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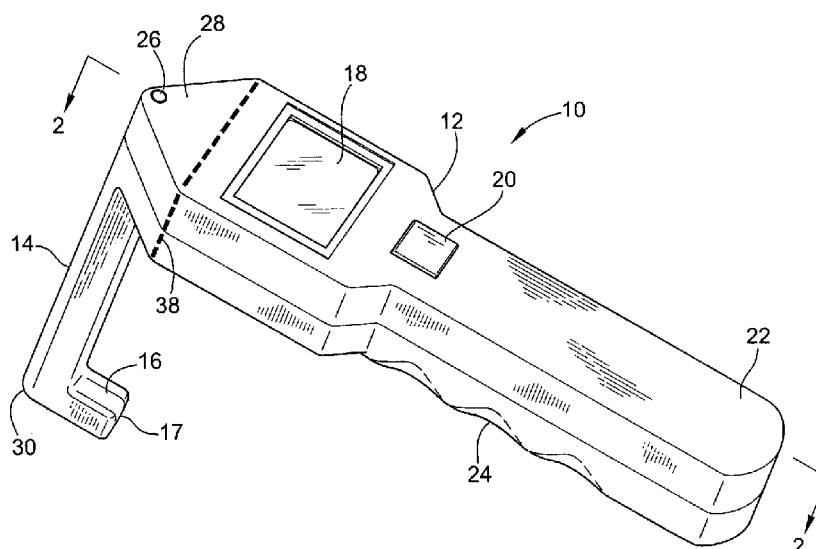


FIG. 1

(57) **Abstract:** An intramedullary nail targeting device (10) for detecting the precise location and position of an opening in an intramedullary nail (60) in a bone (100) or similar object within a body of tissue is described. The device includes a body (12) with a handle end (22) and a drill sleeve end, an activation button (20), a drill sleeve (14) connected to the drill sleeve end and having a first proximal end and a second distal end. The drill sleeve has a length necessary to extend through the depth of the tissue and includes a sensor foot (16) and a drill guide (26). The sensor foot includes a sensor array (34) for sensing the opening in the intramedullary nail when the sensor foot is placed on or near the surface of the bone.

INTRAMEDULLARY NAIL TARGETING DEVICE

Alfred A. Durham

REFERENCE TO RELATED APPLICATIONS

5 This is a continuation-in-part to U.S. Patent Application serial number 10/679,166, in the name of Szakelyhidi, Jr. et al., entitled "Magnetic Targeting Device" filed October 3, 2003 (hereinafter "Szakelyhidi et al.") and claims priority to U.S. Provisional Patent Application serial number 61/190,709, filed September 2, 2008, both of which are incorporated herein by reference in their entirety.

FIELD OF THE INVENTION

10 The present invention is directed to a targeting device in general and specifically relates to an intramedullary nail ("IMN") targeting device and method for positioning locking screws for intramedullary nails.

DESCRIPTION OF THE PRIOR ART

15 The use of magnetic targeting to locate hidden holes or openings in orthopedic hardware has been tried in many forms. However, the distances involved make sensing the magnetic fields difficult. Even the fields of the strongest magnets diminish to that of the earth's magnetic field at distance of about 10 cm.

20 The earliest successful magnetic targeting was accomplished by Durham et al. and was described in a succession of patents covering a mechanical magnetic targeting system using a mechanically balanced cannulated magnet (U.S. Patents Nos. 5,049,151; 5,514,145; 5,703,375; and 6,162,228). Hollstien et al. (U.S. Patent No. 5,411,503) followed with an electrically based system of stacked flux finders connected to a PC display.

25 The magnetic targeting devices to date target a magnet to accurately position a drill bit for insertion in an opening in intramedullary nails. However, these devices operate at the level of the skin, and the magnet may not be strong enough to accurately position the drill bit. As a result, all of these systems are subject to interference and slow response time and are yet to be practical in surgical use. The key component lacking in these systems is a way to target the surface of the bone where the magnetic field strengths are the highest.

SUMMARY OF THE INVENTION

The present invention solves the issue of diminished magnetic strength by placing the magnetic sensors of the magnetic targeting device directly on the bone. This is accomplished by affixing the magnetic sensors directly on the drill bit cannula at the area of the IMN opening. The configuration resembles a foot wherein the system of sensors is in the area of the toe portion.

The present invention is specifically directed to an intramedullary nail targeting device for detecting the precise location and position of an opening in an intramedullary nail in a bone or similar object within a body of tissue having a depth, comprising a body including a handle end, a drill sleeve end, a power source and processing circuits for sensing the correct orientation of the device with respect to the bone; an activation button; and an extended drill sleeve connected to the drill sleeve end and having a first proximal end and a second distal end, wherein the drill sleeve has a length necessary to extend through the depth of the tissue. The drill sleeve includes a sensor foot at the distal end of the drill sleeve, a drill guide extending from the proximal end of the drill sleeve to the distal end, and a sensor array within the sensor foot for sensing the opening in the intramedullary nail when the sensor foot is placed on or near the surface of the bone. The intramedullary nail targeting device further includes display means to determine the correct orientation of the device with respect to the bone when the sensor foot is placed on or near the surface of the bone.

The present invention is also directed to a system for detecting the precise location and position of an opening in an intramedullary nail in a bone or similar object within a body of tissue having a depth. The system comprises an intramedullary nail targeting device, as described in the preceding paragraph, and an intramedullary nail, comprising at least one locking screw opening traversing the intramedullary nail, a magnet in association with the opening wherein the sensor array detects the magnetic flux lines of the magnet.

The present invention is also directed to a method for detecting the location and position of interlocking transverse screw openings within an intramedullary nail for the internal fixation of long bones within a limb, wherein the intramedullary nail includes a longitudinal opening and interlocking screw openings. The steps include placing the intramedullary nail in the marrow of the bone, wherein the intramedullary nail includes a magnet positioned at a reproducible distance from the opening; positioning an intramedullary nail targeting device, described above, near the general location of the

opening, inserting the drill sleeve of the device in the limb such that the sensor foot touches the surface of the bone; activating the device to zero the sensor array; and positioning the device such that the display means determines the correct orientation of the device with respect to the bone for drilling.

5 The primary advantage of the current system is accuracy and the ability to use a magnet of considerably less strength. In addition, because the sensing array is now very close to the magnet, it is much more accurate.

Other advantages of the device of the present invention are as follows:

- 10 • At least parts of the device are disposable and thus not subject to reuse which can cause health issues.
- The device is conveniently handheld and portable.
- The device is not affected by environmental metal or electrical interference.
- The device can be used in any plane.
- The device can self-zero and self-adjust to the strength of the targeting magnet.
- 15 • The device can be used with cannulated or non-cannulated nails.
- The sensor foot is small enough to be used percutaneously.

The objects and advantages of the invention will appear more fully from the following detailed description of the preferred embodiments of the invention made in conjunction with the accompanying drawings.

20

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a perspective view of the IMN targeting device of the present invention.

FIG. 2 is a cross-sectional view of the IMN targeting device of **FIG. 1** taken along lines 2 – 2 of **FIG. 1**.

25 **FIG. 3** is a cross-sectional view of the foot of the IMN targeting device of **FIG. 1** taken along lines 3 – 3 of **FIG. 2**.

FIGS. 4A and **4B** are partial side plan views of the IMN targeting device of **FIG. 1**, illustrating an alternative embodiment of the sensor foot of the present invention.

30 **FIG. 5** is a side plan view of the IMN targeting device illustrating its operation with respect to a long bone.

FIG. 6 is a top view of the intramedullary nail of the present invention.

FIG. 7 is a top plan view of the IMN targeting device of **FIG. 1** with the cover (i.e., upper body portion) removed.

FIG. 8 is a block diagram illustrating the operation of the IMN targeting device of the present invention.

FIG. 9 is a top plan view of the IMN targeting device of **FIG. 1** illustrating the display window.

5 **FIG. 10** is a diagram illustrating the amplitude output of the sensors.

FIG. 11 is a diagram illustrating the flux density of density of the sensors.

DETAILED DESCRIPTION OF THE NEW EMBODIMENT TO THIS INVENTION

Referring now to **FIG. 1**, the present invention is directed to an IMN targeting
10 device **10** which includes a body **12** with a handle portion **22**, a drill sleeve **14**, an activation or on/off button **20**, a sensor foot **16** connected to the distal end of the drill sleeve **14**, a display window **18**, and a drill guide **26** extending through the interior of the drill sleeve **14**.

Advantageously, the IMN targeting device **10** places the sensor foot **16** of the drill sleeve
15 **14** directly on the bone **100**, illustrated in **FIG. 5**, for more accurate reading.

Body 12

The body **12** can be made of a variety of materials known to the medical arts, including plastic and metal as appropriate for durability and reusability of the device **10**. As illustrated in **FIG. 1**, the body **12** is designed to be handheld and comfortable with
20 finger grips **24** in the handle **22**. The body **12** also holds the batteries **32**, the electronic process circuits **86** and the display window **18** as illustrated in **FIGS. 2, 5** and **7**. Typically, the device **10** can operate on two AAA batteries. Alternatively, the battery system can be rechargeable cells or the device **10** could be wired for electrical operation.

The body **12** of the device **10** is amenable to several non-limiting, non-mutually
25 exclusive design variations, each with various advantages. First, the body **12** and sensor-drill sleeve **14** may be provided as separate units and may be separable, for example, at line **38** (see **FIGS. 1** and **2**). Connecting elements are known to the art for joining the drill sleeve **14** to the body **12** in a manner to enable the electrical connection between the two units. The body **12**, which contains the electronic circuitry (such as the comparator
30 circuit module **86**), could be placed in a sterile bag (not illustrated) and would not have to be sterilized prior to use. During use, the plastic bag containing the body **12** could be perforated by the sensor-drill sleeve portion **14** of the device to connect to the electronic circuitry in the body **12** to render the device **10** ready for use. Having the drill sleeve **14** and the body **12** as separate units also allows for different interchangeable sensor-drill

sleeve **14** options for the same body **12**. The advantage of having different sensor-drill sleeve **14** combinations is that they can be used for different applications such as humeral or tibial nail-locking, which might use smaller diameter locking screws. Additionally, shorter drill sleeves **14** would allow more efficient use of the device **10** when deep soft
5 tissues do not have to be avoided. The ability to use different sensor-drill sleeve **14** combinations therefore prevents the necessity of making a different targeting device **10** for each application. Finally, the separate drill sleeve **14** can be made of disposable materials for simple disposal after use.

In a second design variation, the electronics can be gas sterilizable so that the drill
10 sleeve **14** section could be attached to the body **12** at line **38** and used without a sterile bag. All of the advantages above would be realized.

In a third design variation, the electronics could be made to withstand any other form of sterilization: autoclaving, CIDEX® disinfecting solutions (Johnson & Johnson Corporation, New Brunswick, NJ) or similar chemical soaks, gas sterilization or any
15 equivalent.

In a fourth design variation, the targeting device **10** can be connected wirelessly between the sensor foot **16** and the display window **18** to transfer targeting or display information wherever needed. The sensing information could be transmitted by radio, infrared or equivalent from the sensor handle to the display window **18**. The display
20 window **18** may be separate from the body **12** and can comprise any medium, including virtual projections, heads-up glasses, or a personal computer or television screen. Such a display window **18** can be made from any compatible non-magnetic material.

In a fifth design variation, the body **12** may be separable along line **39**, shown in **FIG. 2**, to divide the body **12** into an upper body portion **12A** and a lower body portion
25 **12B**. The upper and lower body portions **12A** and **B**, may be connected by screws **13A** that insert into threaded holes **13B**, the latter of which extend from the lower body portion **12B** into the upper body portion **12A**. Other means of connecting the upper and lower body portions **12A** and **B** may be used. The ability to separate the upper and lower body portions **12A** and **B** allows the user to access internal parts of the device **10**, such as the
30 battery **32** and the comparator circuit **86**.

Window Display 18

While the display window **18** can operate in the manner described with respect to Szakelyhidi et al., the display window **18** is preferably graphical in nature and provides a crosshair **92** in combination with a moving icon or target dot **90**, illustrated in **FIG. 9**, to

indicate the amount of misalignment of the sensor array **34** with respect to the magnet **70** on the IMN **60**. Referring to **FIGS. 5** and **9**, when the targeting dot **90** is centered on the crosshair **92**, the drill guide **26** is centered over the opening **64** or **65** in the IMN **60**, and the bone **100** may be drilled through the drill guide **26**. An advantage of this type of display is that it has sub-millimeter resolution. The determination of the “targeter centered” condition is entirely up to the surgeon, and does not depend on software in the device **10** to provide a “green light, drill here” indication. Such an indication would have to be determined by a software engineer who typically has no medical experience, and this is inappropriate at best. Although the device **10** may include such program logic, the device **10** preferably does not include such program logic because it would replace to some extent the judgment of the surgeon.

Activation or On/Off Button 20

The activation button **20** is provided generally on the top surface of the body **12** at a convenient location for the surgeon to power and calibrate the device **10**. The button **20** is positioned for comfortable use. There may be a button **20** on either side of the handle **22** activating the same function, to allow for left- or right-handed use.

Drill Guide 14 and Sensor Foot 16

The preferred design of the present invention includes a drill sleeve **14** about 10 cm in length. While the length of the drill sleeve **14** is variable, a length of 10 cm incorporates most distal femoral soft tissue sleeves. For tibial and humeral applications, the drill sleeve **14** can be as short as 3-4 cm.

The sensor foot **16** is incorporated as part of the molded drill sleeve **14**. The sensor foot **16** resembles a foot wherein the toe portion **17** contains the system of sensors **34A, B, C, D**, as illustrated in **FIG. 3**. A smaller sized sensor foot **16** on the drill sleeve **14** is more practical to use. Alternatively, the sensor foot **16** could be made separate and incorporated into a metal drill sleeve **14**.

In an alternative version of the sensor foot **16**, as shown in **FIGS. 4A** and **4B**, the sensor foot **16** can be a swivel design wherein it is hingedly attached to the drill sleeve **14** by means of a hinge unit **40**. This configuration eases insertion of the sensor foot **16** into the soft tissues at the point of insertion. The hinge unit **40** can be made of a number of materials and designs to incorporate the swivel functioning of the unit. Prior to insertion into an opening in a limb for positioning next to a bone **100**, the sensor foot **16** is rotated by means of the hinge **40** and pointed in parallel alignment with the drill sleeve **14** for ease of movement toward the bone **100**, as illustrated in **FIG. 4A**. As the toe portion **17**

comes in contact with the bone **100**, the foot **16** will rotate in an arc approximating arrow **42** until the foot portion **16** rests on the bone **100** approximately perpendicular to the drill sleeve **14**, as illustrated in **FIG. 4B**.

Sensor Array **34**

5 As illustrated in **FIG. 3**, the sensor foot **16** is preferably fitted with four magnetic sensors **34A, B, C, D** arranged in a generally rectangular sensor array. The sensor array **34** is connected to the main electronics **86** in the body **12** by printed circuit wiring or wires **36** extending within the drill sleeve **14** beside the drill guide **26** (see **FIG. 2**). It is within the scope of the present invention to use different magnet shapes and materials as long as the sensor array **34** used to target them is adjusted to match the flux field of the magnet. It must also provide the desired flux field for feedback of discriminate targeting in all required planes. An electro-magnet may be used to achieve a similar field if
10 desired. A preferred example of a magnet which may be used in the sensor array **34** is a Honeywell HMC 1052 (Morristown, NJ) magneto resistive magnet. Magneto resistive magnets advantageously have an internal magnetic reset function that can reverse the magnetizing effect of a permanent magnet when brought too close to the sensor array **34**. This feature works well and is used to reset the sensors **34** upon every calibration operation (described below). The sensor reset driver pushes a large current pulse through all sensors at once to perform the reset.

20 Essentially the same sensor array **34** as described with respect to Szakelyhidi et al. is used in the present invention to indicate the correct placement of the drill guide **14** over the IMN **60**. However, rather than requiring eight sensors as suggested in Szakelyhidi et al., it is within the scope of the present invention to employ fewer sensors, e.g., four sensors **34A, B, C, D**. The sensors **34A, B, C, D** are affixed directly on the
25 sensor foot **16** of the drill sleeve **14** at the area of the drill guide **26**.

The sensors **34A, B, C, D** are preferably sized and configured such that, at 10 cm **80**, the sensor array **34** detects one or fewer flux lines **78** at a time, as shown in **FIG. 11**. Only by positioning the sensors **34** very near to the magnet **70** on the surface of the IMN **60** can the flux density be translated into targeting information by detecting multiple flux
30 lines **78** (see reference **82** in **FIG. 11**). Placing the sensor array **34** on the foot **16** allows the sensor array **34** to be at the surface of the bone **100** in close proximity to the magnet **70**. As illustrated in **FIG. 11**, the very high flux density at the surface of the bone **100** (see reference number **82**) ensures that each sensor **34** detects multiple flux lines **78** in spite of the relatively small size of the sensors **34**. This allows for greater resolution in

targeting. As a non-limiting example, the sensors **34A, B, C, D** are preferably 1-2 mm square and are arranged in an array **34** about 5-8 mm across and 2-5 mm thick. The center of the magnetic field can be as little as 6-10 mm from the center axis of the hole to be drilled. Targeting will be more accurate when the distance from the sensor magnet **70** to the center axis of the hole **64** or **66** in the IMN **60** is minimized.

The sensor array **34** may be molded into a plastic drill sleeve **14** with the wires **36** from the sensor **34** ascending the drill sleeve **14** to the comparator circuit **86**, as linked to an LCD window display **18**, as illustrated in **FIGS. 2 and 7**.

Intramedullary Nail 60

Referring to **FIG. 5**, the device is illustrated in association with a long bone **100**, such as a broken femur, tibia or humerus bone. Within the bone **100**, there is illustrated an intramedullary nail (IMN) **60**, known to the art. Examples of IMNs are prevalent in the prior art. For example, reference is made to U. S. Patent 6,503,249 to Krause and the patents to Durham (cited herein), the contents of which are incorporated herein for a description of IMNs and manners of use. The IMN **60** is an elongated metal rod typically having a hollow body portion or shaft **62**, although, as described with respect to the IMN **60** in **FIG. 6**, the IMN **60** may also be a solid body. The IMN **60** typically includes a first locking screw opening **64** and a second, more distal locking screw opening **66**. While the screw openings **64, 66** of typical IMNs **60** are transverse, i.e., positioned at a ninety degree angle in relation to the long axis of the nail **60**, as illustrated in **FIGS. 5 and 6**, it is within the scope of the present invention to have non-transverse or oblique screw openings, i.e., openings at other than ninety degrees in relation to the length of the IMN **60**. It is also within the scope of the present invention to have screw openings **68** placed along the circumferential axis (e.g., 90 degrees) relative to screw openings **64, 66**, as illustrated in **FIG. 6**. Prior to placement of the IMN **60**, a reaming rod known to the art is worked through the medullary cavity **101** of a long bone **100**, such as a broken femur, tibia or humerus bone. The IMN **60** is then placed within the medullary cavity **101** for securing within the bone **100** by means of cross-locking screws or bolts positioned through the screw openings **64, 66**.

Magnet 70

Reference is made to Szakelyhidi et al. for a complete description of the magnetic field and its use with respect to the present invention. All magnets obey the inverse square rule, where the strength of the magnetic field drops off at the square of the

distance. For example, doubling the distance decreases the magnetic field strength by 25%. If the distance is 10 cm, the magnetic field is 0.01 times (1%) the strength and field density it would be at 1 cm from the magnet. Conversely, the strength of the magnetic field at 1 cm from the magnet would be 100 times stronger than the same magnetic field measured at 10 cm. Early magnetic sensors were about 5 mm in size and prevented construction of a foot plate of practical size. New magnetic sensors are as small as 1-2 mm and make practical the construction of a magnetic sensing drill sleeve for placing directly on the bone.

As illustrated in **FIGS. 5** and **6**, the IMN **60** advantageously has the magnet **70** embedded directly on the surface of the IMN **60**. Therefore, the IMN **60**, as illustrated in **FIG. 6** can be a solid IMN. It is also within the scope of the present invention to feed a removable magnet through the center of the IMN **60** as disclosed in Szakelyhidi et al. It is also within the scope of the invention to place a magnetic ring around the periphery of the opening **64, 66** or to place the magnet **70** in the center of the opening **64, 66**, as a displaceable "bull's-eye." The magnet **70** can be alternatively located at the opening **64, 66** and be on a swivel that retracts when the drill enters the opening **64, 66** of the IMN **60**. In thin-wall IMNs, the magnet is centered within the IMN by a circular spring mechanism or equivalent. In thick-wall IMNs, the magnets are small enough to be within the diameter of the guide wire cannulation in the IMN.

20 **Flux Density**

With the magnet **70** residing on or near the surface of the IMN **60**, there is a close positioning of the sensor array **34** and the magnet **70** as shown in **FIG. 5**. This permits exposure of the sensor array **34** to greater flux density (see **FIG. 11**). All magnets, such as magnet **70**, obey the inverse square rule, i.e., double the distance and the magnetic field is one-fourth the strength. As described above, if the distance between the sensor array **34** and the magnet **70** is 10 cm, the magnetic field is 1% the strength and field density of a sensor array **34** 1 cm from the magnet **70**. A preferred distance between sensor array **34** and the magnet **70** in the present invention is a distance of about 1.5 cm, typically the average thickness of the side of the bone **100**. At that distance, the field density is about 30 times the density at a distance of 10 cm.

As illustrated in **FIG. 11**, a magnetic field sensed at 10 cm (reference number **80**) has spread so thin that the flux lines **78** are too far apart to accurately locate the center of a 5 mm hole. At about 1.5 cm from the magnet (reference number **82**), the flux density is

high enough to detect sufficient positional information for accurate targeting of the device **10**.

To date the most difficult distal targeting goal has been the distal femur. The working distances from the center line of the IMN **60** is typically no more than 3 cm at the surface of the distal femur and is usually 1-2 cm. This makes targeting nearly any other bone: tibia, humerus, or any other long bone even easier because of smaller cortex to nail distances.

Internal Operation of Device 10

Reference is now made to **FIGS. 7 and 8** for a description of the internal operation of the device **10**. In action, the microcontroller **102** powers a single sensor **34A, B, C, or D** in turn, using the switch **103** to connect it to the high gain amplifier **104**. The microcontroller **102** then sets the digital voltage generator **106** to a predetermined value. The microcontroller **102** waits for the sensor **34A, B, C, or D** and amplifier **104** to settle and then reads the voltage from the amplifier **104**. This voltage is proportional to the applied magnetic field but also contains some environmentally generated noise and noise which is inherent in the sensors **34A, B, C, or D**. The microcontroller **102** selects the four sensors **34A, B, C, or D** in sequence, measuring their outputs and saving them for targeting computations. A complete set of measurements is made typically 20 to 50 times per second. As with any high gain sensor system, small errors can be multiplied by factors of 1000 or more, resulting in huge problems making the required measurements. The sensors **34A, B, C, or D** are no different and have offset errors in their outputs that make measurements difficult without some adjustment. The amplifier **104** introduces errors as well. The digital voltage generator **106** is used during the calibration process to null out these errors.

When the device **10** is powered on by the activation button **20**, the device **10** immediately begins a calibration sequence. This involves selecting each sensor **34A, B, C, and D** in turn and determining the value from the digital voltage generator **106** that is required to bring the amplifier **104** into its linear amplifying region of operation. This operation takes only a couple seconds. Thereafter, as each sensor **34A, B, C, and D** is selected, the digital voltage generator **106** is loaded with the particular value for that sensor **34A, B, C, or D**, resulting in nullification of static errors for that sensor's measurement. The circuit also features a two-step amplifier gain selection, though the software may use only the high gain setting. Such a system allows use of the device **10** for various thicknesses of human bone **100** without software changes. This design uses

one amplifier **104** and an inexpensive commodity solid state switch **103** to select which sensor **34A, B, C, or D** to read. Another feature not shown is that the microcontroller **102** does not leave all sensors **34A, B, C, or D** powered continuously, but rather turns them on in sequence, saving power consumption.

5 The microcontroller **102** uses a vector algorithm to determine how to position the target icon **90** on the window display **18**. The position of each sensor **34A, B, C, or D** is assigned a vector direction depending on its position in the array **38**. The amplitude of the output of each sensor **34A, B, C, or D** provides the magnitude of each vector. Addition of the magnitudes of the vectors provide a resultant vector that determines the
10 position of the device **10** relative to the magnet, which is represented as a two-dimensional position of a targeting dot **90** on the window display **18** (see **FIG. 9**). **FIG. 10**, for example, shows a center box representing the magnet **70** and four other boxes representing the magnetic sensors **34A, B, C, or D**. The vector lines **35A, B, C, and D** attached to each sensor **34A, B, C, and D**, respectively, indicate the strength of the field at
15 each sensor. Vector line **71** is the resultant vector, which indicates the direction the sensor array **34** should be moved to center it over the magnet **70**. The magnet **70** in **FIG. 10** corresponds with the targeting dot **90** in **FIG. 9**.

Referring back to **FIG. 8**, the thermal cutoff **108** is present in case the device **10** is accidentally run through a sterilizer cycle. The thermal cutoff **108** activates at 82° Celsius
20 and disables operation of the device **10** permanently. Without the thermal cutoff **108**, it is likely that the device **10** would work somewhat after being exposed to such heat, but reliable operation could not be guaranteed. A low battery indicator is implemented that warns the user of low batteries **32** on the window display **18** and also prevents the device **10** from operating.

25 **User Operation**

The single activation button **20** is used to turn the device **10** on, and the device **10** immediately performs a calibration cycle. If the button **20** is pressed briefly thereafter, another calibration cycle is initiated. The window display **18** indicates to the user that calibration is in progress. It is not possible to turn the device **10** on without initiating a
30 calibration cycle. To turn the device **10** off, the button **20** is held down for a couple seconds until the display goes off. The device **10** also powers off after two minutes to prevent the batteries from draining.

To perform targeting, the device **10** is held in the same orientation as it will be used. The device **10** is raised 10-12 inches above the targeting magnet **70** and the button

20 is pressed to start a calibration cycle. It is important that the device **10** be oriented approximately as it will be used in order to properly null the magnetic field of the earth. Once the device **10** completes its calibration operation, it is lowered to the work area and moved to achieve an on-target indication.

5 **Performance**

If the device **10** has some difficulty detecting the magnetic field of the targeting magnet **70**, the display **18** will show an error indication. There could be reasons for the error indication:

- The magnetic field of the earth: This field is about 0.5 gauss. The device **10** must
10 be used with a targeting magnet **70** and at a distance such that the earth's magnetic field is not a significant factor.
- Alternating current (AC) power and radio frequency (RF) noise sources: These are minimized through device shielding.
- Sensor noise: The sensors **34A, B, C, D** produce electronic noise as a byproduct
15 of their operation. This noise is added to the voltage output of each sensor caused by the magnetic field. The device **10** must be used with a targeting magnet **70** and at a distance such that the sensor noise is not a significant factor.
- Proximate ferromagnetic objects: The earth's magnetic field bends where it
20 approaches magnetically attractive objects, such as chairs, metal tables, rebar in concrete, etc. However, if the earth's magnetic field is not an issue, then proximate ferromagnetic objects will not be an issue, either.

Placing the sensor array **36** proximal to the magnet **70** (e.g., against the bone), as occurs with the present device **10**, successfully avoids the above problems by ensuring a strong magnetic field at the sensors when the device is in use.

25 **Operation of Device 10**

In order to locate the general location of the openings **64** and **66** in the IMN **60**, the IMN **60** is placed in the marrow of the bone **100** and urged through the bone **100** as described in Szakelyhidi et al. The openings **64, 66** in the IMN **60** to be targeted has a magnet **70** placed at a reproducible distance from the openings **64, 66** with the magnetic
30 field oriented to the magnetic sensor array **34** in the foot **16** of the IMN targeting device **10**. A handle wand extension, known in the art, which is the same length as the IMN **60** and attached to the IMN **60**, is urged over the exterior of the limb along the same direction as the IMN **60**.

When the IMN **60** is fully positioned in the bone **100**, the end of the handle wand extension indicates both the end of the IMN **60** and the approximate location of the openings **64, 66** in the IMN **60** in the bone **100**. The magnet **70**, situated on the surface of the IMN **60**, as illustrated in **FIG. 6**, corresponds to the distance of the openings **64, 66**.

5 Alternatively, a magnet can be placed in the hollow shaft of the IMN **60**, carried down the cannulation of the IMN **60** by a handle wand, and locked at a corresponding distance of the opening **64** or **66** to be targeted or placed into the wall of the IMN **60**.

An incision is made in the limb. An oval trochar can be used to make a path for the drill sleeve **14** down to the surface of the bone **100**. Once placed on the surface of the bone **100**, the display window **18** is activated by the action of the on/off button **20**. A signal is sent to the sensor array **34** to zero the sensors **34A, B, C, D**. When the sensor array **34** is moved across the surface of the bone **100**, the sensor information appears on the display window **18**, generally in the form of a targeting dot **90** on a targeting grid **92** as illustrated in **FIG. 9**. The placement of the targeting dot **90** in the center of the targeting grid **92** indicates correct placement of the device **10** for drilling. In addition to moving the targeting dot **90** centrally with respect to the targeting grid **92**, more accurate information could be attained in the form of enlarging the dot **90** in response to the strength of the magnetic field being sensed. **FIG. 9** illustrates the device **10** wherein the targeting dot **90** is misaligned with respect to the targeting grid **92**, indicating that the drill sleeve **14** placement is incorrect with respect to the IMN **60** within the bone **100**. Once the targeting dot **90** is centered on the targeting grid **92**, this "maximum size" ensures that the targeting device **10** has not been sensing a symmetrical set of field lines around the magnet **70** or a flux pattern created between two or more magnets **70** embedded into the side of a solid IMN **60**. The same information could be displayed in any equivalent fashion such as a variable LED, audio output, color change or similar signaling device.

Typically, the drill bit **96**, illustrated in **FIG. 5**, used in distal targeting drills the minor diameter of the screw to be inserted in the IMN **60**. This gives more room for the targeting to be accurate than if the major diameter were to be drilled first. A star point drill prevents the drill from "walking" on the slippery curved surface of the bone and is therefore preferred.

The drill bit **96** is then inserted into the drill guide **26** at the upper drill sleeve opening **28** while this information is obtained. The lower opening **30** of the drill guide **26** is placed directly on the bone **100**. This is accomplished by affixing the sensors **34**

directly on the drill guide **26** at the area of the opening **64** or **66**. The drill sleeve **14** is inserted into the bone **100** at the location of the opening **64** or **66**.

The sensor array **34** is activated to locate the magnet **70**, which then determines the location of the opening **64** or **66**. The information sent to the comparator circuit **86** is processed and displayed on an LCD screen **106** that moves a target dot to the center of a cross hair that represents the center of the magnetic field of the targeting magnet.

The drill **96** with a star point is located adjacent to the target magnet **70** and is parallel to the IMN **60** when the magnetic sensor **34** is balanced over the magnetic field. Because the sensor array **34** is proximal to the openings **64** or **66**, when the field is balanced, the drill **96** is free to pass through the opening **64** or **66** in the IMN **60** to the opposite cortex. As soon as the target dot **90** aligns at the center of the targeting grid **92**, the drill **96** is drilled through the opening **64** or **66** to the opposite cortex. The process could be repeated for any additional holes to be targeted.

To target openings **68** in the IMN **60** as shown in **FIG. 6**, the magnetic wand can be rotated **90** degrees. If the magnets **70** are implanted in the side wall of the IMN **60**, they are available for targeting even after locking the IMN **60** proximally.

An aiming device is always more accurate if it has two references in space to align it. The first reference to assist the accuracy of the device **10** comes from determining an entry point on the skin directly over the opening to be targeted in the IMN **60**. The easiest way to determine this point is with a wand that extends from the handle that holds the IMN **60** for insertion. The wand reproduces the curvature of the IMN **60** and has markings corresponding to the length of each IMN **60**. The wand shows the correct entry point over each opening so that when the drill sleeve **14** is inserted at that point, the soft tissues help to stabilize the device perpendicular to the axis of the IMN **60**. The importance of being able to rest the device **10** on the surface of the bone **100** during use cannot be over emphasized. The accuracy needed is on the order of 1 mm. A device **10** held in space cannot be as accurate while simultaneously using a drill.

In most applications it is advantageous to insert the screw through the lumen in the cannula after the opening in the IMN **60** has been magnetically targeted. The device **10** would be used in the standard fashion to drill the minor diameter of the locking screw. A calibration on the drill measures the depth of the drilled hole at the upper drill sleeve **14** opening **28** of the device **10**. Alternatively, after drill removal, the device **10** can remain against the bone **100**. When the drill guide **26** is removed, a depth gauge is used to measure the length of the screw to be inserted. Once measured, the screw of the

appropriate length is loaded onto a screw driver and inserted across the opening **64, 66** of the IMN **60**. Self tapping screws are used in the preferred embodiment.

It is understood that the invention is not confined to the particular construction and arrangement of parts herein illustrated and described, but embraces such modified
5 forms thereof as come within the scope of the following claims.

I Claim:

1. An intramedullary nail targeting device for detecting the precise location and position of an opening in an intramedullary nail in a bone or similar object within a body of tissue having a depth, comprising:
 - a. a body comprising a handle end, a drill sleeve end, a power source and processing circuits for sensing the correct orientation of the device with respect to the bone;
 - b. an activation button;
 - c. an extended drill sleeve connected to the drill sleeve end and having a first proximal end and a second distal end, wherein the drill sleeve has a length necessary to extend through the depth of the tissue, the drill sleeve including:
 - i) a sensor foot at the distal end of the drill sleeve, and
 - ii) a drill guide extending from the proximal end of the drill sleeve to the distal end, and
 - iii) a sensor array within the sensor foot for sensing the opening in the intramedullary nail when the sensor foot is placed on or near the surface of the bone; and
 - d. display means to determine the correct orientation of the device with respect to the bone when the sensor foot is placed on or near the surface of the bone.
2. The device of claim 1 wherein the display means is an LCD display window.
3. The device of claim 1 wherein the display means is a remote projector.
4. The device of claim 1 wherein the power source is selected from the group consisting of batteries, rechargeable cells, or an electrical source.
5. The device of claim 1 wherein the body and the drill sleeve are one unit.
6. The device of claim 1 wherein the drill sleeve is releasably connected to the body.
7. The device of claim 6 wherein the drill sleeve is made of disposable material.
8. The device of claim 1 wherein the length of the drill sleeve corresponds with the depth of the tissue.
9. The device of claim 1 wherein the sensor array is connected to the processing circuits by wires extending the length of the drill sleeve.
10. The device of claim 1 wherein the sensor array is wirelessly connected to the processing circuits in the body of the device.
11. The device of claim 1 wherein the drill sleeve has a length of between about 3 cm and 12 cm.

12. The device of claim 1 wherein the drill sleeve has a length of about 10 cm.
13. The device of claim 1 wherein the sensor foot is fixedly attached to the distal end of the drill sleeve.
14. The device of claim 1 wherein the sensor foot is hingedly attached to the distal end of the drill sleeve.
15. The device of claim 1 wherein the sensor array comprises four magnetic sensors arranged in a generally rectangular array.
16. The device of claim 15 wherein the magnetic sensors are selected from the group consisting of electro-magnets and magneto resistive magnets.
17. The device of claim 15 wherein the magnetic sensors are magneto resistive magnets.
18. The device of claim 15 wherein the sensors are approximately 1-2 mm square.
19. The device of claim 1 wherein the generally rectangular array of magnetic sensors intersects at a center and wherein a center of the rectangular array is between about 6 and about 10 mm from the center axis of the opening in the intramedullary nail.
20. A system for detecting the precise location and position of an opening in an intramedullary nail in a bone or similar object within a body of tissue having a depth, comprising:
- a. an intramedullary nail targeting device, comprising
 - i) a body comprising a handle end, a drill sleeve end, a power source and processing circuits for sensing the correct orientation of the device with respect to the bone;
 - ii) an activation button,
 - iii) an extended drill sleeve connected to the drill sleeve end and having a first proximal end and a second distal end, wherein the drill sleeve has a length necessary to extend through the depth of the tissue, the drill sleeve including:
 - (1) a sensor foot at the distal end of the drill sleeve,
 - (2) a drill guide extending from the proximal end of the drill sleeve to the distal end, and
 - (3) a sensor array within the sensor foot for sensing the opening in the intramedullary nail when the sensor foot is placed on or near the surface of the bone; and
 - iv) display means to determine the correct orientation of the device with respect to the bone when the sensor foot is placed on or near the surface of the bone; and
 - b. an intramedullary nail, comprising:

- i) at least one locking screw opening traversing the intramedullary nail,
 - ii) a magnet in association with the opening wherein the sensor array detects the magnetic flux lines of the magnet.
21. The system of claim 20 wherein the magnet is embedded on the surface of the intramedullary nail.
22. The system of claim 20 wherein the magnet includes the following characteristics: a) a magnetic field of sufficient strength to match the sensor array of the device, and b) a magnetic field of sufficient shape to provide the desired feedback necessary to discriminate targeting in all required planes.
23. The system of claim 20 wherein the sensor array is positioned a distance of about 1.5 cm from the magnet to detect the magnetic flux lines of the magnet.
24. The system of claim 20 wherein the drill sleeve is releasably connected to the body.
25. The device of claim 20 wherein the drill sleeve has a length of about 10 cm.
26. A method for detecting the location and position of interlocking transverse screw openings within an intramedullary nail for the internal fixation of long bones within a limb, wherein the intramedullary nail includes a longitudinal opening and interlocking screw openings, comprising:
- a. Placing the intramedullary nail in the marrow of the bone, wherein the intramedullary nail includes a magnet positioned at a reproducible distance from the opening;
 - b. Positioning an intramedullary nail targeting device near the general location of the opening, wherein the intramedullary nail targeting device comprises:
 - i) a body comprising a handle end, a drill sleeve end, a power source and processing circuits for sensing the correct orientation of the device with respect to the bone;
 - ii) an activation button,
 - iii) a extended drill sleeve connected to the drill sleeve end and having a first proximal end and a second distal end, wherein the drill sleeve has a length necessary to extend through the depth of the tissue, the drill sleeve including:
 - (1) a sensor foot at the distal end of the drill sleeve,
 - (2) a drill guide extending from the proximal end of the drill sleeve to the distal end, and

- (3) a sensor array within the sensor foot for sensing the opening in the intramedullary nail when the sensor foot is placed on or near the surface of the bone;
- iv) display means to determine the correct orientation of the device with respect to the bone when the sensor foot is placed on or near the surface of the bone;
- 5 the bone when the sensor foot is placed on or near the surface of the bone;
- c. inserting the drill sleeve of the device in the limb such that the sensor foot touches the surface of the bone;
- d. activating the device to zero the sensor array; and
- e. positioning the device such that the display means determines the correct
- 10 orientation of the device with respect to the bone for drilling.

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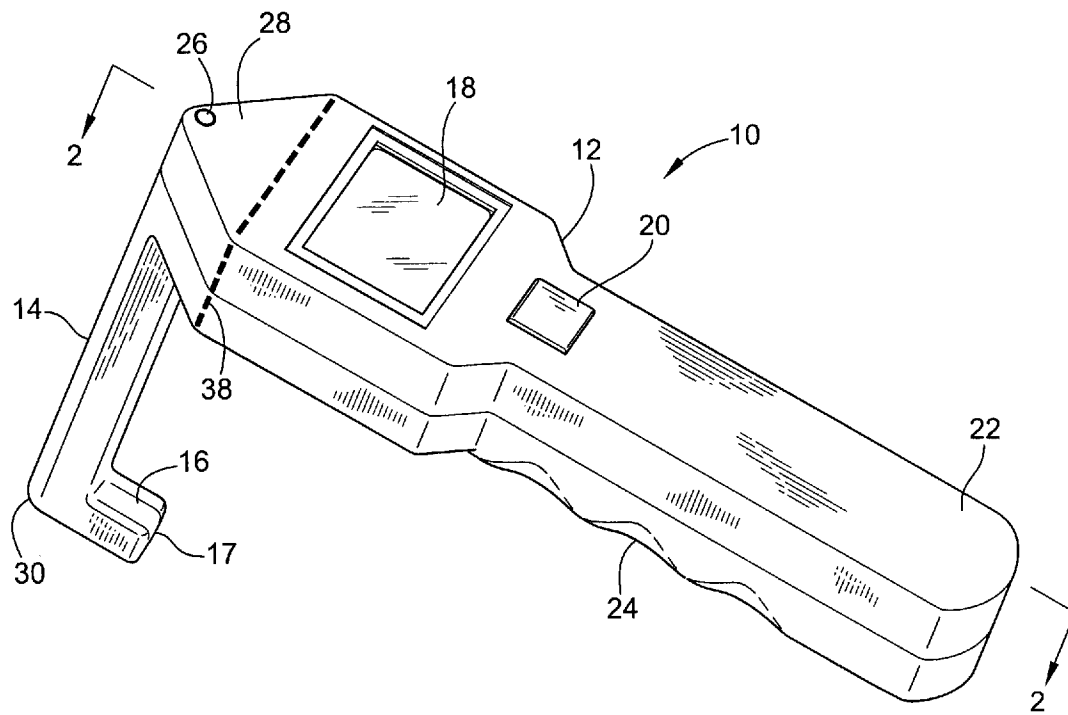


FIG. 1

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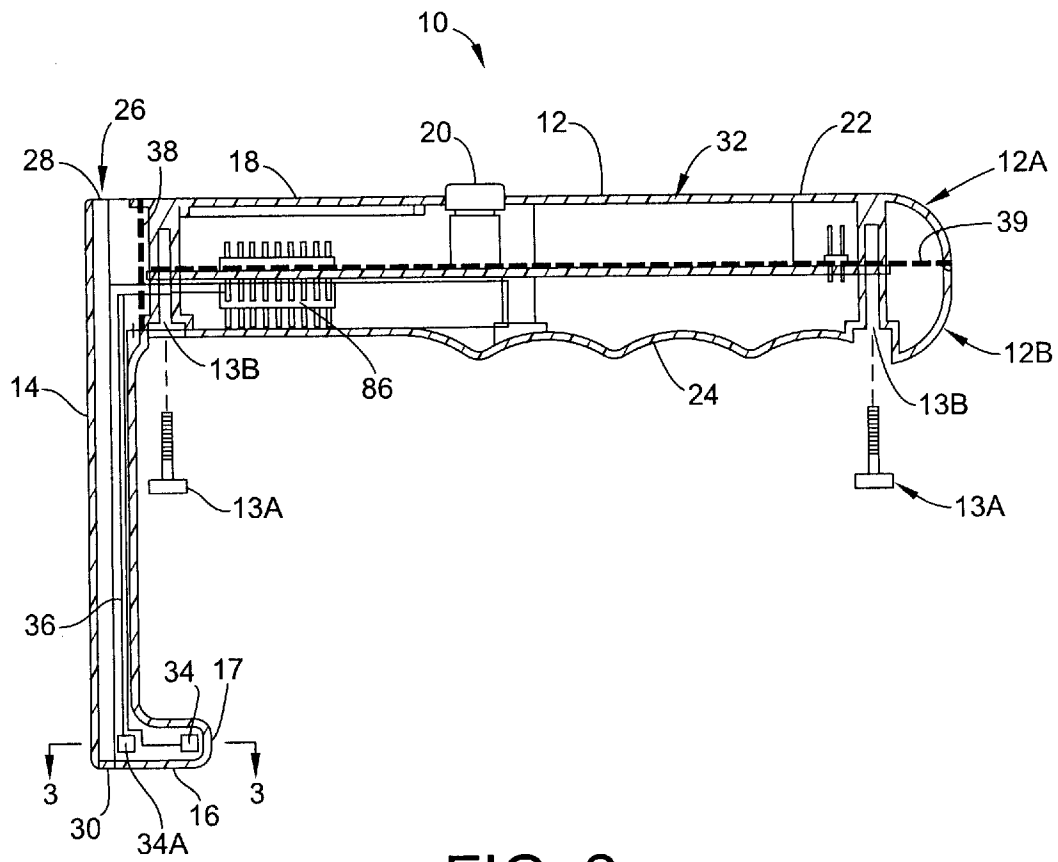


FIG. 2

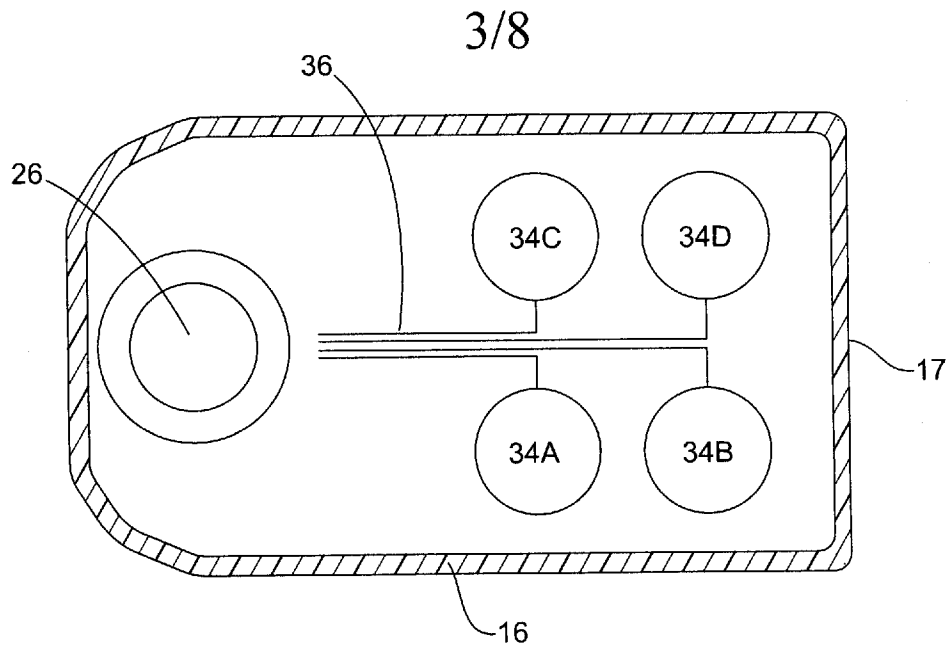


FIG. 3

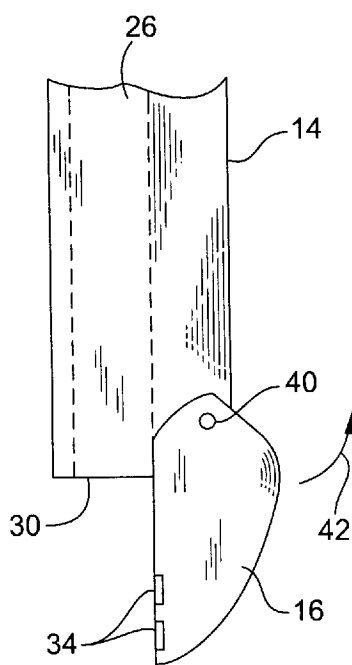


FIG. 4A

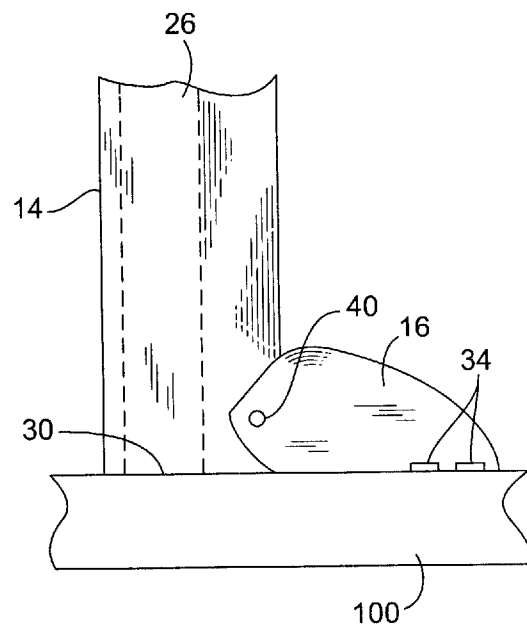


FIG. 4B

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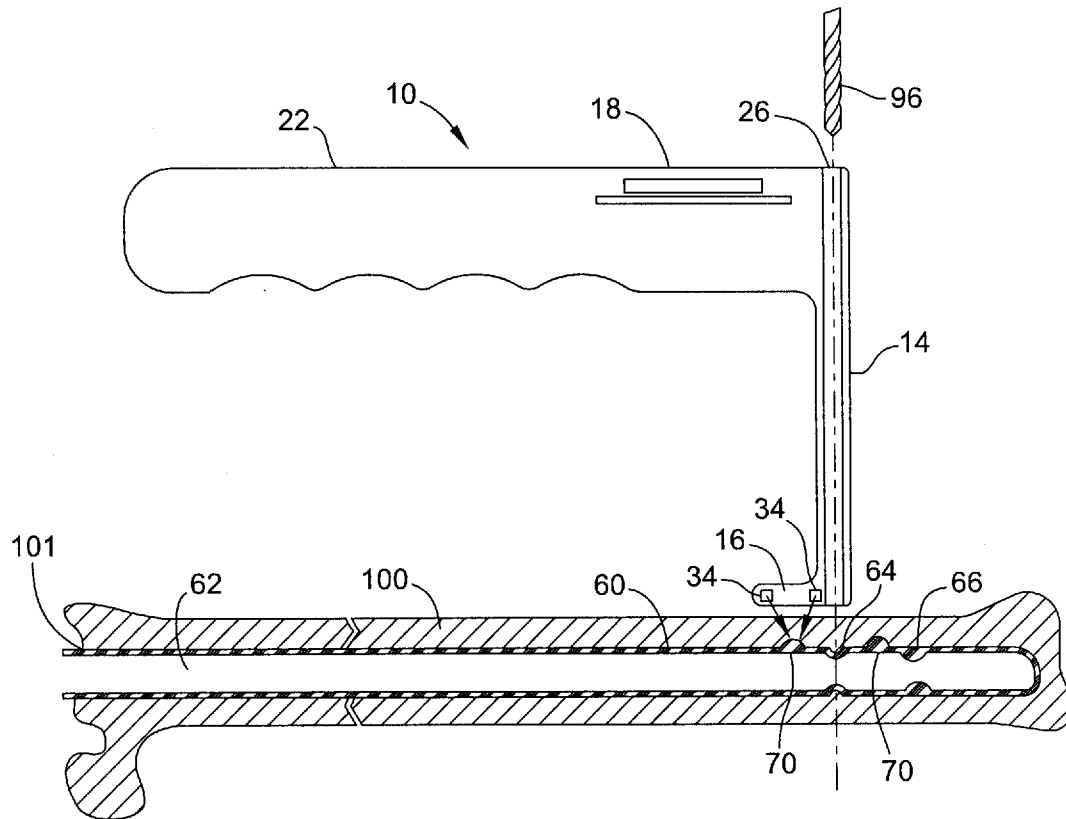


FIG. 5

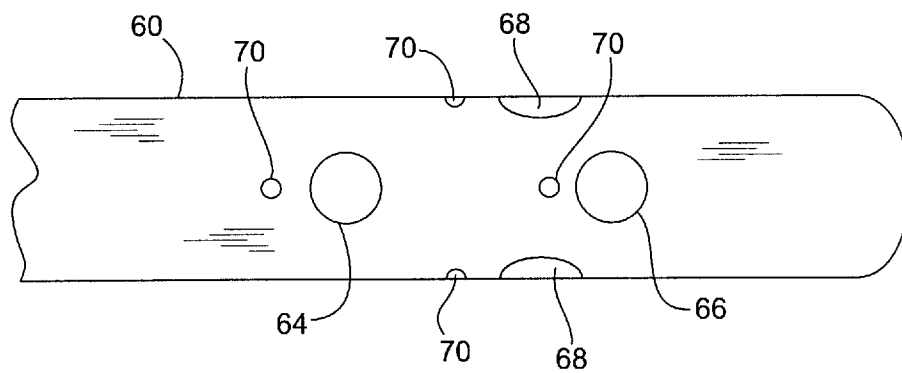


FIG. 6

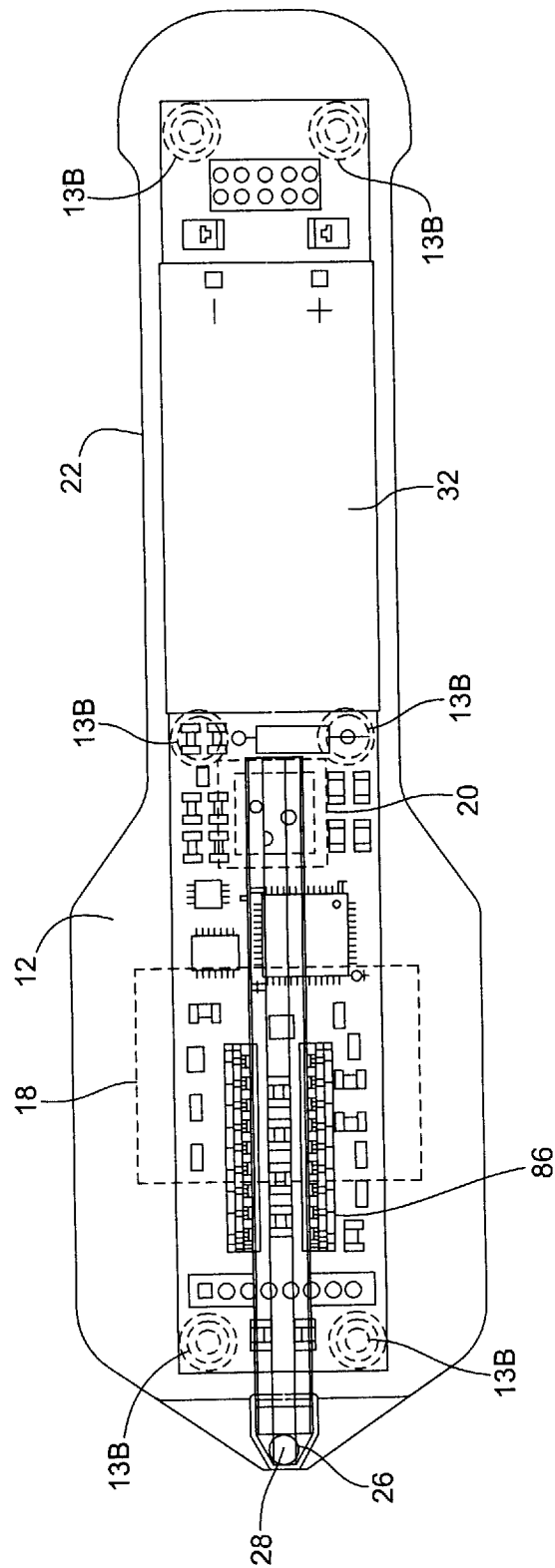


FIG. 7

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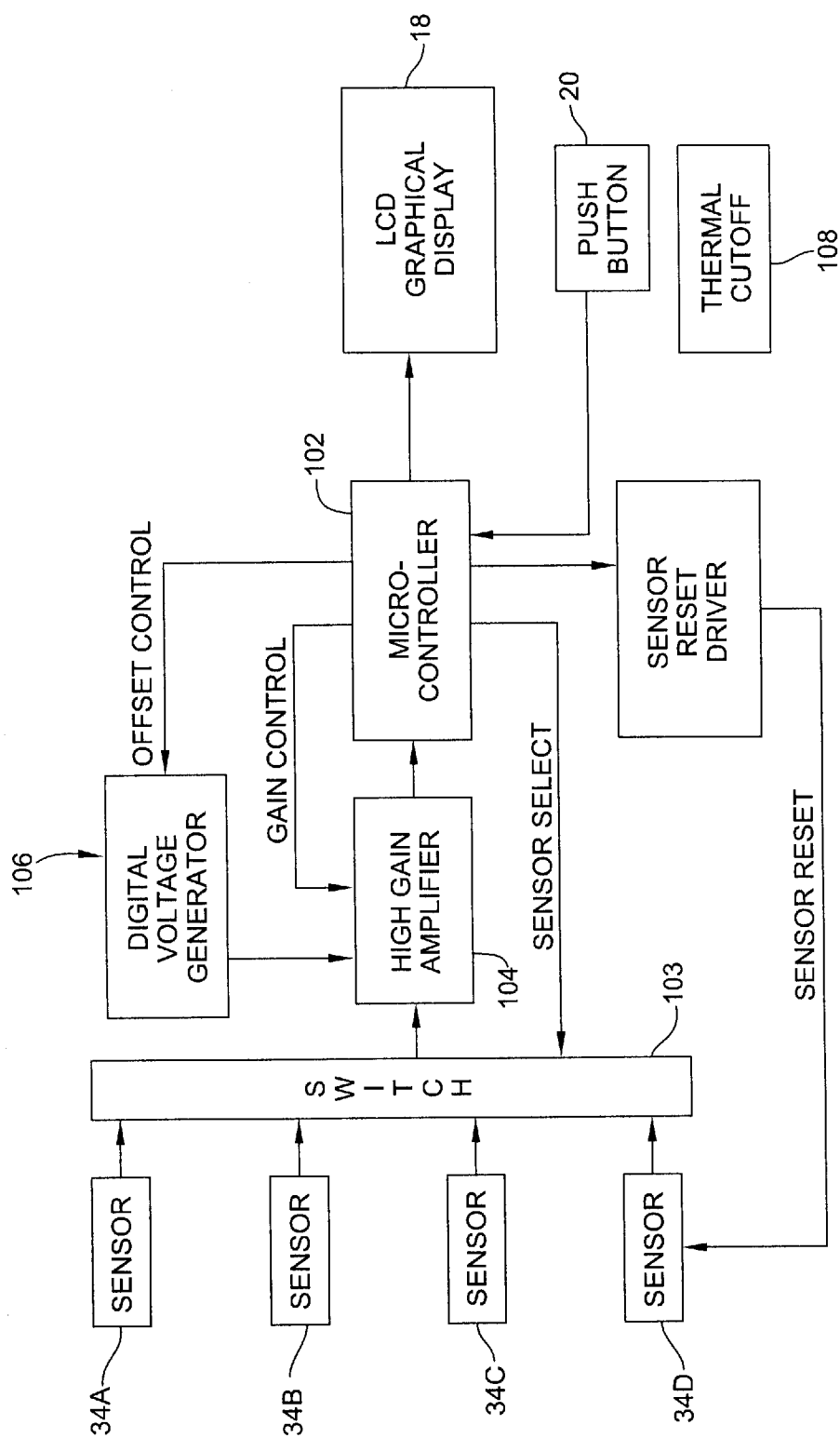


FIG. 8

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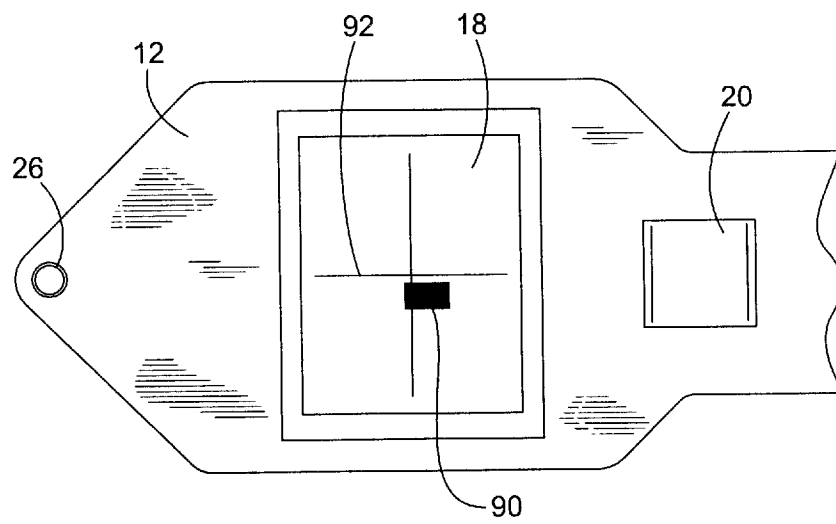


FIG. 9

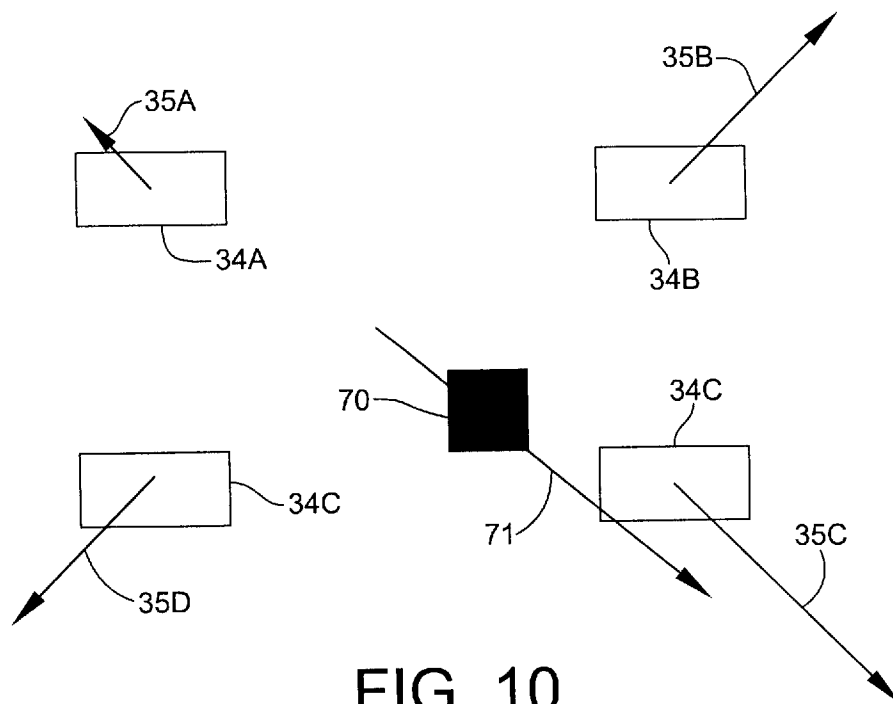


FIG. 10

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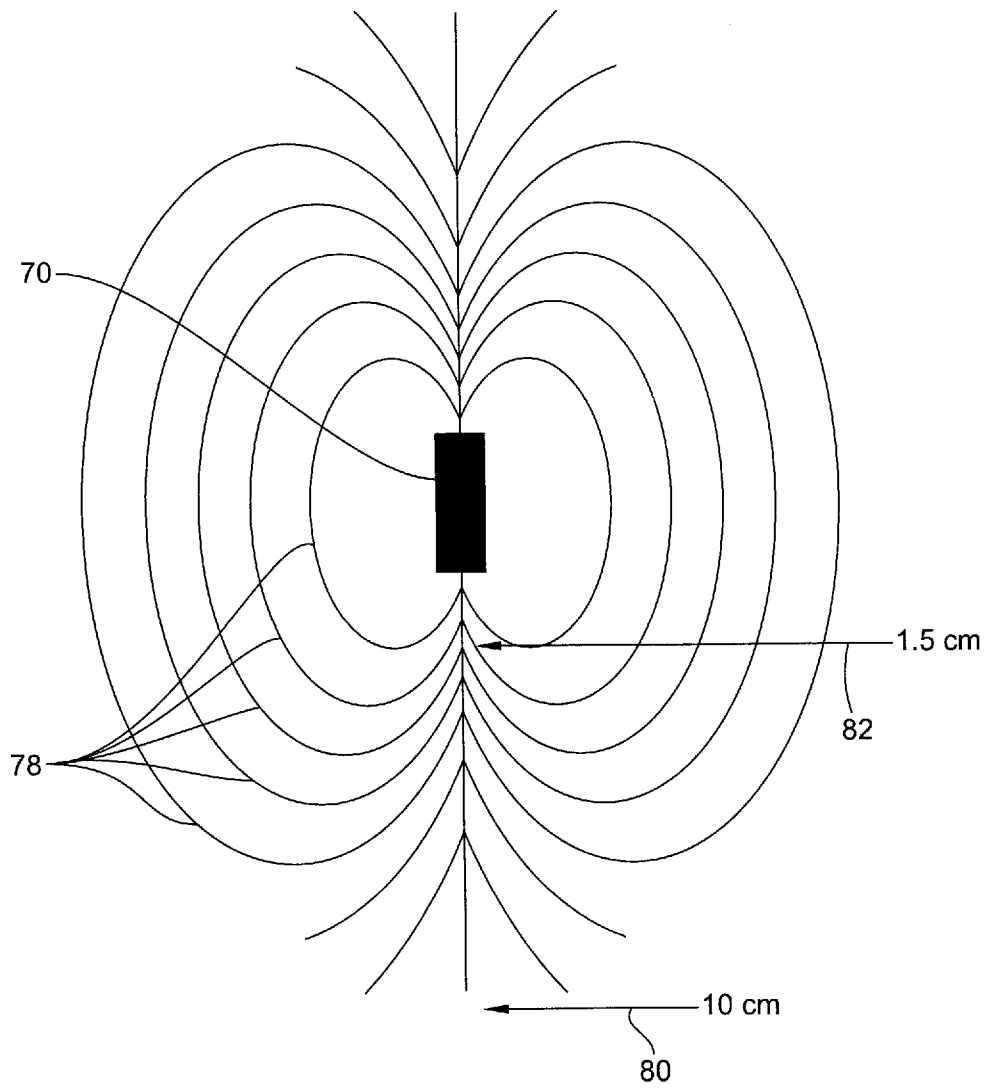


FIG. 11

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2009/055734

A. CLASSIFICATION OF SUBJECT MATTER

INV. A61B17/17
ADD. A61B17/72

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
A61B

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

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

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 411 503 A (HOLLSTIEN STEVEN B [US] ET AL) 2 May 1995 (1995-05-02) column 4, line 38 - column 5, line 55; figure 1	1,20
A	EP 0 589 592 A (ORTHOFIX SRL [IT]) 30 March 1994 (1994-03-30) page 4, line 41 - line 54; figure 1	1,20

☐ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

* Special categories of cited documents

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

18 December 2009

Date of mailing of the international search report

08/01/2010

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Authorized officer

Hagberg, Åsa

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2009/055734

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

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

1. ☒ Claims Nos.: 26
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

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

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

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

Remark on Protest

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

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2009/055734

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5411503	A	02-05-1995	AU 7057194 A WO 9500085 A1	17-01-1995 05-01-1995
EP 0589592	A	30-03-1994	AU 664113 B2 AU 4736293 A CN 1084382 A JP 6277229 A MX 9305807 A1 NO 933360 A TR 27301 A	02-11-1995 31-03-1994 30-03-1994 04-10-1994 31-03-1994 23-03-1994 29-12-1994