



(19) **United States**

(12) **Patent Application Publication**
SWOYER et al.

(10) **Pub. No.: US 2007/0219608 A1**

(43) **Pub. Date: Sep. 20, 2007**

(54) **HIGH EFFICIENCY NEUROSTIMULATION LEAD**

Publication Classification

(76) Inventors: **John M. SWOYER**, Andover, MN (US); **Jeffrey S. GAGNON**, Champlin, MN (US)

(51) **Int. Cl.**
A61N 1/05 (2006.01)

(52) **U.S. Cl.** **607/116; 607/118**

Correspondence Address:
CRAIG TAYLOR LAW OFFICE, PLLC
774 RANDY AVE.
SHOREVIEW, MN 55124-2905

(57) **ABSTRACT**

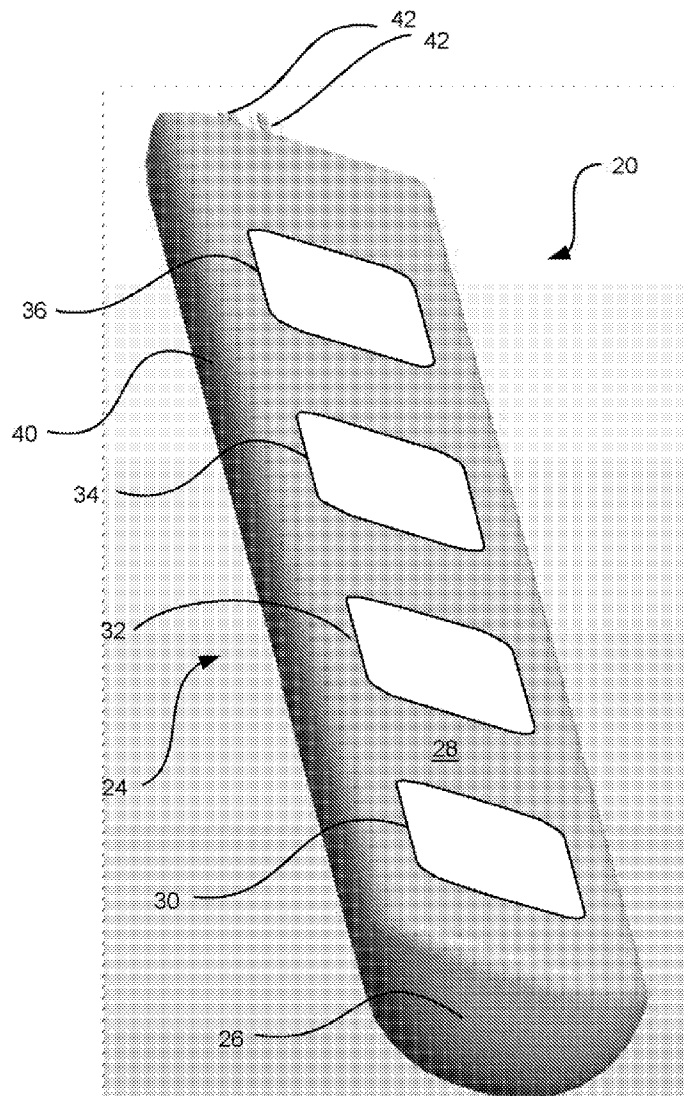
Devices and methods for stimulating nerves, such as peripheral nerves. Some devices can include small cross-section lead bodies having one or more electrodes in the distal portion, the electrodes having a substantially flat surface and being suitable for placement through a 12 gauge needle or smaller. Some lead electrodes are covered with a hydrophilic coating at least about 0.001 inch thick. The hydrophilic coating may include a steroid and/or a GABBA agonist which can elute or diffuse over time, away from the electrode.

(21) Appl. No.: **11/686,322**

(22) Filed: **Mar. 14, 2007**

Related U.S. Application Data

(60) Provisional application No. 60/782,819, filed on Mar. 16, 2006.



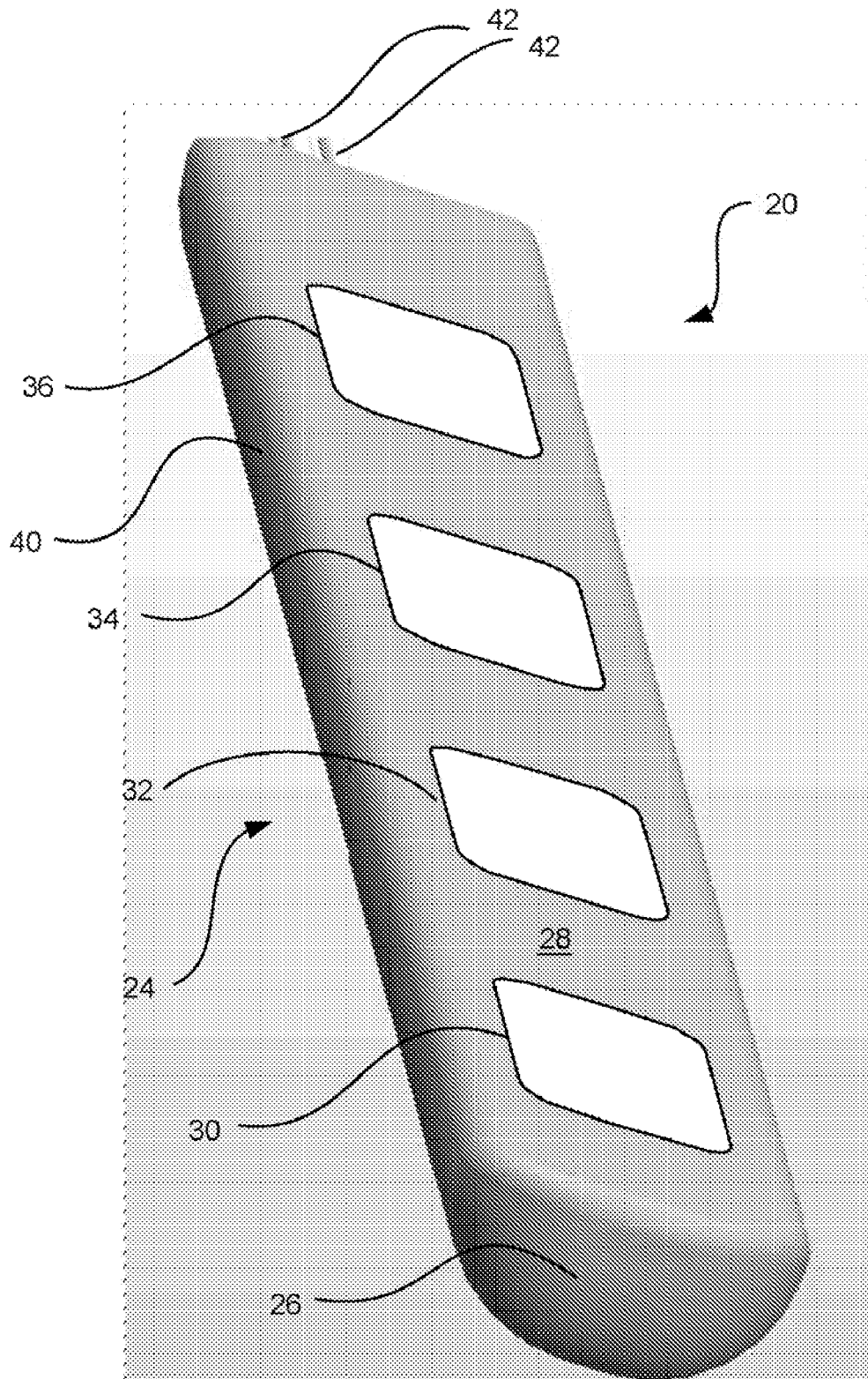


FIG 1

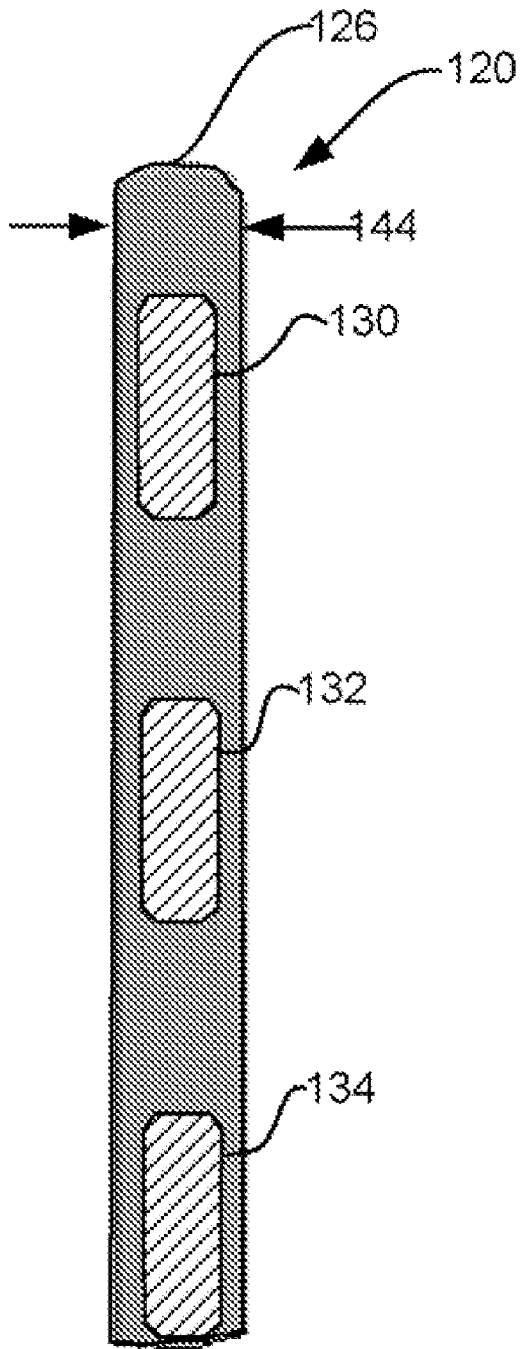


FIG 2

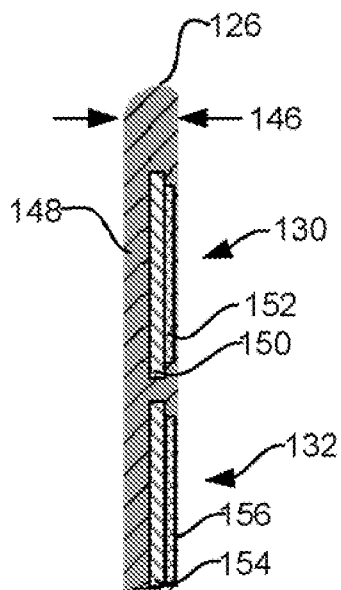


FIG 3

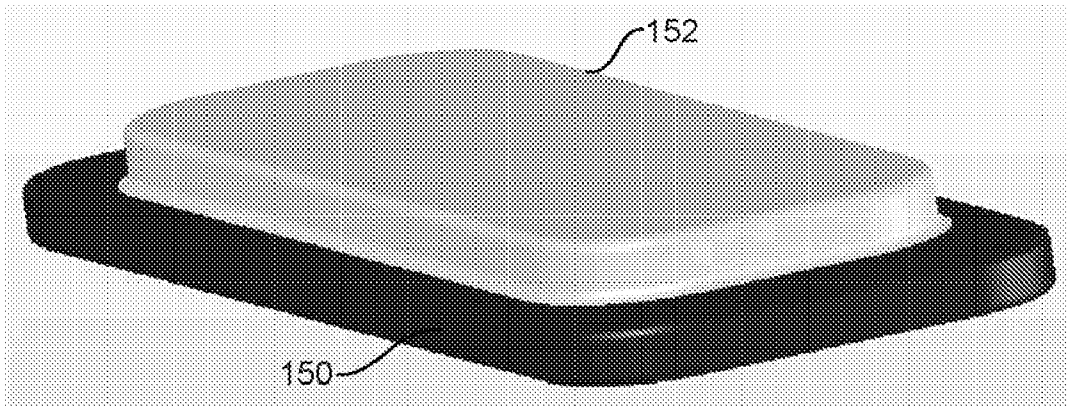


FIG 4

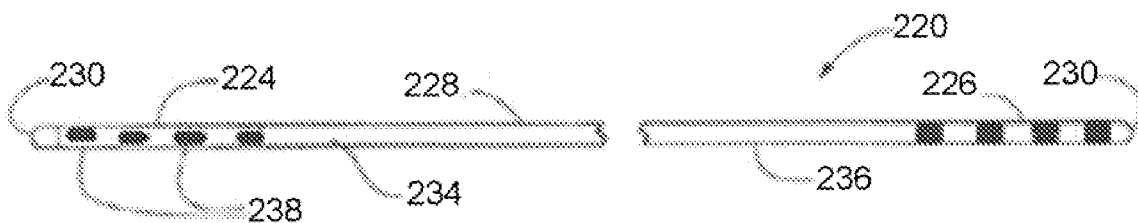


FIG 5

HIGH EFFICIENCY NEUROSTIMULATION LEAD

RELATED APPLICATIONS

[0001] The present application is a nonprovisional of U.S. Provisional Patent Application No. 60/782,819, filed Mar. 16, 2006, titled HIGH EFFICIENCY NEUROSTIMULATION LEAD.

TECHNICAL FIELD

[0002] The present invention is related generally to medical devices. More specifically the present invention is related to neurostimulation leads.

BACKGROUND

[0003] Implantable leads, typically having externally exposed ring or band electrodes can be used to deliver electrical stimulation to surrounding tissue and/or to sense electrical energy produced by the surrounding tissue. Such leads are often implanted, for example, within the epidural or intrathecal spaces of the spinal column, along peripheral nerves, within the brain, and about the heart. Electrical stimulation of the spinal cord has been shown to be effective in relieving intractable pain in some patients. Such electrical stimulation can reduce or eliminate the use of pain relieving drugs. Examples of some leads may be found in U.S. Pat. Nos.; 6,721,604; 6,981,314; 6,216,045; and 5,483,022, herein incorporated by reference.

[0004] One such lead is formed of polymeric material, for example, polyurethane or silicone. The lead can be nominally 1 mm in outer diameter and about 20 cm in length. A typical lead may have a series of electrodes formed as bands or rings disposed in a spaced apart relationship in a lead distal region. The distal region of the lead can be introduced, for example, into the epidural region for use in stimulation of the spinal column. The lead proximal region may have a corresponding set of band or ring connectors or terminals, one for each corresponding electrode in the distal region. Each proximal region terminal can thus be connected to one distal electrode in a typical configuration.

[0005] The terminals can be used to couple the proximal end of the lead to a lead extension, which can in turn be coupled to an implantable pulse generator (IPG). The lead extension can provide added length to extend the reach of the lead to a more distantly placed IPG. In some embodiments, the lead extension is between about 20 and 50 cm in length.

[0006] The lead typically has a lumen extending from the proximal end through to the distal region, with the lumen being dimensioned to accept a stiffening member or stylet. The lead, commonly formed of a polymeric material and being very small in cross section, is typically very floppy and not pushable. With a stylet or stiffening member inserted, the lead gains the needed pushability, and can be advanced into and up the spinal column to the desired location.

[0007] Current neurostimulation leads often use polished platinum electrodes having relatively large surface areas. Leads are described in U.S. Pat. Nos. 5,103,837; 5,324,324; 5,345,933; 4,044,774; and 5,265,608, herein incorporated by reference. Typical percutaneously inserted leads can use ring electrodes that wrap around 360 degrees. This is often wasteful, as energy is delivered to tissue that is not intended

to be stimulated. Such wasted energy may lead to shortened battery life. This can also lead to side effects such as pain in those tissues.

[0008] What would be desirable are leads that can be percutaneously inserted and provide directional stimulation.

SUMMARY

[0009] Some embodiments of the present invention include the use of directional electrodes that can be percutaneously delivered. In some embodiments, the lead has a flat face at the distal end that predominately or only stimulates in one direction. Optionally, the electrodes can be coated with a hydrophilic polymer film, layer, or coating. The polymer absorbs body fluid, which allows the electrical charge to pass through the polymer from the metal substrate to the stimlatable tissue. This minimizes polarization of the electrodes. It may also present a more biocompatible surface to the tissue, minimizing the foreign body response to the implanted electrode.

[0010] Minimizing the response may limit the amount of fibrosis, or scar tissue that forms at the electrode surface. This type of tissue essentially acts as an insulator and increases the energy requirements of the system. Additionally, pharmaceutical agents can be included in the polymer. These agents can elute out of the polymer matrix over time and modify the tissue response to the lead. Pharmaceuticals agents in some embodiments may include steroids, for example, beclamethasone, dexamethosone, etc and their derivatives. These agents minimize the inflammatory response to the implanted foreign body.

[0011] A different class of drug that can be included in some embodiments are GABBA agonists, for example, baclofen. These drugs enhance the ability of the stimulation to generate action potentials in the target nerves.

[0012] The present invention provides an implantable medical electrical lead including, an elongate body having a proximal portion, a distal portion, and at least one electrical conductor extending between the proximal portion and the distal portion: and a first electrode disposed in the distal portion, the first electrode having a substantially flat, planar surface and being electrically coupled to the conductor. The lead elongate body and electrode may be sized small enough in cross-sectional profile to be insertable through a 12 gauge needle. Some leads also include a hydrophilic coating over the flat electrode surface, the coating having a thickness of at least about 0.001 inch or 0.0005 inch, or between about 0.0001 and 0.01 inch, in various embodiments. The hydrophilic coating is swellable in water, in some embodiments. Some coatings include a steroid substance disposed within the hydrophilic coating for diffusion out of the hydrophilic coating. In some embodiments, the steroid is selected from the group consisting of beclamethason, dexamethosone, and their derivatives, and combinations thereof.

[0013] Some embodiment leads include a GABBA agonist substance disposed within the hydrophilic coating for diffusion out of the hydrophilic coating, for example, baclofen.

[0014] In various embodiments, the lead conductor has surface area of less than about 3 square mm., the lead has a thickness of less than about 1 mm, and the lead distal region has a width of less than about 2 mm or less than about 1.5 mm.

[0015] The present invention also provides a method for implanting a medical electrical lead, the method including advancing an implantable medical electrical lead through a

needle smaller than about 12 gauge to a target site. The lead can include an elongate body having a proximal portion and a distal portion, and at least one electrical conductor extending between the proximal portion and the distal portion. The lead may also include a first electrode disposed in the distal portion, the first electrode having a substantially flat, planar surface and being electrically coupled to the at least one conductor, where the lead elongate body and electrode are sized small enough in cross-sectional profile to be insertable through a 12 gauge needle.

[0016] In some methods, the target site is selected for peripheral nerve stimulation. The nerve stimulation target site may be selected from the group consisting of occipital, supra orbital, sub orbital, and pudendal nerve stimulation, and combinations thereof. The target site may also include the spinal cord, stimulated from a lead advanced in the epidural or intrathecal space, depending on the embodiment.

DESCRIPTION OF THE DRAWINGS

[0017] FIG. 1 is a fragmentary, perspective view of a lead distal region in one embodiment, having substantially planar, nominally square shaped electrodes.

[0018] FIG. 2 is a fragmentary, front view of a lead distal region in one embodiment, having substantially planar, nominally rectangular electrodes.

[0019] FIG. 3 is a fragmentary, side, cross-sectional view of the lead distal region of FIG. 2, having a polymeric, hydrophilic layer over the electrode metal surface.

[0020] FIG. 4 is a perspective view of one electrode of FIG. 3, having the polymeric hydrophilic layer disposed over the metallic electrode surface.

[0021] FIG. 5 is a fragmentary, bottom view of a lead according to the present invention having a distal region with four electrodes and a proximal region with four conductor rings.

DETAILED DESCRIPTION

[0022] The present invention provides leads, sized in some embodiments such that it can be delivered via a percutaneously placed needle, for example, smaller than 12 gauge, typically between 13 and 28 gauge. The lead distal end or distal portion can have at least one flat face. Electrodes may be disposed on at least one of the flat faces and can have a flat surface in some embodiments of the invention. The electrode metal substrate can be a corrosion resistant, biocompatible and biostable material, such as platinum, platinum alloys, titanium or titanium alloy, gold, etc.

[0023] The lead can have 1 or more electrodes (preferably 1-32 and more preferably 4-8 electrodes), with electrical conductors connecting the distal and proximal regions and/or ends, with the proximal region and/or end containing the corresponding number of contacts and configured to be compatible to an implanted pulse generator, or other power source. The electrode dimensions are optimized for battery life by minimizing current loss into undesirable tissue.

[0024] The insulation of the lead body, and distal and proximal ends, can be a biocompatible and biostable polymer, such as polyurethane>silicone. polyurethane-silicone hybrid, peek, polyimide, etc.

[0025] FIG. 1 shows one lead 20 according to the present invention, having a polymeric distal region 24 including a housing 40 having four flat surface planar electrodes 30>32>24, and 36 disposed within a substantially flat hous-

ing region 28. Electrical conductors 42 may be seen in a cutout view extending proximally. Lead 20 terminates in this embodiment in a somewhat rounded distal end 26.

[0026] FIG. 2 illustrates another lead embodiment 120 having a width of less than 2 mm. here about 1.3 mm, indicated at 144, and having electrodes 130, 132: and 134 with a surface area of less than about 3 square mm, here about 2 square mm. Lead 120 terminates in a distal tip 126.

[0027] FIG. 3 illustrates lead 120 of FIG. 2 in cross section, having a thickness of less than about 1 mm, here about 0.75 mm, indicated at 146. Housing 148 houses a first electrode 130 having a conductor layer and a surface coating or layer 152>which can include a hydrophilic material, steroid, and/or a GABA agonist. A second electrode 132 having a conductor layer 154 with a coating 156 is also shown.

[0028] FIG. 4 illustrates lead electrode 1330, having a flat metal substrate 150 coated with a hydrophilic layer 152.

[0029] FIG. 5 illustrates a neurological stimulation lead 220 according to the present invention. Lead 220 can incorporate a multiconductor cable. Lead 220 has a distal region 224>a proximal region 226>and an intermediate region 228 disposed between the distal and proximal regions. In a preferred embodiment, the intermediate region is defined to lie between the innermost distal and proximal electrical contacts described below. Lead 220 can be formed of a body or shaft 234 extending between a distal end 230 and a proximal end 232. Lead body 234 has an exterior surface or side wall 236. In some embodiments, the lead body proximal of the distal region has a substantially round cross section, while in other embodiments the lead regions proximal of the distal region are flat on at least one side, similar to the shape of the distal region bearing the electrodes. Lead body 234 is preferably formed of a polymeric material, for example, polyurethane or silicone.

[0030] Lead distal region 224 may include a number of electrodes 238, which may, for example, be cathodes disposed along the bottom of lead body 234 in a spaced-apart configuration. Electrodes 238 may also be described as electrical contacts or contacts. Electrodes 238 are normally adapted to be inserted into the human body, are externally exposed, and can be used for neurological stimulation. One exemplary use of electrodes 328 is the stimulation of the nerves within the spinal cord. Proximal region 226 can include a number of connector bands or connector rings 240 disposed in a spaced-apart configuration. Connectors 240 may also be described as electrical contacts or terminals, and are preferably also externally exposed. Connectors may be circumferential or flat, and may be made from platinum, platinum alloys, stainless steel, nickel alloys, etc. Electrodes 238 and connectors 240 may be formed of Platinum and/or Iridium. Connectors 240 can be used for connecting lead 220 to a lead extension to extend the effective length of the lead. In some uses, connectors 240 may also be used to directly couple lead 222 to an implantable pulse generator.

[0031] Electrodes 238 and connectors 240 can be coupled to each other in a one-to-one arrangement. In some leads, the distal-most electrode is coupled to the distal-most connector, the second-to-distal-most electrode coupled to the second-to-distal-most connector, and so forth. The electrodes and connectors can be coupled through conductors extending between the two. In some leads, the conductors are embedded within the lead while in other leads, the conductors lie within lumens extending the length of the lead. In some

leads, the conductors are disposed within lumens that are later backfilled to substantially fill the lumens with a polymeric material. Some leads have stylet lumens for receiving a stiffening stylet member.

[0032] Lead **220** can be varied in outer diameter and length to suit the application for which it is intended. In some embodiments, lead **220** has a total length of between about 5 cm and about 100 cm. In other embodiments, lead **220** has an outer diameter of less than about 1 mm and a total length of between about 10 cm and 150 cm.

[0033] Uses for the present invention include, but are not limited to: spinal cord stimulation, brain stimulation; any central nervous system stimulation; any peripheral nerve stimulation, including but not limited to occipital, orbital, cranial, sacral, pudendal; vagus; and/or radial nerves; cardiac pacing and/or defibrillation; smooth muscle stimulation (stomach, liver, etc); and skeletal muscle stimulation.

[0034] Leads according to the present invention can be introduced into the epidural space and used to stimulate the spinal cord. In another use, a lead can be introduced into the intrathecal space for spinal cord stimulation. While not wishing to be bound by theory, applicants believe that intrathecal stimulation is not currently used because circumferential electrodes would dump too much current into the highly conductive cerebral spinal fluid. Highly directional electrodes of some embodiments of the present invention, placed in close proximity to the spinal cord, may require much lower current. In some embodiment methods, a GABBA agonist coating or layer on the lead distal region: for example, on the electrode, can be placed much closer to the spinal cord when the lead is placed in the intrathecal space. This close proximity to the nerves may increase the effectiveness of the GABBA agonist. The GABA agonist coating or coating may also be used in peripheral nerve stimulation. Applicants believe peripheral nerve stimulation may also benefit from the close proximity of the electrode and drug to the nerve.

[0035] Various examples and embodiments of the present invention have been presented above., and are intended to illustrate some aspects of the present invention. The scope of the present invention is to be defined by the claims which follow.

What is claimed is:

1. An Implantable medical electrical lead comprising:
 - an elongate body having a proximal portion, a distal portion, and at least one electrical conductor extending between the proximal portion and the distal portion;
 - a first electrode disposed in the distal portion, the first electrode having a substantially flat., planar surface and being electrically coupled to the at least one conductor, where the lead elongate body and electrode are sized small enough in cross-sectional profile to be insertable through a 12 gauge needle.
2. The lead of claim 1, further comprising a hydrophilic coating over the flat electrode surface, the coating having a thickness of at least about 0.001 inch.
3. The lead of claim 2, further comprising a substance disposed within the hydrophilic coating for diffusion out of the hydrophilic coating., where the substance is a steroid.
4. The lead of claim 3, in which the steroid is selected from the group consisting of beclomethason, dexamethosone, and their derivatives, and combinations thereof.

5. The lead of claim 2, further comprising a substance disposed within the hydrophilic coating for diffusion out of the hydrophilic coating, where the substance is a GABBA agonist.

6. The lead of claim 5, in which the GABBA agonist includes baclofen.

7. The lead of claim 2, further comprising a steroid disposed within the hydrophilic coating for diffusion out of the hydrophilic coating

8. The lead of claim 2, further comprising a GABBA agonist disposed within the hydrophilic coating for diffusion out of the hydrophilic coating.

9. The lead of claim 2, further comprising a steroid and a GABBA agonist disposed within the hydrophilic coating for diffusion out of the hydrophilic coating.

10. The lead of claim 2, in which the hydrophilic coating is swellable in water.

11. The lead of claim 2, in which the coating has a thickness of at least about 0.005 inch.

12. The lead of claim 2, in which the coating has a thickness of between about 0.001 and 0.01 inch.

13. The lead of claim 1, in which the lead conductor has surface area of less than about 3 square mm.

14. The lead of claim 1, in which the lead distal region has a width of less than about 2 mm.

15. The lead of claim 1, in which the lead distal region has a width of less than about 1.5 mm.

16. The lead of claim 1, in which the lead distal portion has a thickness substantially perpendicular to the conductor flat surface, where the thickness is less than about 1 mm.

17. A method for implanting a medical electrical lead, the method comprising:

- advancing an implantable medical electrical lead through a needle smaller than about 12 gauge to a target site, the lead including

- an elongate body having a proximal portion, a distal portion, and at least one electrical conductor extending between the proximal portion and the distant portion, and

- a first electrode disposed in the distal portion, the first electrode having a substantially flat, planar surface and being electrically coupled to the at least one conductor, where the lead elongate body and electrode are sized small enough in cross sectional profile to be insertable through a 12 gauge needle.

18. The method of claim 17 in which the target site is selected for peripheral nerve stimulation.

19. The method of claim 18 in which the nerve stimulation target is selected from the group consisting of occipital, supra orbital, sub orbital, and pudendal nerve stimulation, and combinations thereof.

20. The method of claim 17, further comprising eluting a steroid from a hydrophilic coating disposed over the lead conductor.

21. The method of claim 17, further comprising eluting a GABBA agonist from a hydrophilic coating disposed over the lead conductor.

22. The method of claim 21, in which the GABA agonist eluting includes eluting baclofen.

23. The method of claim 17, in which the target site is the epidural space.

24. The method of claim 17, in which the target site is the intrathecal space.

25. The method of claim **24**, further including eluting a GABA agonist within the intrathecal space.

26. An Implantable medical electrical lead comprising: an elongate body having a proximal portion, a distal portion, and at least four electrical conductors extending between the proximal portion and the distal portion; at least four electrodes disposed in the distal portion, the at least four electrodes each having a substantially flat, planar surface and being electrically coupled to at least one of the at least four conductors,

where the lead elongate body and electrodes are sized small enough in cross-sectional profile to be insertable through a 12 gauge needle; and

a hydrophilic coating over the flat electrode surfaces, the coating having a thickness of at least about 0.001 inch.

27. The implantable lead of claim **26**, in which the electrodes have a substance disposed within the hydrophilic coating for diffusion out of the hydrophilic coating, where the substance is a steroid selected from the group consisting of beclamethason, dexamethosone, and their derivatives and combinations thereof.

28. The implantable lead of claim **26**, in which the electrodes have a substance disposed within the hydrophilic coating for diffusion out of the hydrophilic coating, where the substance is a GABBA agonist.

* * * * *