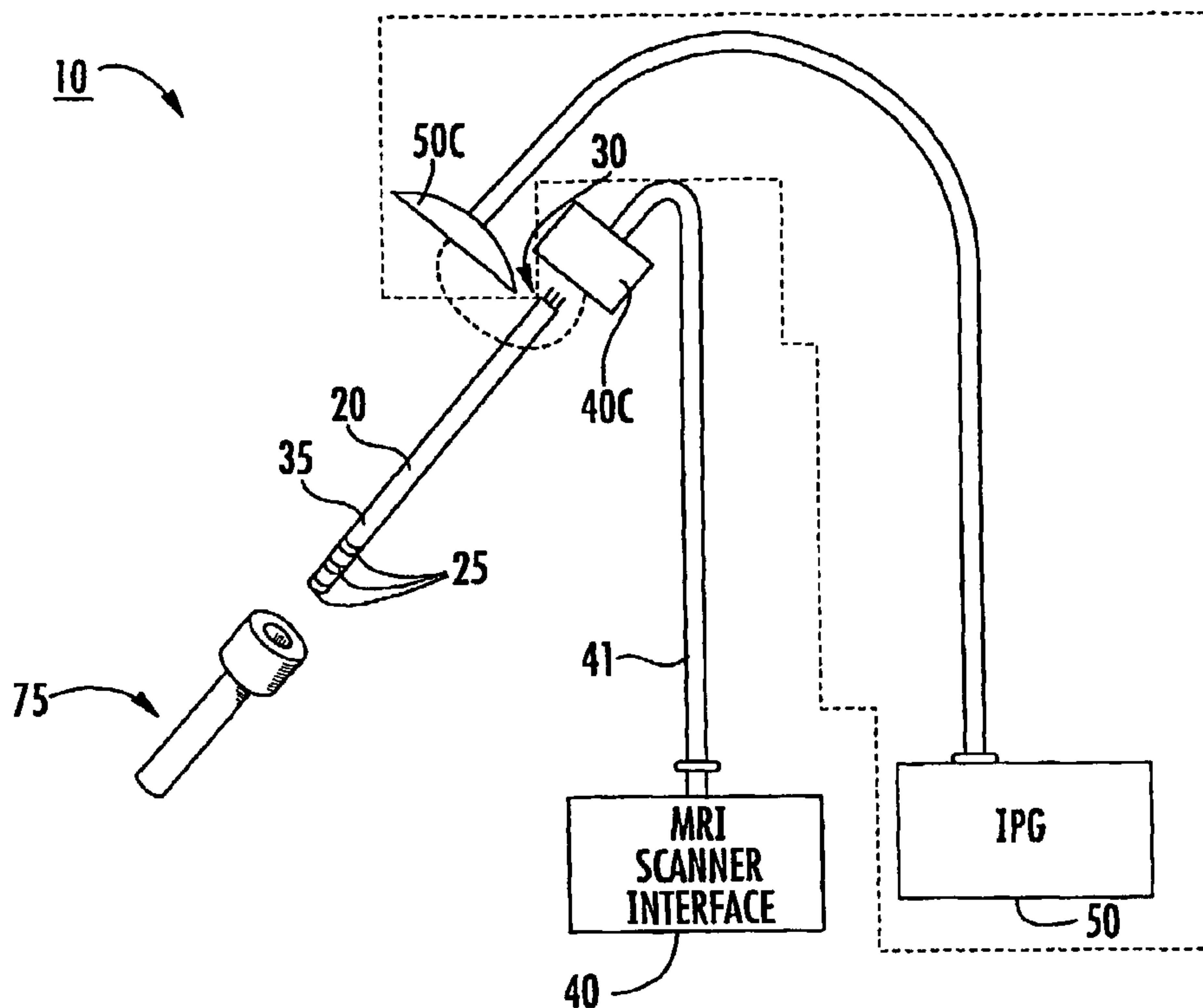




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In vivo medical stimulation probes include an elongate lead having at least one stimulation electrode disposed on a distal portion thereof. The probes may include a plurality of axially spaced apart RF chokes disposed on and/or in an axially extending shielding layer of the lead in advance of the at least one electrode to inhibit induced RF current from forming and/or traveling along the shielding layer.

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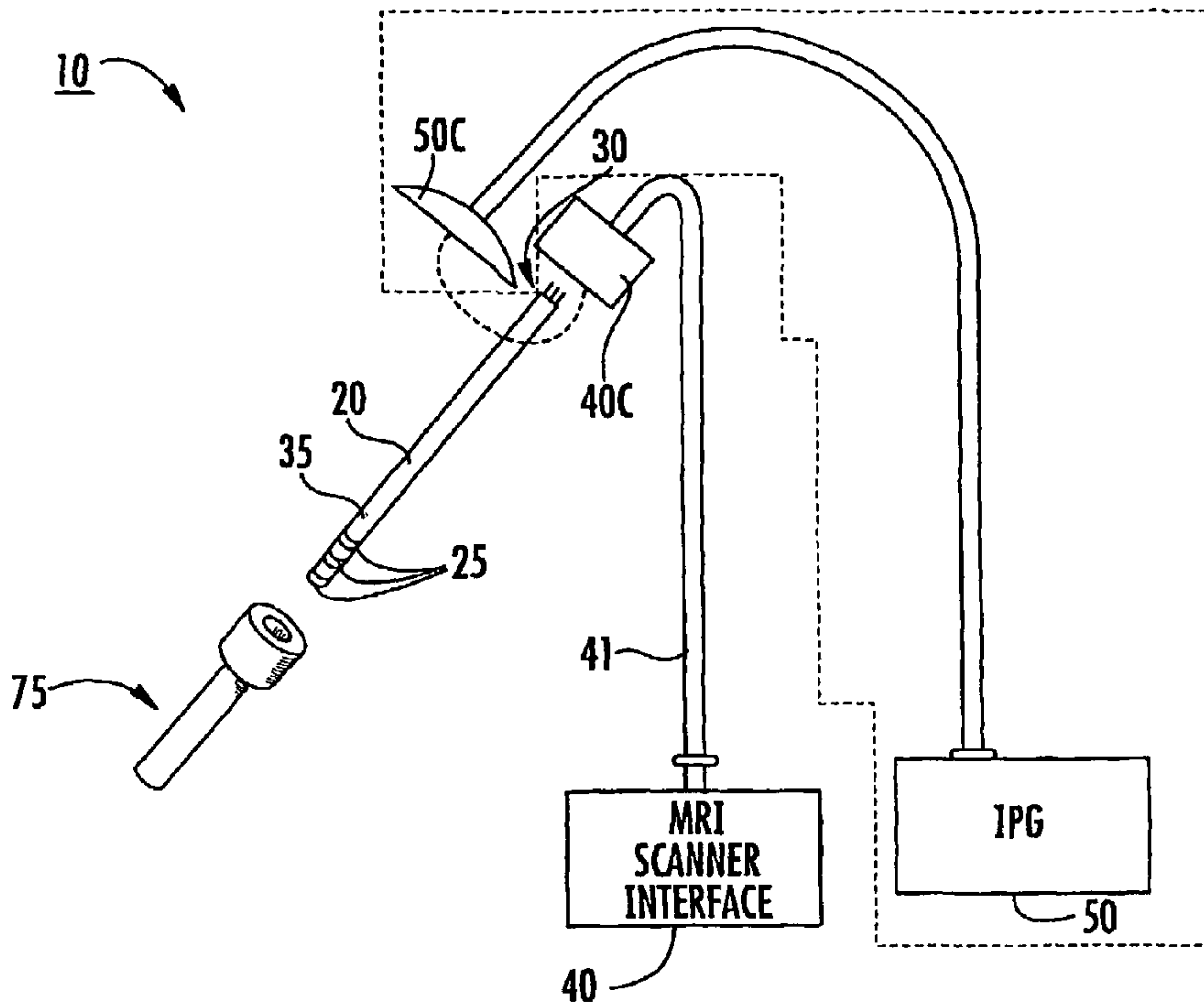
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IMPLANTABLE MRI COMPATIBLE STIMULATION LEADS AND ANTENNAS AND RELATED SYSTEMS AND METHODS

GOVERNMENT FUNDING

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RELATED APPLICATIONS

This application claims the benefit of priority to U.S. Provisional Application Serial Nos. 60/600,020 filed August 9, 2004 and 60/608,195, filed September 9, 2004, the contents of which are hereby incorporated by reference as if recited in full herein.

FIELD OF THE INVENTION

The present invention relates to interventional medical leads and may be particularly suitable for MRI compatible implantable Deep Brain Stimulation ("DBS") and/or implantable sympathetic nerve chain stimulation leads.

5

BACKGROUND OF THE INVENTION

Deep Brain Stimulation (DBS) is becoming an acceptable therapeutic modality in neurosurgical treatment of patients suffering from chronic pain, Parkinson's disease or seizure, and other medical conditions. Other electro-
10 stimulation therapies have also been carried out or proposed using internal stimulation of the sympathetic nerve chain and/or spinal cord, etc.

One example of a prior art DBS system is the Activa® system from Medtronic, Inc. The Activa® system includes an implantable pulse generator stimulator that is positioned in the chest cavity of the patient and a lead with axially
15 spaced apart electrodes that is implanted with the electrodes disposed in neural tissue. The lead is tunneled subsurface from the brain to the chest cavity connecting the electrodes with the pulse generator. These leads can have multiple exposed electrodes

at the distal end that are connected to conductors which run along the length of the lead and connect to the pulse generator placed in the chest cavity.

Functional MRI is an imaging modality that can be used to evaluate cardiac, neurological and/or other disorders. It may be desirable to use MRI for patients with implanted stimulation devices and leads. However, currently available lead systems may be unsuitable to use in a magnetic resonance imaging (MRI) environment. For example, the devices may not be MRI compatible, *i.e.*, they may contain ferromagnetic materials, which may distort the MRI images. Also, currently available lead/probe/cable systems may be susceptible to unwanted induced RF and/or AC current and/or localized heating of the tissue. For example the Medtronic Activa® device recommends that MRI imaging be carried out in a 1.5T magnet without using body coils, *i.e.*, only using head coils for transmission of the RF excitation pulse(s). Also, the problem of unwanted RF deposition may increase as higher magnetic fields, such as 3T systems, become more common for MRI imaging (the RF pulses having shorter wavelengths).

It is believed that the clinical outcome of certain medical procedures, particularly those using DBS, may depend on the precise location of the electrodes that are in contact with the tissue of interest. For example, presently, to treat Parkinson's tremor, the DBS probes are placed in neural tissue with the electrodes transmitting a signal to the thalamus region of the brain. However, it has been reported that in about 30% of the patients implanted with these devices, the clinical efficacy of the device/procedure is less than optimum. This may be attributed to the inaccurate/imprecise placement of the lead/probe/electrode with respect to the cranial or neural tissue.

Other non-MRI applications, such as RF/microwave diathermy treatment, can use leads that employ an alternating current to cauterize tissue. The diathermy alternating current delivered during the therapy can be in the range of between about 1KHz-350 MHz. In certain situations, the lead system may undesirably act as an antenna, picking-up and depositing current in localized tissue where the leads are exposed, thereby potentially increasing the SAR or *specific absorption rate* (a way of measuring the quantity of radiofrequency (RF) energy that is absorbed by the body).

Notwithstanding the above, there remains a need for alternative medical lead configurations.

SUMMARY OF EMBODIMENTS OF THE INVENTION

Some embodiments of the present invention provide therapeutic lead systems. The lead systems may include RF chokes and/or induced current resistant leads. The lead systems may be implantable, MRI compatible multi-purpose lead systems
5 configured to provide an internal MRI receive antenna and at least one stimulating electrode.

The lead can be a relatively long implantable lead having a length in the body of greater than 10 cm. The at least one electrode can be a plurality of electrodes with at least one being a recording and/or a recording and stimulating and/or ablating
10 electrode.

The device can be a multi-purpose lead system that allows for increased precision placement using an integrated MRI receive antenna and stimulation using electrodes. The lead system can have opposing distal and proximal portions with the proximal portion configured to reside external of the subject during lead placement.
15 In some embodiments, the proximal portion can releaseably attach to either an MRI interface or an implantable pulse generator. The proximal portion can merge into a connector. The distal portion comprises a plurality of stimulating electrodes and an MRI antenna.

Certain embodiments are directed to *in vivo* medical stimulation probes that
20 include: (a) an elongate lead having at least one stimulation electrode disposed on a distal portion thereof; and (b) a plurality of axially spaced apart RF chokes disposed on and/or in an axially extending shielding layer of the lead in advance of the at least one electrode to inhibit induced RF current from forming and/or traveling along the shielding layer.

The lead may be a flexible lead and the at least one electrode can be a plurality of spaced apart electrodes. The lead can include a plurality of conductors held in a core of the lead, a respective one for each electrode. The shielding layer can be discontinuous and can be configured to surround the conductors over at least a major length of the lead and terminate at a lead location that is in advance of the electrodes.
25
30 The probe can include an axially extending primary inner shield surrounding the core with the plurality of electrodes. The discontinuous shielding layer can be a second shield layer that is generally cylindrically disposed over an inner primary shielding layer. The primary and second shields both terminate at a location that is before the electrodes.

Other embodiments are directed to chronically implantable deep brain stimulation and MRI imaging probe systems. The systems include: (a) an MRI compatible elongate lead comprising an MRI antenna having an axially extending radially spaced apart first and second shield layer for internally receiving local *in vivo* MRI signals; (b) at least one electrode held on a distal portion of the lead and, in operation, configured to generate a stimulation pulse to deep brain neural tissue; (c) a stimulation circuit in communication with the at least one electrode; (d) a MRI signal receive circuit in communication with the MRI antenna; (e) a splitter circuit in communication with the stimulation and receive circuits for selectively electrically connecting either the MRI receive or stimulation circuit; and (f) means for inhibiting RF induced current on the second shield layer of the lead.

Still other embodiments are directed to MRI compatible deep brain stimulation and imaging probe systems. The systems include: (a) a flexible elongate probe body having opposing proximal and distal portions, the probe body comprising a plurality of electrodes disposed on the distal portion; (b) a plurality of axially extending conductors disposed in a core of the probe body, a respective one for each electrode; (c) an axially extending inner shield surrounding the plurality of conductors for at least a major portion of the length of the conductors; (d) an axially extending second shield radially spaced apart from the inner shield; (e) an axially extending first insulating/dielectric layer disposed intermediate of the inner and second shields; (f) an MRI antenna held by the probe body at the distal portion thereof; (g) an RF transmit decoupling circuit in communication with the MRI antenna; and (h) at least one connector attached to the proximal portion of the probe body, configured to hold a conductor transmission line for each of the electrodes.

Yet other embodiments are directed to medical kits. The kits include an elongate sterilized biocompatible and MRI compatible lead having opposing distal and proximal portions. The lead includes an MRI antenna, a plurality of stimulation electrodes on the distal portion, and a plurality of axially spaced apart RF chokes disposed in a shielding layer of the lead. The lead is configured to have selective operative first and second electrical transmission paths associated with first and second operational modes. The first transmission path connecting the MRI antenna with an MRI scanner and decoupling the electrodes during MRI operation and the second transmission path connecting the electrodes with a stimulation or recording source during electrical stimulation or recording.

Additional embodiments are directed to methods of placing and operating a deep brain stimulation probe. The methods include: (a) inserting a flexible elongate lead comprising an MRI antenna and at least one stimulation electrode on a distal portion thereof into a brain of a subject; (b) connecting the lead to an MRI scanner interface in communication with a splitter circuit having at least two different electric transmission paths, a first for MRI operation and a second for stimulation operation; (c) obtaining MRI signals associated with local neural tissue proximate the MRI antenna from the MRI antenna using the first transmission path; (d) placing the electrode on the lead at a desired location in the brain based on data from the obtaining step; then (e) stimulating neural tissue with the electrode using the second transmission path; and (f) configuring the lead to inhibit the formation and/or transmission of RF induced current. The stimulating and obtaining steps are carried out using the same lead.

Other embodiments are directed to computer program products for operating a multi-purpose MRI compatible stimulation probe with MRI antenna. The computer program product includes a computer readable storage medium having computer readable program code embodied in the medium. The computer-readable program code includes computer readable program code that controllably engages a desired operational mode of at least one of a plurality of different modes including a first or second operational mode for a MRI compatible stimulation probe with at least one electrode and an MRI antenna. The first operational mode having a first transmission path connecting the MRI antenna with an MRI scanner and decoupling the electrodes during MRI operation and the second operational mode having a second transmission path connecting the electrodes with a stimulation or recording source during electrical stimulation or recording.

The computer readable program code may be configured to time the selection of the second operational mode to occur proximate in time but after an MRI signal acquisition by the MRI antenna in the first operational mode. The computer readable program code may be configured to obtain microrecordings of local tissue in substantially real time proximate in time to an MRI signal acquisition by the MRI antenna in the first operational mode. The computer readable program code may be configured to obtain a plurality of MRI signals of local neural tissue proximate the MRI antenna in substantially real time, then obtain a plurality of microrecordings of

the local neural tissue to allow a clinician to track placement of the probe using both MRI data and audio data.

Yet other embodiments are directed to an MRI compatible therapeutic stimulation probe that includes: (a) an elongate flexible probe body having an axially extending internal cavity disposed therein; (b) at least one electrode held by a distal portion of the probe body; and (c) at least one axially extending conductor configured to slidably extend into the cavity of the probe body, the at least one conductor having increased rigidity relative to the probe body. During positioning in a body, the at least one conductor cooperates with the probe body and defines an *in vivo* MRI antenna used to obtain MRI signals for MRI positional guidance. After placement, the at least one conductor can be removed from the probe body, leaving the probe body in position in the body.

A cannula may be configured to be inserted into a burr hole placed in a patient's skull and the stimulation probe and MRI antenna probe may be configured for deep brain placement guided through the cannula.

In some embodiments, the cannula is configured to cooperate with the MRI antenna probe to define an MRI receive antenna when the MRI antenna probe is held inside the cannula. In particular embodiments, the cannula comprises a conductive shielding layer that cooperates with the MRI antenna probe to define an MRI receive antenna during positioning in a body used to obtain MRI signals for MRI positional guidance of the stimulation electrode on the same probe.

These and other embodiments will be described further below.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a schematic illustration of a medical kit with an implantable stimulation lead according to embodiments of the present invention.

Figure 2 is an image of a subject having implanted deep brain stimulation leads which are MRI compatible according to embodiments of the present invention.

Figure 3A is a side view of a stimulation lead according to embodiments of the present invention.

Figure 3B is a section view of the device shown in **Figure 3A**, taken along line **3B-3B**.

Figure 3C is a side schematic view of a probe with linear and/or extendable electrodes according to embodiments of the present invention.

Figure 4 is an electrical schematic diagram of the device shown in **Figure 3A** according to embodiments of the present invention.

Figure 5A is a partially exploded schematic illustration of a multi-purpose probe according to embodiments of the present invention.

5 **Figure 5B** is a section view of the probe shown in **Figure 5A**, illustrating the probe cavity without the center conductor(s).

Figure 5C is a section view of the probe shown in **Figure 5A**, illustrating the probe cavity holding the center conductor(s).

10 **Figure 6** is a block diagram of a bimodal lead operating circuit according to embodiments of the present invention.

Figure 7A is a block diagram of another operating circuit according to embodiments of the present invention.

Figure 7B is yet another block diagram of an operating circuit according to embodiments of the present invention.

15 **Figure 8A** is a schematic illustration of a splitter circuit according to embodiments of the present invention.

Figure 8B is an end view of the circuit shown in **Figure 8A**.

20 **Figure 9** is a schematic illustration of an MRI guidance fixture system configured to insert an MRI compatible stimulation probe into position according to embodiments of the present invention.

Figure 10A is a schematic perspective illustration of a dual probe system according to other embodiments of the present invention.

Figure 10B is a schematic illustration of a dual probe kit according to embodiments of the present invention.

25 **Figure 11** is a block diagram of a data processing system according to embodiments of the present invention.

DETAILED DESCRIPTION OF EMBODIMENTS OF THE INVENTION

30 The present invention will now be described more fully hereinafter with reference to the accompanying drawings, in which embodiments of the invention are shown. This invention may, however, be embodied in many different forms and should not be construed as limited to the embodiments set forth herein; rather, these embodiments are provided so that this disclosure will be thorough and complete, and will fully convey the scope of the invention to those skilled in the art. Like numbers

refer to like elements throughout. It will be appreciated that although discussed with respect to a certain antenna embodiment, features or operation of one lead system embodiment can apply to others.

In the drawings, the thickness of lines, layers, features, components and/or regions may be exaggerated for clarity and broken lines illustrate optional features or operations, unless specified otherwise. In addition, the sequence of operations (or steps) is not limited to the order presented in the claims unless specifically indicated otherwise. It will be understood that when a feature, such as a layer, region or substrate, is referred to as being "on" another feature or element, it can be directly on the other element or intervening elements may also be present. In contrast, when an element is referred to as being "directly on" another feature or element, there are no intervening elements present. It will also be understood that, when a feature or element is referred to as being "connected" or "coupled" to another feature or element, it can be directly connected to the other element or intervening elements may be present. In contrast, when a feature or element is referred to as being "directly connected" or "directly coupled" to another element, there are no intervening elements present. Although described or shown with respect to one embodiment, the features so described or shown can apply to other embodiments.

Unless otherwise defined, all terms (including technical and scientific terms) used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. It will be further understood that terms, such as those defined in commonly used dictionaries, should be interpreted as having a meaning that is consistent with their meaning in the context of the relevant art and this specification and should not be interpreted in an idealized or overly formal sense unless expressly so defined herein.

Probe embodiments of the present invention can be configured to stimulate any desired internal region of the body or object. The object can be any object, and may be particularly suitable for animal and/or human subjects. Some probe embodiments can be sized and configured for brain stimulation, typically deep brain stimulation. Some probe embodiments can be configured to stimulate a desired region of the sympathetic nerve chain. Other embodiments may be directed to other anatomical structures, organs or features including deep tissue, lumens, and the like. For example, the probe systems of the present invention may be configured for treatment of cardiac, gastrointestinal, urinary, or other body regions.

Examples of known stimulation treatments and/or target body regions are described in U.S. Patent Nos. 6,708,064; 6,438,423; 6,356,786; 6,526,318; 6,405,079; 6,167,311; 6,539,263; 6,609,030 and 6,050,992, the contents of which are hereby incorporated by reference as if recited in full herein.

5 Embodiments of the present invention will now be described in detail below with reference to the figures. **Figure 1** illustrates a medical kit **10** that includes at least one stimulation lead or probe **20** with at least one stimulation and/or ablation electrode **25**. The at least one electrode **25** is shown in **Figure 1** as three generally cylindrical, axially spaced apart electrodes. The terms "lead" and "probe" can be used
10 interchangeably to indicate a body used to support the stimulation electrode(s) **25**. Other numbers of electrodes as well as other electrode configurations can be used. **Figure 3A** illustrates four electrodes **25** and **Figure 3C** illustrates that the electrodes **25** may be generally linear electrodes **25** that can translate with respect to the body of the probe to extend out an end portion and/or sidewall portions thereof to provide
15 additional (typically individually selectable) stimulation and/or sensing options. It is contemplated that the electrode(s) can be sized and configured to "fit" the desired internal target, which may be a relatively small region, such as less than about 1-3 mm. Typically, as shown in **Figure 1**, the electrode(s) **25** can be held on a distal portion of the probe body. The connector **30** on the proximal end portion of the probe
20 body can be configured to reside outside of the body during lead placement. The proximal portion of the probe body can be configured to releasably engage either an MRI scanner interface **40** or an implantable pulse generator **50** via respective connectors **40c**, **50c**.

As shown by the broken line, the kit **10** may optionally also include at least
25 one implantable pulse generator **50** with an implantable lead **51** and a connector **50c**. The generator and lead **50**, **51** can also comprise MRI compatible materials and/or components. In addition, optionally, the kit **10** can include an MRI interface lead **41** and/or MRI scanner interface **40**. The MRI interface **40** and/or lead **41** may be standard equipment useable across many procedures with different probes **20**. The
30 lead **41**, connector **40c** and/or MRI interface **40** may be provided as a single-use disposable sterilized component in the kit **10** or sterilized by the clinic between uses. The probe **20** is typically an elongate flexible probe comprising an outer layer of elastomeric material such as a polymer that extends across the outer surface of the probe body while leaving the electrode(s) **25** configured to contact the tissue in

position in the body. The probe **20** includes at least one conductor lead **26** (**Figure 3B**) that electrically connects the electrode **25** to a remote input or output source, such as either the MRI scanner interface **40** or implantable pulse generator **50**. The lead(s) **26** can comprise any suitable material, and may, in some embodiments, comprise a shape memory alloy such as Nitinol. The connector **50c** for the implantable pulse generator **50** may be configured with a smaller profile than that of the MRI lead connector **40c** as it may be configured to be implanted under the skin of the subject for chronic stimulation.

As shown in **Figure 1**, in some embodiments, the kit **10** may also include an MRI compatible generally rigid cannula **75** and/or a cannula **75** with increased rigidity relative to the probe that is configured to slidably receive at least the distal and intermediate portions of the probe body to guide the probe **20** into position. As with the interface lead **41**, the cannula **75** can be single-use and disposable and provided as a sterilized component in the kit **75**, or it may be re-used as a standard component and sterilized by the user/clinic. The cannula **75** can be configured according to a desired body entry location; *e.g.*, for oral entry, the cannula **75** can be formed into a bite block, nasal cavity or ear plug member, and for non-neural uses, such as placement in the spinal column, no cannula may be required.

For MRI compatible uses, the cannula **75**, the probe **20**, the MRI interface cable **41** and connector **40c** can comprise non-magnetic MRI compatible material(s) and be configured to operate in a high magnetic field environment. As noted above, in some embodiments, the implantable pulse generator **50** as well as the implantable lead **51** and connector **50c** may also comprise MRI compatible materials to allow post-placement MRI interrogation of the subject.

In some embodiments, the probe **20** comprises an MRI antenna **35** that is configured to pick-up MRI signals in local tissue during an MRI procedure. The MRI antenna **35** can be configured to reside on the distal portion of the probe **20**. In some embodiments, the antenna **35** has a focal length or signal-receiving length of between about 1-5 cm, and typically is configured to have a viewing length to receive MRI signals from local tissue of between about 1-2.5 cm. The MRI antenna **35** is shown in **Figures 3B** and **4** as comprising a coaxial antenna. However, other antenna configurations can be used, such as, for example, a whip antenna, a coil antenna, a loopless antenna, and/or a looped antenna. *See, e.g.*, U.S. Patent Nos. 5,699,801; 5,928,145; 6,263,229; 6,606,513; 6,628,980; 6,284,971; 6,675,033; and 6,701,176, the

contents of which are hereby incorporated by reference as if recited in full herein. *See also* U.S. Patent Application Publication Nos. US 2003/0050557; US 2004/0046557; and 2003/0028095, the contents of which are also hereby incorporated by reference as if recited in full herein.

5 **Figures 3A-3B** illustrate one embodiment of a probe **20** with a plurality of electrodes **25**. At least one of the electrodes **25** can be a sensing electrode (*i.e.*, micro-electric recording). Typically, one or more of the electrodes **25** may be able to both sense and stimulate (or ablate). For neural uses, different regions in the brain provide different sensed intensities, frequencies and/or pitches (typically readings of between
10 about 1-4 microvolts) which are identifiable.

Figure 3B illustrates that the core of the probe **20** can be configured to hold a plurality of axially extending conductors **26**, typically a respective one for each electrode **25**. In other embodiments, greater or lesser numbers of conductors than electrodes may be used. The conductors **26** may be static and held generally
15 encapsulated in a first insulating dielectric layer **61**. In other embodiments, the conductors **26** may be held in the first dielectric material **61** so that they can translate in the axial and/or generally outward or transverse directions as shown in **Figure 3C**. Referring again to **Figure 3B**, an axially extending first shielding layer **62** can surround the first dielectric layer **61**. A second axially extending insulating dielectric
20 layer **63** can surround the first shielding layer **62**. A second axially extending shielding layer **64** can be electrically connected to the first shield layer **62** (that may also be called a primary shield layer) at a proximal end portion thereof. An outer polymeric insulator layer **65** can surround the inner layers **61-64** while terminating to typically expose the electrodes **25** to allow stronger stimulation contact during
25 operation. The conductors **26** extend from the connector **30** to the respective electrode **25**. The probe **20** includes an electrical ground **68** and the connector **30** connects the ground **68** and each electrode **25**.

Figure 4 illustrates an electrical schematic of the probe **20** shown in **Figures 3A** and **3B**. As shown, the primary or first shield layer **62** axially terminates at a
30 distal portion of the probe in advance of the first electrode **25₁**. The primary shielding **62** may be formed into a coil **62c** at a distal portion of the probe **20**. In other embodiments, the primary shielding **62** can terminate without coiling (not shown). In yet other embodiments, the shielding **62** may be coiled a distance past one or more electrodes **25**, including all the way forward to the distal end portion (not shown). In

some embodiments, a respective one conductor **26** can extend to a corresponding electrode **25**, with the longest conductor **26** corresponding to the more distal electrode **25**. The conductor(s) **26** may be substantially linear along the length in the probe body as shown, or may be coiled. If coiled, the coil for the conductor **26** may be at a distal portion, just before the respective electrode **25**, which may increase signal (not shown).

Each electrode **25** is typically in communication with at least one of the insulated conductors **26**. At the proximal end of the probe **20**, the conductors **26** are connected to a connector **30** so as to be connected to the implantable signal generator **50** or to the interface circuit **40** during MRI guided probe/lead/cable placement. These insulated conductors **26** are typically covered with a polymeric insulator sleeve **61** and a conducting material is cylindrically layered to form the first shielding layer **62** over the insulator. This shielding **62** is terminated proximal to the electrodes and is not in electrical contact with the conductors or the electrodes. A second insulator/polymeric/dielectric layer **63** further insulates this shielding to form multi-core coaxial type cable system with an impedance that is typically between about 10-100 ohms. The RF chokes **64rf** can be integrated or built into the shielding **64** in the form of a second shielding, which is not continuous and has multiple sections each $\lambda/4$ or less in length. As shown in **Figure 4**, at the proximal end, each section or segment **64s** is connected to the primary shielding **62**, and the distal end may not be electrically connected to the primary shielding **62**, or is connected with a capacitance **164** in between the primary and secondary shielding, **62**, **64**, respectively. A top insulator/polymeric layer **65** can be used to insulate the probe body **20b**, except for the electrodes **25**.

As shown by the axial arrow in **Figure 4**, the MRI active portion of the antenna **35** may extend between a location where the primary shield **62** terminates and the first electrode **25₁**. However, as noted above, other antenna **35** configurations may also be used. As shown, the second shield layer **64** comprises a plurality of axially spaced apart RF chokes **64rf**. The term "RF chokes" refers to a shielding layer configuration that provides an electrical length of less than or equal to $\lambda/4$ (from the perspective of external electromagnetic waves) to inhibit the formation and/or propagation of RF induced current or standing waves in an AC (alternating current, e.g., diathermy applications) or RF exposure environment. The physical length that provides the electrical wavelength may vary depending on the materials used in

fabricating the probe (such as dielectric constant) and the magnetic field in which it is used. In some embodiments, the probe **20** has a physical length that is greater than 10 cm, typically between about 20 cm to about 150 cm. In some embodiments, the implantable lead segment **50** can also include RF chokes **64rf** formed along target regions or along substantially the entire implantable length. In the embodiment shown in **Figure 4**, the RF chokes **64rf** comprise a plurality of disconnects of the shield **64** and/or discrete electrically isolated second shield segments. In other embodiments, the RF chokes **64rf** can include a series of axially spaced apart Balun circuits or other suitable circuit configurations. *See, e.g.*, U.S. Patent No. 6,284,971, the contents of which are hereby incorporated by reference as if recited in full herein, for additional description of RF inhibiting coaxial cable that can inhibit RF induced current.

As shown in **Figure 4**, the second shield layer **64** may be coupled to the first shielding layer **62** at opposing ends of the segments **64s**. As shown, one end (typically the proximal end portion) of the disconnected segment **64s** is directly coupled to the shielding layer **62** and the other end (typically the distal end portion) is capacitively coupled to the first shielding layer **62**. Each segment **64s** may be configured to engage the first shield layer **62** in the same manner or in an opposing different electrical manner (not shown).

Figures 5A-5C illustrate another embodiment of a probe **20**. In this embodiment, the probe body **20b** is configured to define an axially extending cavity **27** (**Figure 5B**) that is sized and configured at least one conductor **26** therein. **Figure 5B** illustrates the probe body **20b** without the conductor **26** with the cavity **27** open and **Figure 5C** illustrates the conductor **26** in position. The conductor **26** can be a single conductor **26** as shown or a plurality of (typically structurally bound or bundled) conductors. The conductor **26** can be configured similar to a guidewire that can be released from the probe body **20** once the probe body is in position and/or can provide structural rigidity to the probe **20** during placement. As shown in **Figure 5C**, in position, the conductor **26** can be configured to cooperate with the other components held on/in the probe **20** to define the MRI antenna **35** and to allow for MRI guidance. The probe **20** can provide a ground while the conductor **26** can provide the (+) electrical path. The conductor **26** can reside in the probe body during insertion or be extended through a distal end portion thereof. The conductor **26** can be packaged in position in the probe **20** or may be assembled on site. Instead of using

the internal conductor(s) to connect the electrodes, the probe body **20b** can include electrical conductive traces or wires **25t** (typically subsurface and under an insulation layer **65**) that connect the electrodes **25** to the input/output source(s). Although shown as having several dielectric and conductive shield layers in **Figures 5B** and **5C**, other probe body configurations can be used to provide the desired electrode and/or MRI antennas.

The probe body cavity **27** and/or conductor **26** can be configured with matable structures that allows for angular adjustment in the cavity. For example, one outer perimeter portion of the conductor **26** may be planarized, flattened or roughened.

Although not shown, in some embodiments, the probe **20** can be configured with one or more lumens and exit ports that deliver desired cellular, biological, and/or drug therapeutics to the target area, such as the brain. The probe **20** may also incorporate biopsy and/or injection needles and/or ablation means.

Embodiments of the present invention provide a multi-function MRI safe lead or probe **20** that can operate at least bimodally: namely, during MRI procedures to obtain MRI signal from local tissue *in vivo* and to stimulate and/or ablate the target tissue during and/or after an MRI procedure. In some embodiments, the probe **20** is chronically implantable and can operate in at least three modes to: (1) selectively sense microelectric activity in local tissue, (2) stimulate local tissue, and (3) obtain MRI signals of local tissue. Each of the operations are typically performed serially or independently of the others. The probe **20** can be configured for use in any suitable MRI scanner, such as low field magnets (typically about 0.5-1.0 T fields), to a conventional 1.5T magnet or higher, such as 2T, 3T or even higher. MRI scanners are well known to those of skill in the art and include, but are not limited to, SIEMENS and GE MRI systems.

Configuring a probe **20** to function both as an MRI antenna **35** and a stimulation probe may reduce the time needed to place the electrodes in the desired location, provide for increased accuracy in location and/or reduce the number of times a device is inserted into the brain or other target region.

Figure 6 illustrates a circuit **100** that can provide the bimodal operation of the probe **20**. As shown, the circuit **100** includes a splitter circuit **102** that is in communication with an electrode stimulation circuit **110** that provides the stimulation to the electrode(s) **25**. The splitter circuit **102** is also in communication with an RF transmit decoupler circuit **115** that is in communication with an MRI antenna RF

receive circuit **120** and the antenna **35**. Certain components or all of the components can be held in the MRI scanner interface **40**. In other embodiments, certain or all of the components of the circuit **100** can be held in the connector **30**.

Generally stated, the interventional probe system can have a plurality of
5 different operational modes such that the probe is associated with two or more operational transmission paths. In some embodiments, the probe can have two primary operational modes with different electric transmission paths, which are electrically directed using the splitter circuit **102**. In operation, during an MRI procedure, an RF excitation pulse is transmitted to a subject. The MRI antenna is
10 decoupled during RF transmission, then operative during a receive cycle to receive signal from local tissue. The stimulation electrodes **25** are typically isolated via the splitter circuit **102** so that only the MRI antenna **35** is active. The MRI interface **40** (**Figure 1**) communicates with the MRI scanner and may be configured with a supplemental port to allow the implantable pulse generator to connect thereto, thereby
15 allowing the IPG to stimulate the electrodes without decoupling the interface during the placement procedure (confirming proper placement). In some embodiments, the MRI interface **40** can include a stimulation and/or sensing mode that operates the electrodes independently of the IPG. In other embodiments, the connector **30** can be disconnected from the MRI interface after initial placement and connected to the IPG,
20 and reconnected to the MRI interface **40** as desired for additional MRI data and/or images.

During MRI guided clinical implantation of the probe **20**, the probe **20** can be used as an MRI antenna **35** to collect MRI or NMR data. Optionally, the probe **20** may be used to generate high-resolution images of the target internal anatomy (such
25 as neural tissue) and/or to locate the position of the electrodes **25** in the body by obtaining MRI signals and (whether local signal or images). The MRI signal can be acquired by the external coils and/or internal MRI antenna **35**. The electrodes **25** can also be used to assess location via acquiring electrical signals from the target (neural) anatomy.

Figures 7A and **7B** illustrate different circuits **100** that may be used to provide
30 the different operational modes of the probe **20**. **Figure 7A** illustrates an MRI antenna receive circuit **135** that receives the MRI responsive signal from local tissue and an RF transmit decoupler circuit **135D** that can decouple the antenna **35** and the electrodes during RF transmission. The circuit **100** also includes an electrode

stimulation circuit **125** that provides the stimulation pulses to the electrodes **25** and can include an electrode pulse filtering circuit **225** and a recording electrode circuit **226** used to gather local microelectric signals. **Figure 7B** illustrates that the circuit **100** can include a controller **200**, such as a digital signal processor, that can form part of the splitter circuit **102** (**Figure 6**) to direct operation of the MRI antenna mode or the electrode mode.

Figure 8A is a schematic illustration of an exemplary splitter circuit **102** that provides different transmission paths for signals operating in the imaging (MR signal) mode and in the sensing microelectrical mode according to some embodiments of the present invention. **Figure 8A** illustrates that the circuit **102** can have two sides, **102A**, **102B**, respectively that substantially overlies each other as shown in **Figure 8B** with a ground plane therebetween. Side A **102A** is the MRI antenna **35** active path with matching and tuning components including decoupling capacitors **127**, conductor connections **126** (to respective conductors **26**), an input (shown as a BNC input) to the MRI scanner **131**, an input to a multi-pin connector for an electrode pulse signal **132** (EP signal) a PIN diode **128**, a matching tuning inductor **129** and a matching/tuning circuit capacitor **130**. Side B is the electrode operational circuit configured to act as a high pass filter. As shown, the respective electrical transmission paths to the conductors **26** include capacitors **138** (shown as 1000 pF capacitors) and 64 MHz RF blocking inductors **139**. The blocking inductors **139** can be changed to block the frequency of the MRI system in use (higher frequencies for higher field magnets, *i.e.*, for proton imaging, 96 MHz for 2T, 128 MHz for 3T). It is noted that components of the exemplary circuits are shown with respect to side A or B for ease of discussion, but certain of the circuits (or the entire circuit) may reside on a different side than that shown (and are not required to be on one side).

In some embodiments, the probe **20** can be placed in the brain, such as in the subthalamic nucleus or other deep brain target via a burr hole formed in the skull. MR imaging using the probe **20** can guide an increased accurate placement in the thalamus or other desired anatomies. Further, the electrical signals from the local tissue can be analyzed and evaluated to determine a final location of the electrodes **25**. During this time, the probe can be connected to the MRI scanner interface **40** that can include a matching-tuning decoupling circuitry, and a splitter circuit to separate MR signal from the electrical signals generated by the local target tissue. Once the probe system is appropriately located in the desired anatomy, the stimulator can be

connected for physiological confirmation of the function. If satisfactory, the proximal section of the probe can be routed subcutaneously and connected to the signal generator **50** implanted in the chest cavity. A telescopic system to lengthen or shorten the lead may be implemented in the proximal section of the probe, since
5 diameter/profile may not be a significant concern in this region.

Figure 9 schematically illustrates a stereotaxis guidance system **400** that can be used to implant the probes **20**. Other guidance or implantation systems can also be used as will be known to those of skill in the art. In the embodiment shown, the system **400** comprises a generally circular ring **401** which is fixed to the patient's head.
10 This ring **401** has one or more rotating semicircular rings **402** that are configured to advance the probe **20** in substantially straight trajectory. The ring **401** has a locator arm **403** which has MRI fiducial markers **404** to identify the long axis plane in which the probe **20** will be advanced into the brain. MR imaging is carried out to identify the location of the distal end of the probe **20**. Once this is done a suitable long axis
15 plane is identified and the locator/directing arm **403** is fixed in this plane. The probe **20** is used as an MRI antenna and tracked during advancing into the desired anatomy, collecting electrical signals generated by the cranial anatomy in substantially real-time. Once the distal electrodes are located in the appropriate anatomy the stereotaxis guidance system **400** is removed and the probe **20** remains in location as an implanted
20 DBS lead.

The probe **20** may have one or more lumens configured to deliver cellular and/or biological therapeutics to the desired neural tissue. The lumens may receive extendable needles that may exit the probe from the distal end or from the sides, proximal, distal, or even, through the electrodes to precisely deliver cellular/biological
25 therapeutics to the desired anatomy target. This delivery configuration may be a potential way to treat patients, where the cellular/biological therapeutics are delivered into the desired anatomy and the neurotransmitter/signal generator paces the cells to modify their function. In this way, even if the signal generator fails, the cells (stem cells) may differentiate and take over the function. MRI can be effectively used to
30 monitor the efficacy of the therapy in the brain.

Figures 10A and **10B** illustrate a dual probe system according to other embodiments of the present invention. In this embodiment, an MRI antenna probe **120a** and a stimulation probe **120b** can be sized and configured to serially enter a common cannula **75**. The antenna probe **120a** and the stimulation probe **120b** can

each include at least one sensing electrode. Each probe **120a**, **120b** can have a graduated scale or coordinate system that allows the antenna probe **120a** to be used to obtain MRI imaging data used to locate the target *in vivo* location. The cannula **75** can include MRI fiducial markers (not shown). The antenna probe **120a** can then be removed and replaced with the stimulation probe **120b** that can be automatically advanced in the same trajectory to the same position based on the data provided by the antenna probe **120a** and the controlled insertion to the location defined by the antenna probe **120a**, typically to a high degree of precision. The two probes **120a**, **120b** can be sized and configured to have substantially the same cross-sectional area. In some embodiments, a non-conductive elastomeric sleeve (not shown), coating or other configuration can be used to size the probes **120a**, **120b** to snugly fit the cannula **75** as desired. In other embodiments, an insert can be used to adjust the size of the cannula **75** to correspond to that of the probe in use (also not shown). The cannula **75** and both probes **120a**, **120b** are MRI compatible and at least the probes **120a** and **120b** may include the RF chokes **64rf** (**Figure 4**).

In some embodiments, the antenna probe **35** can define a relatively small MRI receiver length "L," such as less than about 5 cm, typically between about 1-2.5 cm as noted above. As before, the antenna **35** can be any suitable type and is not limited to a coaxial cable type (including, for example, a dipole or loopless antenna as discussed above). The cannula **75** can form a shielding layer. In some embodiments, the cannula **75** may comprise a polymer and may include MRI compatible conductive material, such as Nitonal.

In some embodiments, the cannula **75** can be configured to cooperate with the antenna probe **120a** to define an antenna **35**. The cannula **75** can provide a ground and positive signal path. With reference again to **Figure 4**, the cannula **75** can provide one or more insulating layers **61**, **63** or shielding layers **62**, **64** with the antenna probe **120a** providing at least one conductor **26** and potentially one or more of the insulating layer **61** or shielding layers **62**, **64**. In particular embodiments, the cannula **75** provides the secondary shield layer **64** and may include RF chokes **64rf**.

Figure 10B illustrates that a kit **10'** can comprise the two probes **120a**, **120b** and, optionally, the cannula **75**. The antenna probe **120a** can be configured to connect with the MRI interface **40** (**Figure 1**) while the stimulation probe **120b** can be configured to connect to the implantable pulse generator **50**, each of which (along with respective leads **41**, **51**) may also form part of the medical kit **10'**.

Figure 11 is a block diagram of exemplary embodiments of data processing systems that illustrates systems, methods, and computer program products in accordance with embodiments of the present invention. The data processing systems may be incorporated in a digital signal processor in either the implantable pulse generator **50** and/or MRI scanner interface **40** and/or be in communication therewith. The processor **410** communicates with the memory **414** via an address/data bus **448**. The processor **410** can be any commercially available or custom microprocessor. The memory **414** is representative of the overall hierarchy of memory devices containing the software and data used to implement the functionality of the data processing system. The memory **414** can include, but is not limited to, the following types of devices: cache, ROM, PROM, EPROM, EEPROM, flash memory, SRAM, and DRAM.

As shown in **Figure 11**, the memory **414** may include several categories of software and data used in the data processing system: the operating system **452**; the application programs **454**; the input/output (I/O) device drivers **458**; the MRI Antenna operation or Electrode Operation Module **450**; and data **456**.

As will be appreciated by those of skill in the art, the operating system **452** may be any operating system suitable for use with a data processing system, such as OS/2, AIX, DOS, OS/390 or System390 from International Business Machines Corporation, Armonk, NY, Windows CE, Windows NT, Windows95, Windows98, Windows2000 or other Windows versions from Microsoft Corporation, Redmond, WA, Unix or Linux or FreeBSD, Palm OS from Palm, Inc., Mac OS from Apple Computer, LabView, or proprietary operating systems. The I/O device drivers **458** typically include software routines accessed through the operating system **452** by the application programs **454** to communicate with devices such as I/O data port(s), data storage **456** and certain memory **414** components. The application programs **454** are illustrative of the programs that implement the various features of the data processing system and can include at least one application, which supports operations according to embodiments of the present invention. Finally, the data **456** represents the static and dynamic data used by the application programs **454**, the operating system **452**, the I/O device drivers **458**, and other software programs that may reside in the memory **414**.

While the present invention is illustrated, for example, with reference to the Module **450** being an application program in **Figure 11**, as will be appreciated by

those of skill in the art, other configurations may also be utilized while still benefiting from the teachings of the present invention. For example, the Module 450 may also be incorporated into the operating system 452, the I/O device drivers 458 or other such logical division of the data processing system. Thus, the present invention
5 should not be construed as limited to the configuration of Figure 11 which is intended to encompass any configuration capable of carrying out the operations described herein. Further, the Module 450 can communicate with other components, such as an MRI scanner.

The I/O data port can be used to transfer information between the data
10 processing system, the product pusher, and the clipper mechanism or another computer system or a network (*e.g.*, the Internet) or to other devices controlled by the processor. These components may be conventional components such as those used in many conventional data processing systems, which may be configured in accordance with the present invention to operate as described herein.

15 The computer-readable program code can include computer readable program code that controllably engages a first or second operational mode for a MRI compatible stimulation probe with at least one electrode and an MRI antenna. The first operational mode having a first transmission path connecting the MRI antenna with an MRI scanner and decoupling the electrodes during MRI operation and the
20 second operational mode having a second transmission path connecting the electrodes with a stimulation or recording source during electrical stimulation or recording.

The computer readable program code may be configured to time the selection of the second operational mode to occur proximate in time but after an MRI signal acquisition by the MRI antenna in the first operational mode. The computer readable
25 program code may be configured to obtain microrecordings of local tissue in substantially real time proximate in time to an MRI signal acquisition by the MRI antenna in the first operational mode. The computer readable program code may be configured to obtain a plurality of MRI signals of local neural tissue proximate the MRI antenna in substantially real time, then obtain a plurality of microrecordings of
30 the local neural tissue to allow a clinician to track placement of the probe using both MRI data and audio data.

The flowcharts and block diagrams of certain of the figures herein illustrate the architecture, functionality, and operation of possible implementations of the present invention. In this regard, each block in the flow charts or block diagrams

represents a module, segment, or portion of code, which comprises one or more executable instructions for implementing the specified logical function(s). It should also be noted that in some alternative implementations, the functions noted in the blocks may occur out of the order noted in the figures. For example, two blocks
5 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.

In the drawings and specification, there have been disclosed embodiments of the invention and, although specific terms are employed, they are used in a generic
10 and descriptive sense only and not for purposes of limitation, the scope of the invention being set forth in the following claims. Thus, the foregoing is illustrative of the present invention and is not to be construed as limiting thereof. Although a few exemplary embodiments of this invention have been described, those skilled in the art will readily appreciate that many modifications are possible in the exemplary
15 embodiments without materially departing from the novel teachings and advantages of this invention. Accordingly, all such modifications are intended to be included within the scope of this invention as defined in the claims. In the claims, means-plus-function clauses, where used, are intended to cover the structures described herein as performing the recited function and not only structural equivalents but also equivalent
20 structures. Therefore, it is to be understood that the foregoing is illustrative of the present invention and is not to be construed as limited to the specific embodiments disclosed, and that modifications to the disclosed embodiments, as well as other embodiments, are intended to be included within the scope of the appended claims. The invention is defined by the following claims, with equivalents of the claims to be
25 included therein.

THAT WHICH IS CLAIMED IS:

1. An *in vivo* medical stimulation probe, comprising:
an elongate lead having at least one stimulation electrode disposed on a distal
5 portion thereof and an axially extending shielding layer; and
a plurality of axially spaced apart RF chokes disposed on and/or in the
shielding layer in advance of the at least one electrode to inhibit induced RF current
from forming and/or traveling along the shielding layer.
- 10 2. A medical stimulation probe according to Claim 1, the lead further
comprising a recording/sensing electrode disposed at the distal portion.
3. A medical stimulation probe according to Claim 1, wherein the lead is
a flexible lead, and wherein the at least one electrode is a plurality of spaced apart
15 electrodes, the lead further comprising a plurality of conductors held in a core of the
lead, a respective one for each electrode, and wherein the shielding layer is
discontinuous and is configured to surround the conductors over at least a major
length of the lead and terminate at a lead location that is in advance of the electrodes.
- 20 4. A medical stimulation probe according to Claim 3, further comprising
an axially extending primary inner shield surrounding the core with the plurality of
electrodes, wherein the discontinuous shielding layer is a second shield layer that is
generally cylindrically disposed over the inner shielding layer, and wherein the
primary and second shielding layers both terminate at a distal location of the lead that
25 precedes the electrodes.
5. A medical stimulation probe according to Claim 3, further comprising
a first insulating dielectric layer disposed between the conductors and the primary
inner shield, and a second insulating dielectric layer disposed between the primary
30 inner shield and the secondary shield, wherein the conductors, the first and second
insulating layers and the primary inner and second shield layers comprise MRI
compatible materials, and wherein the RF chokes are formed in the secondary shield
to provide an electrical length that is about $\lambda/4$ or less.

6. A medical stimulation probe according to Claim 5, the lead further comprising a bio- and MRI-compatible outer polymeric layer.

5 7. A medical stimulation probe according to Claim 1, wherein the lead is MRI-compatible and further comprises an MRI antenna that is configured to collect and transmit signal data in response to an applied RF excitation pulse in an MRI scanner system.

10 8. A medical stimulation probe according to Claim 7, wherein the MRI antenna is a coaxial antenna having a signal receiving length of between about 1-4 cm.

15 9. A medical stimulation probe according to Claim 1, wherein the lead is chronically implantable, and wherein the at least one electrode is a plurality of electrodes that are sized and configured to apply deep brain stimulation.

20 10. A medical stimulation probe according to Claim 7, further comprising a connector disposed at a proximal end portion thereof, the connector configured to releasably serially attach to both an implantable pulse generator and an MRI scanner interface, thereby allowing bimodal operation of the lead.

25 11. A medical stimulation probe according to Claim 7, wherein the lead is configured to have at least two operational modes, including a first MRI operational mode wherein the lead receives MRI signals from the MRI antenna and a second therapeutic operational mode wherein the lead delivers a stimulation pulse to the at least one electrode.

30 12. A medical stimulation probe according to Claim 11, wherein the at least one electrode is a plurality of electrodes, at least one of which is configured as a recording electrode, and wherein the lead is configured to provide trimodal operation with a third operational mode wherein the lead receives micro-electrical signals from local tissue from the recording/sensing electrode.

13. A medical stimulation probe according to Claim 11, wherein the MRI antenna is configured to receive signal from local tissue over about a distance of about 2.5 cm or less.

5 14. A medical stimulation probe according to Claim 13, wherein the MRI antenna is axially spaced apart from the closest electrode a distance of about 1-4 cm.

15. A medical stimulation probe according to Claim 9, wherein the lead has a length that is greater than 10 cm.

10

16. A chronically implantable deep brain stimulation and MRI imaging probe system, comprising:

an internal MRI-compatible elongate lead comprising an MRI antenna having axially extending, radially spaced apart first and second shielding layers and at least one core conductor, for internally receiving local *in vivo* MRI signals;

15

at least one electrode held on a distal portion of the lead that, in operation, is configured to generate a stimulation pulse to deep brain tissue;

a stimulation circuit in communication with the at least one electrode;

a MRI signal receiver circuit in communication with the MRI antenna;

20

a splitter circuit in communication with the stimulation and receiver circuits for electrically connecting either the MRI receive or stimulation circuit; and

means for inhibiting RF induced current on the second shield layer of the lead.

17. A deep brain stimulation and MRI imaging probe according to Claim 16, further comprising a decoupling circuit to electrically decouple the MRI antenna during an MRI RF transmit operation.

25

18. A deep brain stimulation and MRI imaging probe according to Claim 16, wherein the means for inhibiting comprises a plurality of axially spaced apart discontinuities in the second shield layer.

30

19. A deep brain stimulation and MRI imaging probe according to Claim 16, wherein the means for inhibiting comprises axially spaced apart RF choke circuits in communication with the second shield layer.

20. A deep brain stimulation and MRI imaging probe according to Claim 19, wherein the RF choke circuits comprise Balun circuits.

5 21. A deep brain stimulation and imaging probe according to Claim 17, further comprising a microrecording electrode for receiving microelectric signals associated with neural tissue.

10 22. A deep brain stimulation and imaging probe according to Claim 17, wherein the lead merges into a connector at a proximal portion thereof, the connector configured to serially interchangeably engage an MRI scanner interface and an implantable pulse generator.

15 23. An MRI compatible deep brain stimulation and MRI signal acquiring probe system comprising:

a flexible elongate probe body having opposing proximal and distal portions, the probe body comprising a plurality of electrodes disposed on the distal portion;

a plurality of axially extending conductors disposed in a core of the probe body, a respective one of the conductors associated with each electrode;

20 an axially extending inner shield surrounding the plurality of conductors for at least a major portion of the length of the conductors;

an axially extending second shield radially spaced apart above the inner shield;

25 an axially extending first insulating/dielectric layer disposed intermediate of the inner and second shields;

an MRI antenna member disposed at least partially on and/or in the probe body to be able to collect MRI signals at the distal portion thereof;

an RF transmit decoupling circuit in communication with the MRI antenna member; and

30 at least one connector attached to the proximal portion of the probe body, configured to hold a conductor transmission line for each of the electrodes.

24. A system according to Claim 23, wherein the connector is configured to attach to an MRI scanner having a splitter circuit for selectively operating the electrodes or the MRI antenna member.

5 25. A system according to Claim 23, further comprising an MRI interface that holds the RF transmit decoupler circuit, the MRI interface configured to connect the lead to an MRI scanner with the decoupler circuit decoupling the MRI antenna member during an MRI RF excitation transmission.

10 26. A system according to Claim 23, wherein the RF transmit decoupler circuit is held in the connector.

15 27. A system according to Claim 25, wherein the decoupler circuit comprises a matching and tuning decoupling circuit that engages an MRI scanner and decouples the electrodes.

28. A system according to Claim 23, further comprising a splitter circuit in communication with the RF transmit decoupler circuit and the stimulation circuit.

20 29. A system according to Claim 28, wherein the stimulation circuit comprises a high pass filter disposed in a transmission path intermediate a stimulation source and the electrodes.

25 30. A system according to Claim 29, wherein the stimulation circuit is configured to block RF signals at a resonant frequency associated with a magnetic field strength of a MRI system used to operate an MRI antenna defined at least partially by the antenna member and/or probe.

30 31. A system according to Claim 23, further comprising a splitter circuit having selective operative first and second electrical transmission paths associated with first and second operational modes, the first transmission path connecting the MRI antenna member with the MRI scanner and decoupling the electrodes during MRI operation and the second transmission path connecting the electrodes with a

stimulation or recording source during electrical stimulation or recording, respectively.

5 32. A system according to Claim 31, wherein the splitter circuit is configured to act as a high pass filter during MRI operation to electrically isolate the second transmission path from the first transmission path.

10 33. A system according to Claim 31, wherein the splitter circuit is held in an MRI interface that is releaseably attachable to the connector.

34. A system according to Claim 33, wherein the splitter circuit is held in the connector.

15 35. A system according to Claim 27, wherein the lead is configured to serially engage with an MRI scanner and an implantable pulse generator.

36. A system according to Claim 35, wherein the implantable pulse generator is MR-compatible.

20 37. A system according to Claim 27, wherein the lead comprises at least one lumen and associated port configured to emit a therapeutic fluid to neural tissue.

25 38. A system according to Claim 27, wherein the lead is configured to slidably enter a cannula with the connector held outside the cannula during deep brain placement.

39. A system according to Claim 38, wherein the cannula is MRI-compatible.

30 40. A system according to Claim 38, wherein the system is configured to operate with an MRI-compatible stereotactic guidance system during deep brain placement.

41. A medical kit, comprising:
an elongate sterilized biocompatible and MRI-compatible lead having
opposing distal and proximal portions, the lead comprising an MRI antenna, a
plurality of stimulation electrodes on the distal portion, and a plurality of axially
5 spaced apart RF chokes disposed in a shielding layer of the lead, the lead configured
to have selective operative first and second electrical transmission paths associated
with at least first and second operational modes, the first transmission path connecting
the MRI antenna with an MRI scanner and decoupling the electrodes during MRI
operation and the second transmission path connecting the electrodes with a
10 stimulation, ablation or recording source during electrical stimulation, ablation or
recording, respectively.

42. A medical kit according to Claim 41, wherein the lead is configured and
sized to be chronically implantable for deep brain stimulation.

15

43. A medical kit according to Claim 41, wherein the lead comprises a
connector at the proximal end portion thereof, the connector configured to engage a
chronically implantable pulse generator in the second operational mode.

20 44. A medical kit according to Claim 41, wherein the connector is configured
to be in communication with an MRI scanner in the first operational mode.

45. A medical kit according to Claim 44, wherein the connector is configured
to connect to an MRI interface with a splitter circuit and RF transmit decoupler circuit
25 with the MRI interface connecting to the MRI scanner.

46. A method of placing and operating a deep brain stimulation probe,
comprising:
inserting a flexible elongate lead comprising an MRI antenna and at least one
30 stimulation electrode on a distal portion thereof into a brain of a subject;
connecting the lead to an MRI scanner interface in communication with a
splitter circuit having a first electric transmission path for MRI operation and a second
electric transmission path for stimulation operation;

obtaining MRI signals associated with local neural tissue proximate the MRI antenna from the MRI antenna using the first transmission path;

placing the electrode on the lead at a desired location in the brain based on the obtaining step; then

5 stimulating neural tissue with the electrode using the second transmission path; and

configuring the lead to inhibit the formation and/or transmission of RF induced current,

10 wherein the stimulating and obtaining steps are carried out using the same lead.

47. A method according to Claim 46, further comprising:

implanting the lead in the brain with the electrode held at the desired location in the brain;

15 connecting the lead to an implantable pulse generator; and

stimulating the neural tissue with the at least one electrode based on a stimulation signal transmitted from the implantable pulse generator to provide a therapeutic treatment.

20 48. A method according to Claim 46, wherein the placing step comprises inserting the lead through a bore of a MRI-compatible cannula body having increased rigidity relative to the lead, the cannula body extending through the skull into neural tissue.

25 49. A method according to Claim 46, wherein the placing step is carried out using a stereotaxis guidance system with MRI fiducial markers.

50. A method according to Claim 46, wherein the obtaining step is repeated and carried out in substantially real-time to track the location of the lead in the brain.

30

51. A method according to Claim 46, wherein the lead comprises an axially extending lumen with at least one exit port, the method further comprising releasing a therapeutic fluid into the brain.

52. A method according to Claim 46, wherein the obtaining step is repeated a plurality of times before the placing step, the method further comprising recording microelectric audio signals sensed from the at least one electrode and transmitting the audio signals to an external audio device intermediate at least some of the obtaining
5 steps.

53. A method according to Claim 52, wherein the MRI antenna and at least one electrode are decoupled during RF transmission in an MRI imaging session.

10 54. A method according to Claim 46, wherein the at least one electrode is a plurality of electrodes, at least one of which is both a recording and stimulation electrode, and wherein the MRI antenna has a viewing length of between about 1-4 cm.

15 55. A method according to Claim 46, further comprising splitting the operation of the lead into an MRI operation and a stimulation operation, wherein the electrodes have an electrical circuit transmission path that is different during stimulation and MRI operation.

20 56. A method according to Claim 46, wherein the at least one electrode is a plurality of electrodes, the lead having an axially extending electrical conductor held in a core for each of the electrodes, each conductor generally extending from a respective electrode to a common connector, wherein the method further comprises selectively attaching the connector to an MRI system or to an implantable pulse
25 generator, depending on the desired operational mode.

57. A computer program product for operating a multi-purpose MRI compatible stimulation probe and MRI antenna, the computer program product comprising:

30 a computer readable storage medium having computer readable program code embodied in said medium, said computer-readable program code comprising:

computer readable program code that controllably engages one of a plurality of operation modes including a first or second operational mode for an MRI-compatible stimulation probe with at least one electrode and an MRI antenna, the first

operational mode having a first transmission path connecting the MRI antenna with an MRI scanner and decoupling the electrodes during MRI operation and the second operational mode having a second transmission path connecting the electrodes with a stimulation or recording source during electrical stimulation or recording.

5

58. A computer program product according to Claim 57, wherein the computer readable program code is configured to time the selection of the second operational mode to occur proximate in time but after an MRI signal acquisition by the MRI antenna in the first operational mode.

10

59. A computer program product according to Claim 57, further comprising computer readable program code that is configured to obtain microrecordings of local tissue in substantially real time proximate in time to an MRI signal acquisition by the MRI antenna in the first operational mode.

15

60. A computer program product according to Claim 57, further comprising computer readable program code that is configured to obtain a plurality of MRI signals of local neural tissue proximate the MRI antenna in substantially real time, then obtain a plurality of microrecordings of the local neural tissue to allow a clinician to track and/or guide placement of the probe.

20

61. An MRI compatible therapeutic stimulation probe comprising:
an elongate flexible probe body having an axially extending internal cavity disposed therein;

25

at least one electrode held by a distal portion of the probe body; and
at least one axially extending conductor configured to slidably extend into the cavity of the probe body, the at least one conductor having increased rigidity relative to the probe body,

30

wherein, during positioning in a body, the at least one conductor cooperates with the probe body and defines an *in vivo* MRI antenna used to obtain MRI signals for MRI positional guidance, and wherein, after placement, the at least one conductor can be removed from the probe body, leaving the probe body and electrode in position in the body.

62. An MRI probe according to Claim 61, wherein the at least one electrode is a plurality of electrodes, at least one of which configured as a sensing and stimulation electrode.

5 63. An MRI probe according to Claim 62, wherein the at least one conductor is a plurality of attached insulated conductors.

64. An MRI probe according to Claim 61, wherein the probe body comprises a shielding layer with a plurality of axially spaced apart RF chokes.

10

65. An MRI probe according to Claim 64, wherein the probe body comprises a first insulating layer, a first shielding layer, a second insulating layer and a second shielding layer, and wherein the RF chokes are disposed in the second shielding layer.

15

66. An MRI antenna and stimulation probe system, comprising:
an MRI compatible cannula having an axially extending bore; and
an elongate flexible stimulation and internal MRI signal acquisition probe having at least one electrode held by a distal portion of the probe;
wherein the stimulation and antenna probe are configured to slidably extend
20 through the cannula bore and wherein, in operation, the cannula and antenna probe cooperate to define components of at least one deep brain MRI signal receive antenna.

25

67. An MRI antenna probe and stimulation probe system according to Claim 66, wherein the cannula is configured to be inserted into a burr hole placed in a patient's skull, and wherein the stimulation and MRI antenna probe are configured for deep brain placement.

30

68. An MRI antenna probe and stimulation probe system according to Claim 66, wherein the cannula comprises a plurality of generally concentric tubular members configured to define an axially extending shield disposed over an inner conductive core, with the shield and core being insulated from each other.

69. An MRI antenna probe and stimulation probe system according to Claim 66, wherein the cannula comprises a conductive shielding layer that cooperates with

the stimulation and MRI antenna probe to define an MRI receive antenna during positioning to obtain MRI signals for MRI positional guidance.

5 70. An MRI antenna probe and stimulation probe system according to Claim 69, wherein the stimulation probe is configured to remain implanted in the body.

10 71. An MRI antenna probe and stimulation probe system according to Claim 66, wherein the stimulation and MRI antenna probe comprises a plurality of axially spaced apart RF chokes.

15 72. An MRI antenna probe and stimulation probe system according to Claim 66, the system further comprising a controlled placement system that is configured to determine the positional location of the stimulation and antenna probe when held at a target region in the brain.

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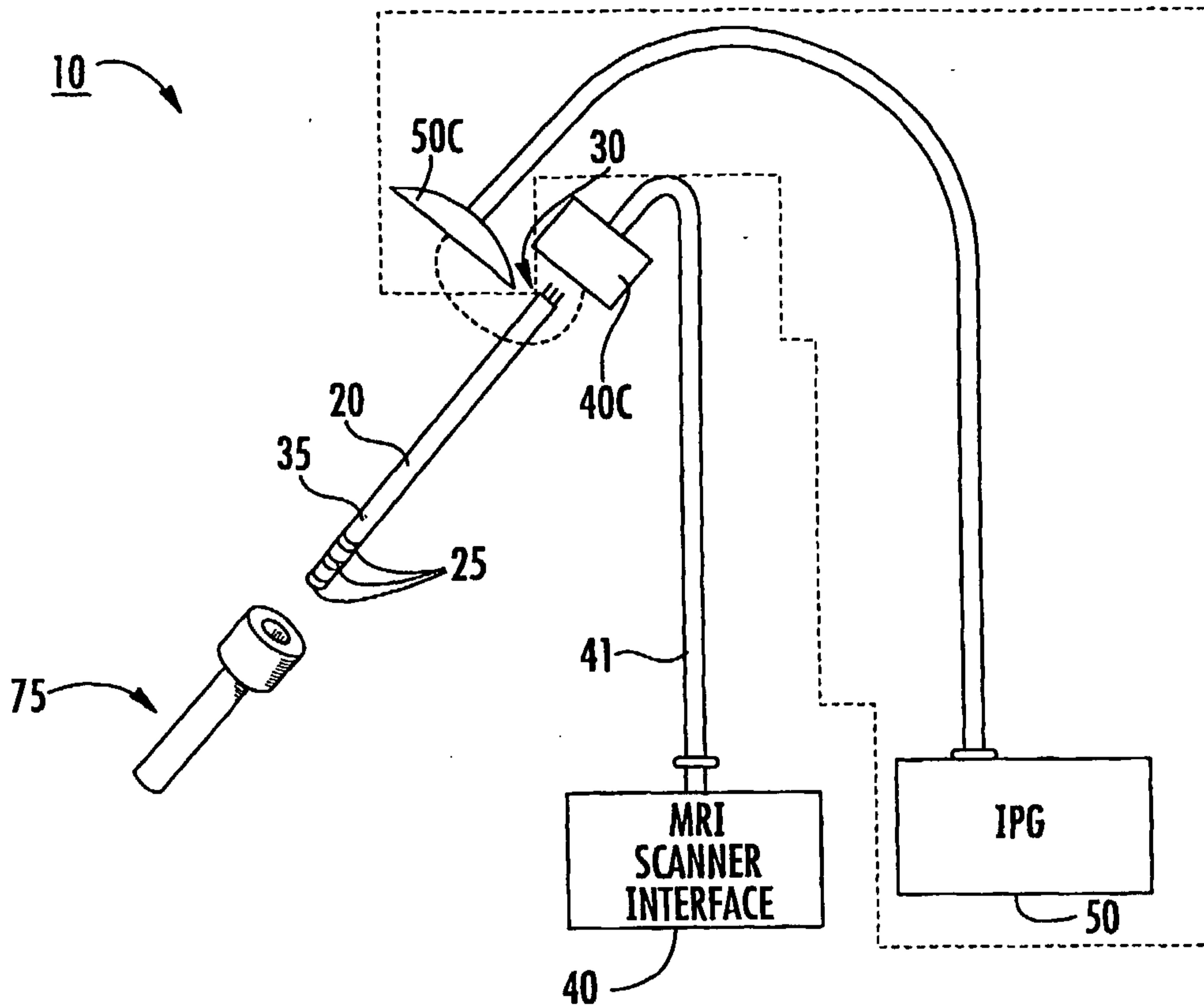


FIG. 1

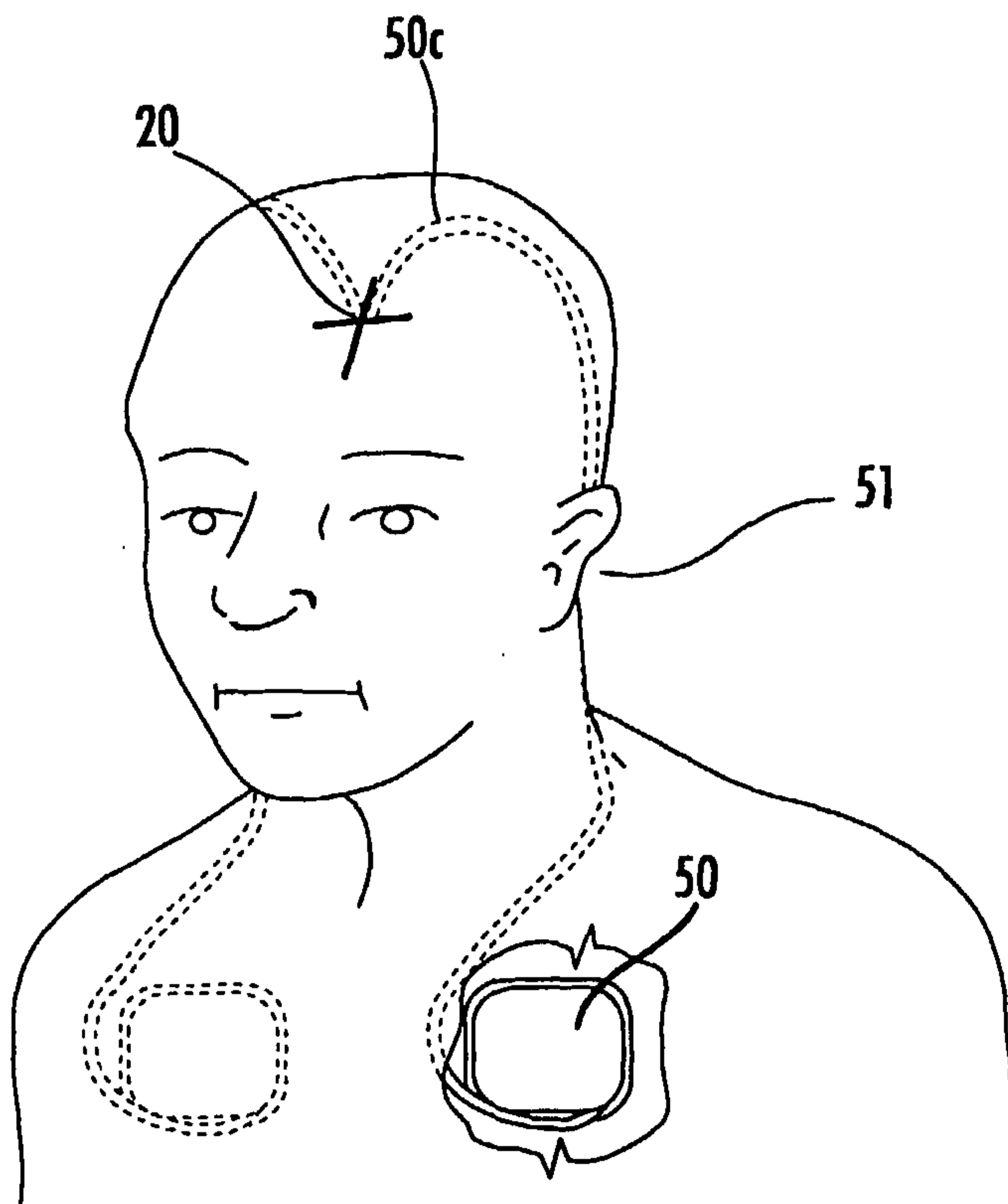


FIG. 2

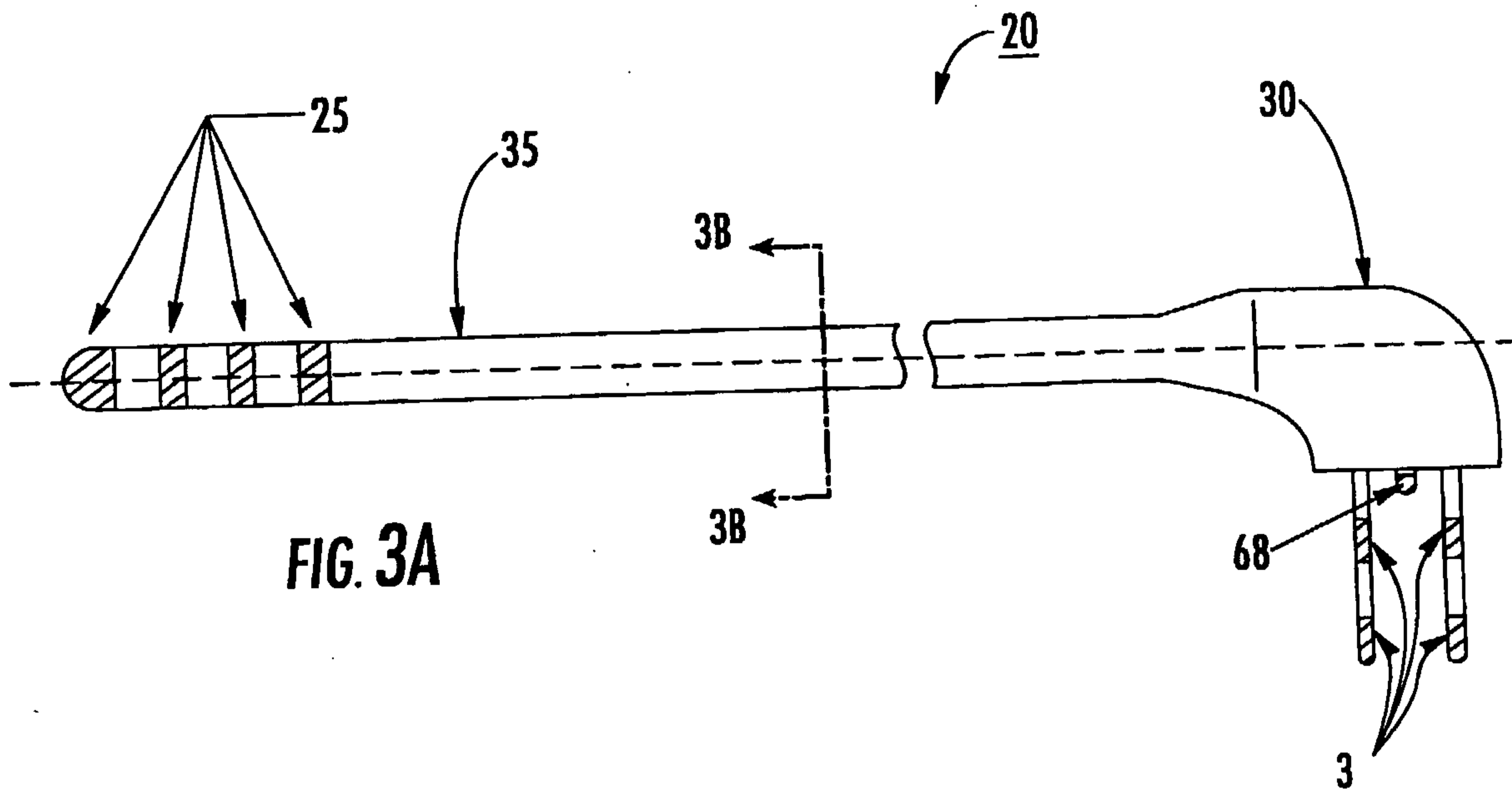


FIG. 3A

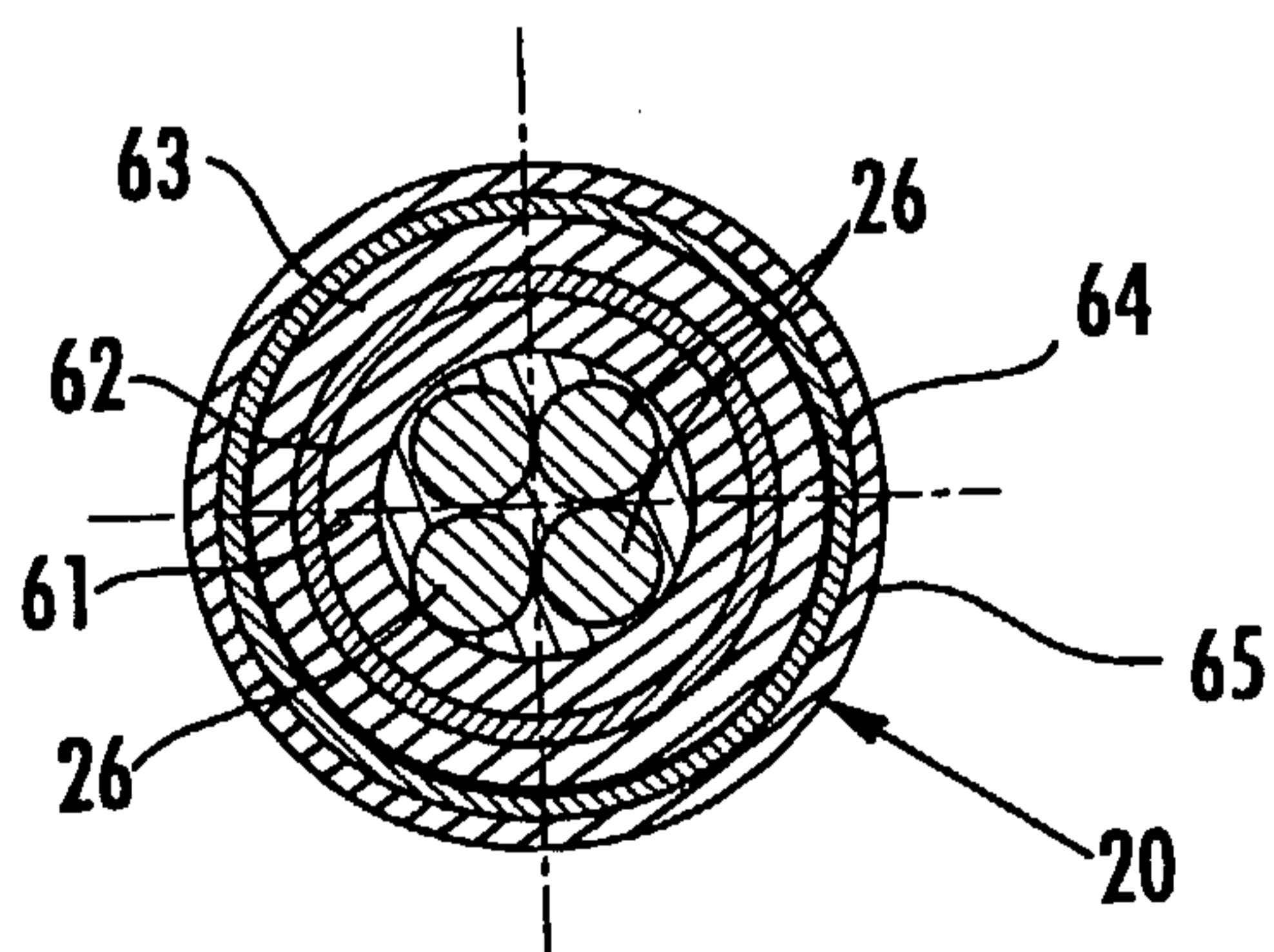
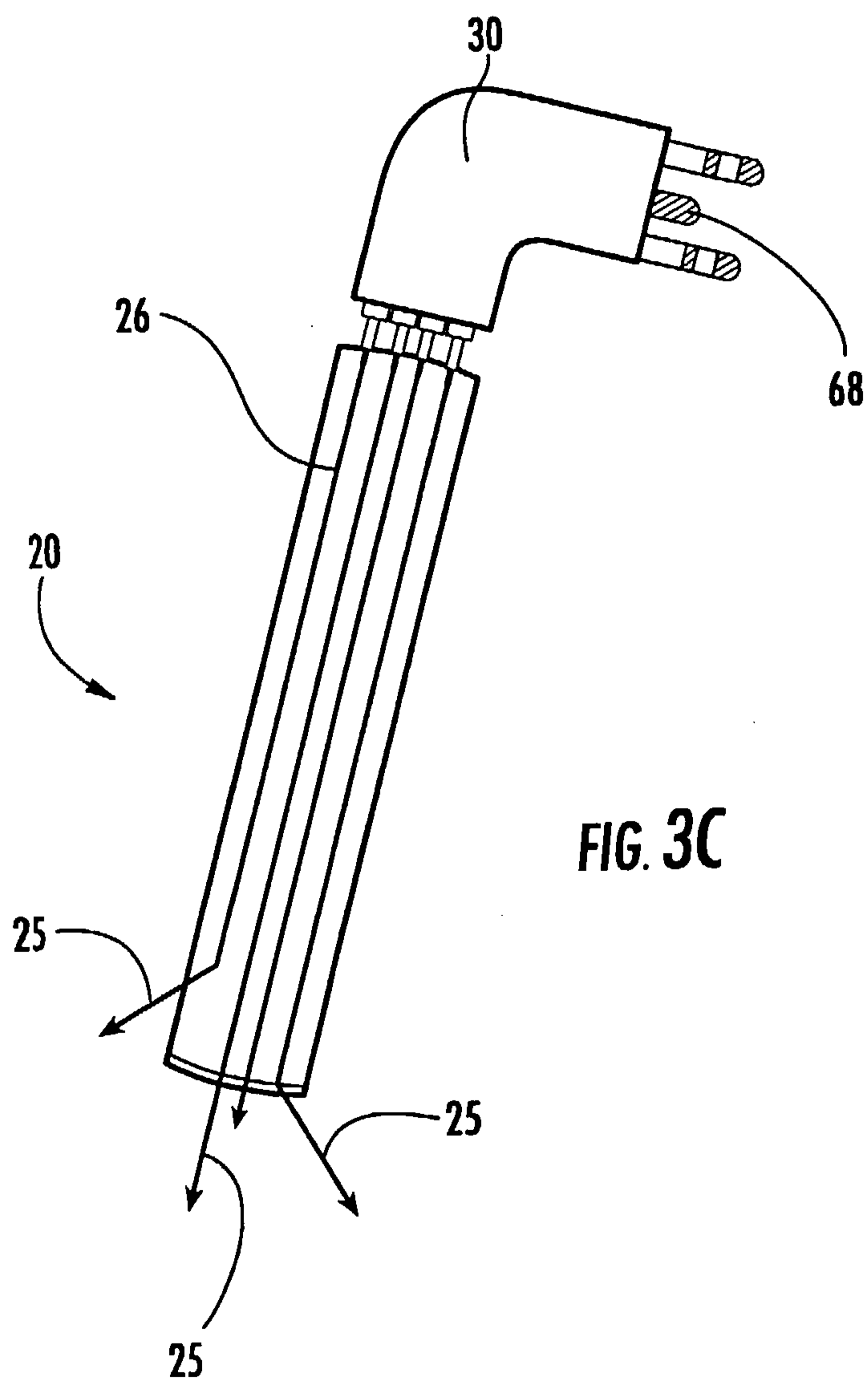


FIG. 3B



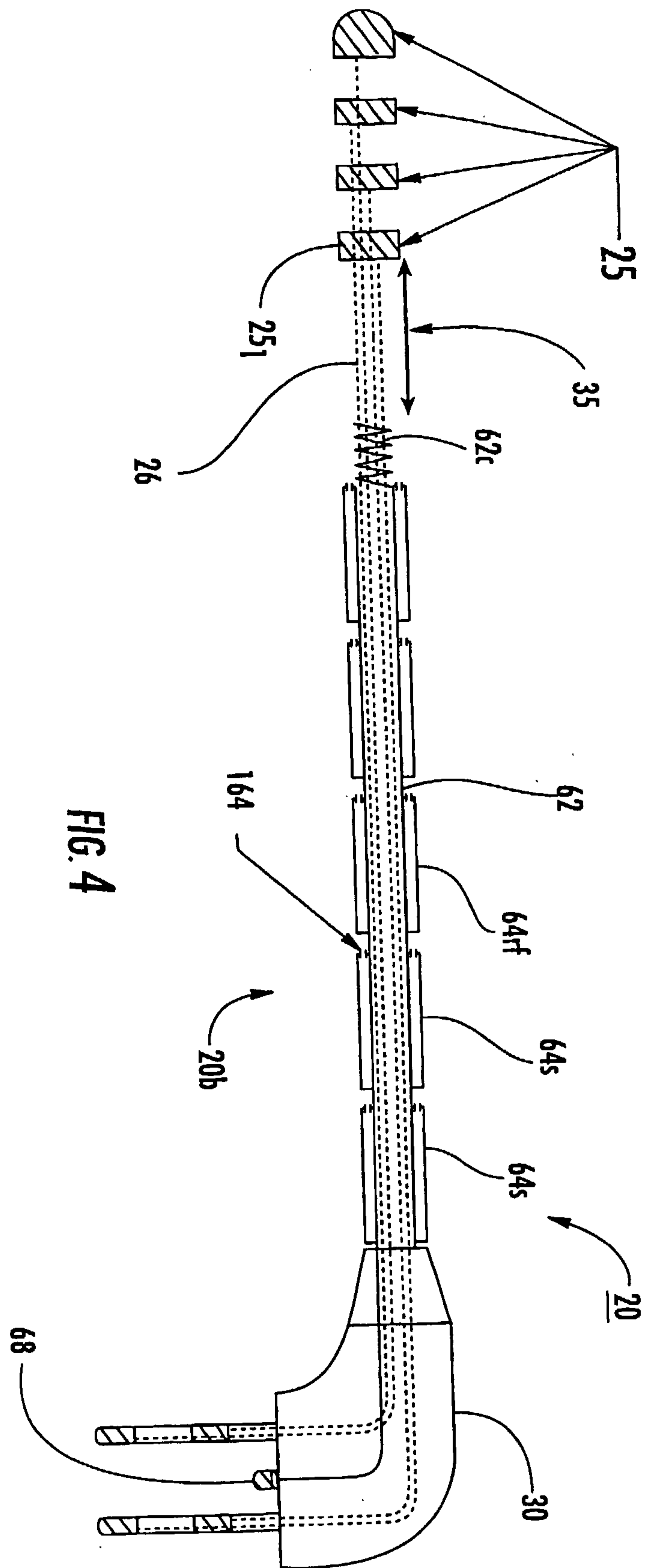


FIG. 4

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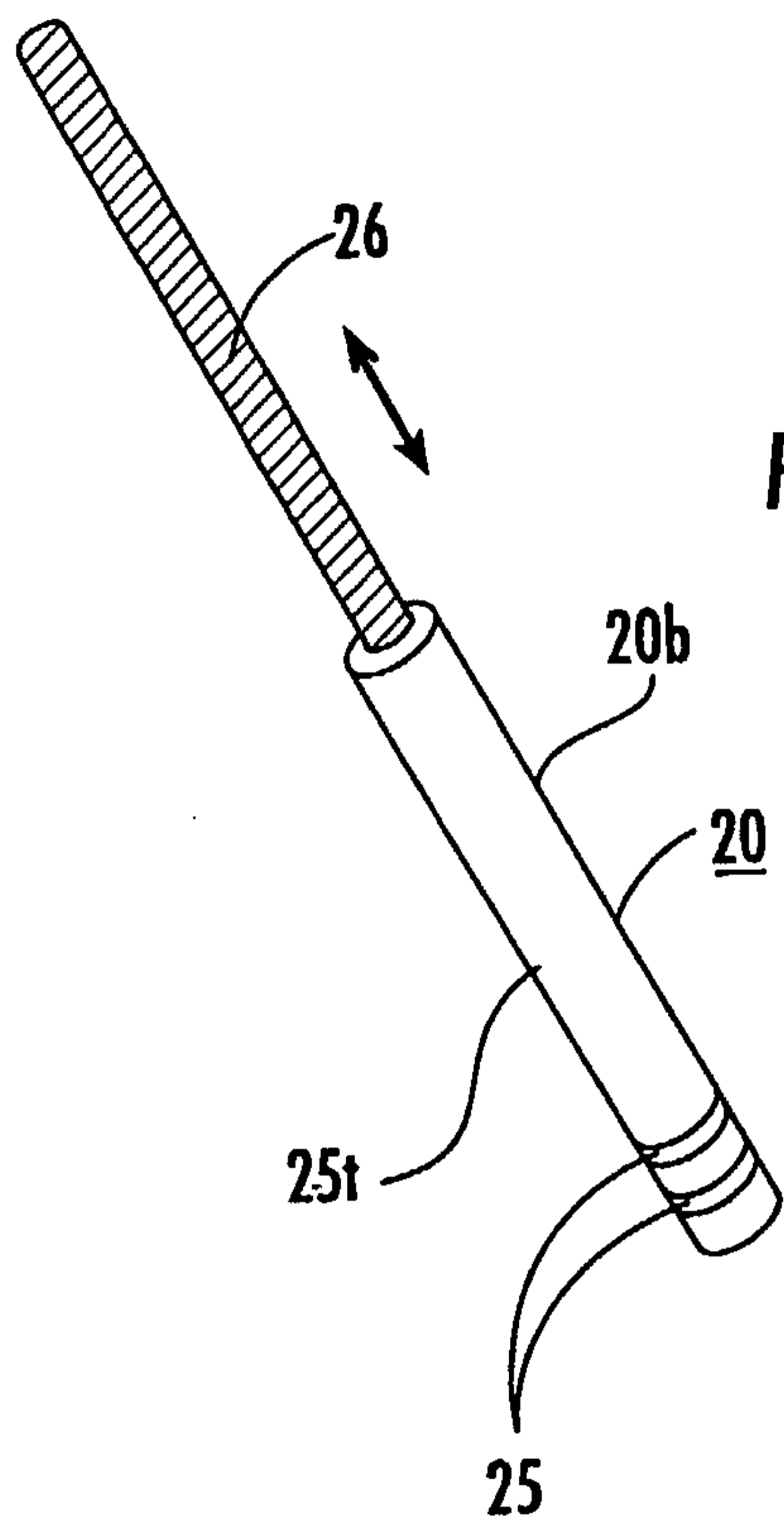


FIG. 5A

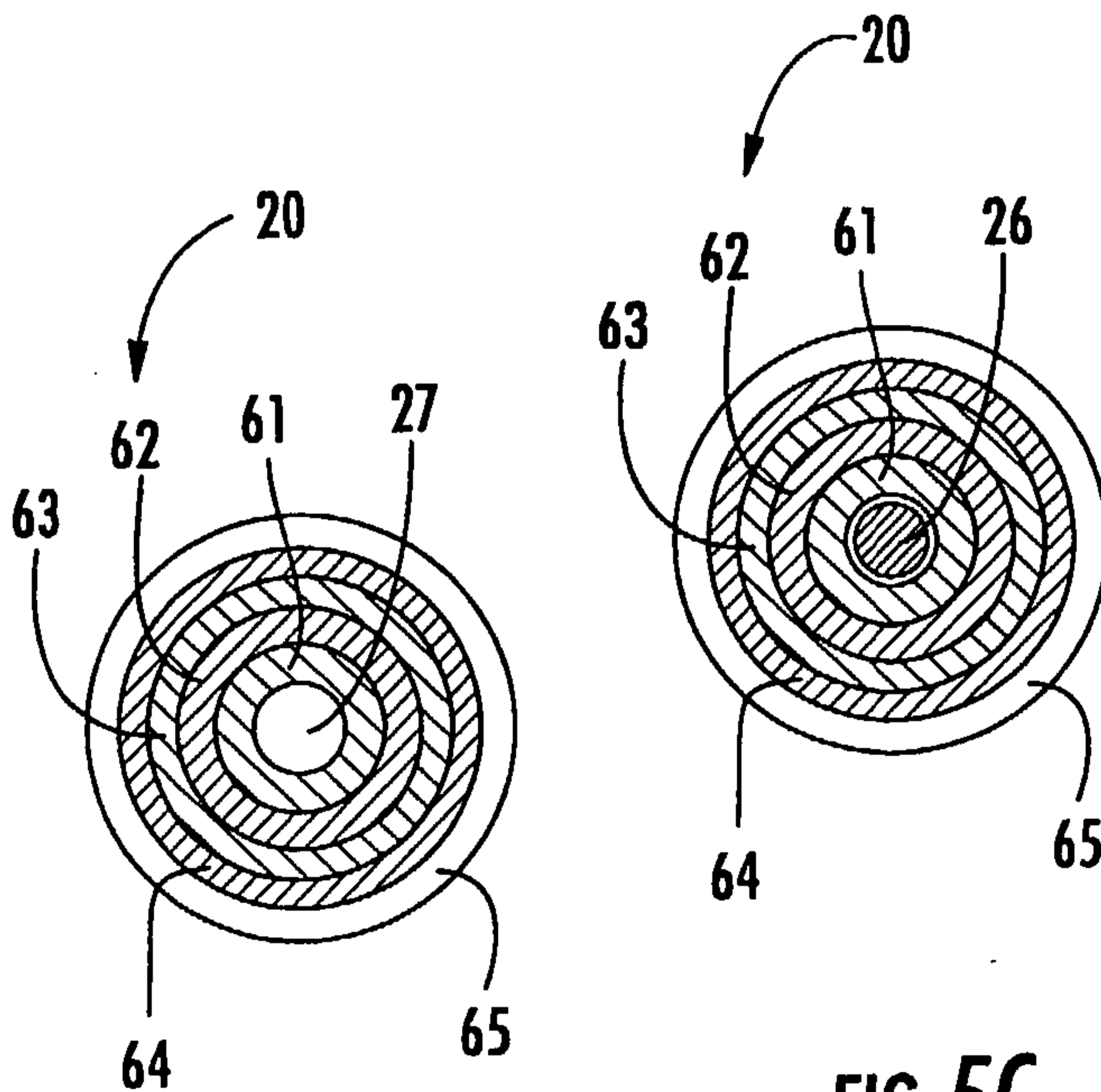


FIG. 5B

FIG. 5C

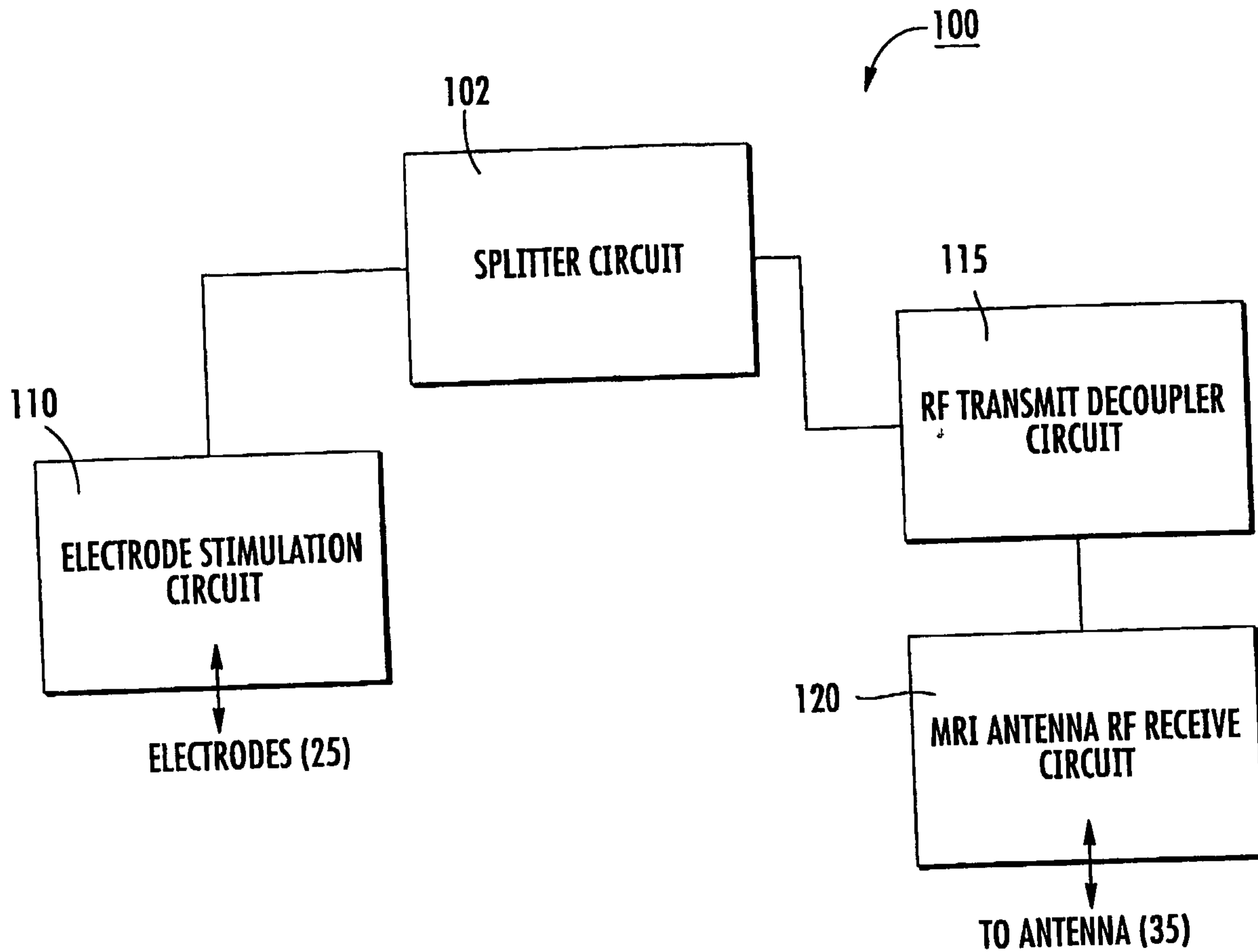


FIG. 6

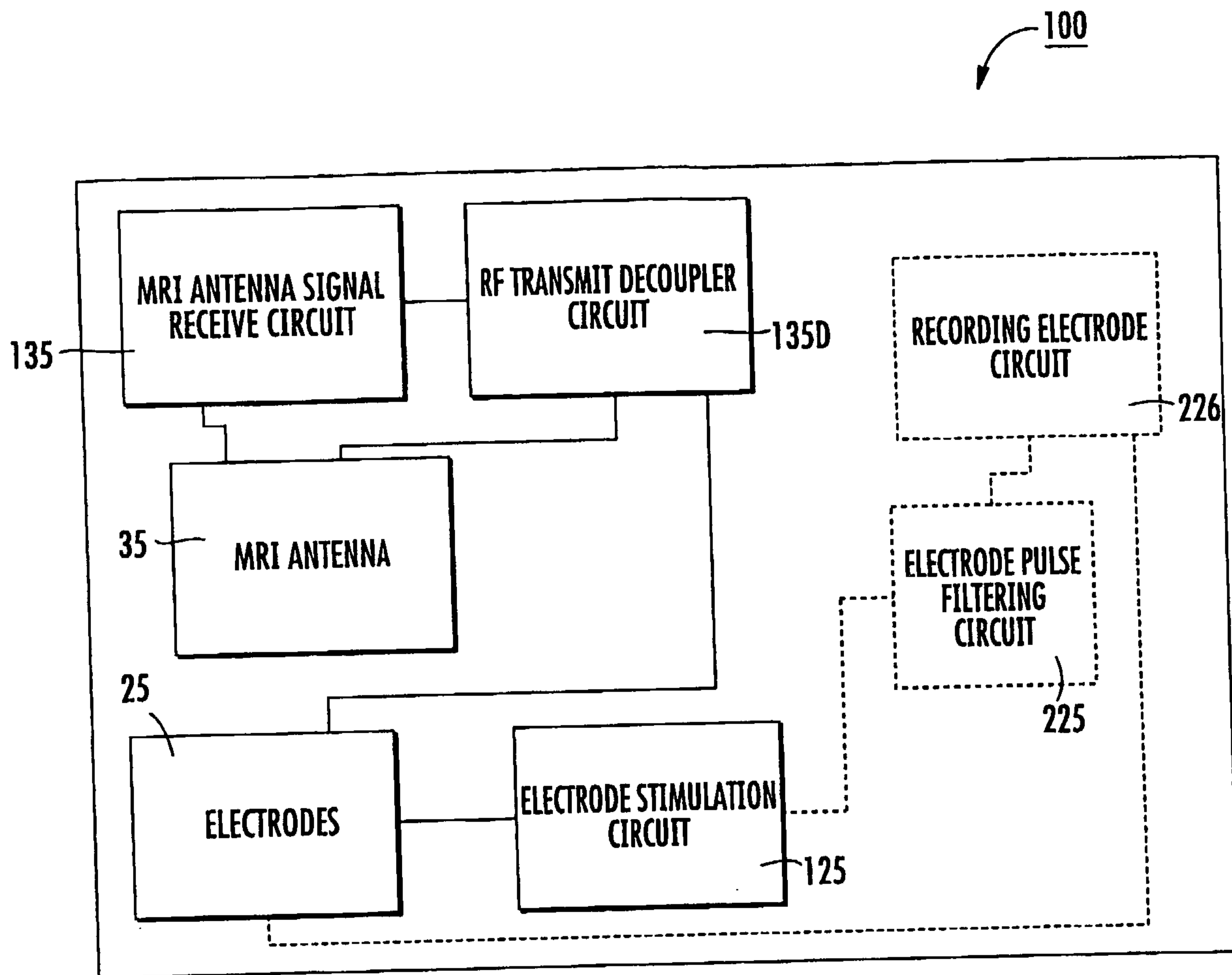


FIG. 7A

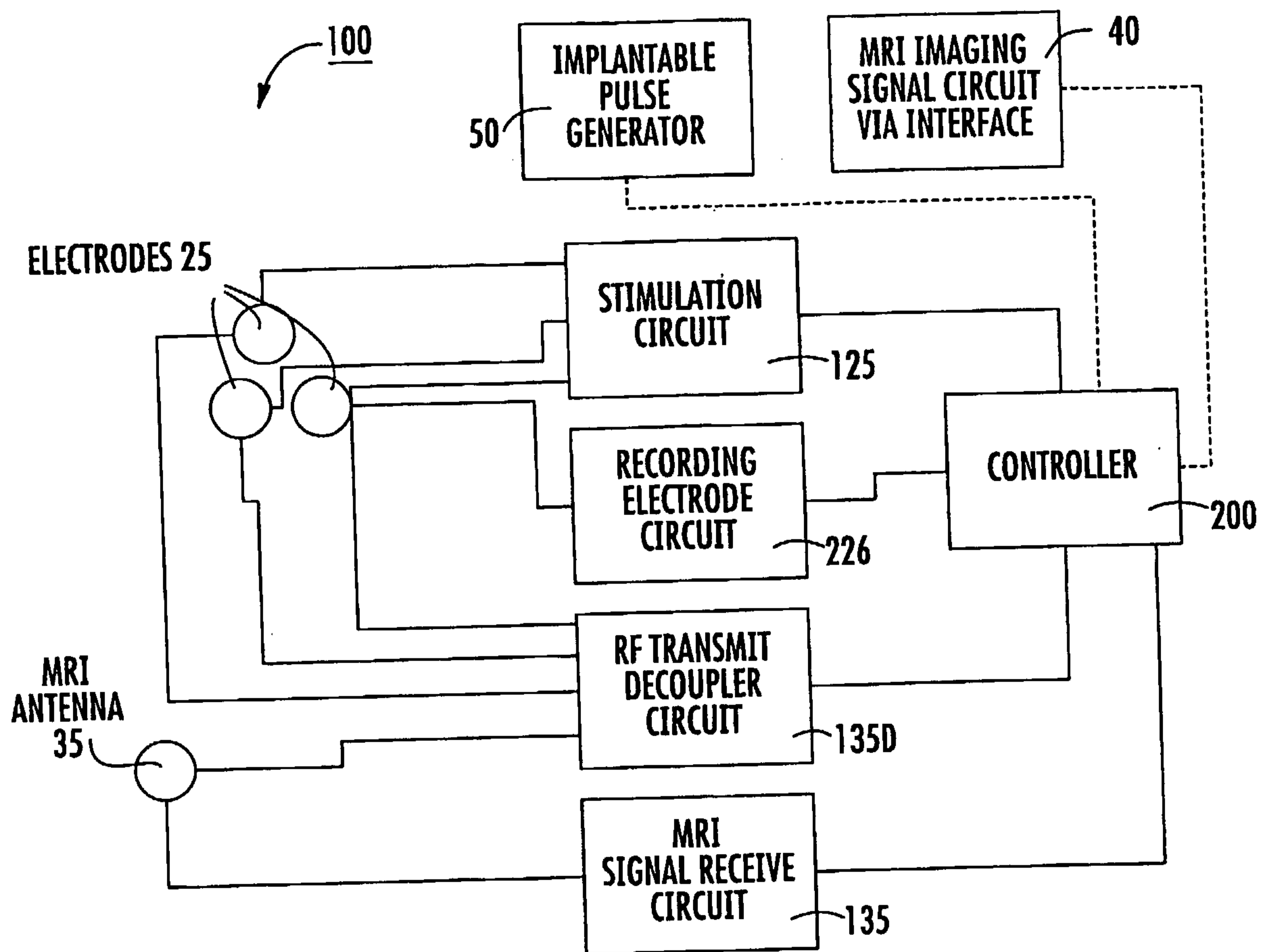
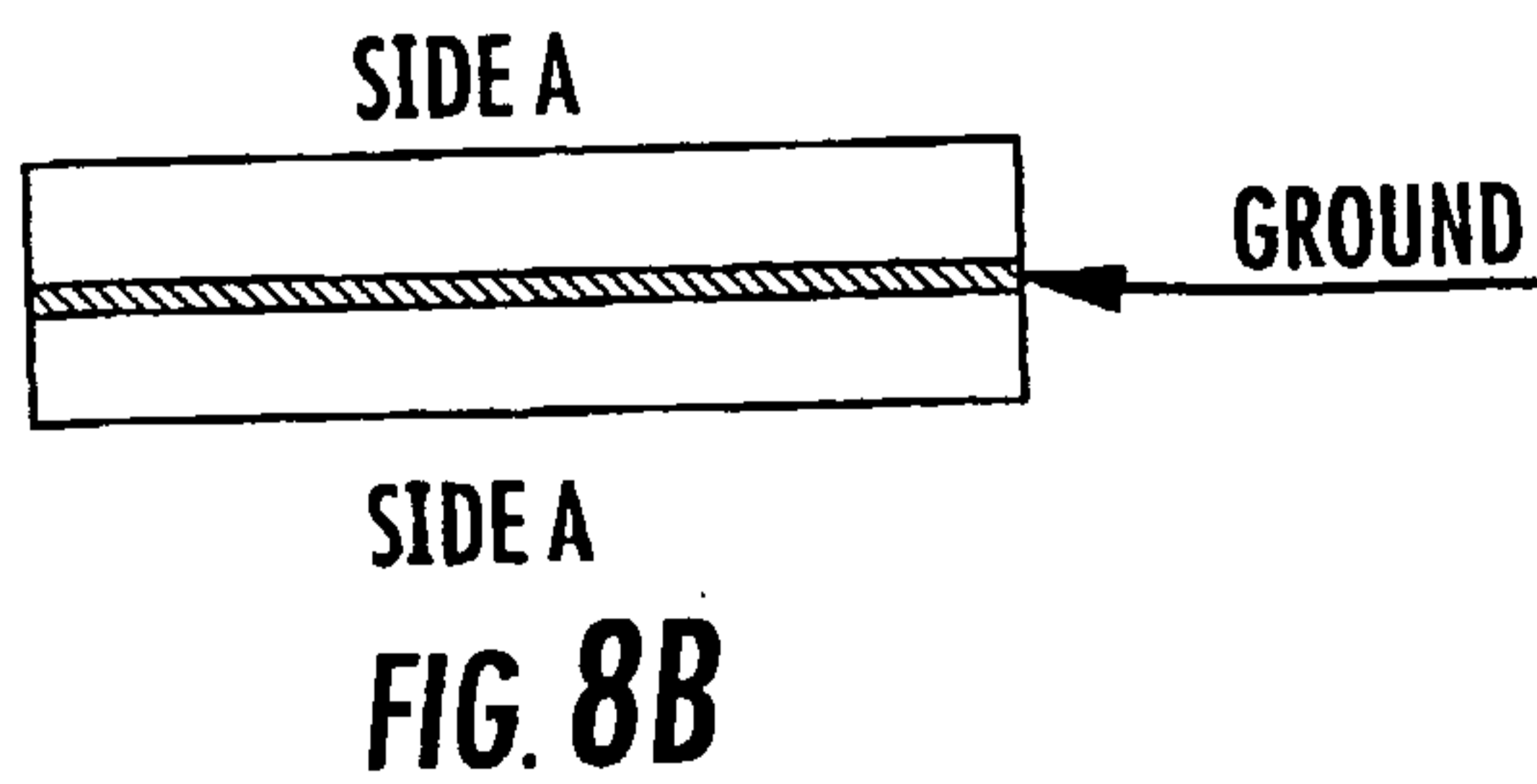
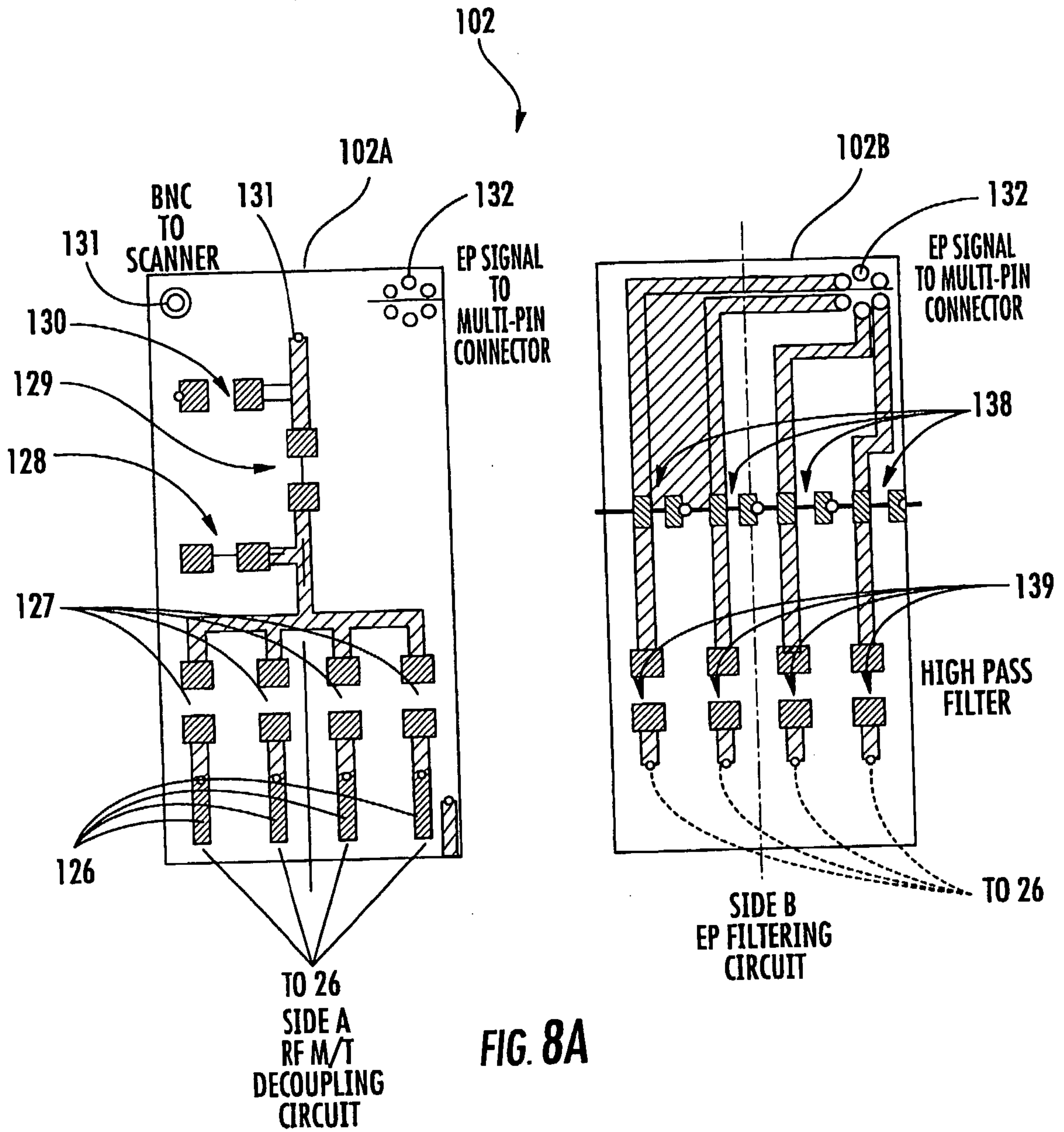


FIG. 7B

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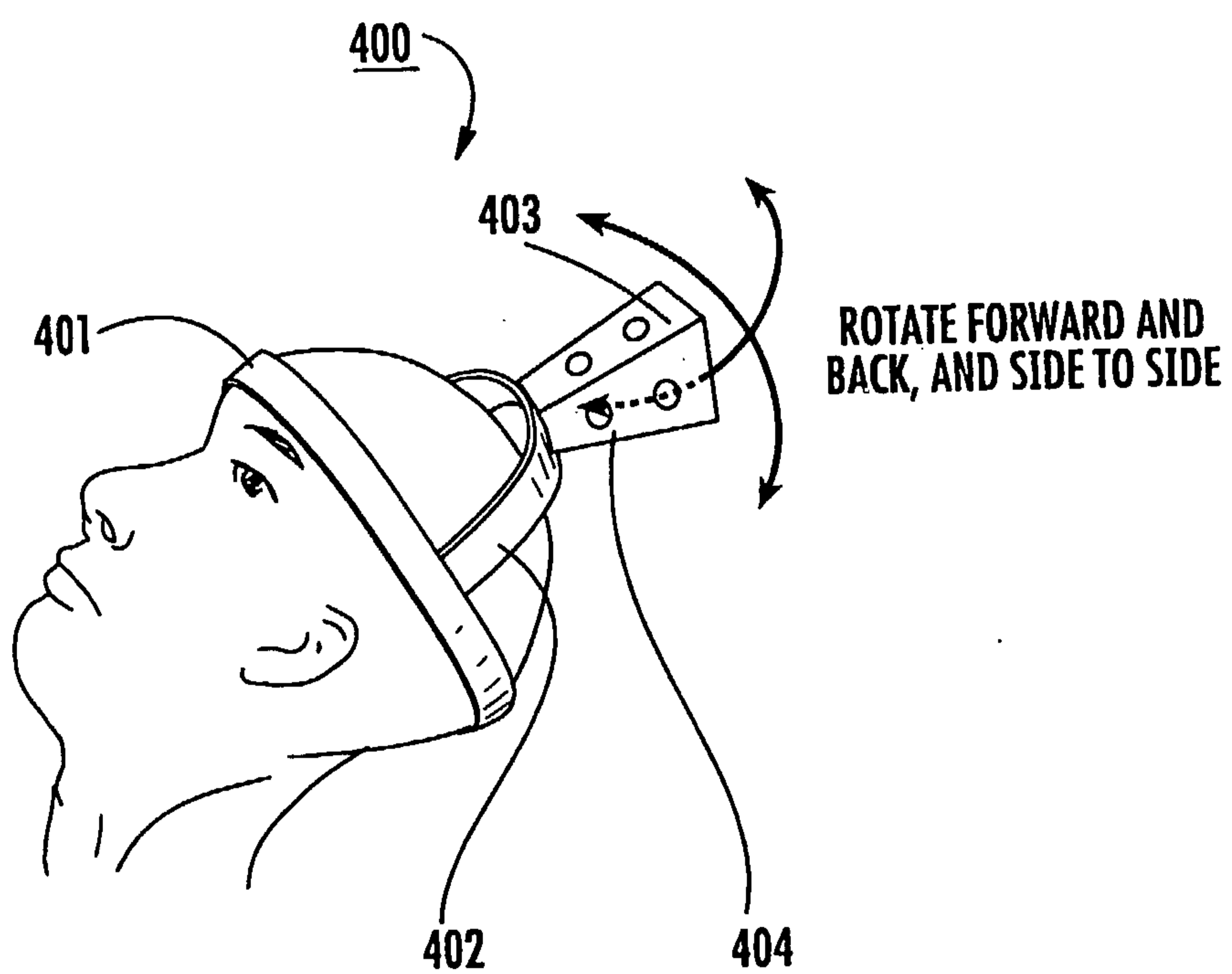


FIG. 9

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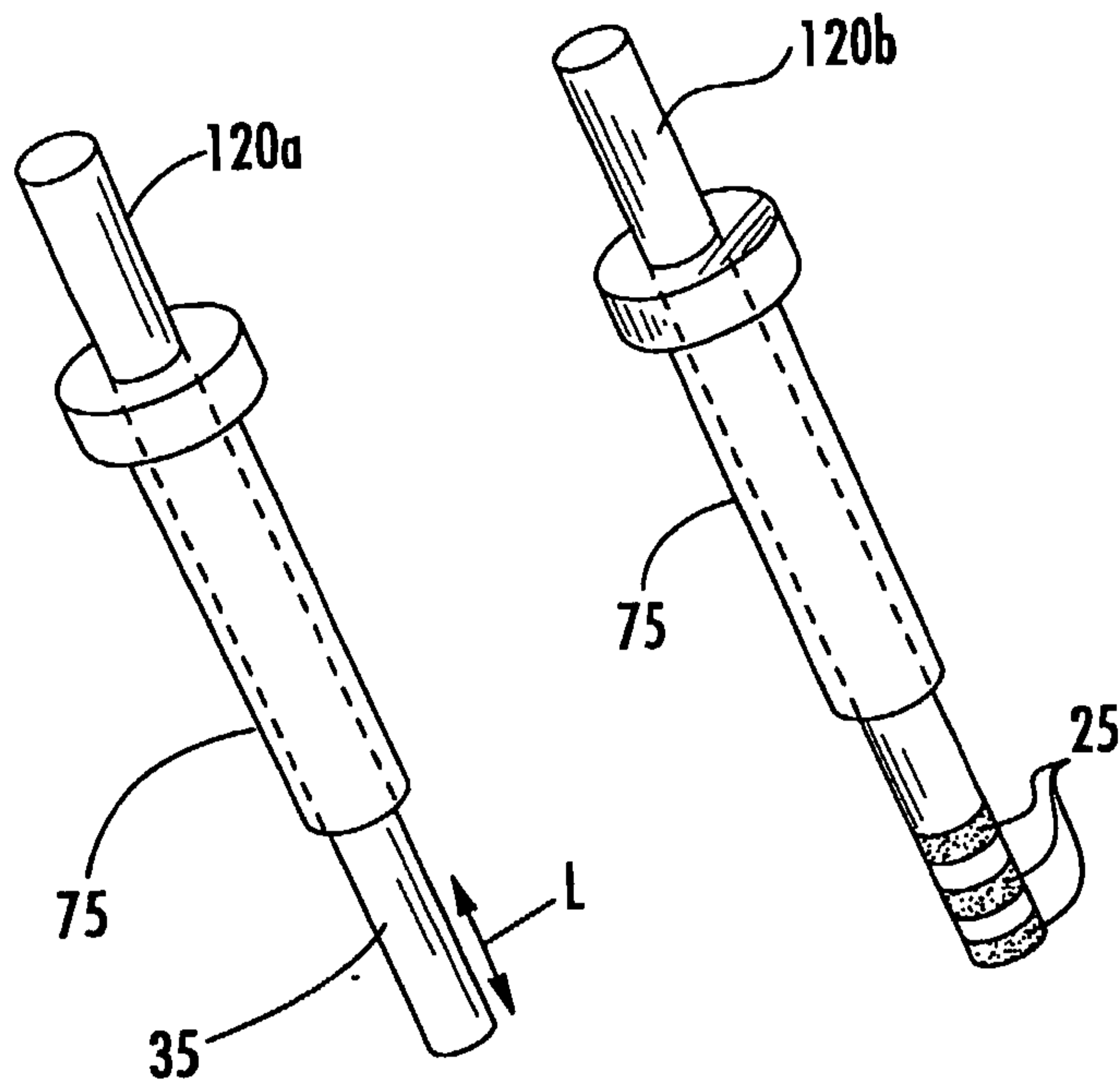


FIG. 10A

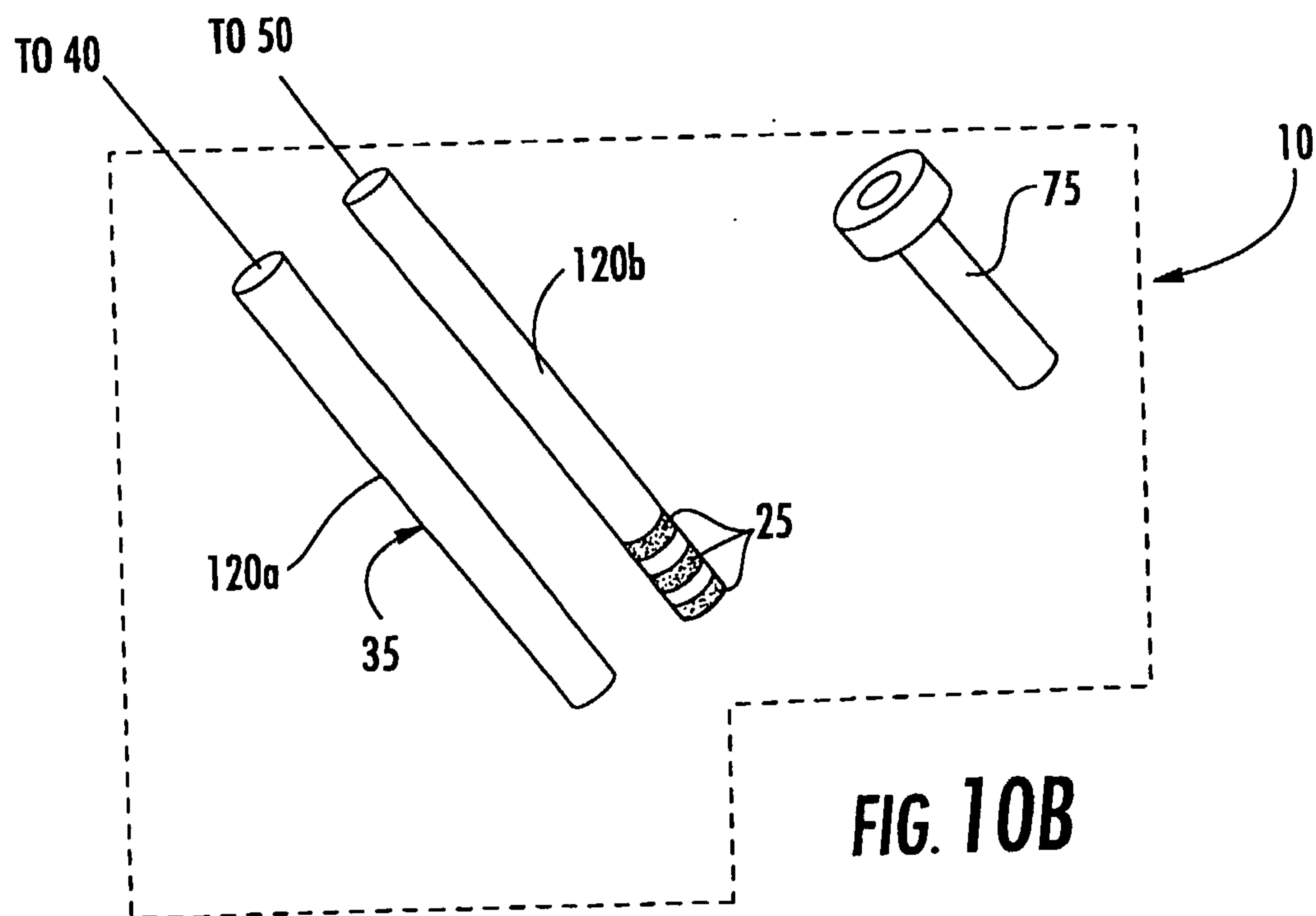


FIG. 10B

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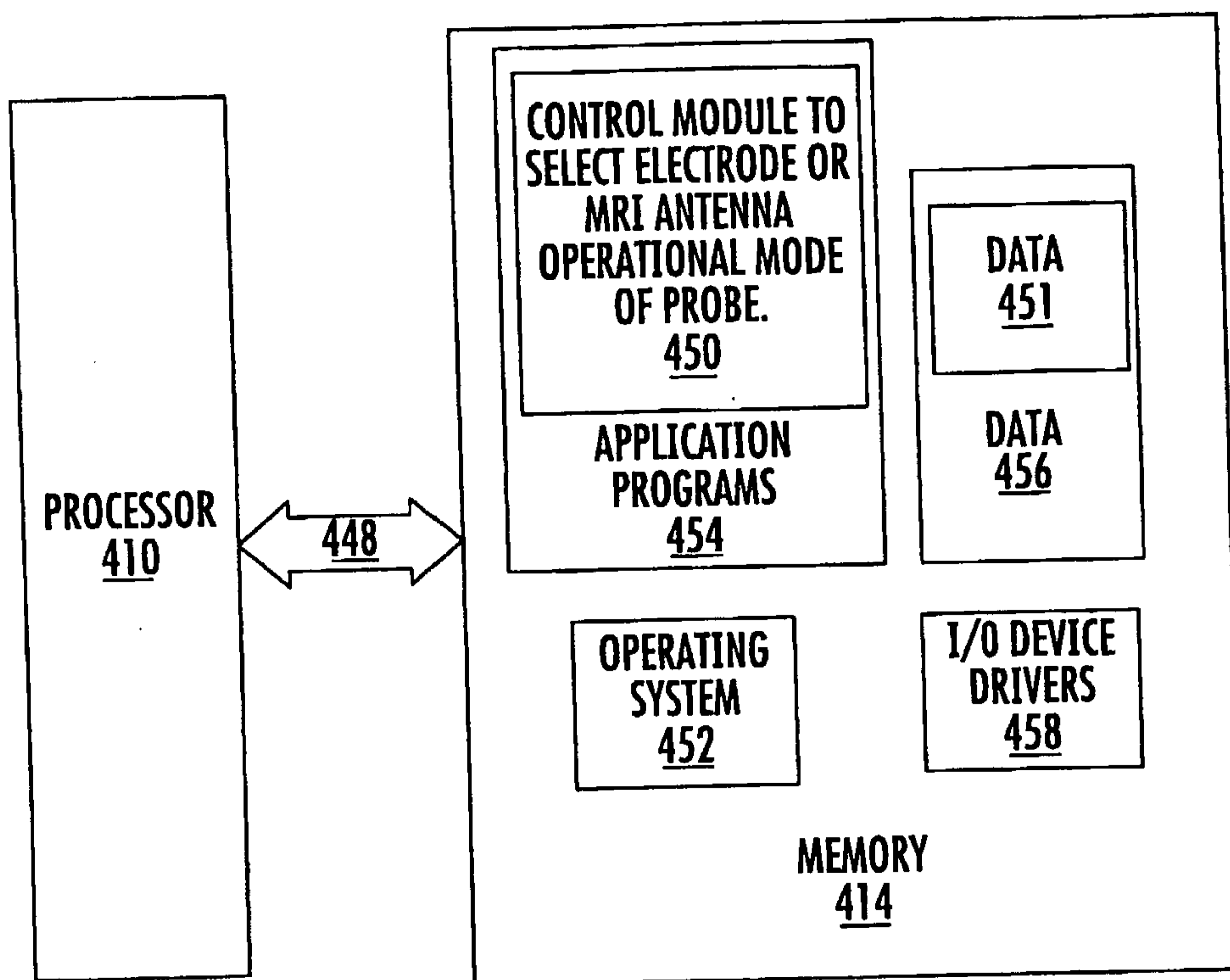


FIG. 11

