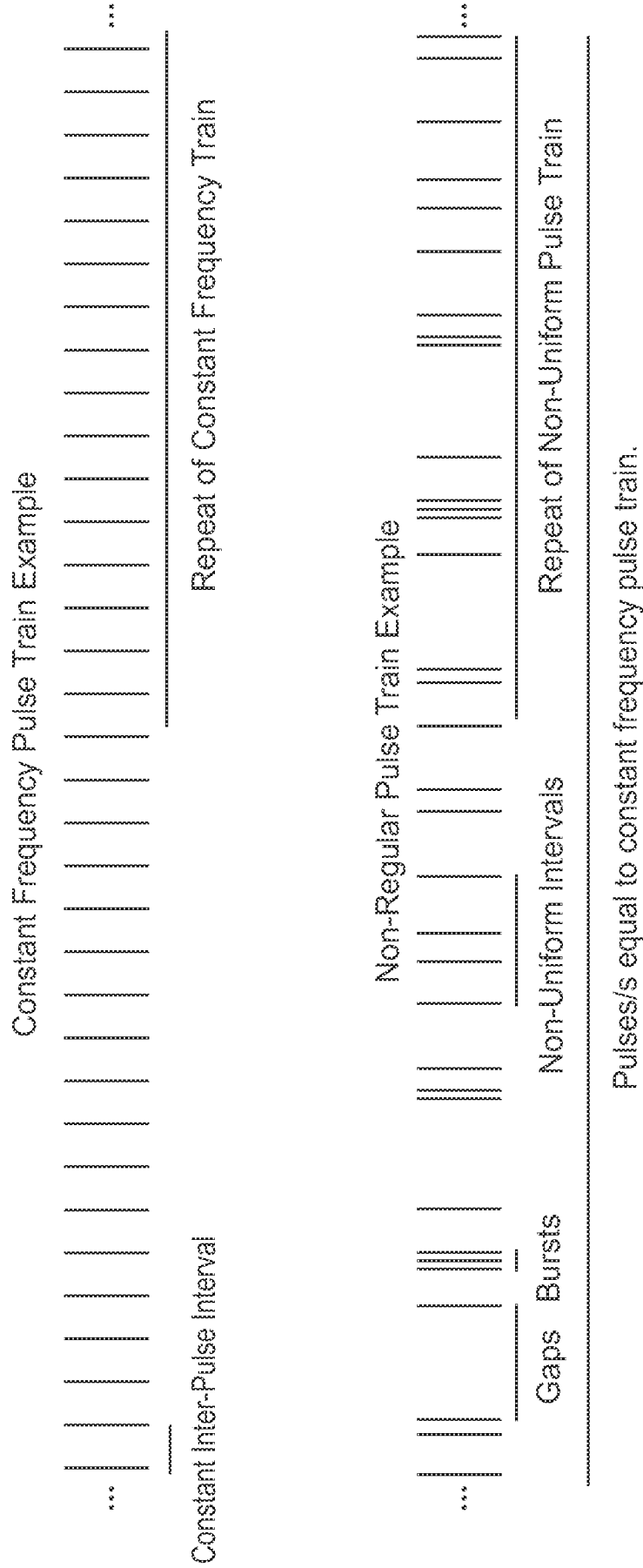


Fig. 12

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2014/025423**A. CLASSIFICATION OF SUBJECT MATTER****A61N 1/36(2006.01)i, A61N 1/08(2006.01)i**

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

A61N 1/36; A61N 1/05; A61N 1/34; A61N 1/18; A61N 1/08

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

Korean utility models and applications for utility models

Japanese utility models and applications for utility models

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

eKOMPASS(KIPO internal) & Keywords: pulse generator, electrodes, electrical stimulation, optimizing, pattern, spinal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2002-0052634 A1 (KEITH L. MARCH) 2 May 2002 See abstract; paragraphs [0037] and [0038]; claims 1-5, 38, 30, and 37.	15-24, 28, 29
A		25-27
A	US 2007-0265679 A1 (KERRY BRADLEY et al.) 15 November 2007 See abstract; claims 16 and 22.	15-29
A	US 2010-0191307 A1 (ZI-PING FANG et al.) 29 July 2010 See abstract; claim 80.	15-29
A	US 2013-0006331 A1 (JEFF A. WEISGARBER et al.) 3 January 2013 See abstract; claims 11 and 43.	15-29
A	US 2011-0087309 A1 (PAUL H. STYPULKOWSKI) 14 April 2011 See abstract; claims 1-3.	15-29

 Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

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

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

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

"&" document member of the same patent family

Date of the actual completion of the international search

29 July 2014 (29.07.2014)

Date of mailing of the international search report

30 July 2014 (30.07.2014)

Name and mailing address of the ISA/KR

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Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: 1-14
because they relate to subject matter not required to be searched by this Authority, namely:
Claims [1-14] pertain to methods for treatment of the human and thus relate to a subject-matter which this International Searching Authority is not required to search (PCT Article 17(2)(a)(i) and PCT Rule 39.1(iv)).
2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of any additional fees.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2014/025423

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2002-0052634 A1	02/05/2002	EP 1330193 A2	30/07/2003
		EP 1330193 A4	22/03/2006
US 2007-0265679 A1	15/11/2007	EP 1848493 A1	31/10/2007
		EP 1848493 B1	31/07/2013
		EP 1868681 A2	26/12/2007
		US 2005-0203588 A1	15/09/2005
		US 2005-0209655 A1	22/09/2005
		US 2005-0245987 A1	03/11/2005
		US 2008-0071325 A1	20/03/2008
		US 7146223 B1	05/12/2006
		US 7881805 B2	01/02/2011
		US 7991482 B2	02/08/2011
		US 8065013 B2	22/11/2011
		US 8233991 B2	31/07/2012
		WO 2006-073393 A1	13/07/2006
WO 2006-112852 A2	26/10/2006		
US 2010-0191307 A1	29/07/2010	EP 2396075 A1	21/12/2011
		US 8255057 B2	28/08/2012
		US 8509906 B2	13/08/2013
		WO 2010-088417 A1	05/08/2010
US 2013-0006331 A1	03/01/2013	EP 2540340 A1	02/01/2013
		US 8571667 B2	29/10/2013
US 2011-0087309 A1	14/04/2011	EP 1467795 A1	20/10/2004
		EP 1467795 B1	21/05/2008
		US 2003-0135248 A1	17/07/2003
		US 2006-0190061 A1	24/08/2006
		US 7050856 B2	23/05/2006
		US 7873418 B2	18/01/2011