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(54) **SYSTEMS AND METHODS FOR
INSTALLING AN ORTHOPEDIC IMPLANT**

(71) Applicant: **MicroPort Orthopedics Holdings Inc.,**
Tiel (NL)

(72) Inventors: **Paul Stemniski**, Arlington, TN (US);
Sarah Lancianese, Orlando, FL (US);
Richard Obert, Poway, CA (US)

(73) Assignee: **MICROPORT ORTHOPEDICS
HOLDINGS, INC.**, Tiel (NL)

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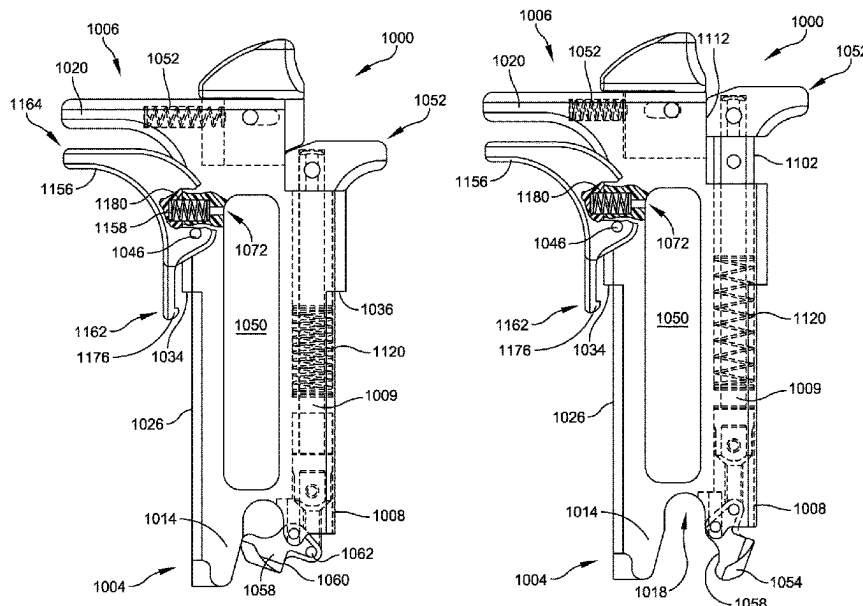
Primary Examiner — Sameh R Boles

(74) *Attorney, Agent, or Firm* — DUANE MORRIS LLP

(57) **ABSTRACT**

A system includes a cartridge having an elongate body
extending from a first end to a second end and having a top
side and a bottom side. The cartridge defines a first hole
adjacent to the first end that extends through the cartridge
from the bottom side to the top side. The top side of the
cartridge defines a pair of parallel slots that extend perpen-
dicular with respect to a longitudinal axis of the cartridge.
Each slot of the pair of parallel slots is equidistant from a
central axis defined by the first hole.

16 Claims, 68 Drawing Sheets



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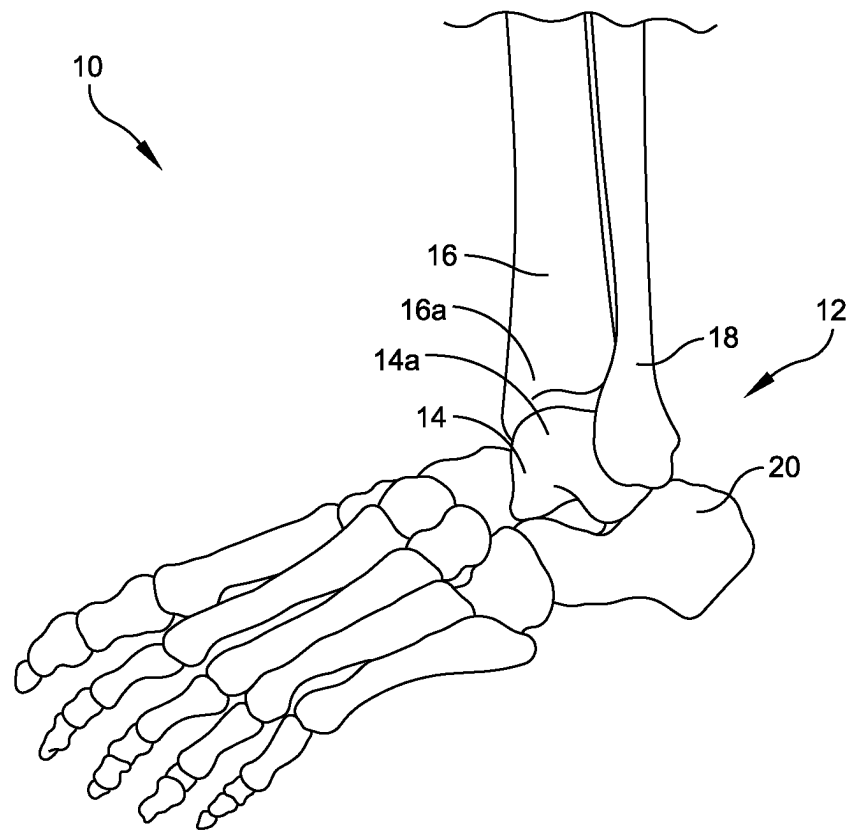


FIG. 1

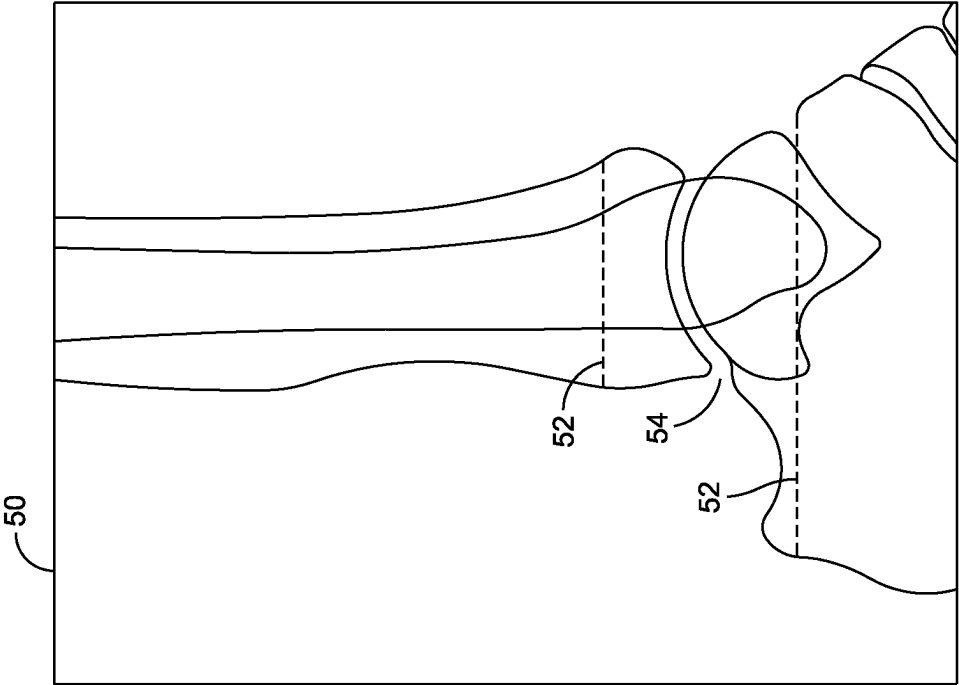


FIG. 2B

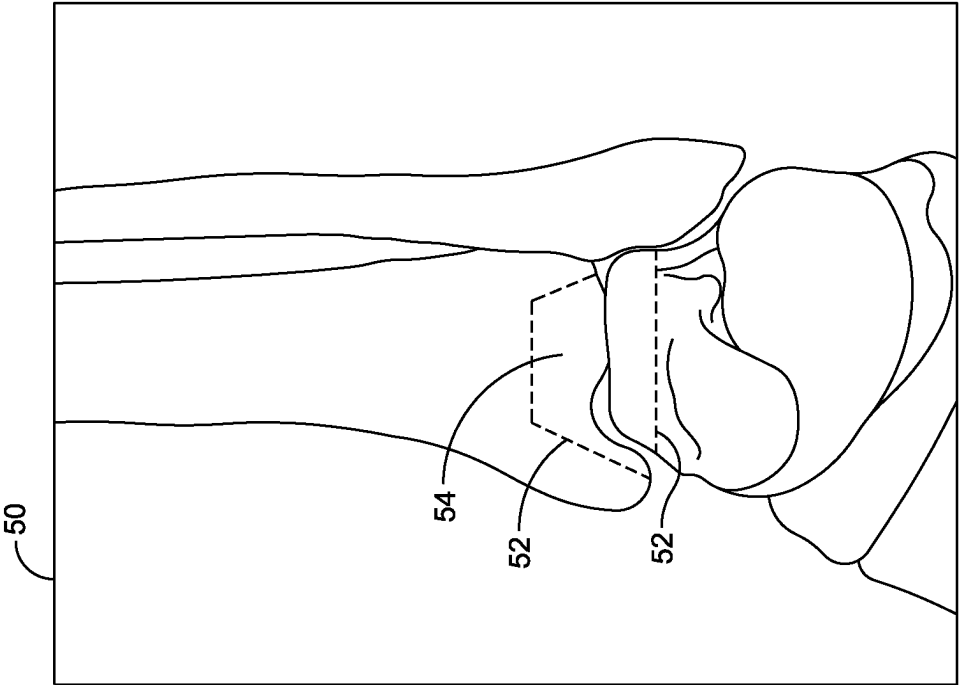


FIG. 2A

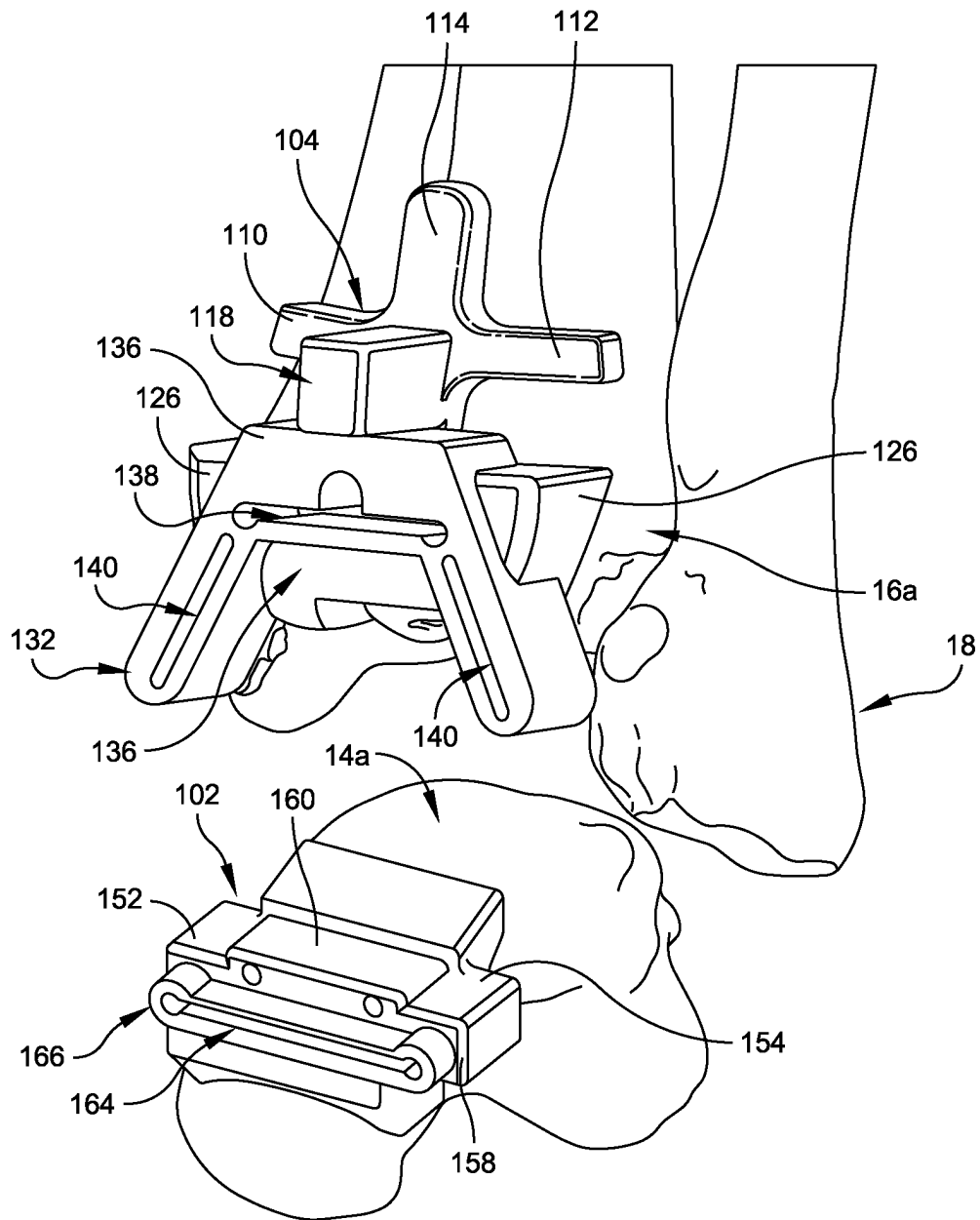


FIG. 3

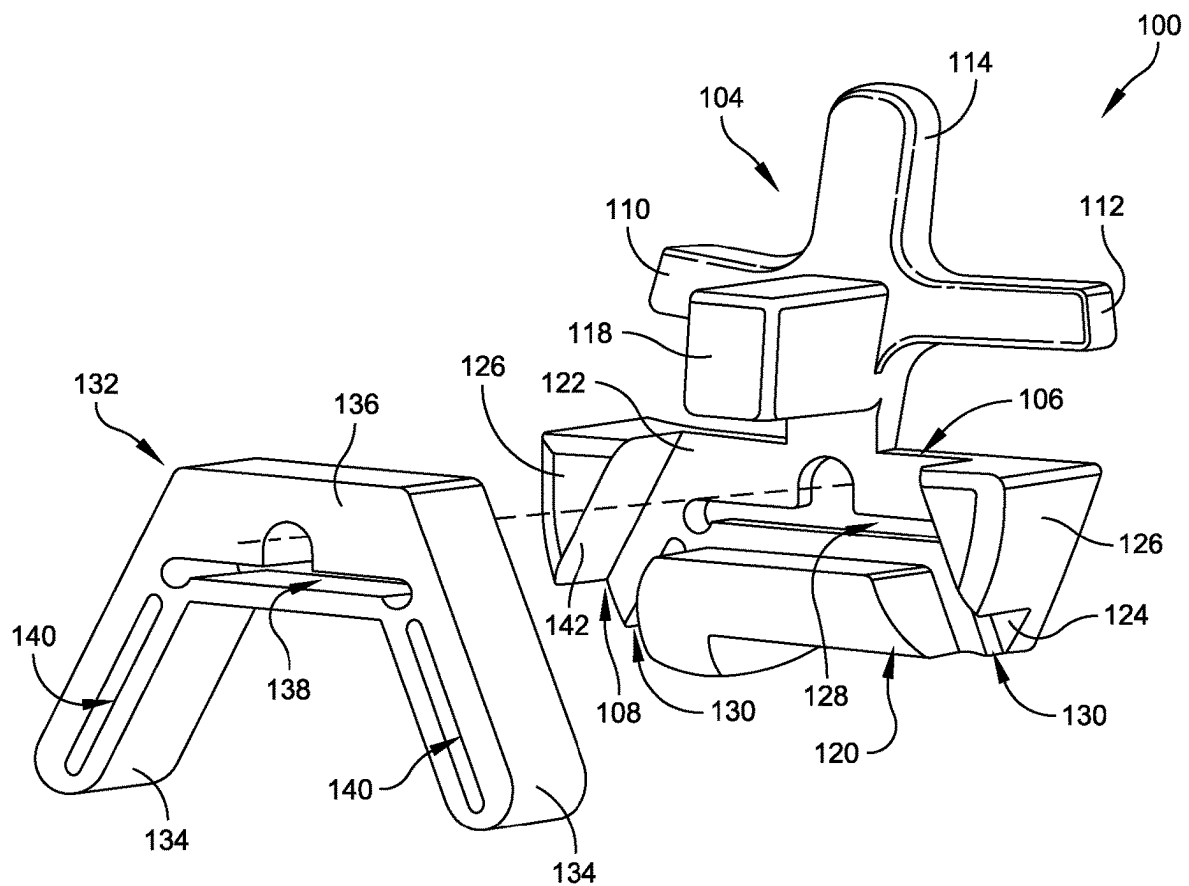


FIG. 4

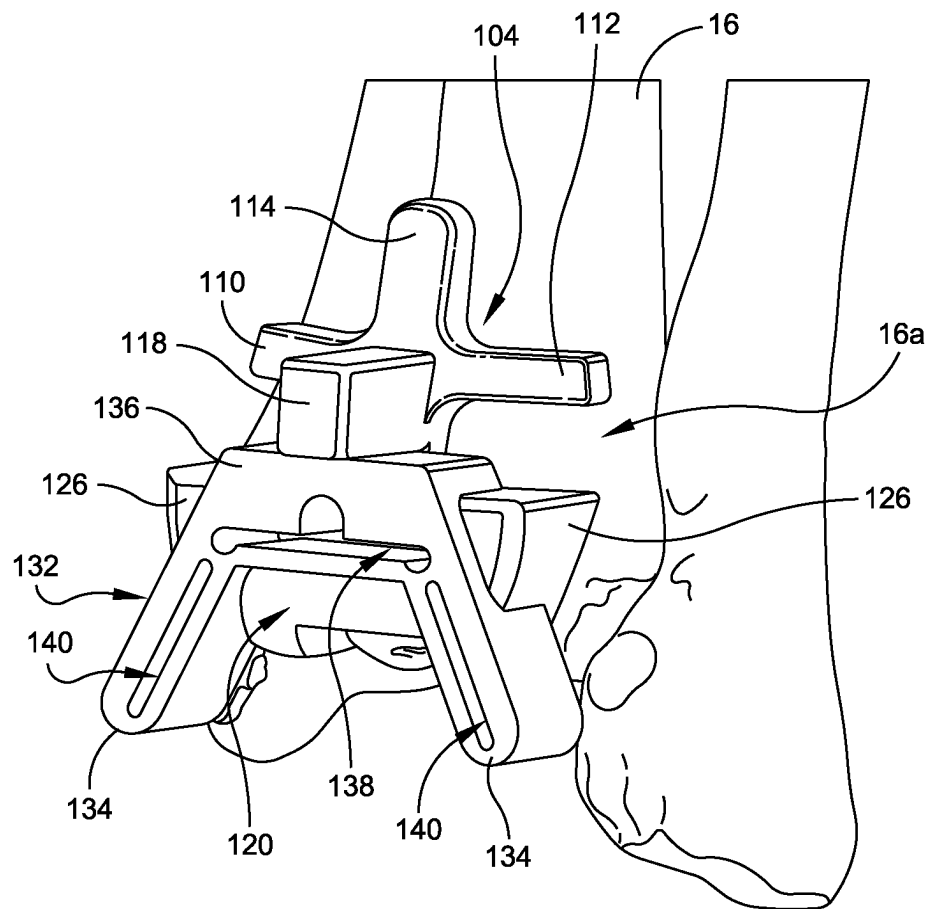


FIG. 5

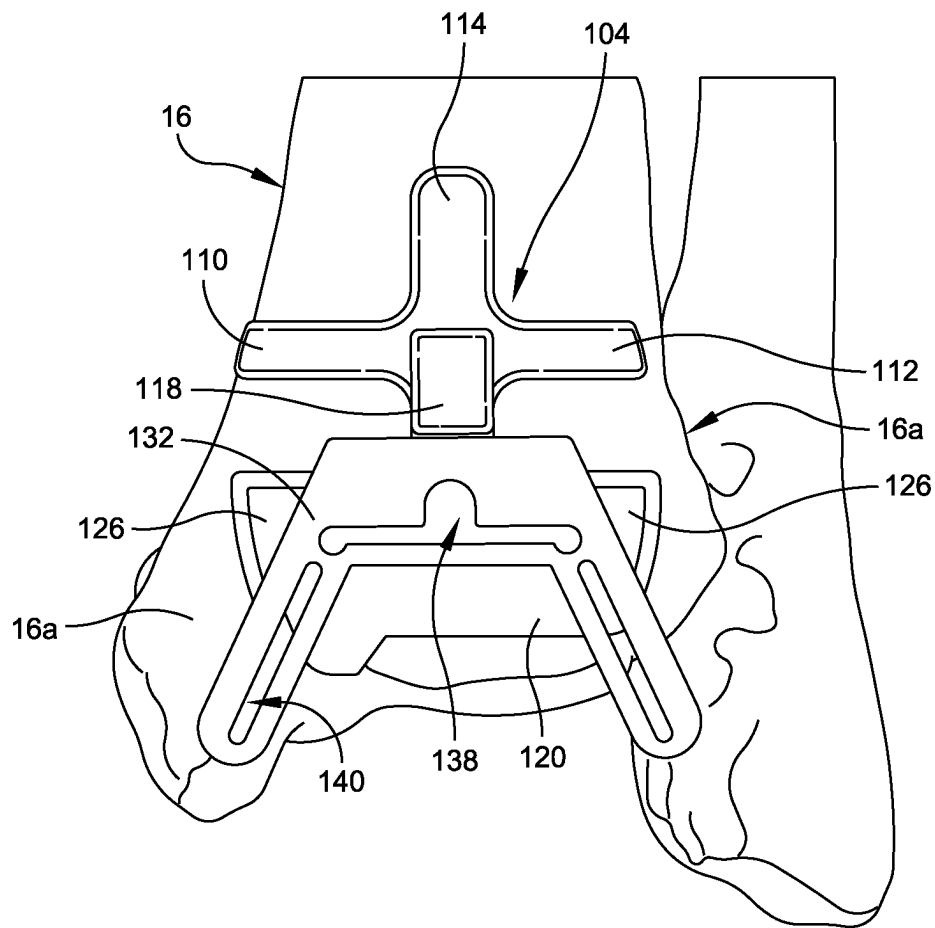


FIG. 6

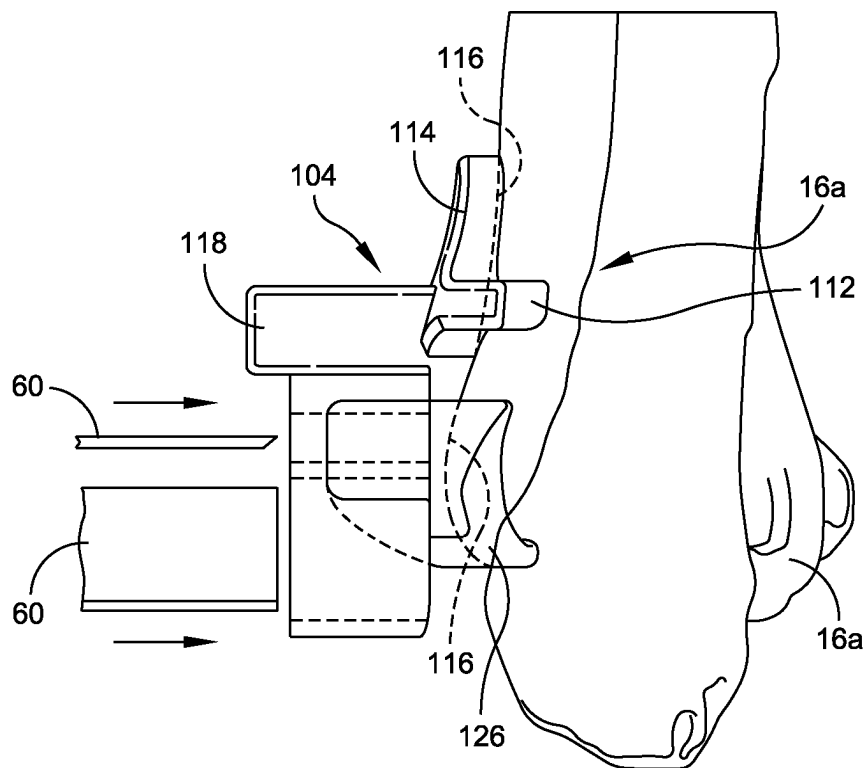


FIG. 7

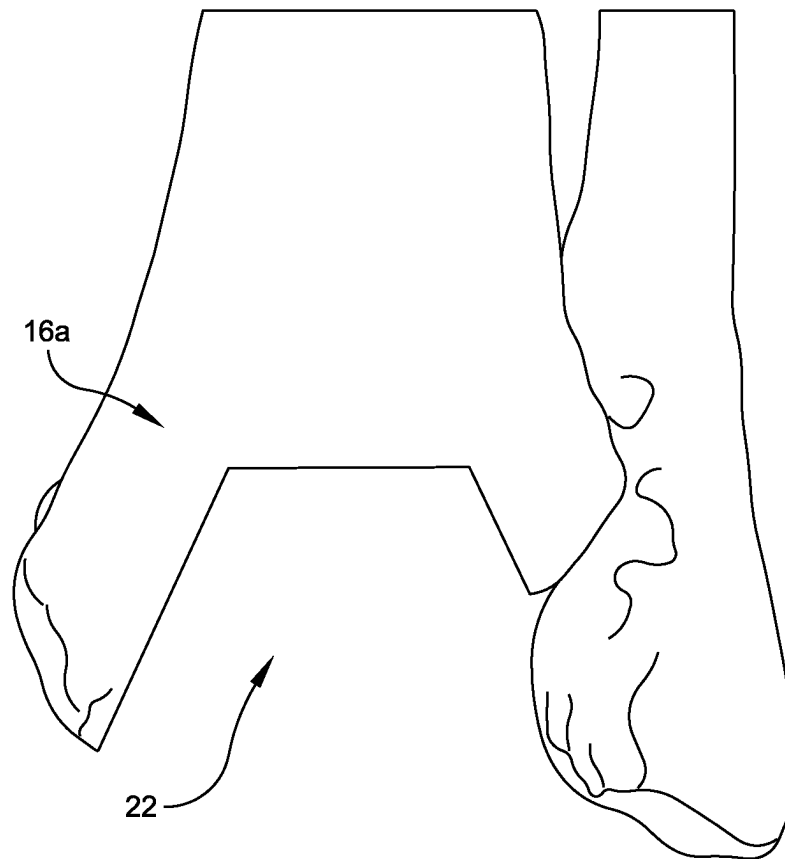


FIG. 8

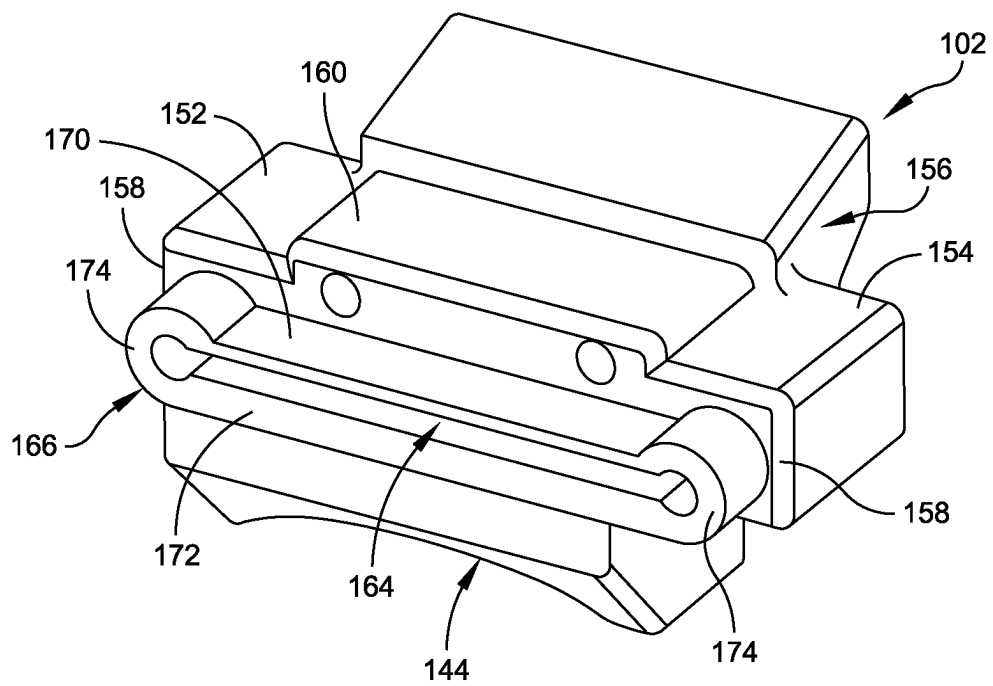


FIG. 9

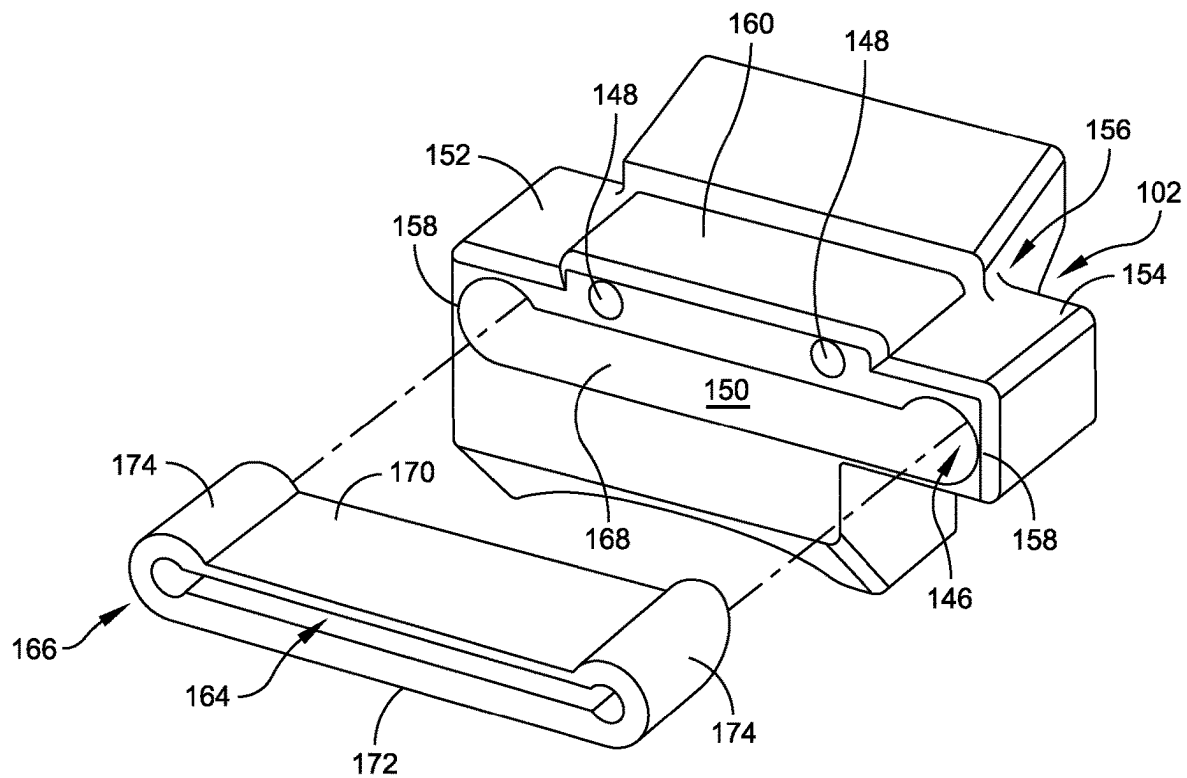


FIG. 10

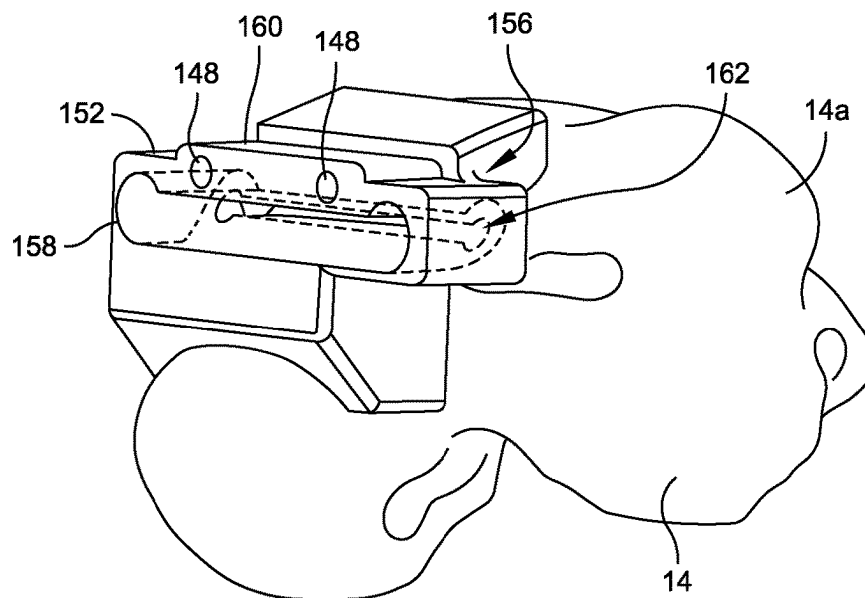


FIG. 11

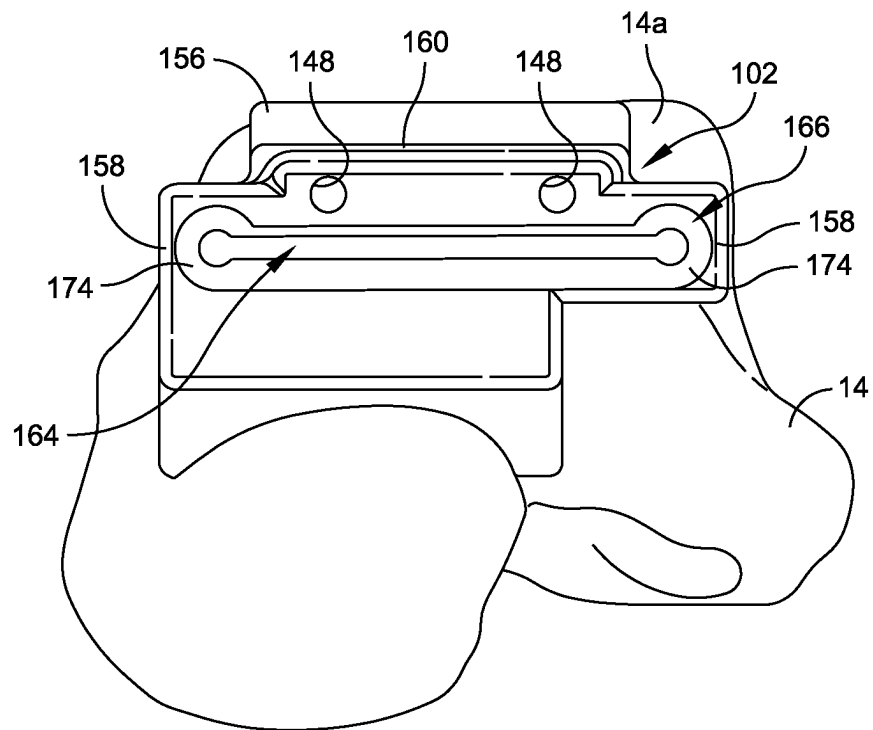


FIG. 12

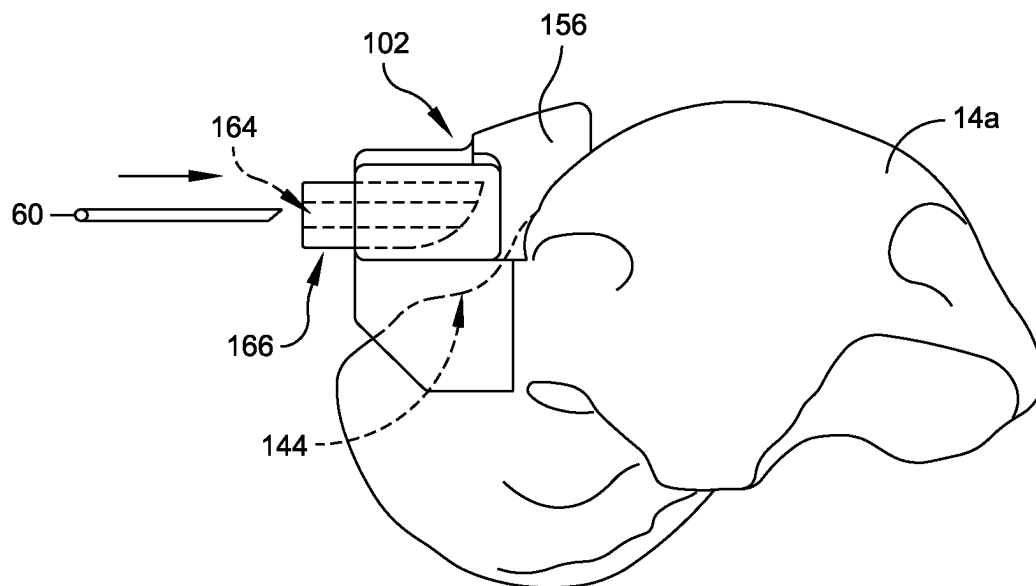


FIG. 13

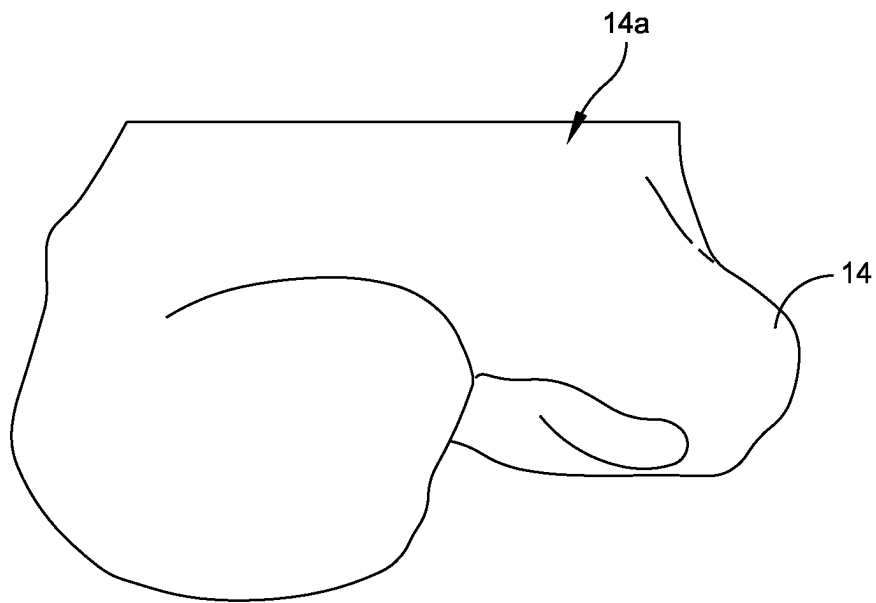


FIG. 14

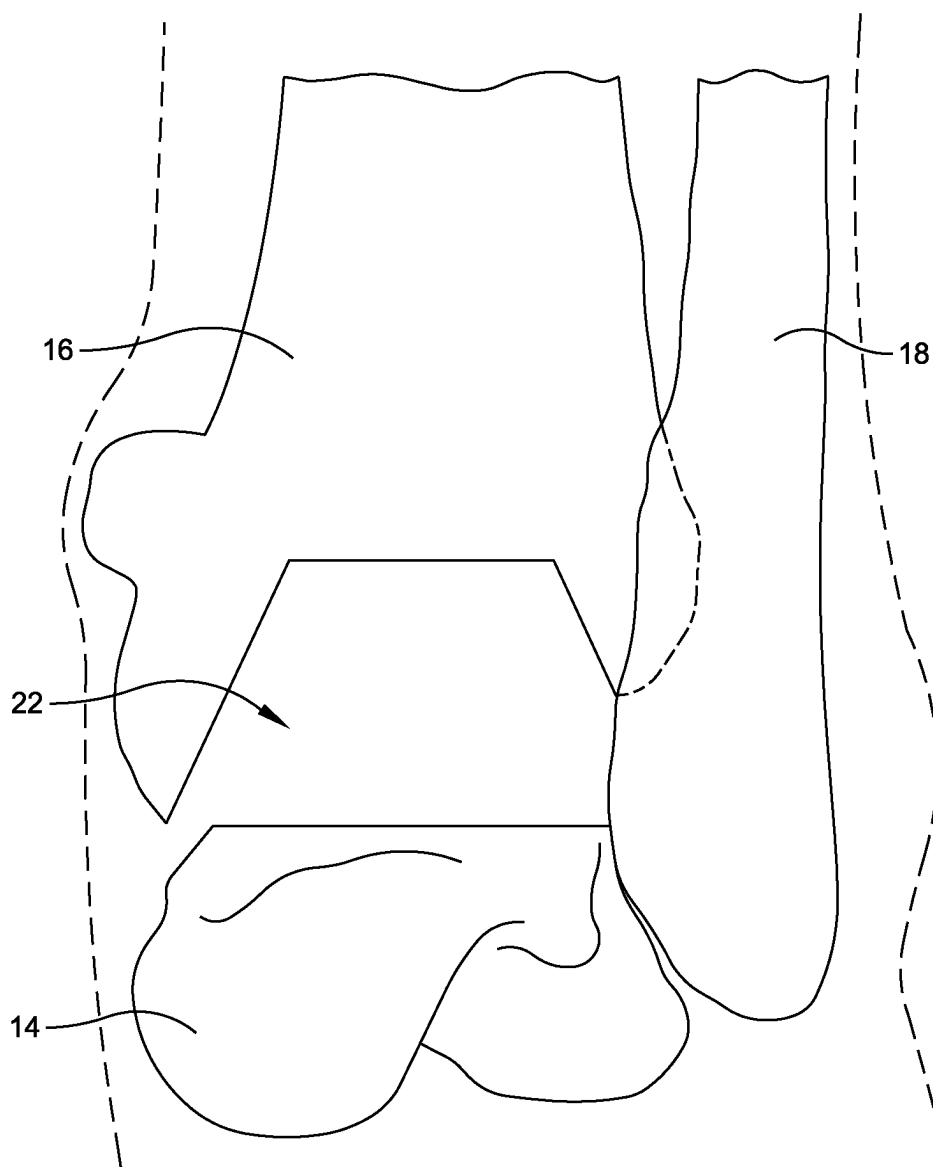


FIG. 15

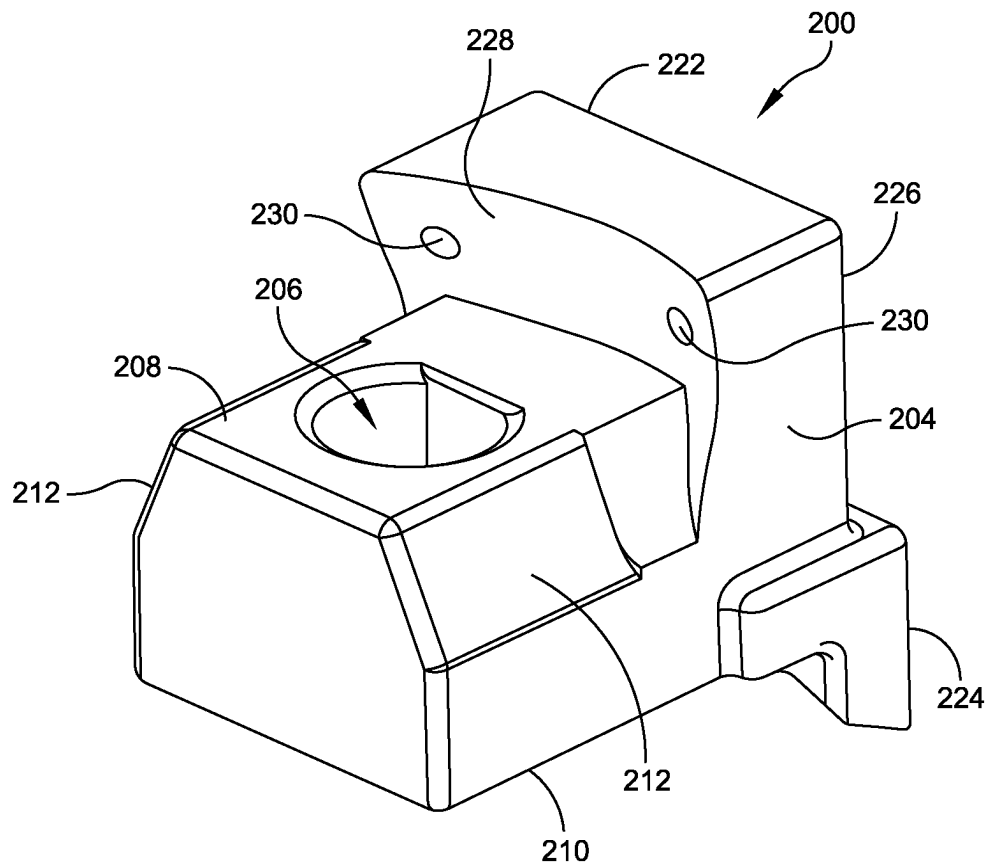


FIG. 16

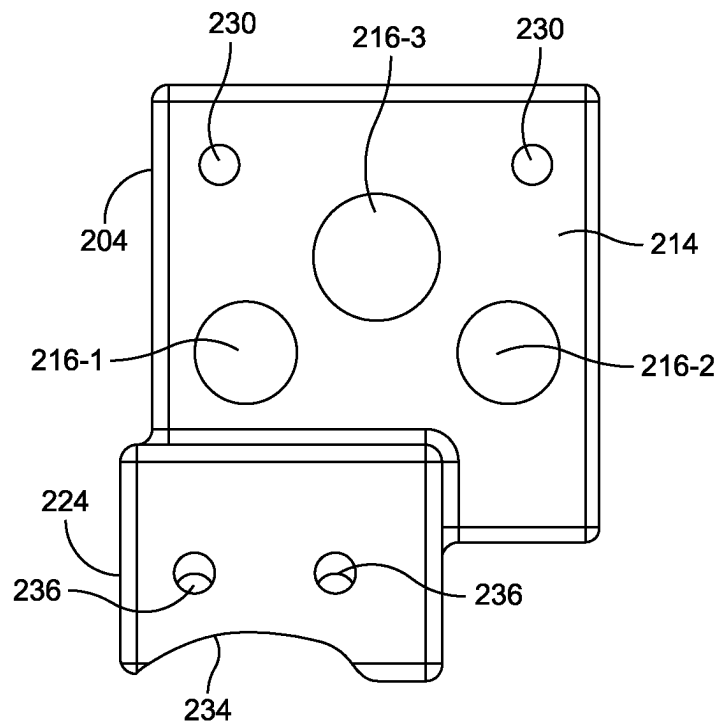


FIG. 17

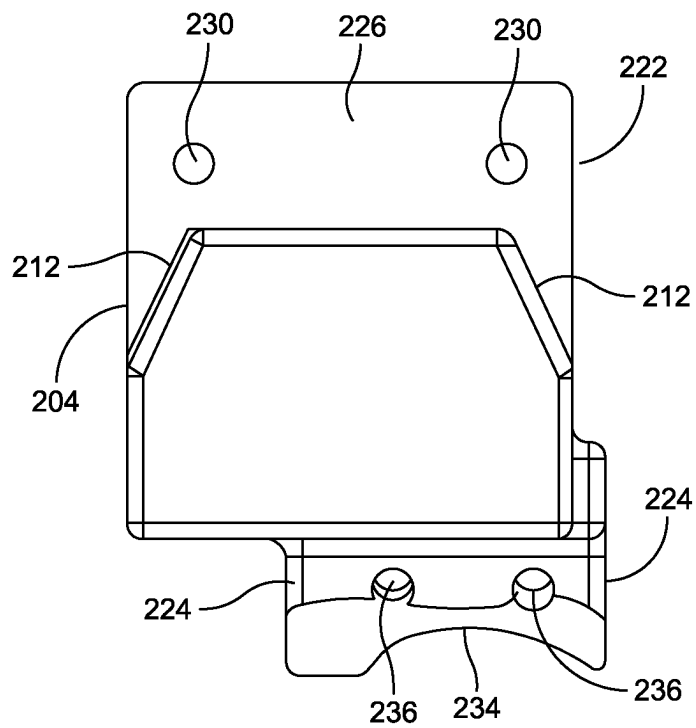


FIG. 18

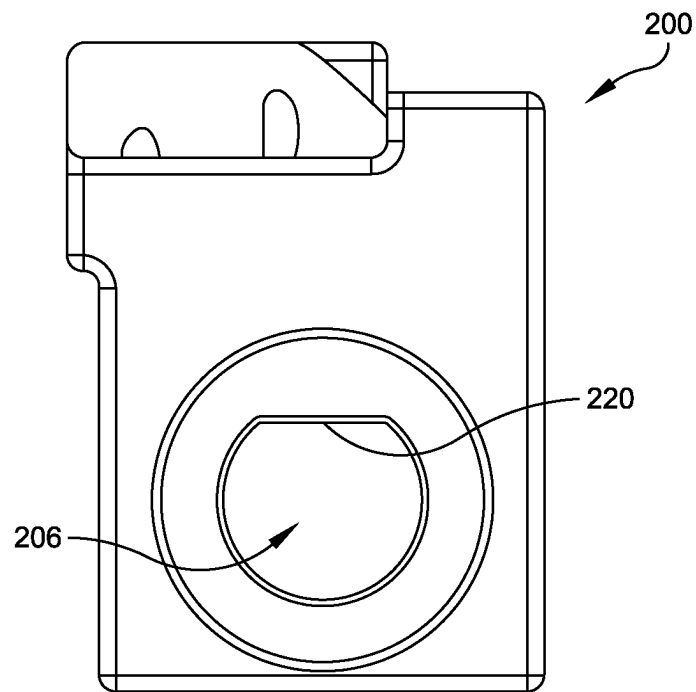


FIG. 19

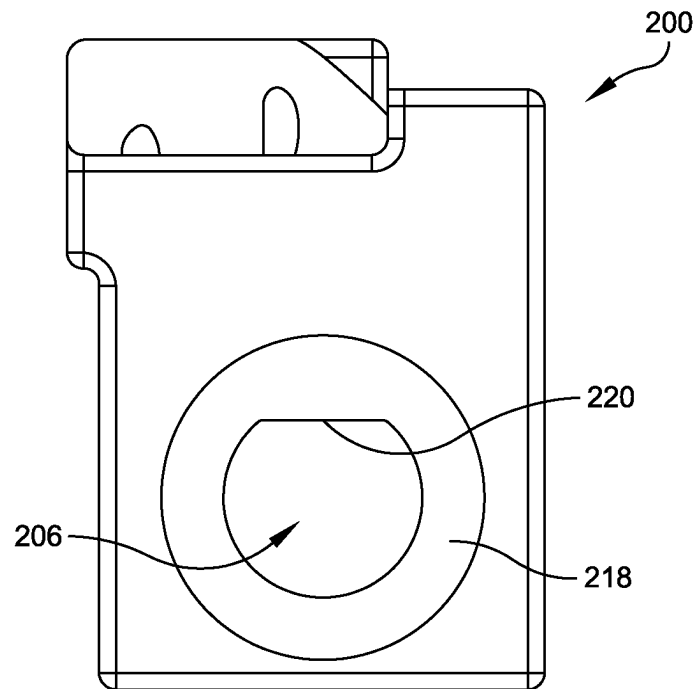


FIG. 20

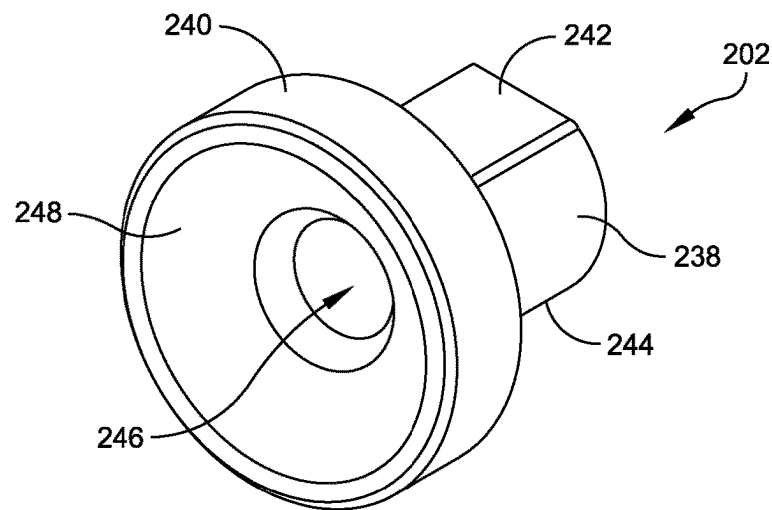


FIG. 21

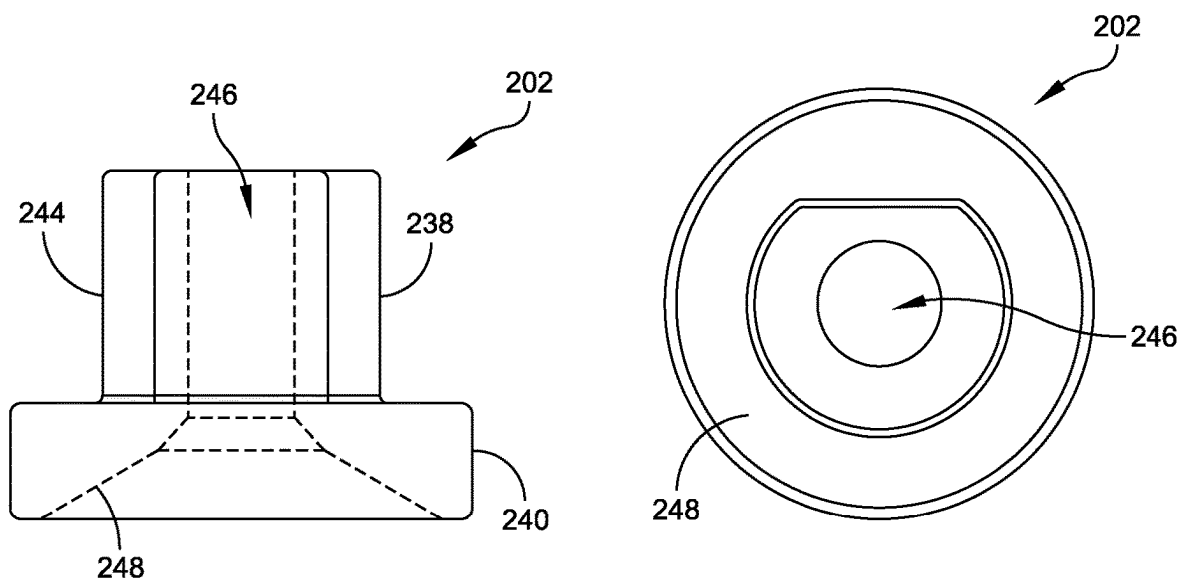


FIG. 22

FIG. 23

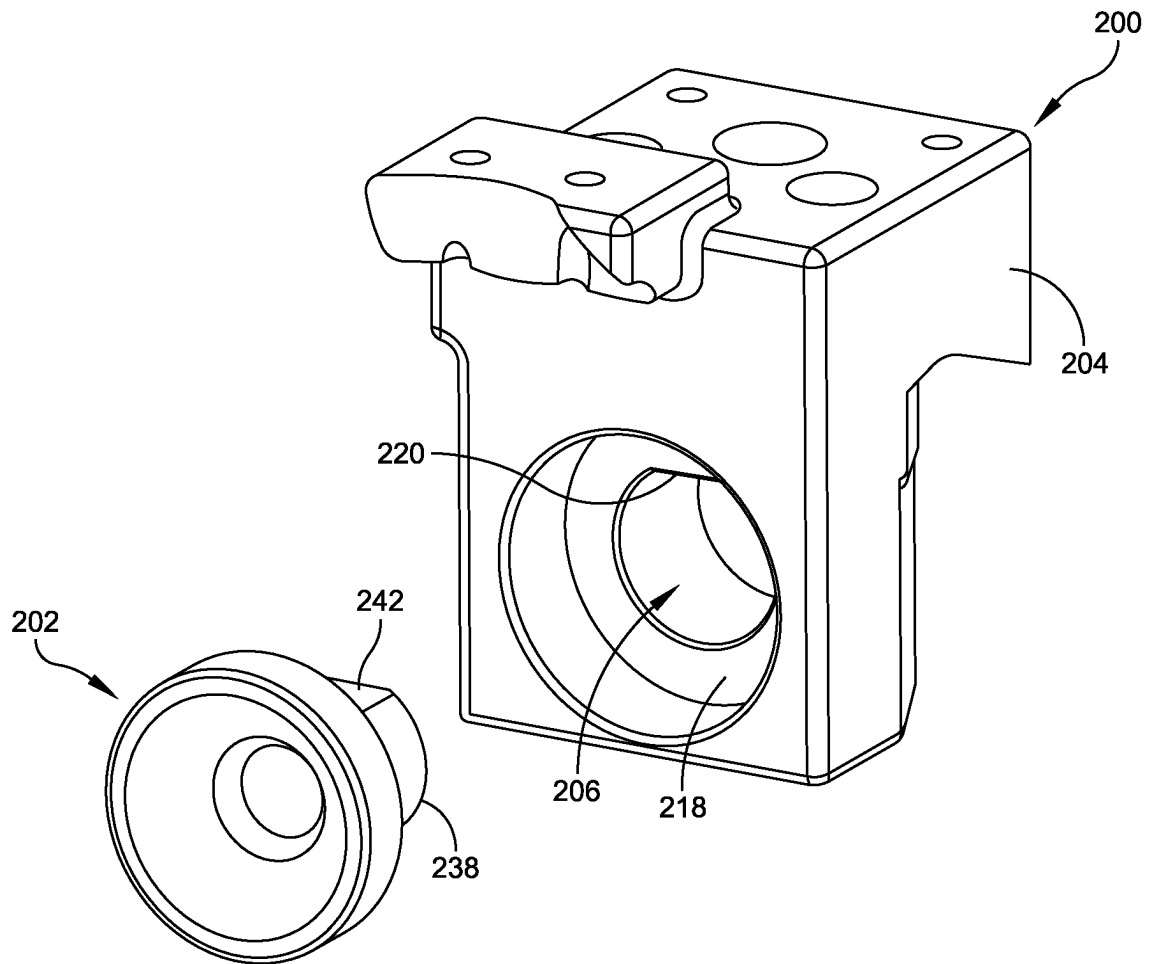


FIG. 24

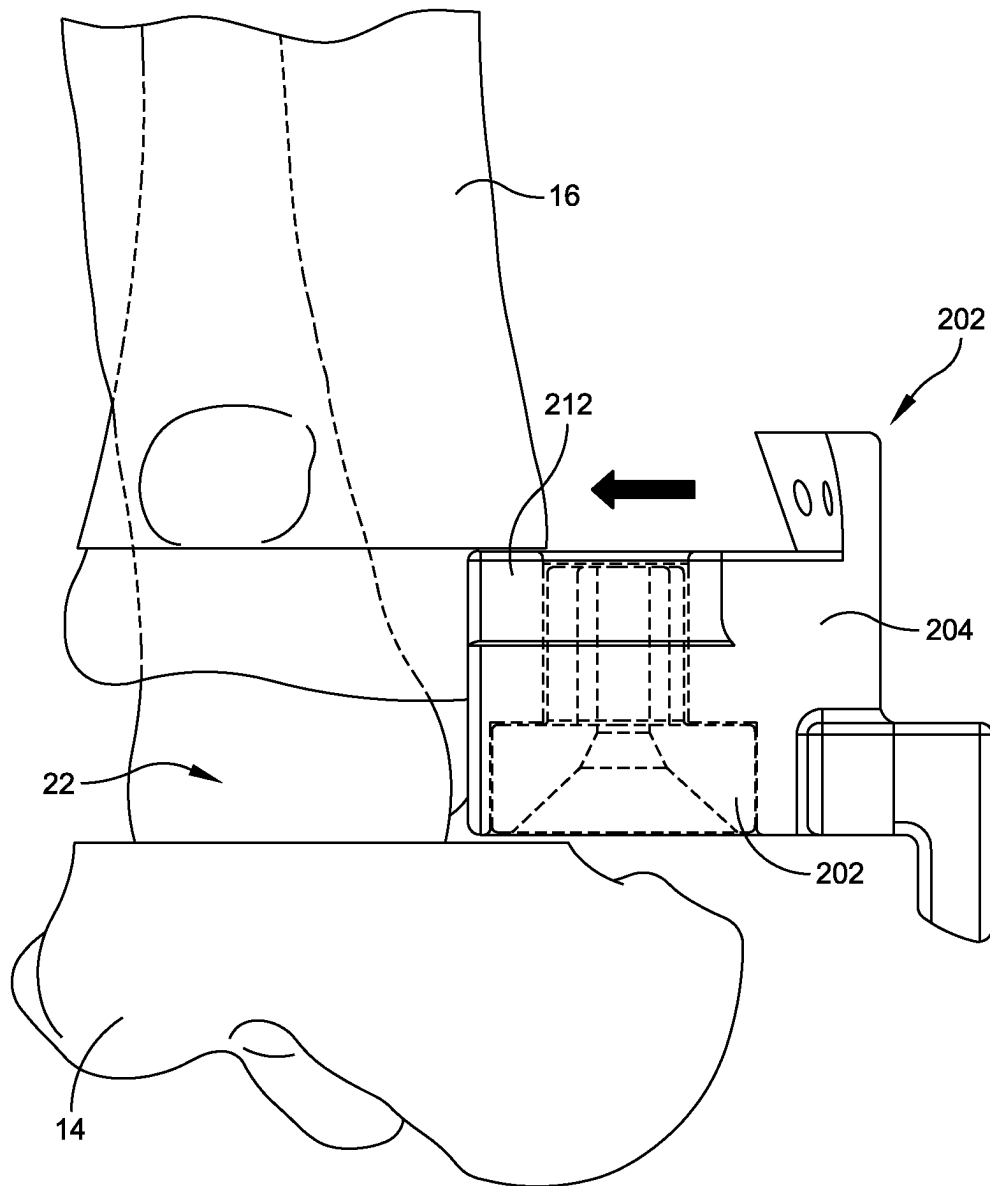


FIG. 25A

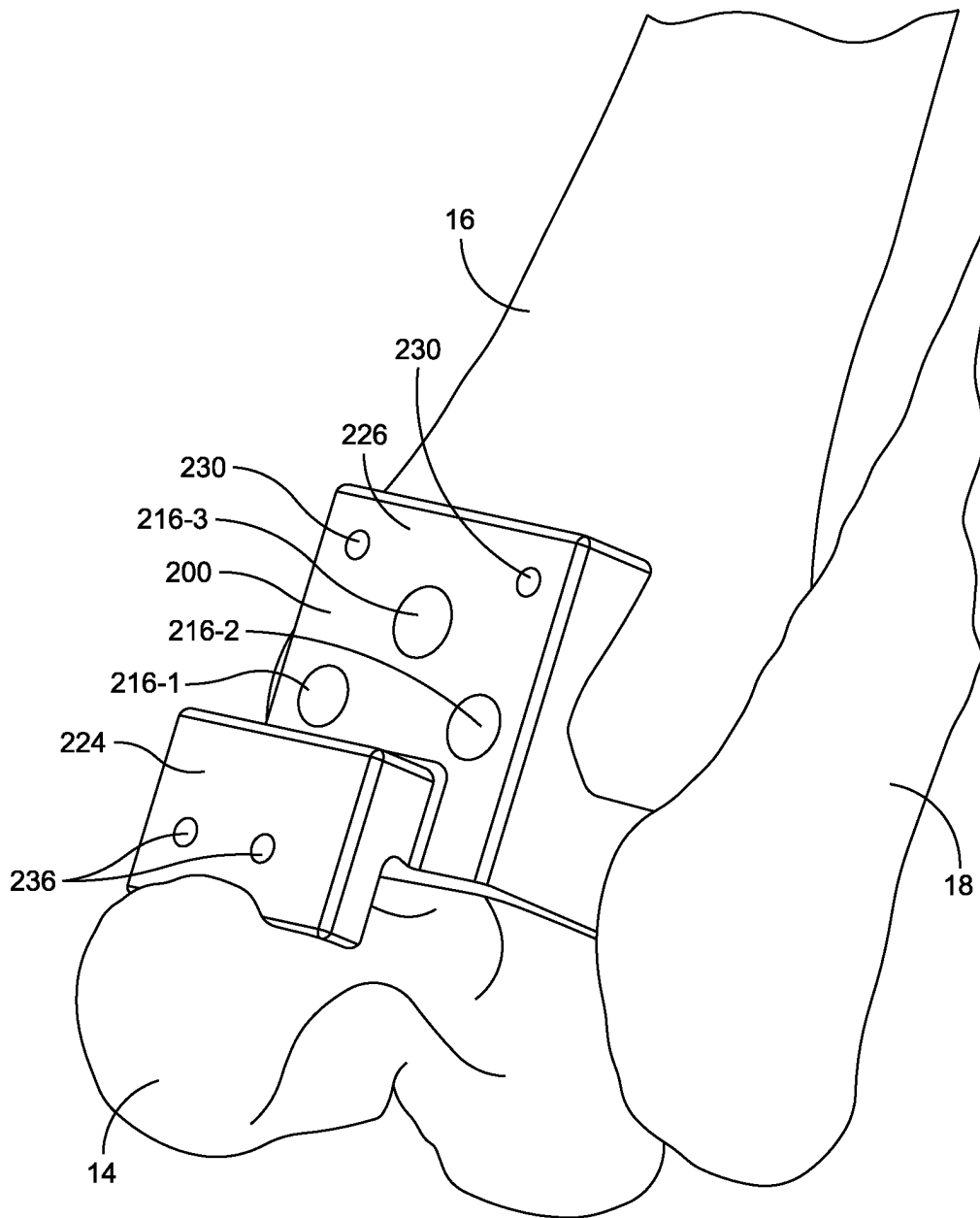


FIG. 25B

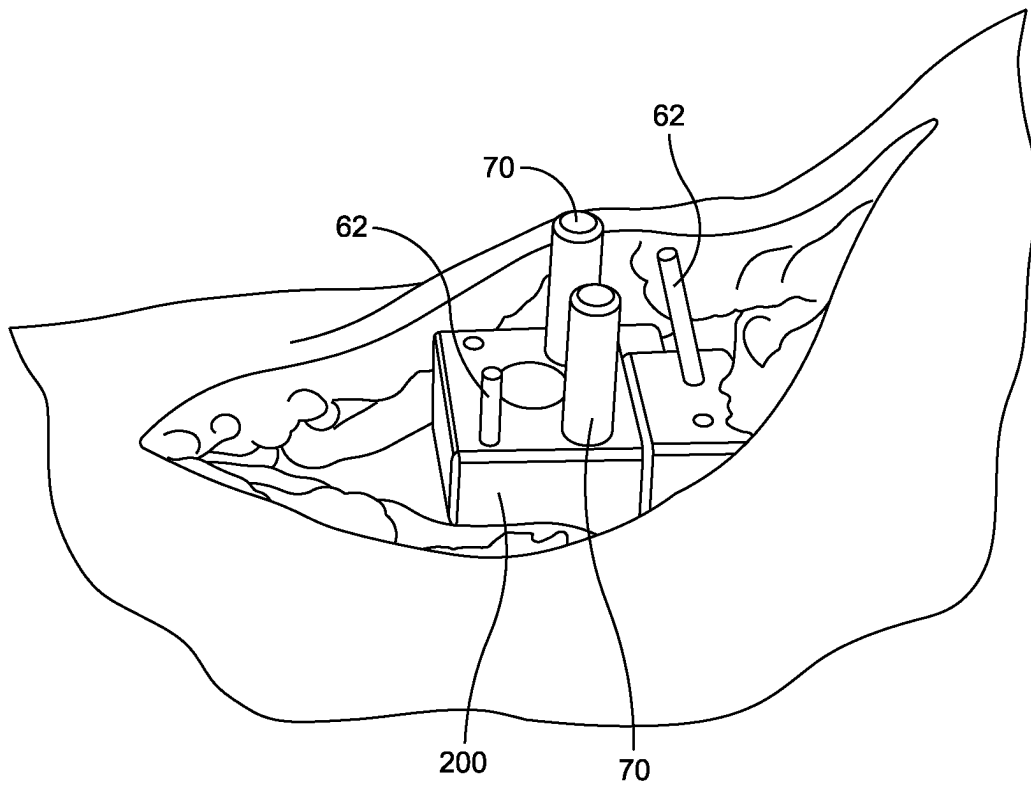


FIG. 25C

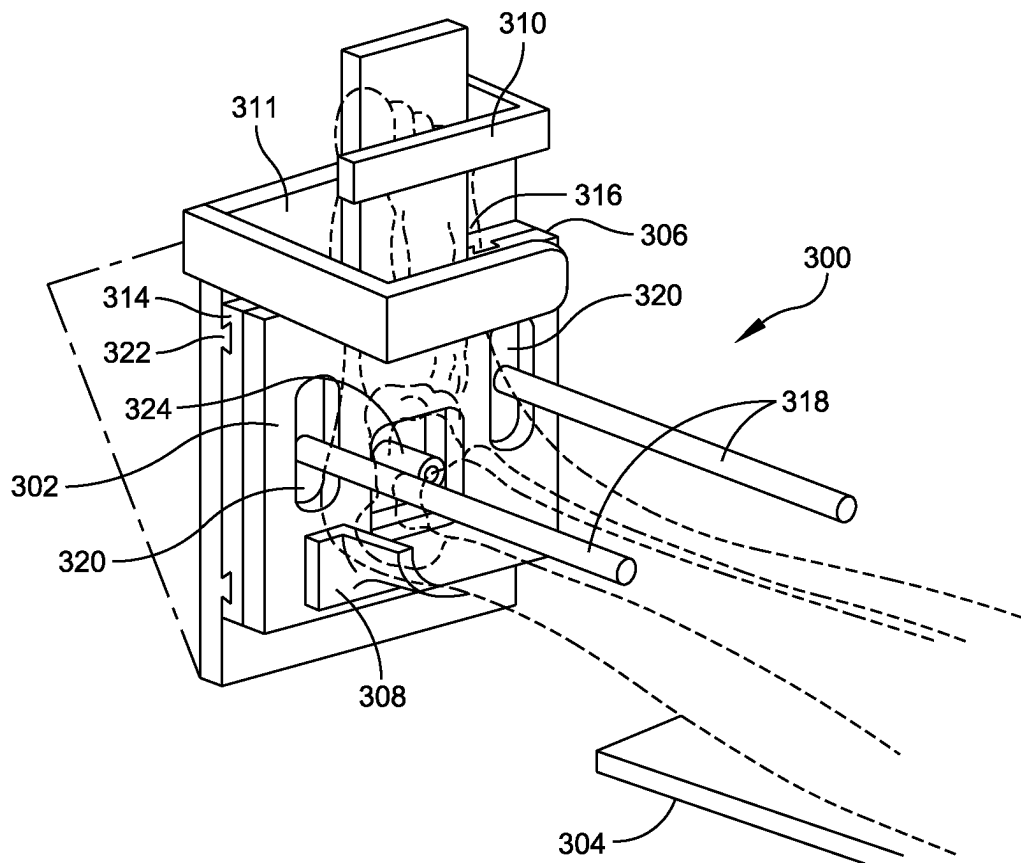


FIG. 26

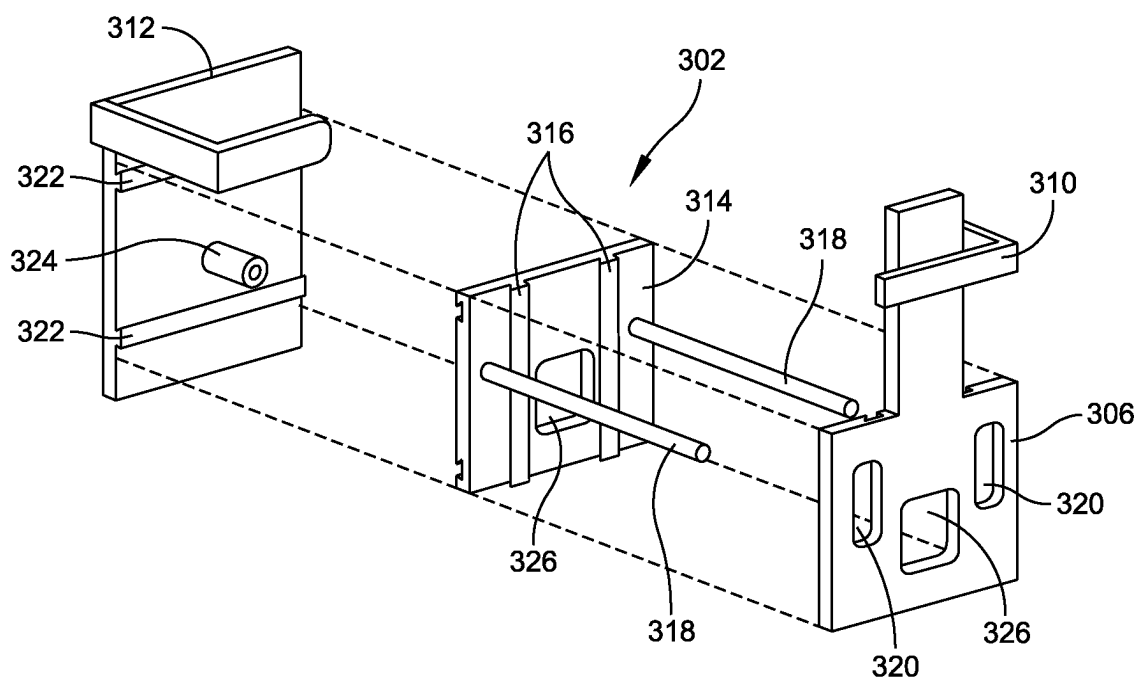


FIG. 27

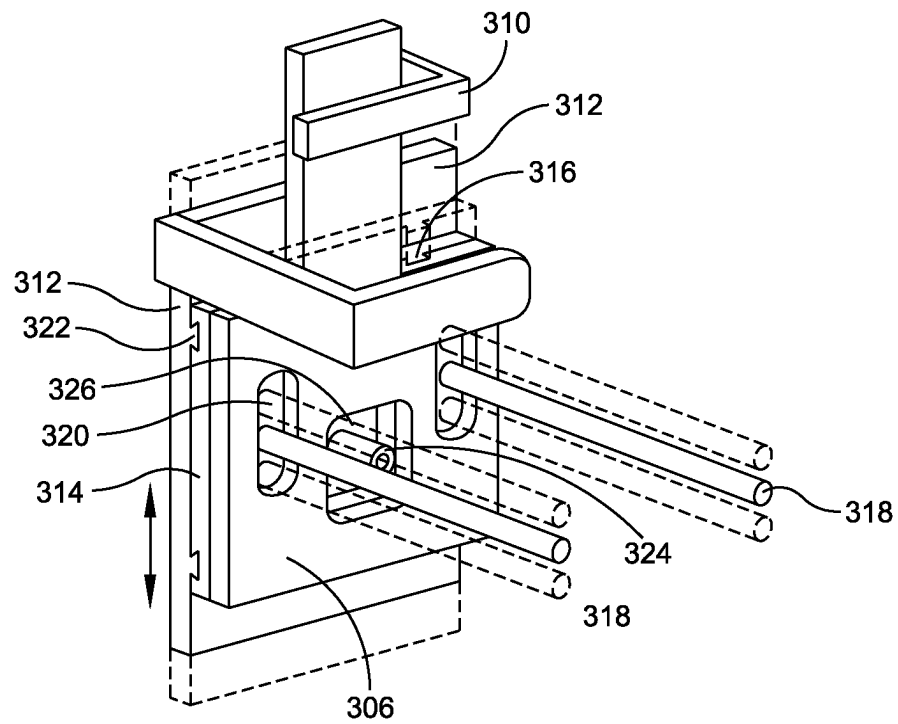


FIG. 28A

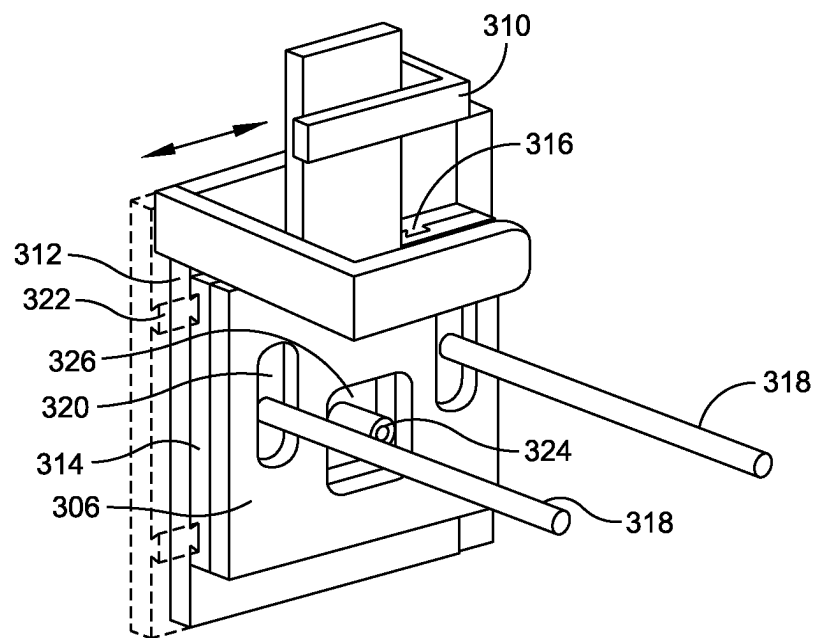


FIG. 28B

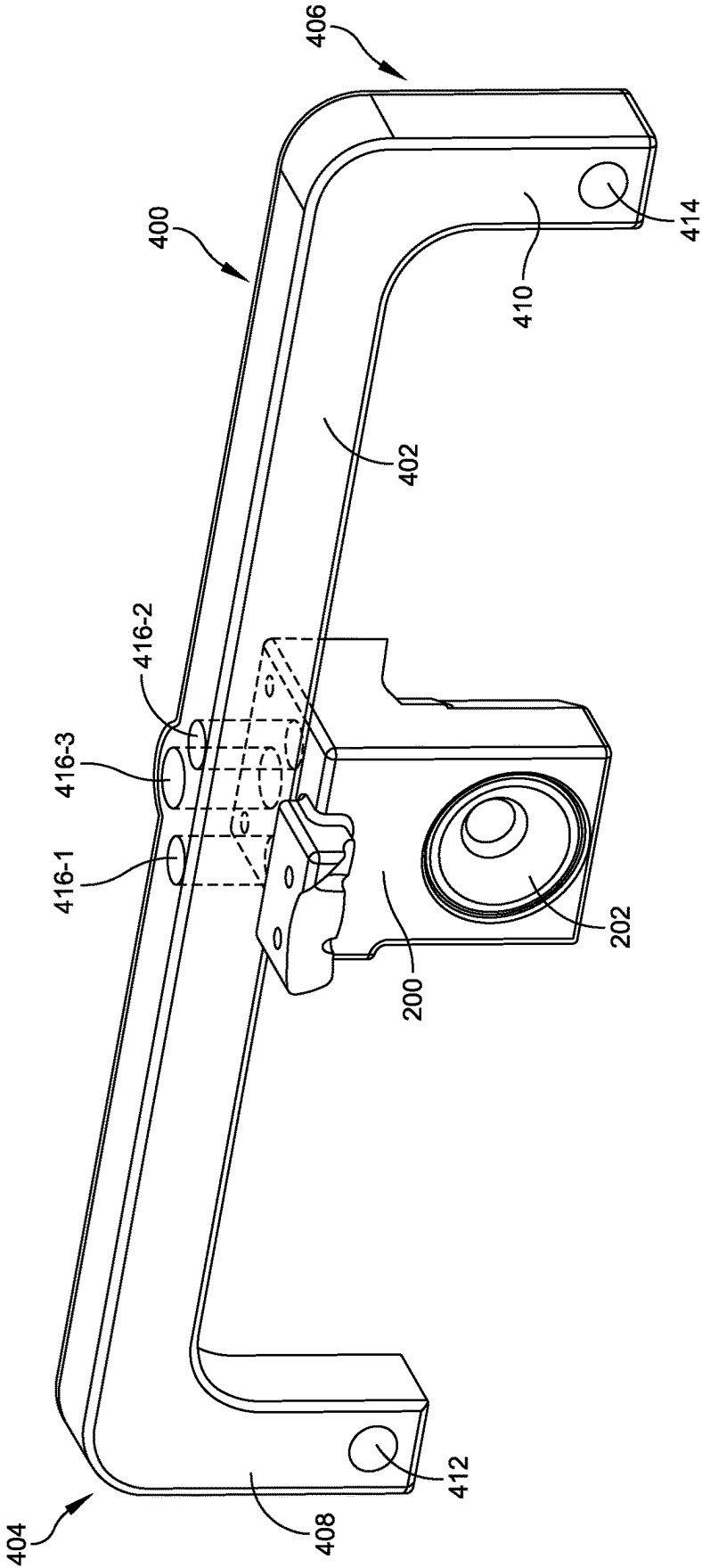


FIG. 29

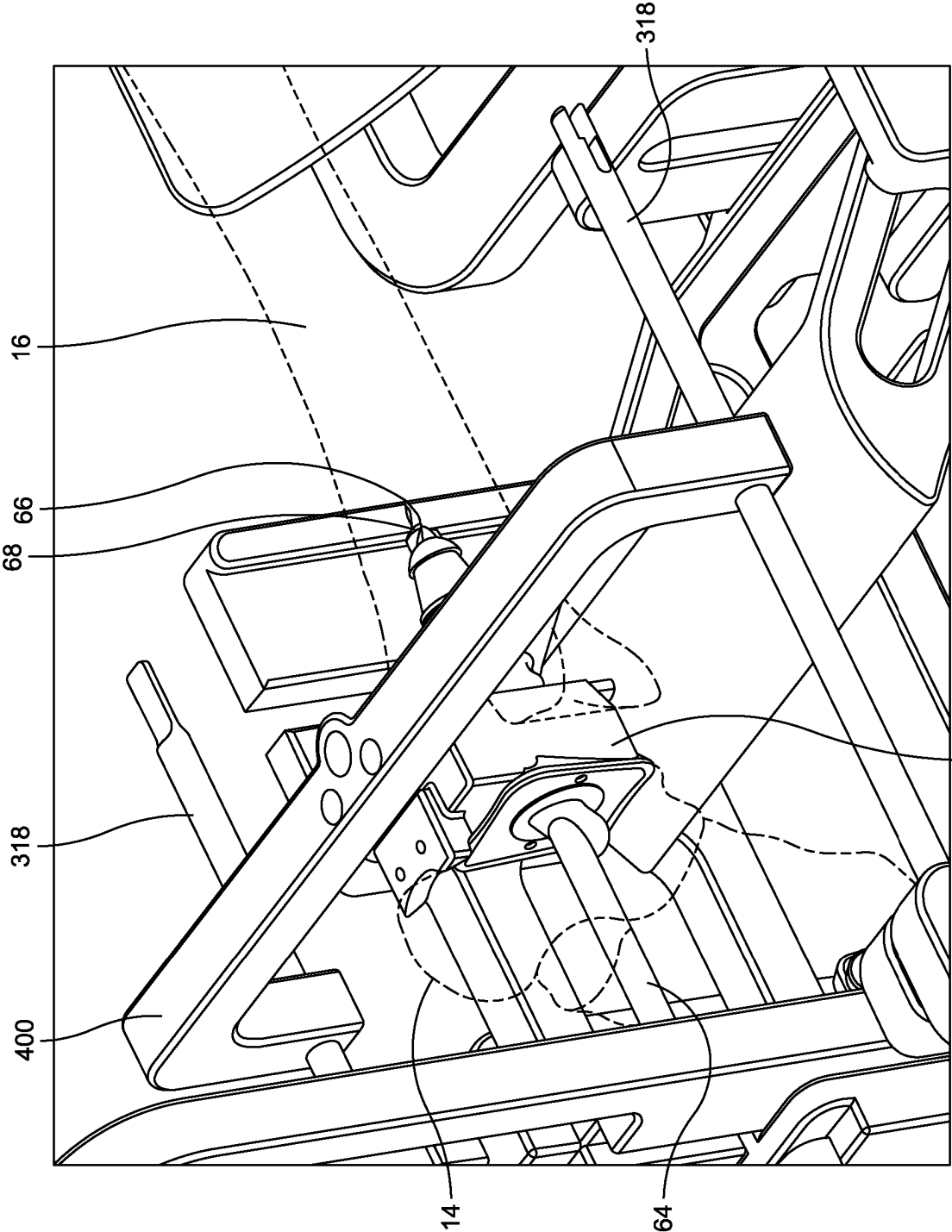


FIG. 30

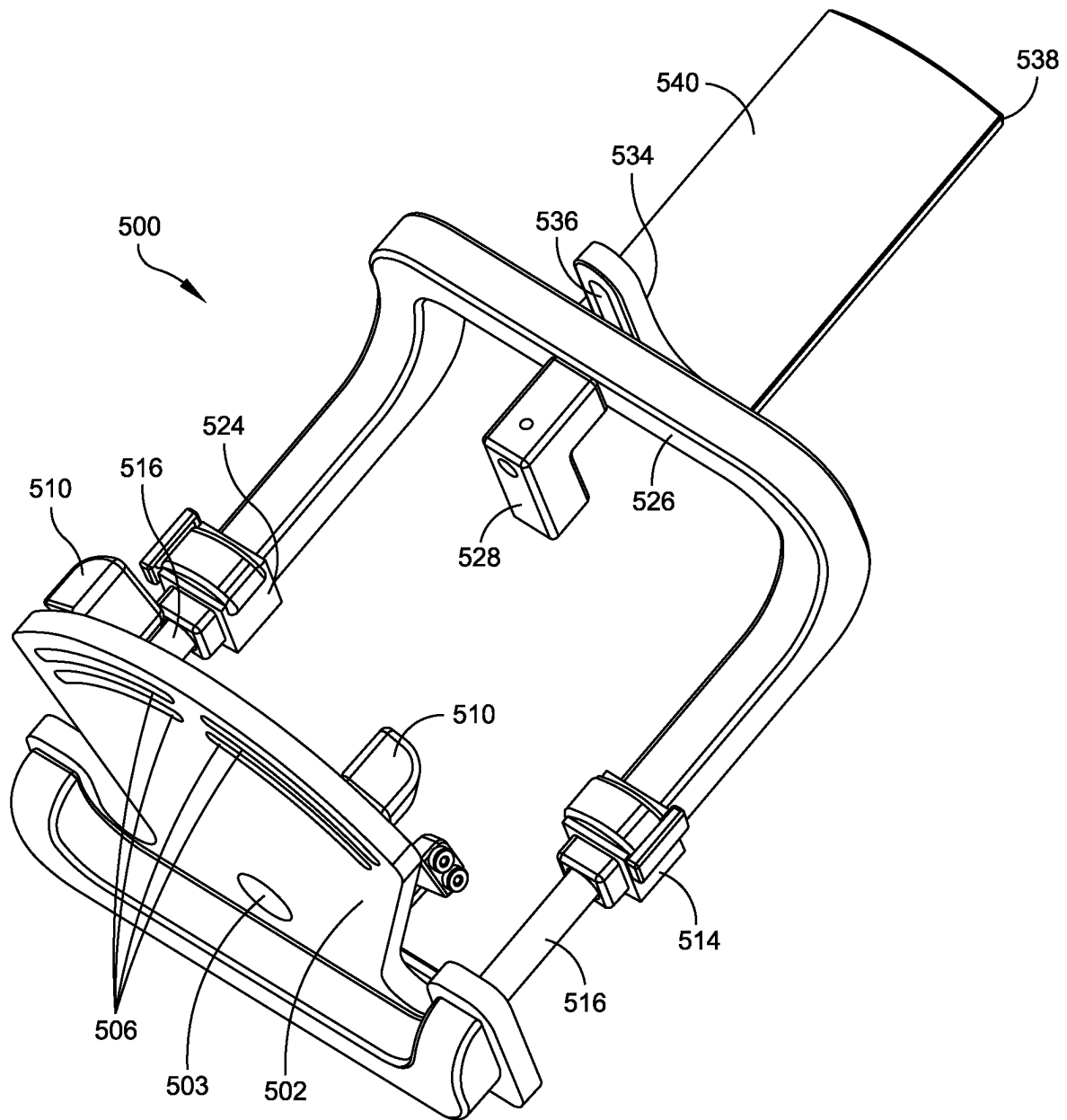


FIG. 31

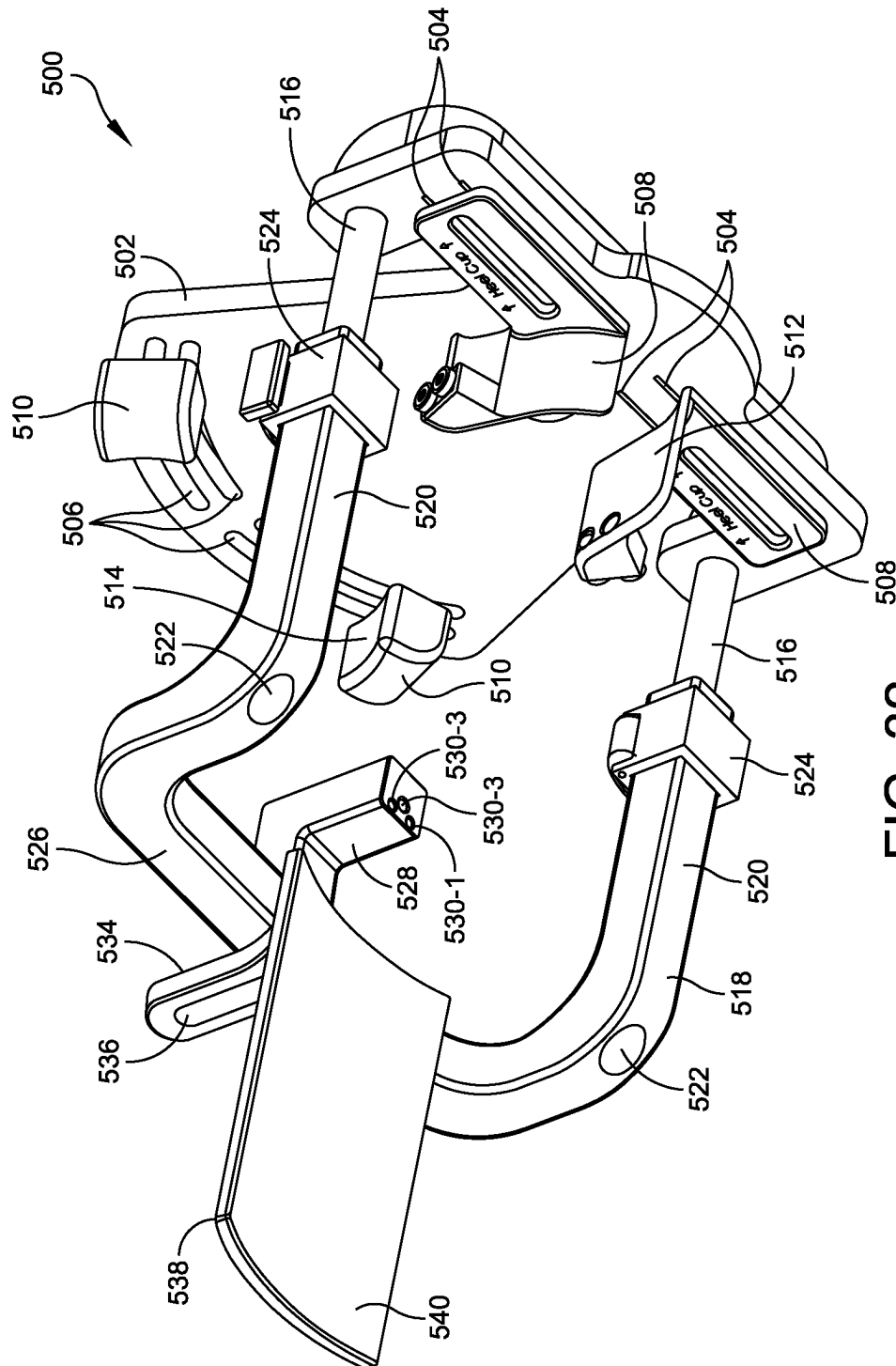


FIG. 32

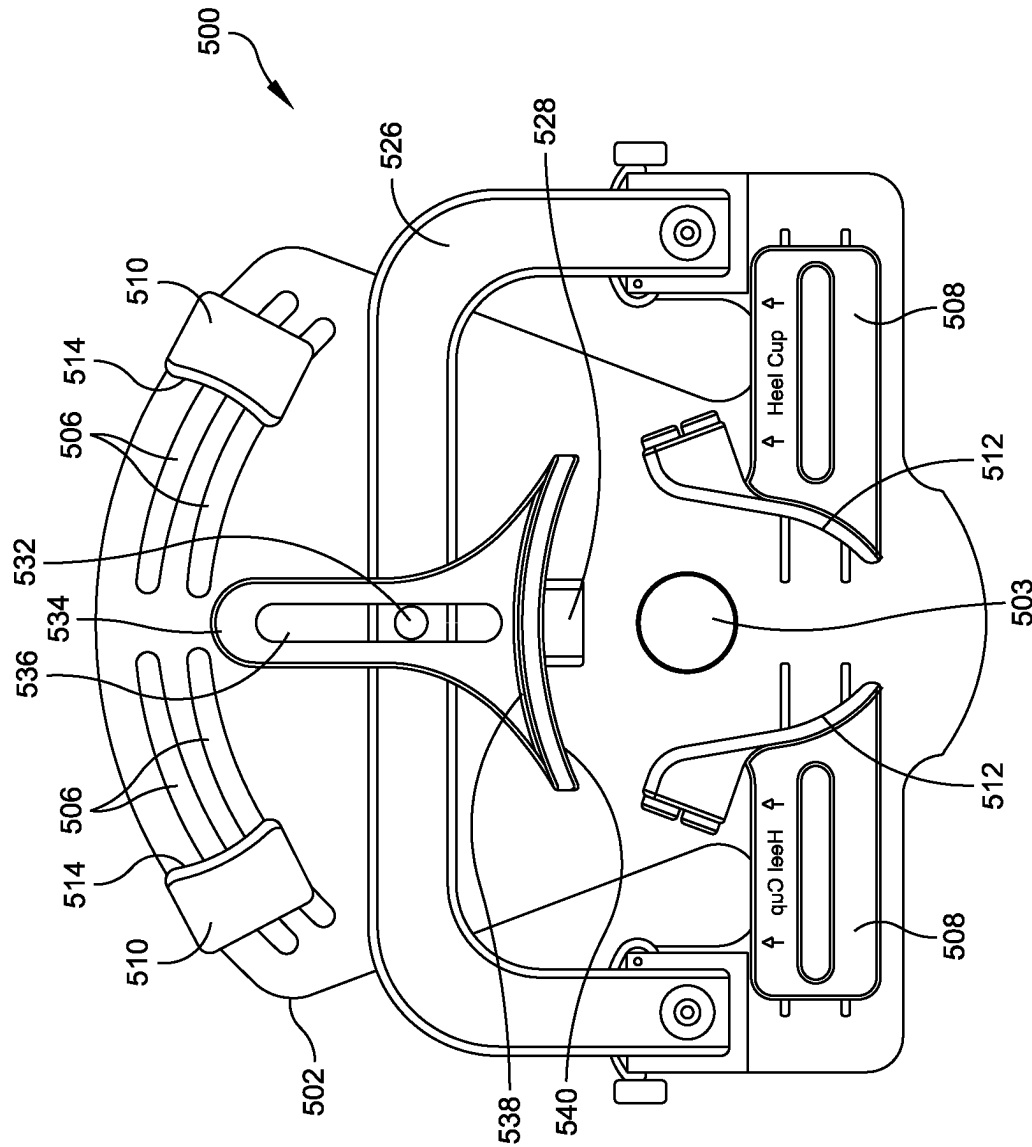


FIG. 33

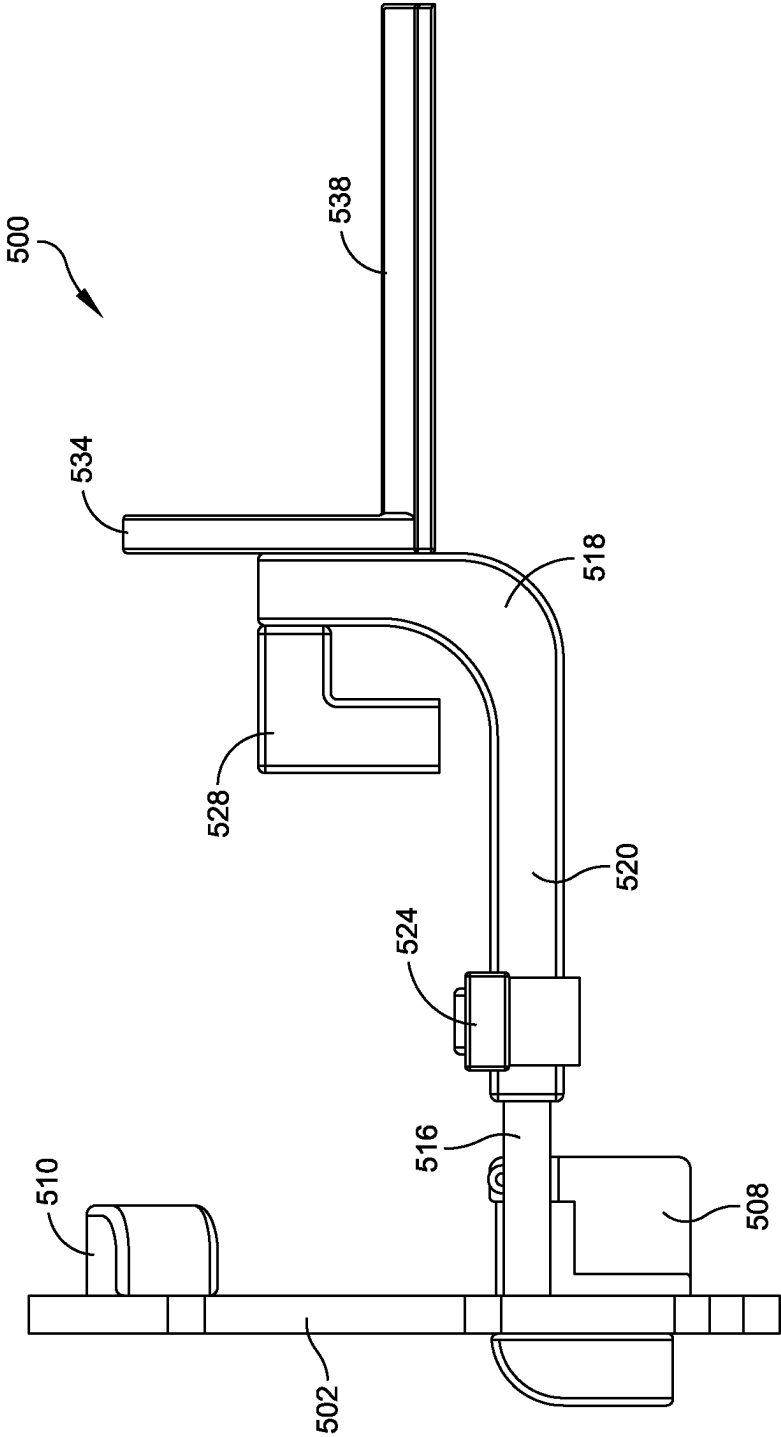
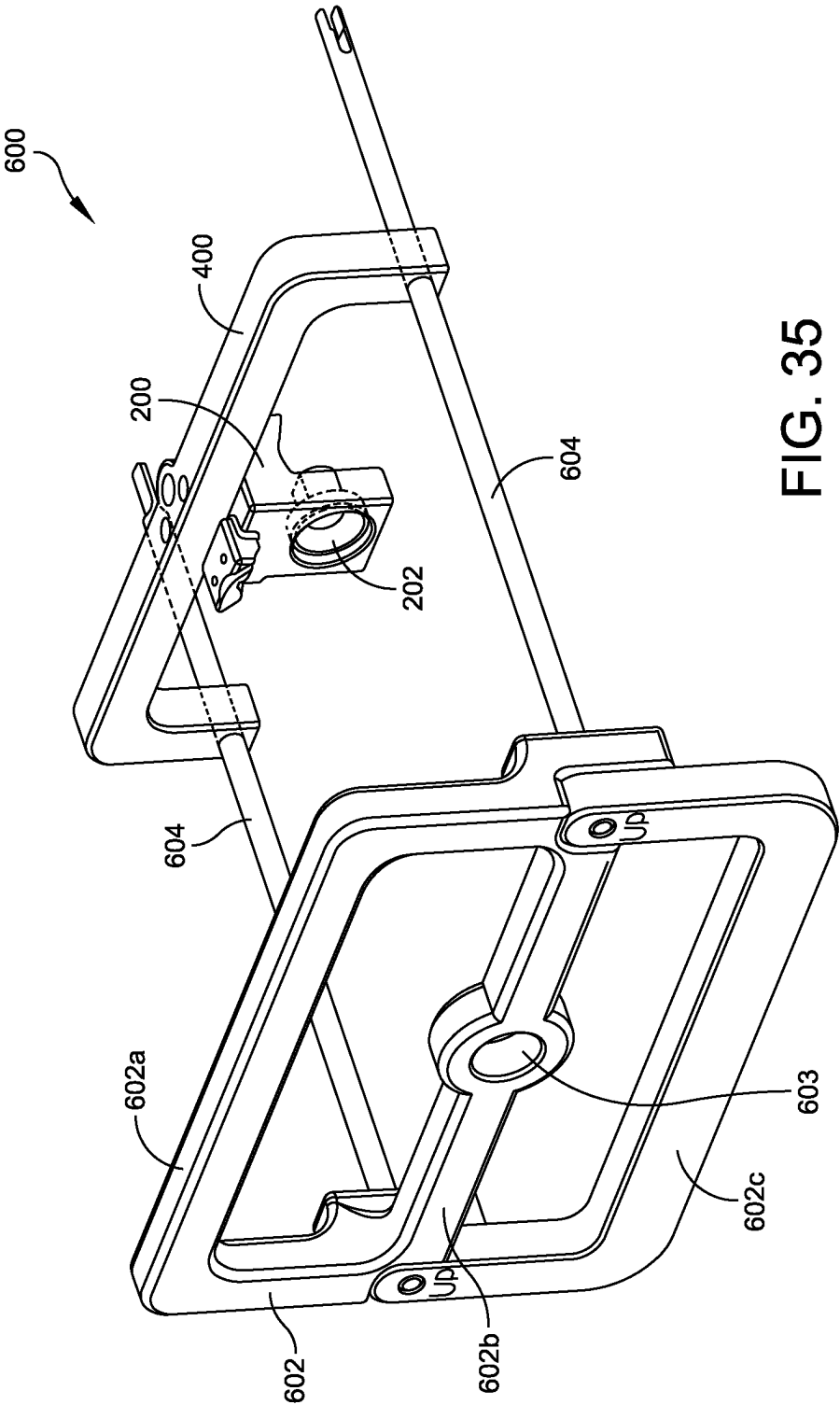


FIG. 34



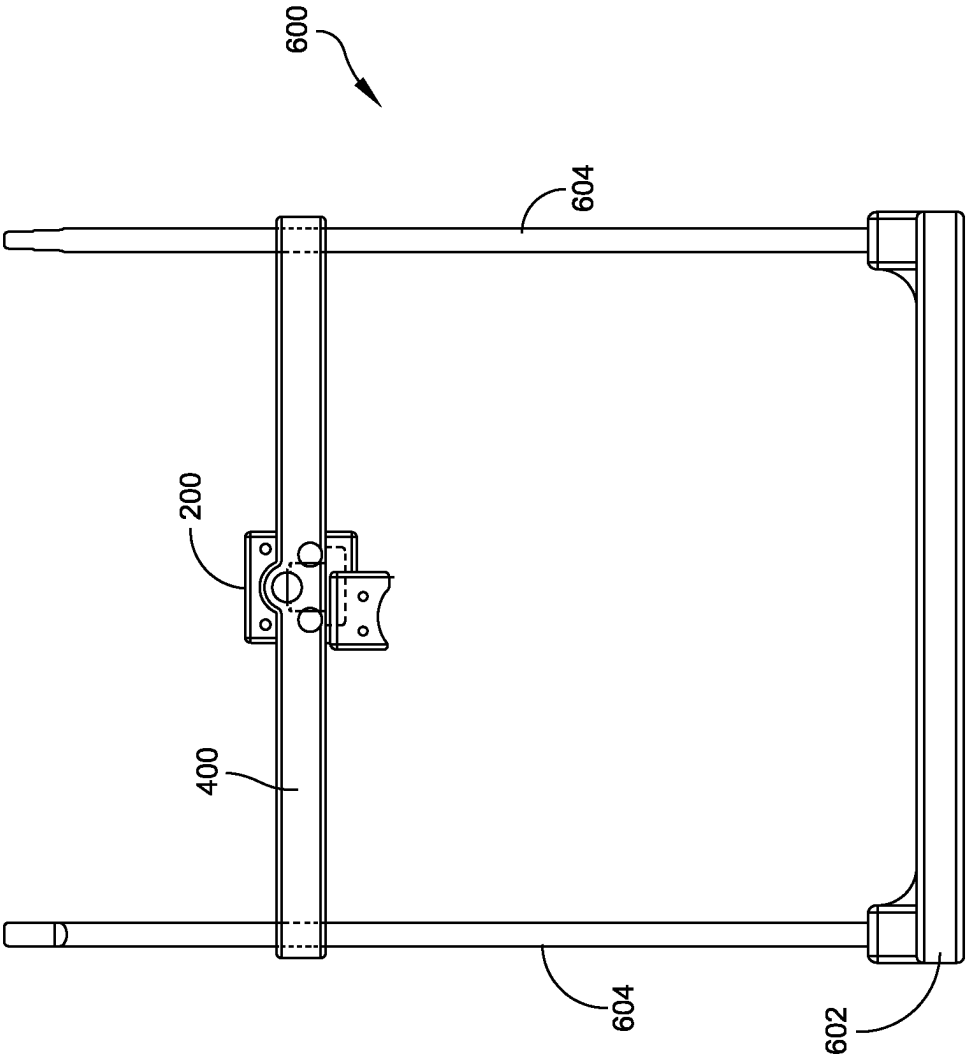


FIG. 36

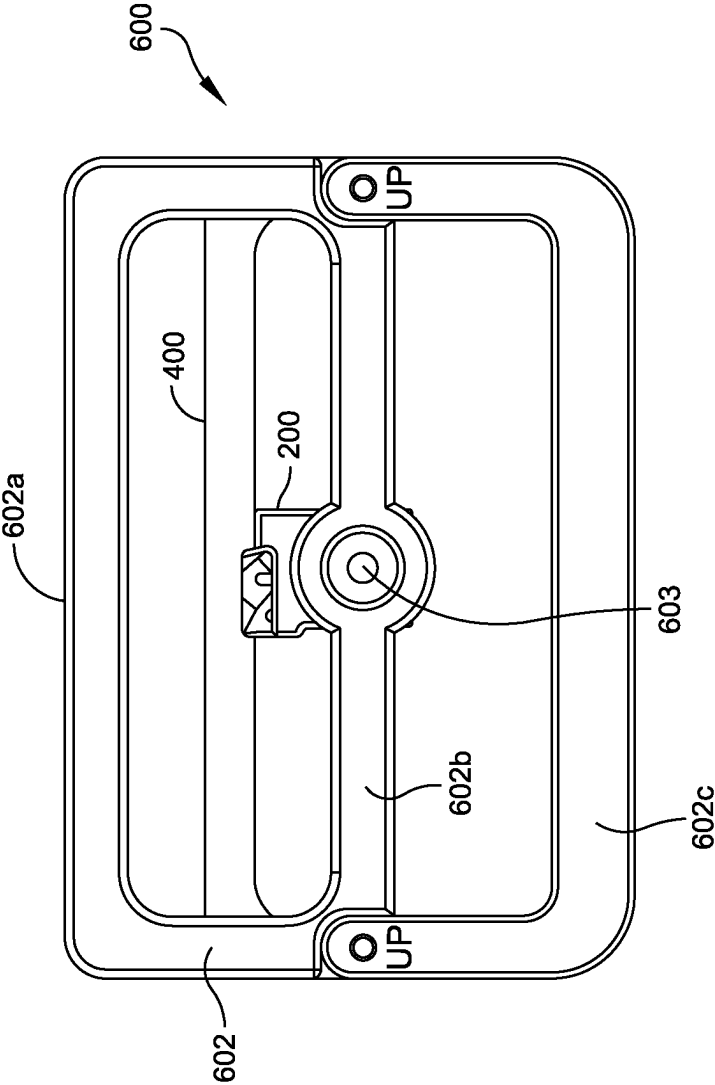


FIG. 37

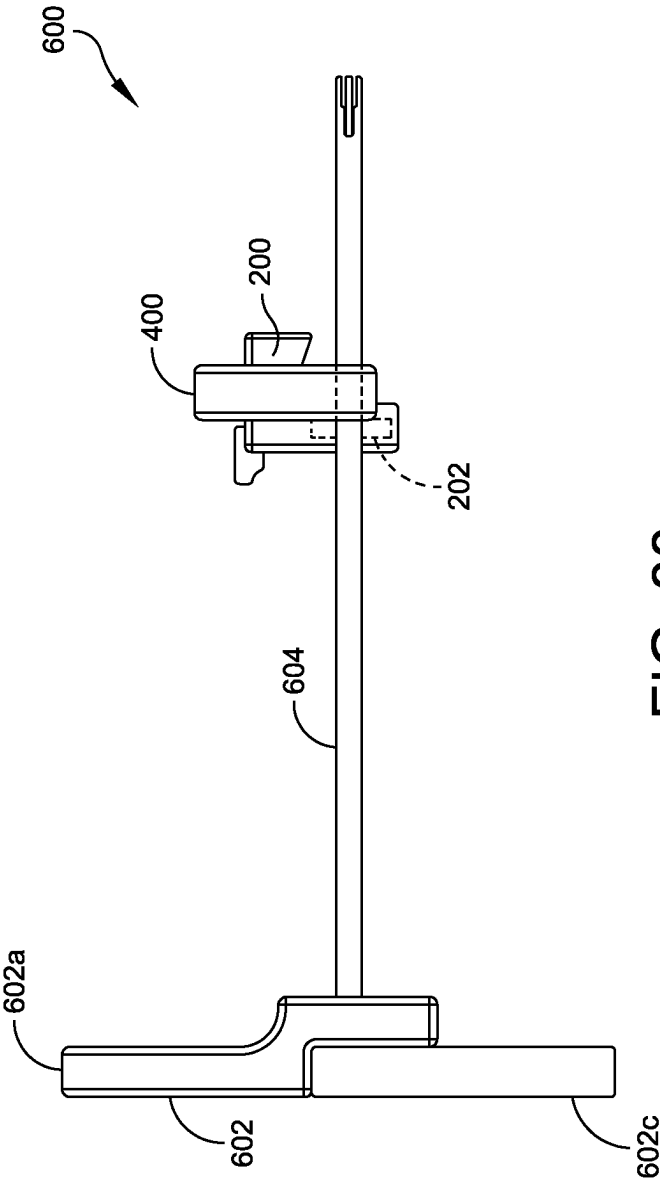


FIG. 38

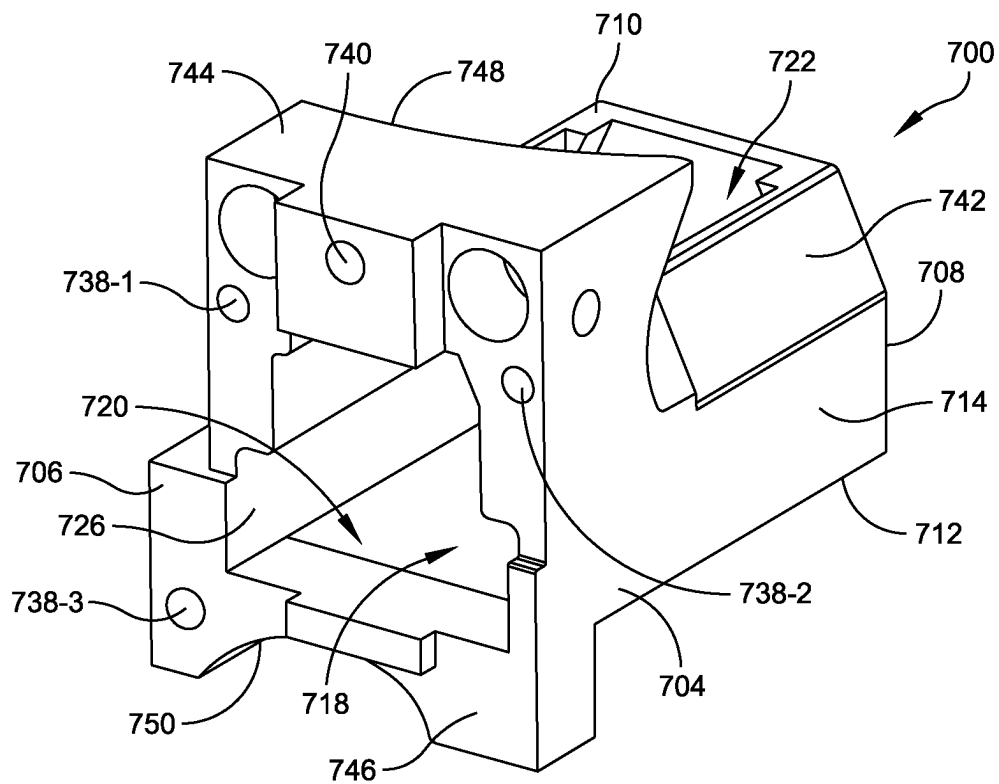


FIG. 39

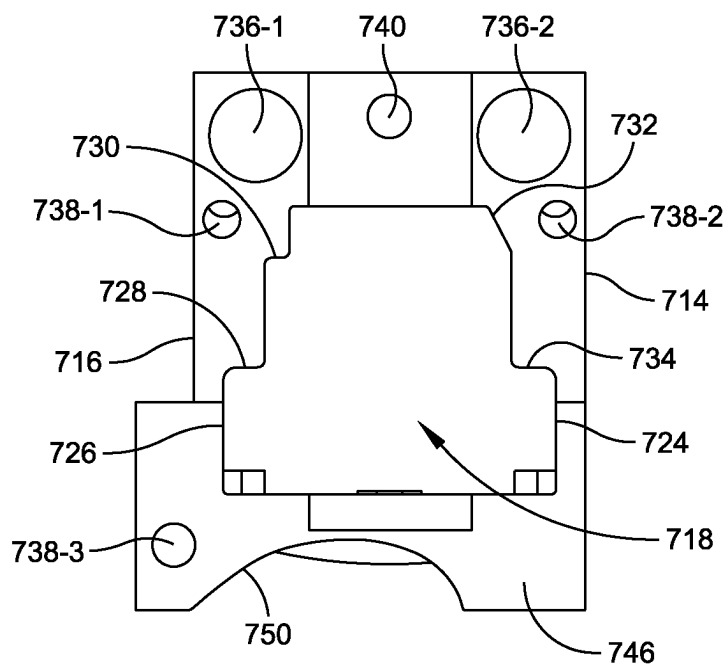


FIG. 40

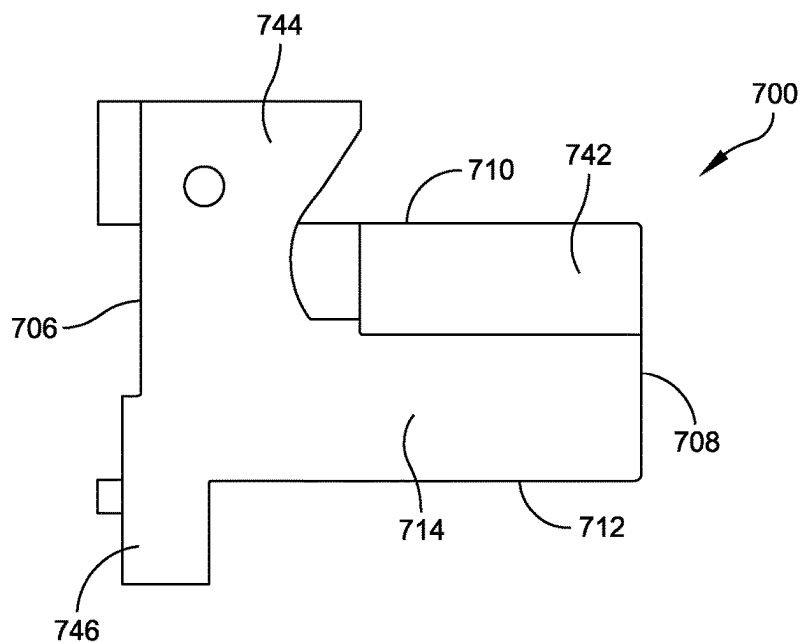


FIG. 41

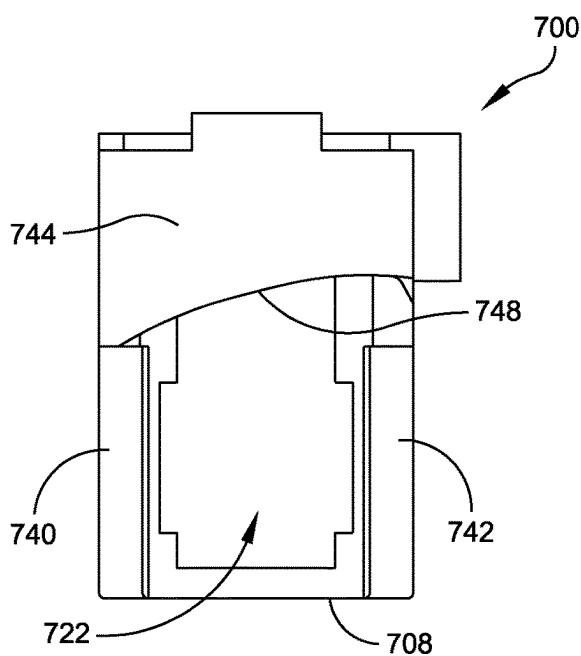


FIG. 42

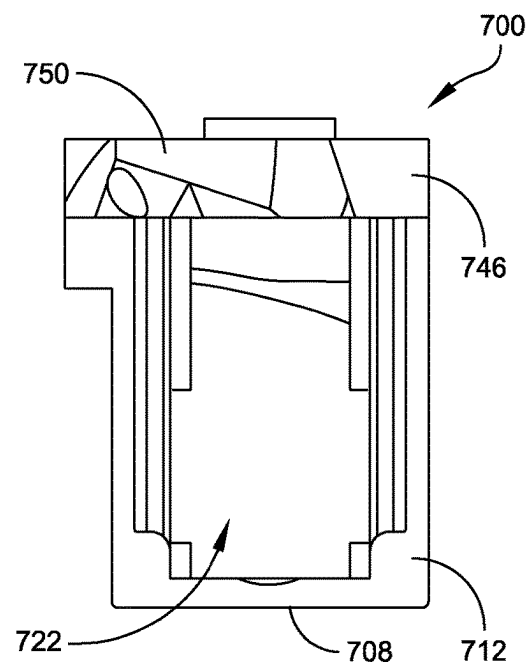


FIG. 43

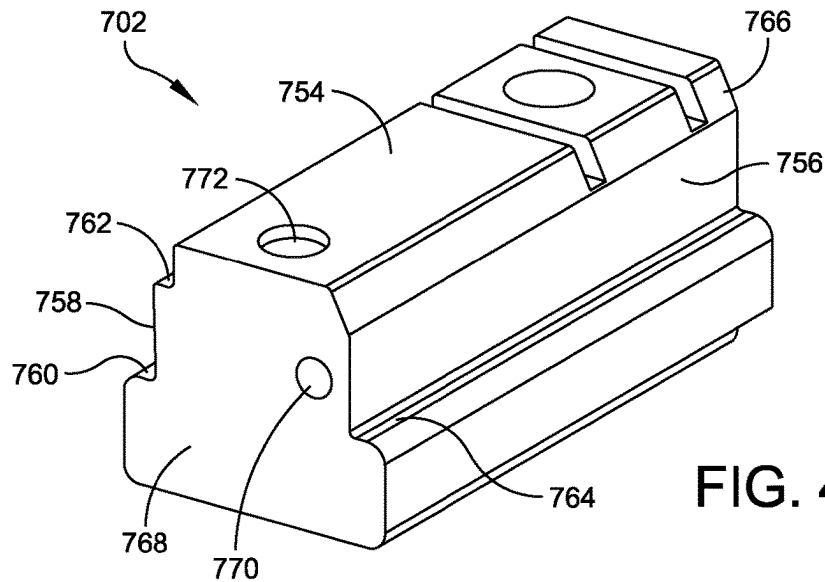


FIG. 44

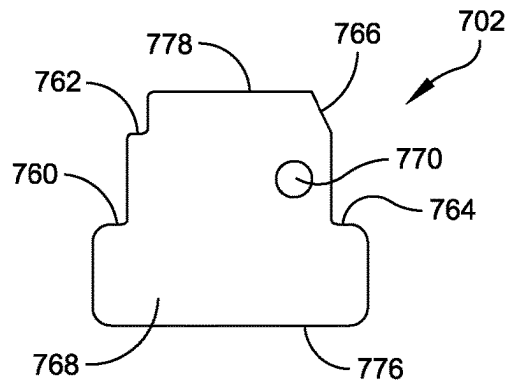


FIG. 45

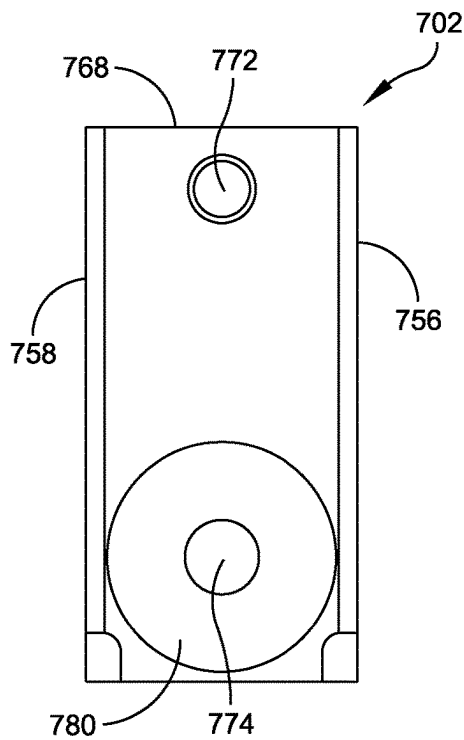


FIG. 46

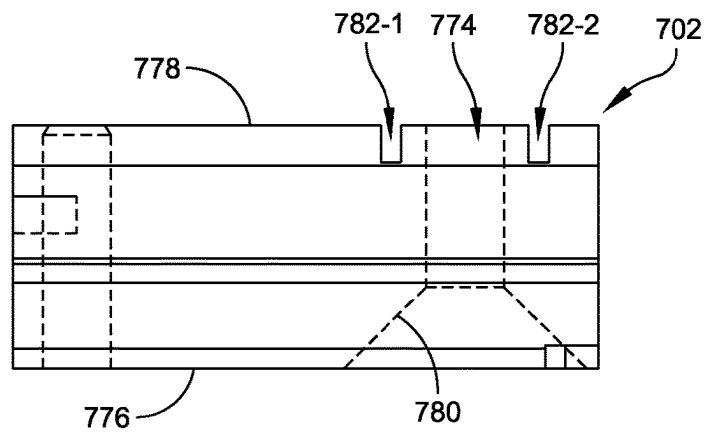
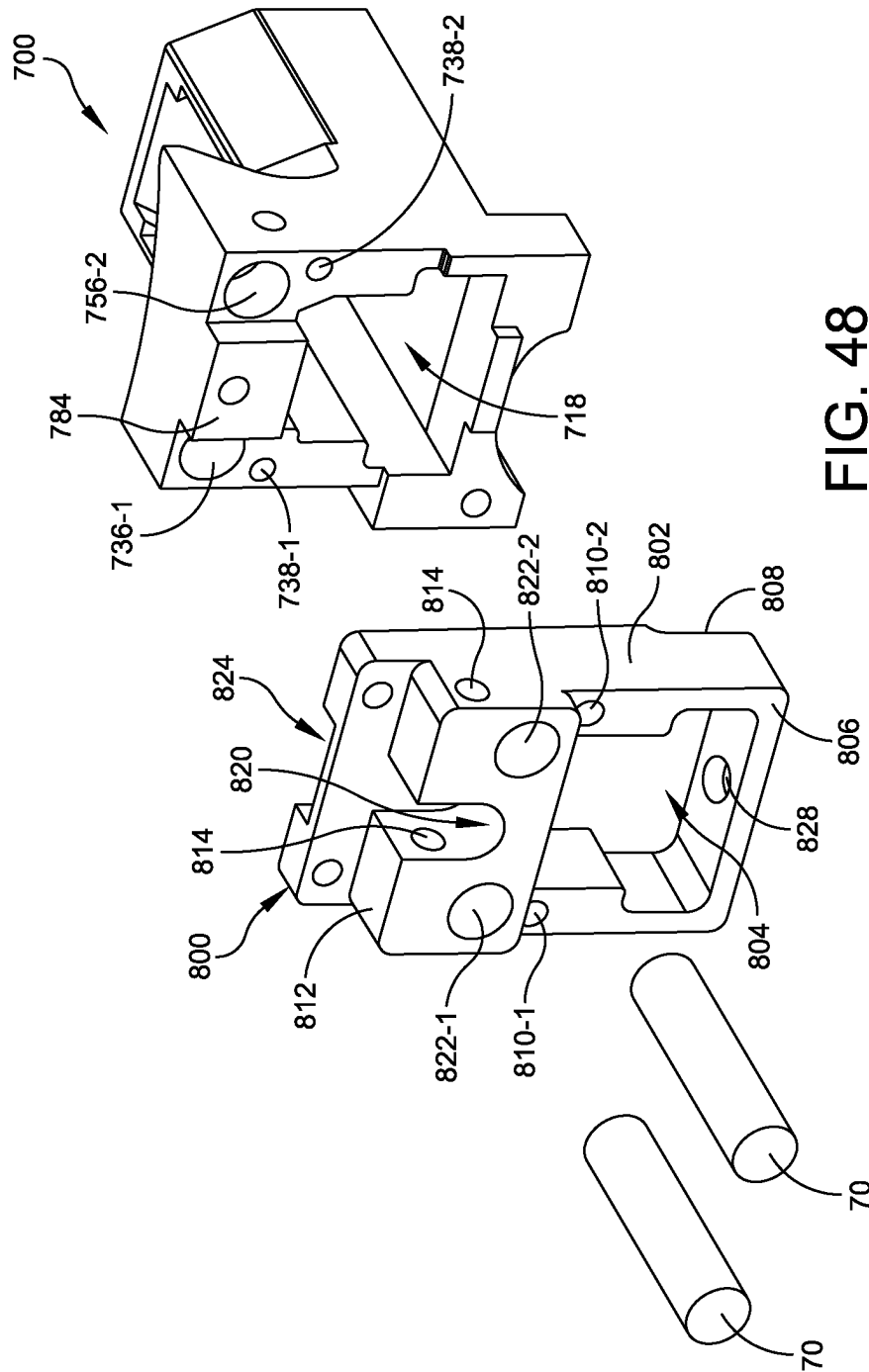


FIG. 47



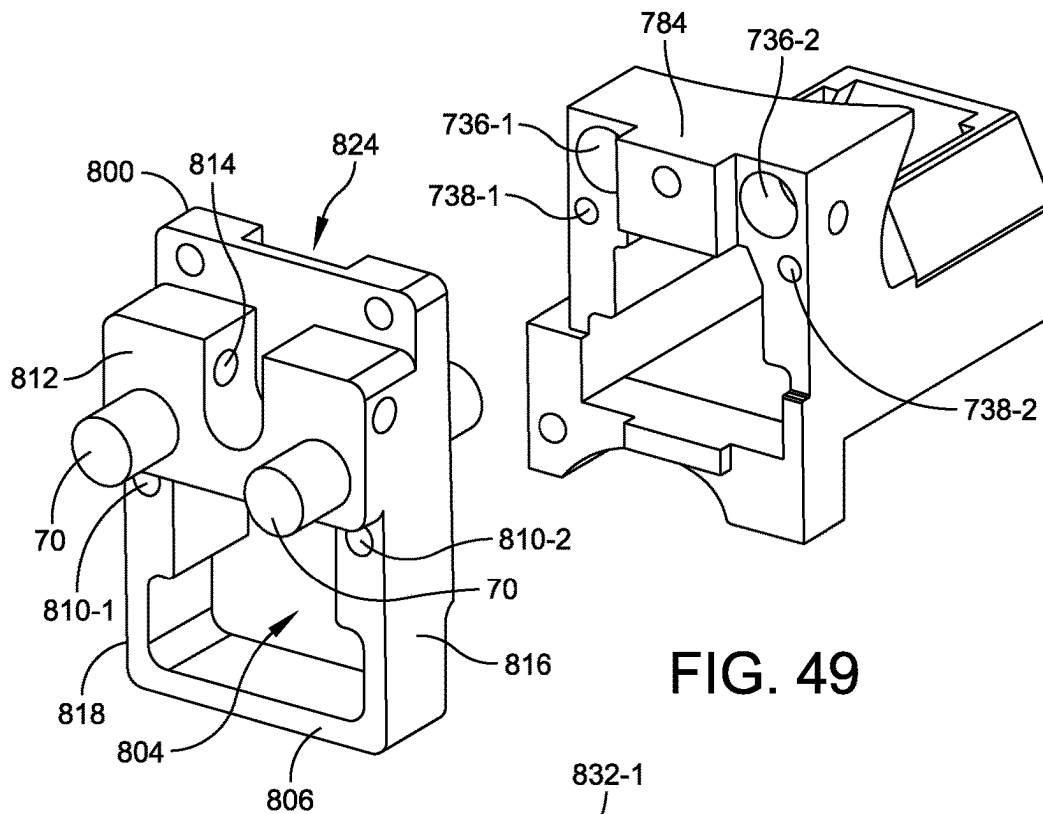


FIG. 49

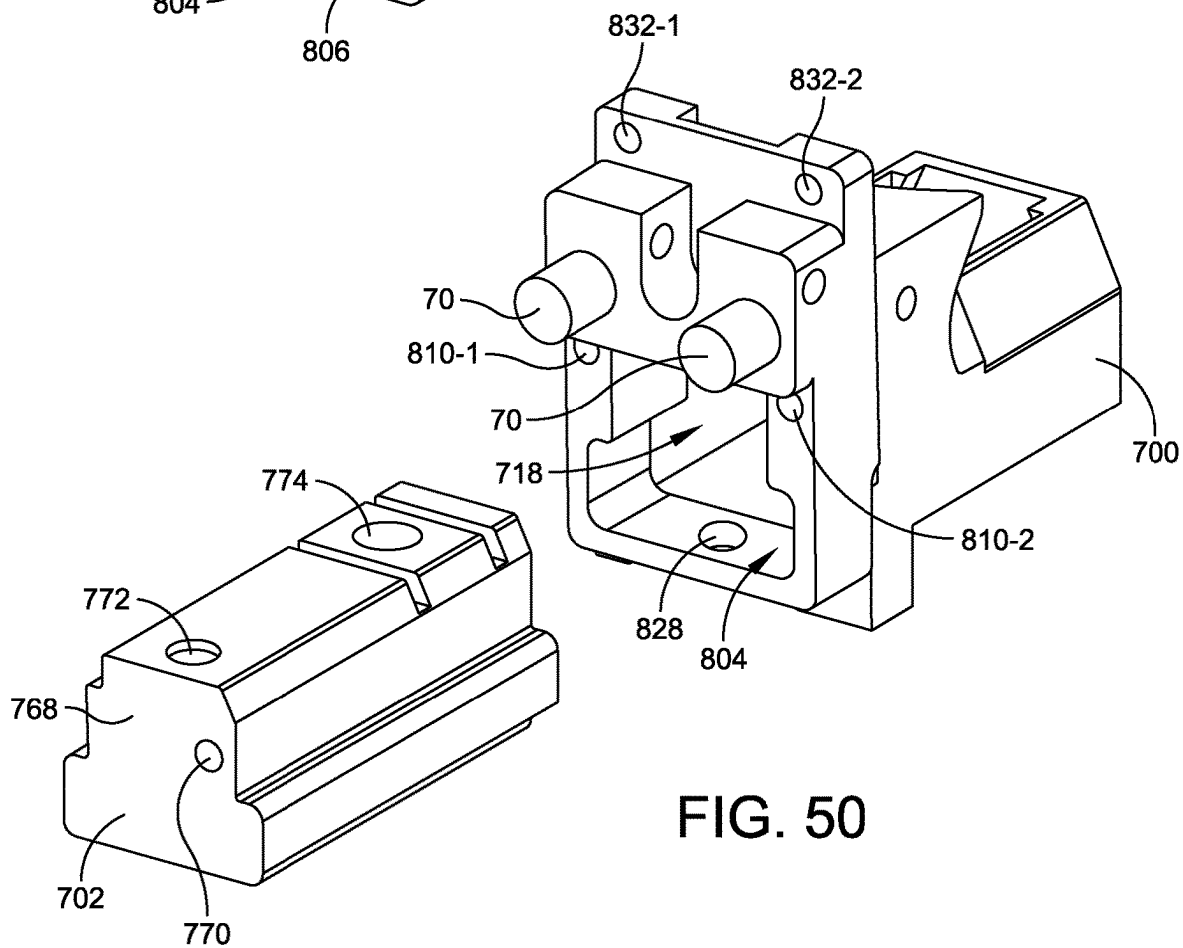


FIG. 50

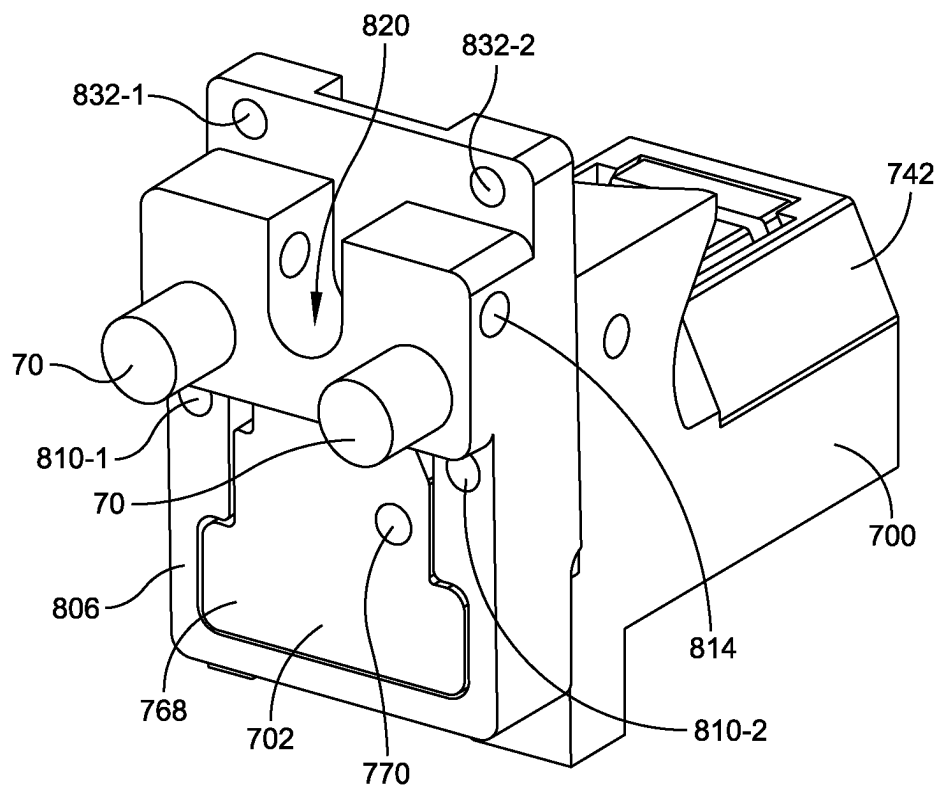


FIG. 51

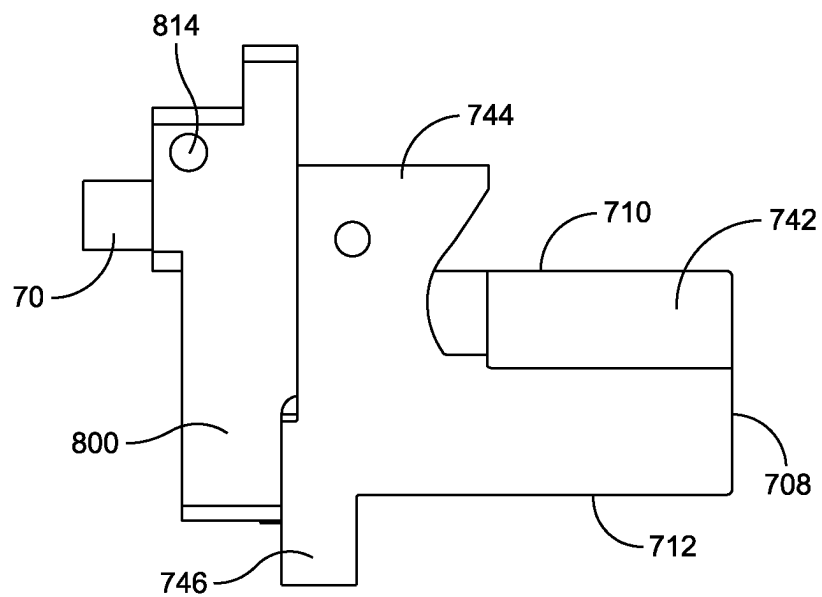


FIG. 52

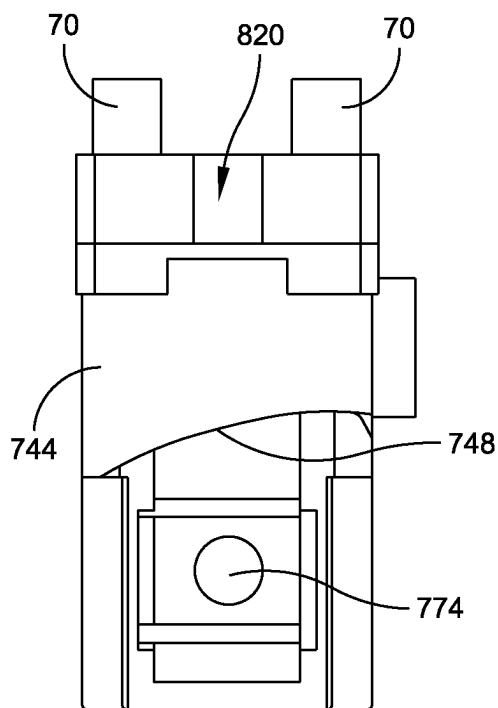


FIG. 53

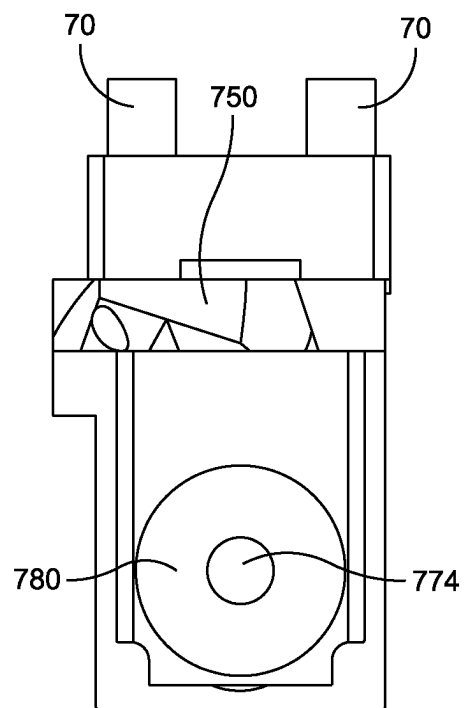


FIG. 54

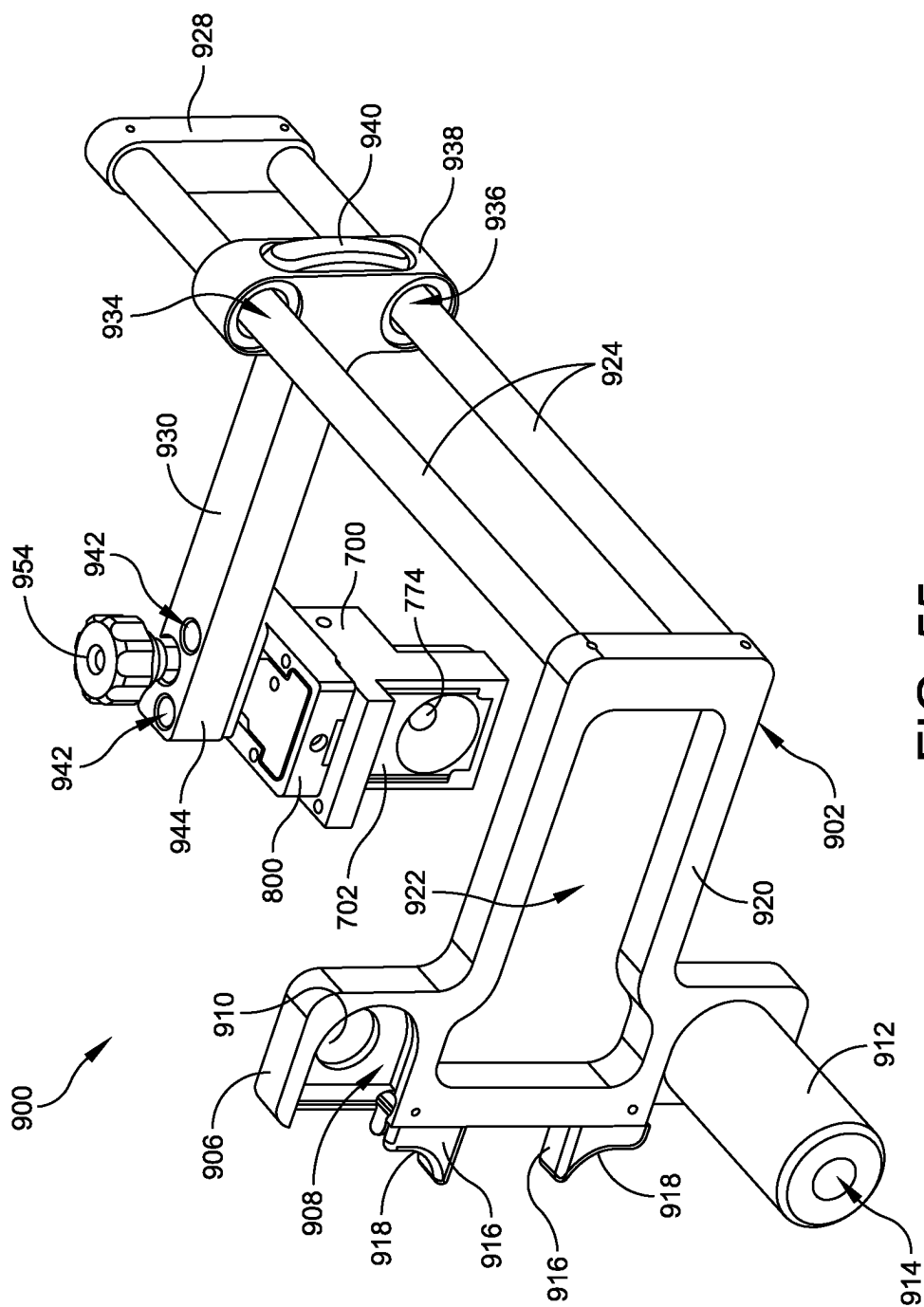
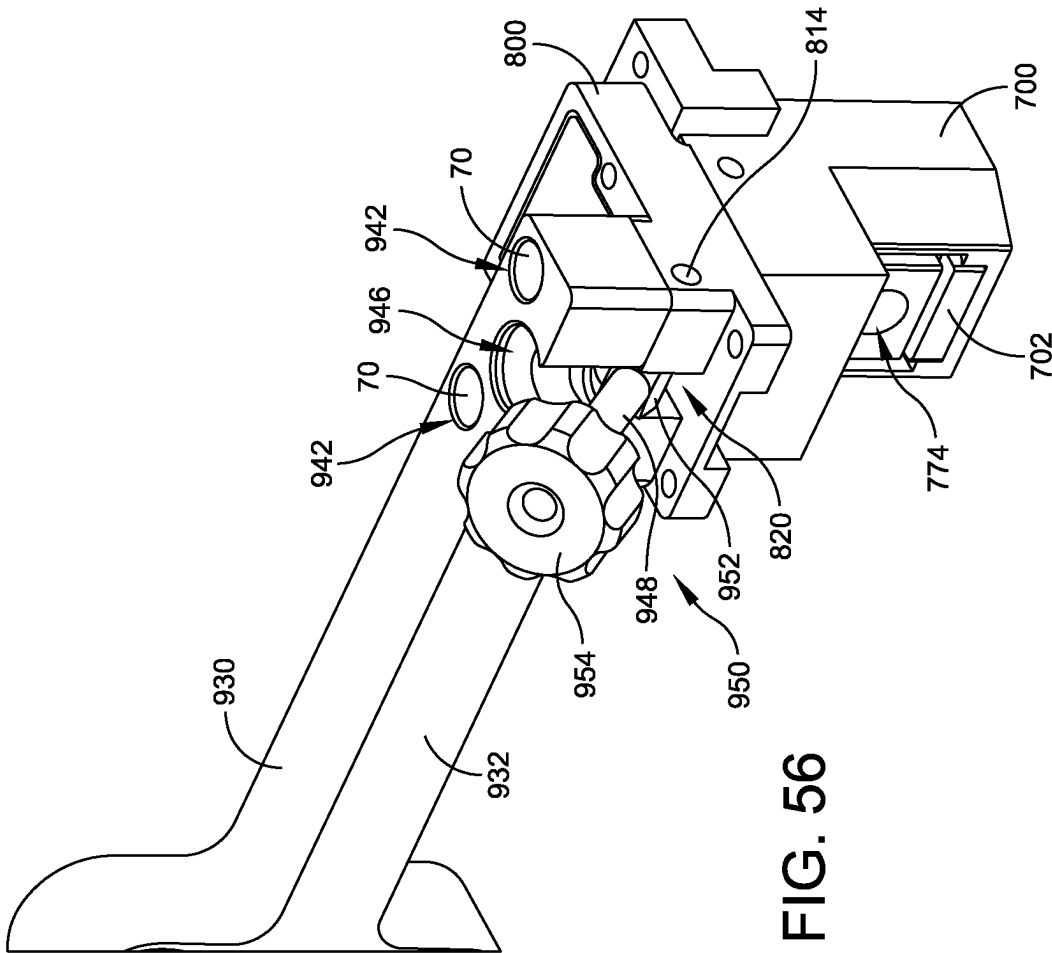


FIG. 55



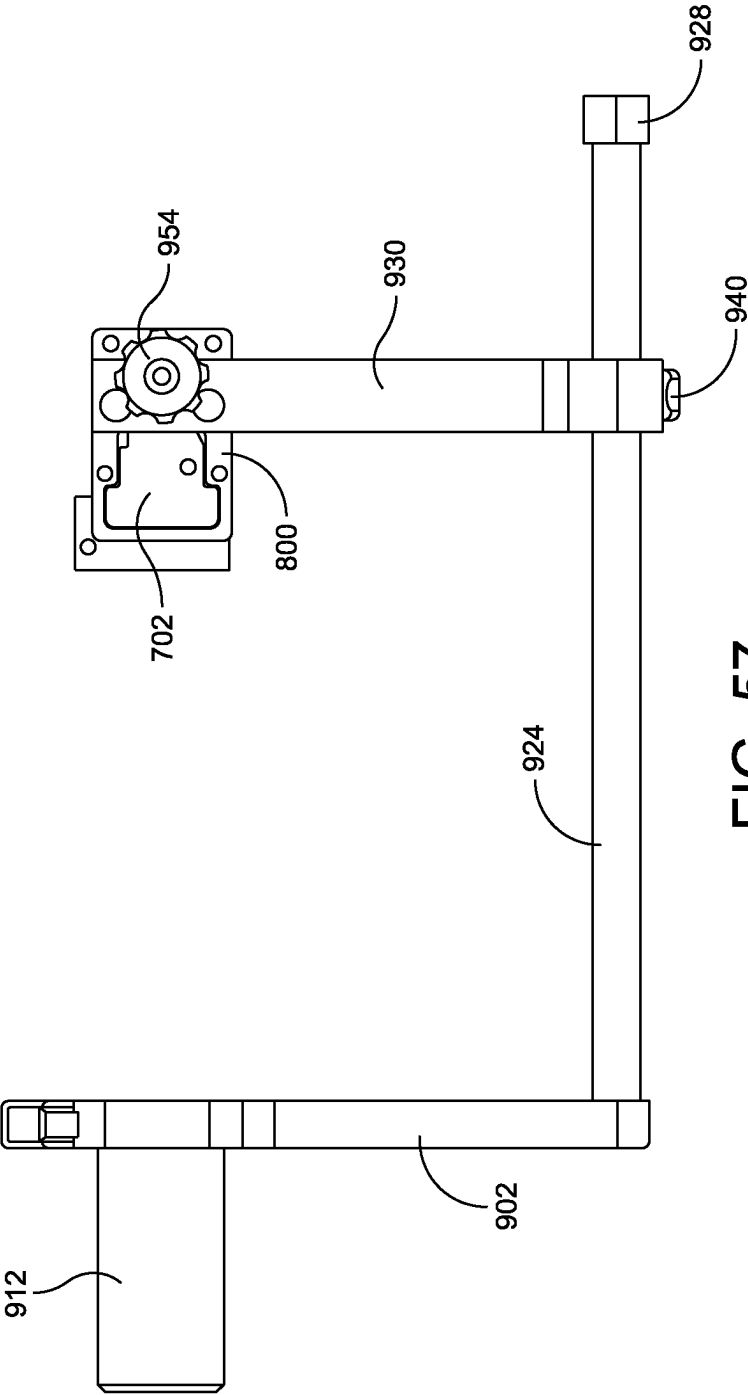


FIG. 57

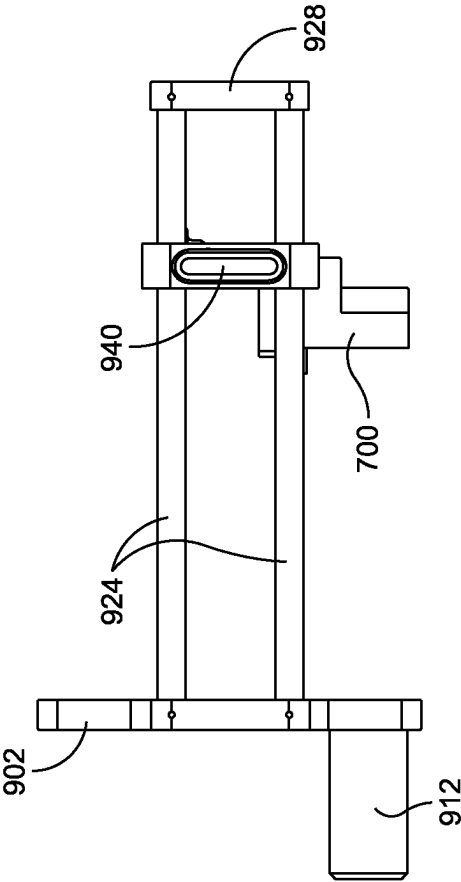


FIG. 58

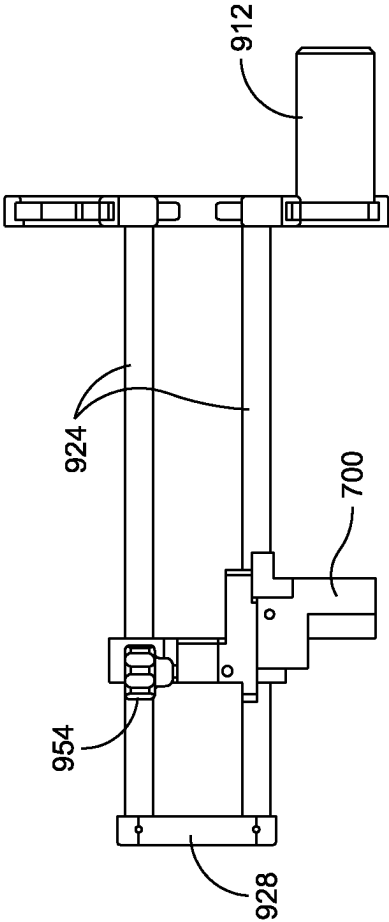


FIG. 59

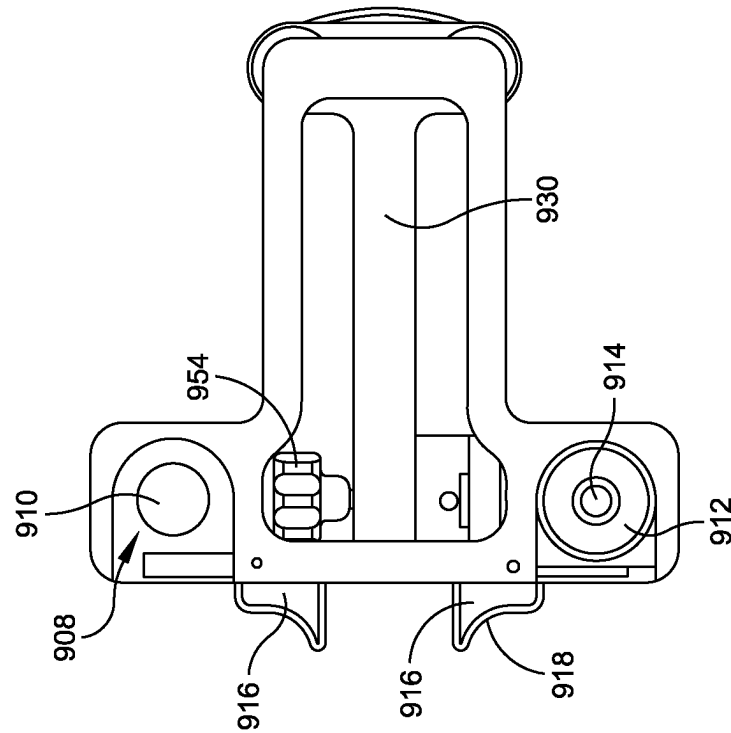


FIG. 61

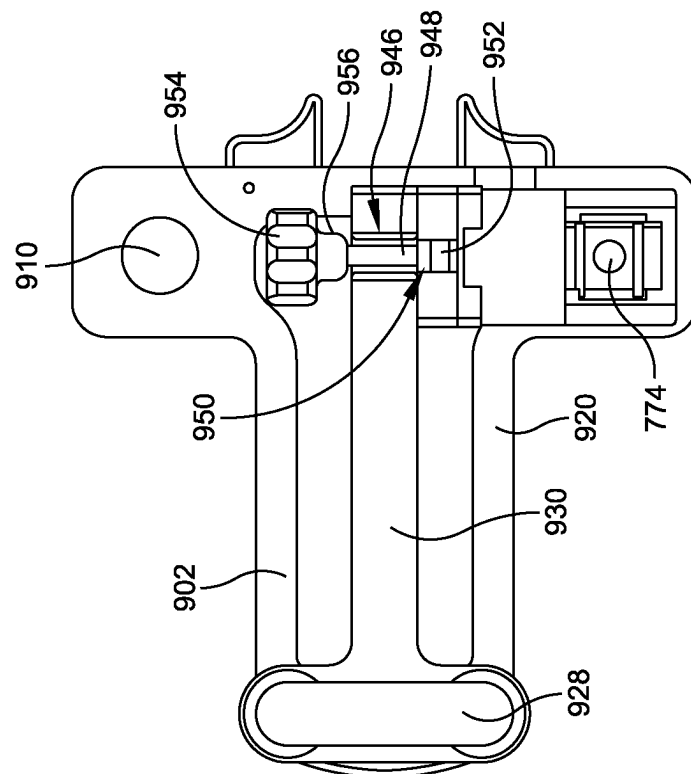


FIG. 60

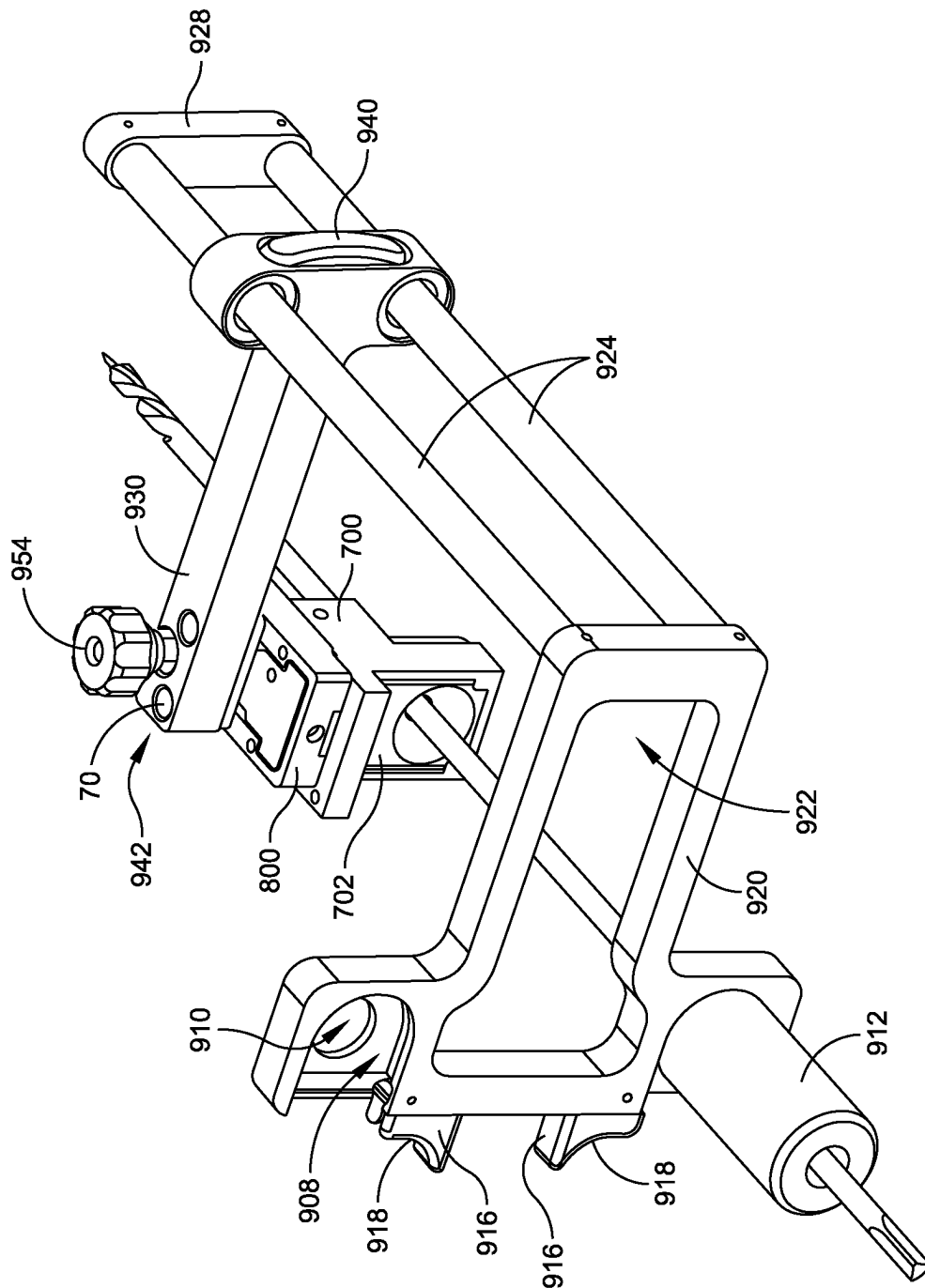


FIG. 62

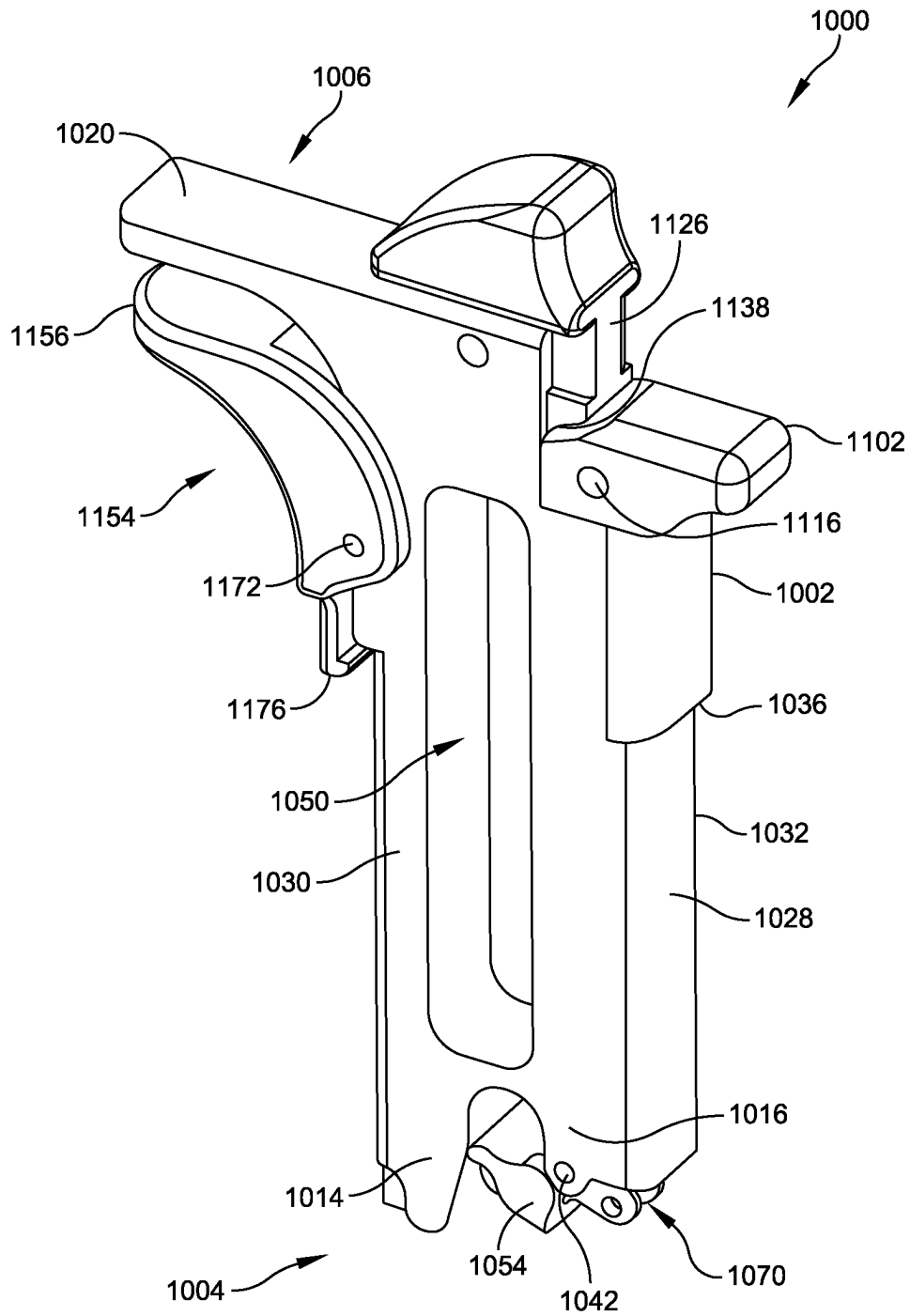


FIG. 63

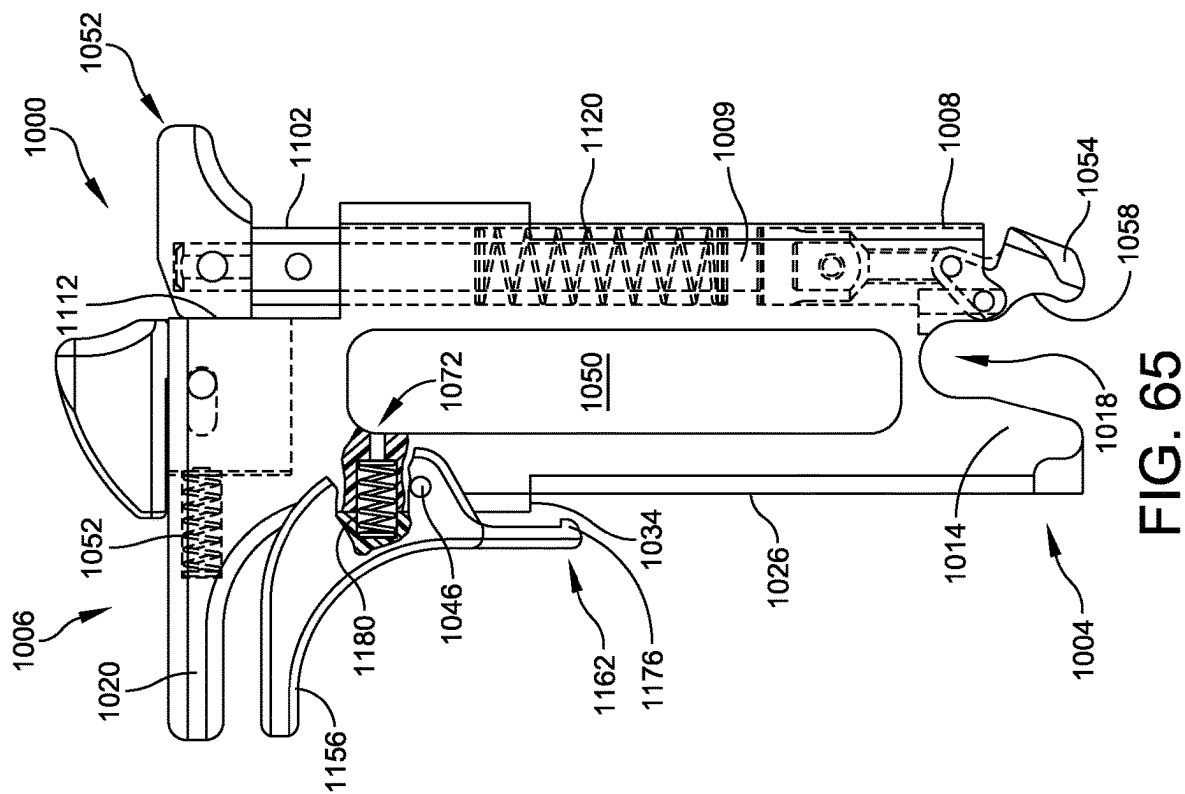


FIG. 65

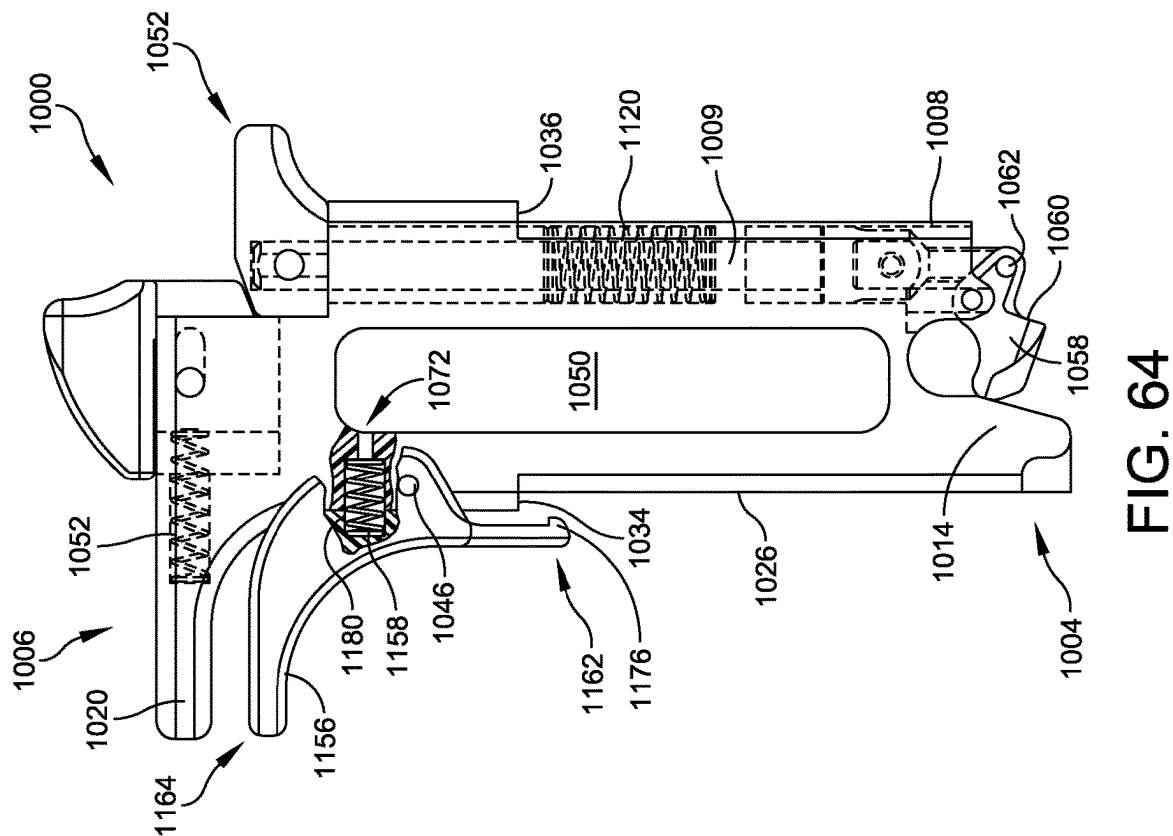


FIG. 64

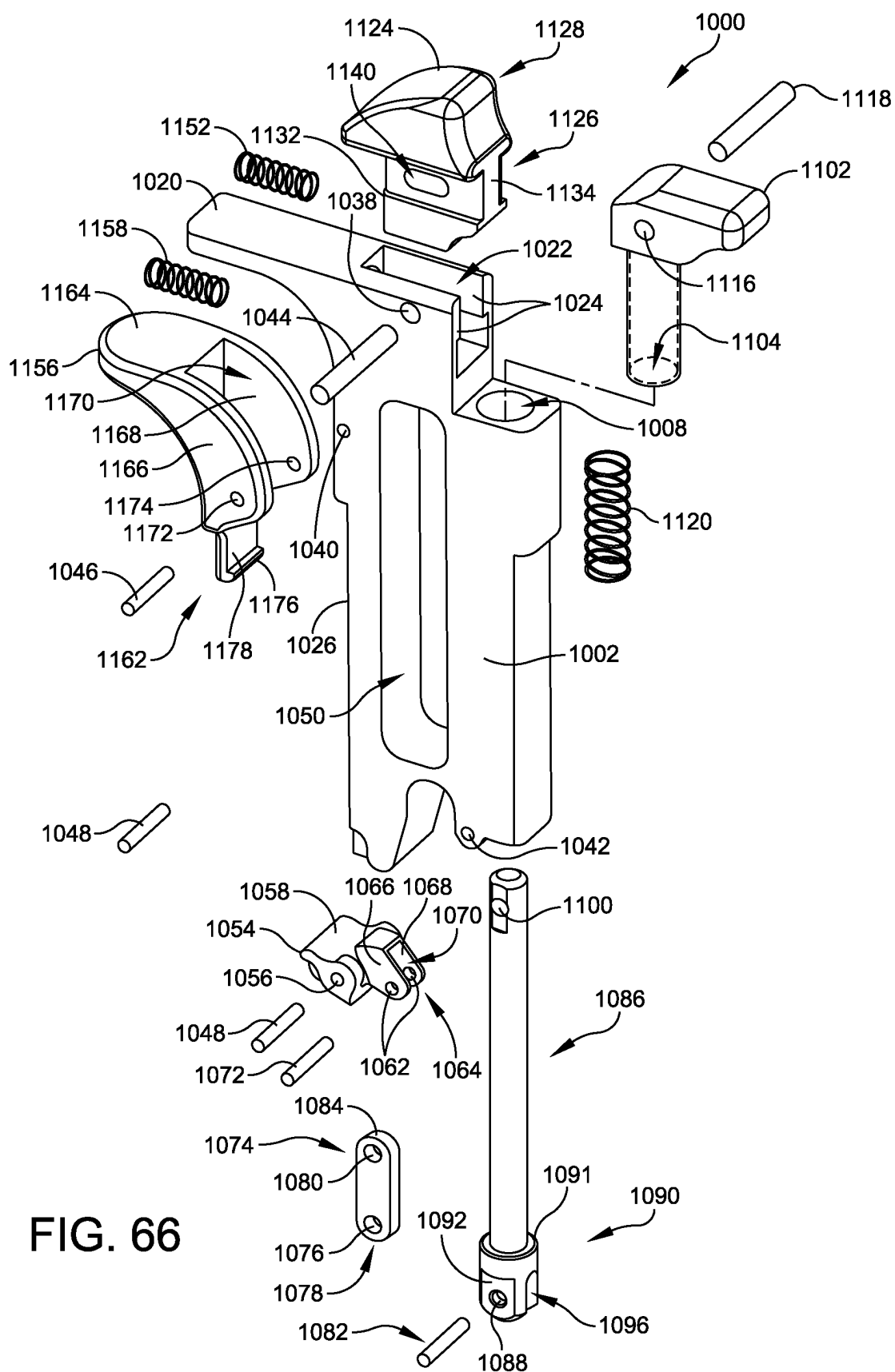


FIG. 66

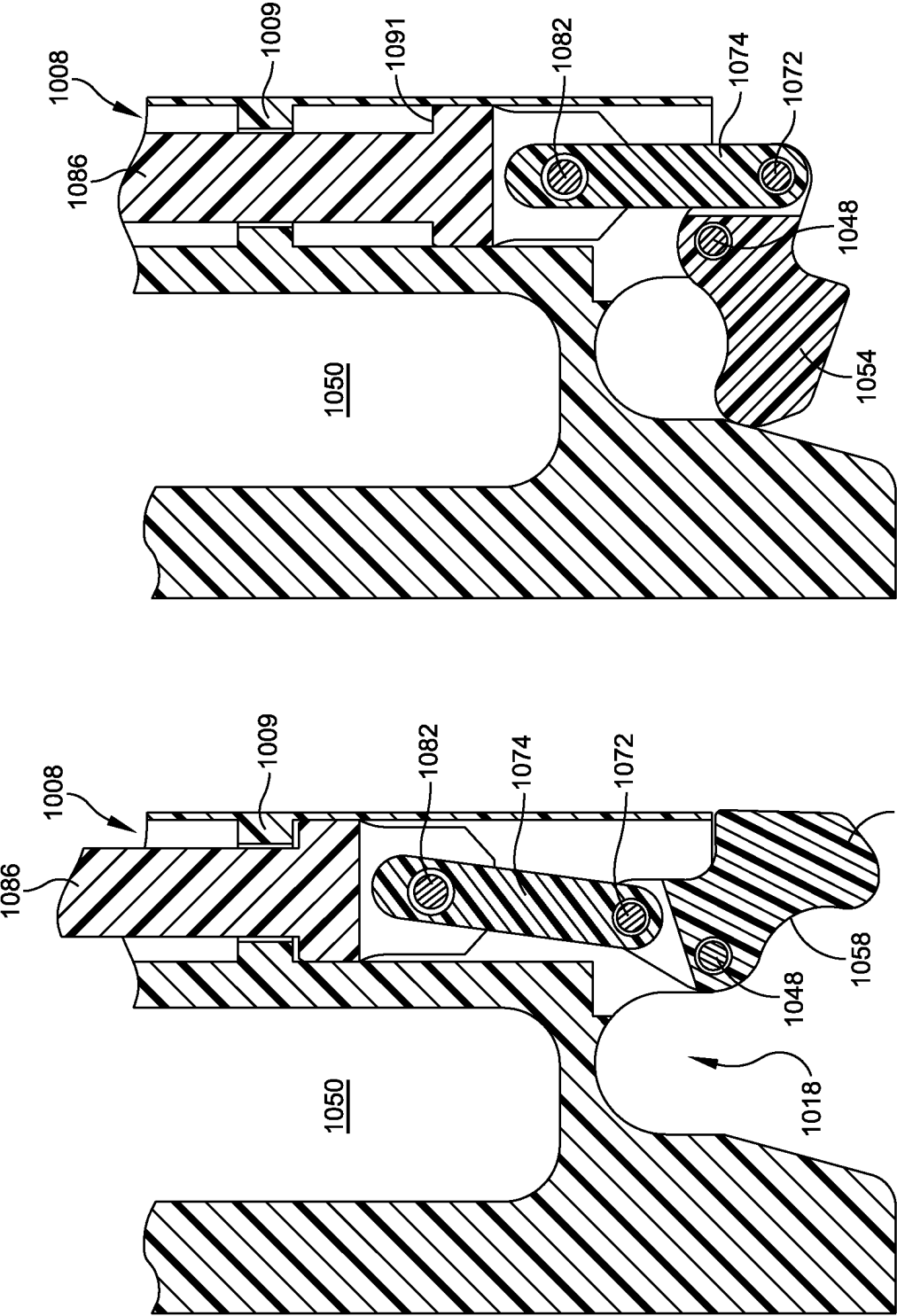


FIG. 68

FIG. 67

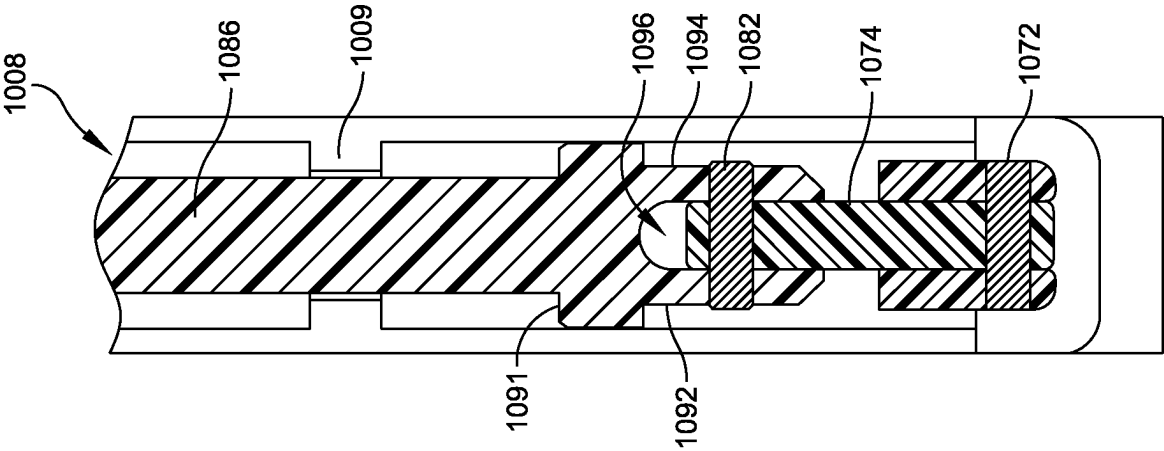


FIG. 69

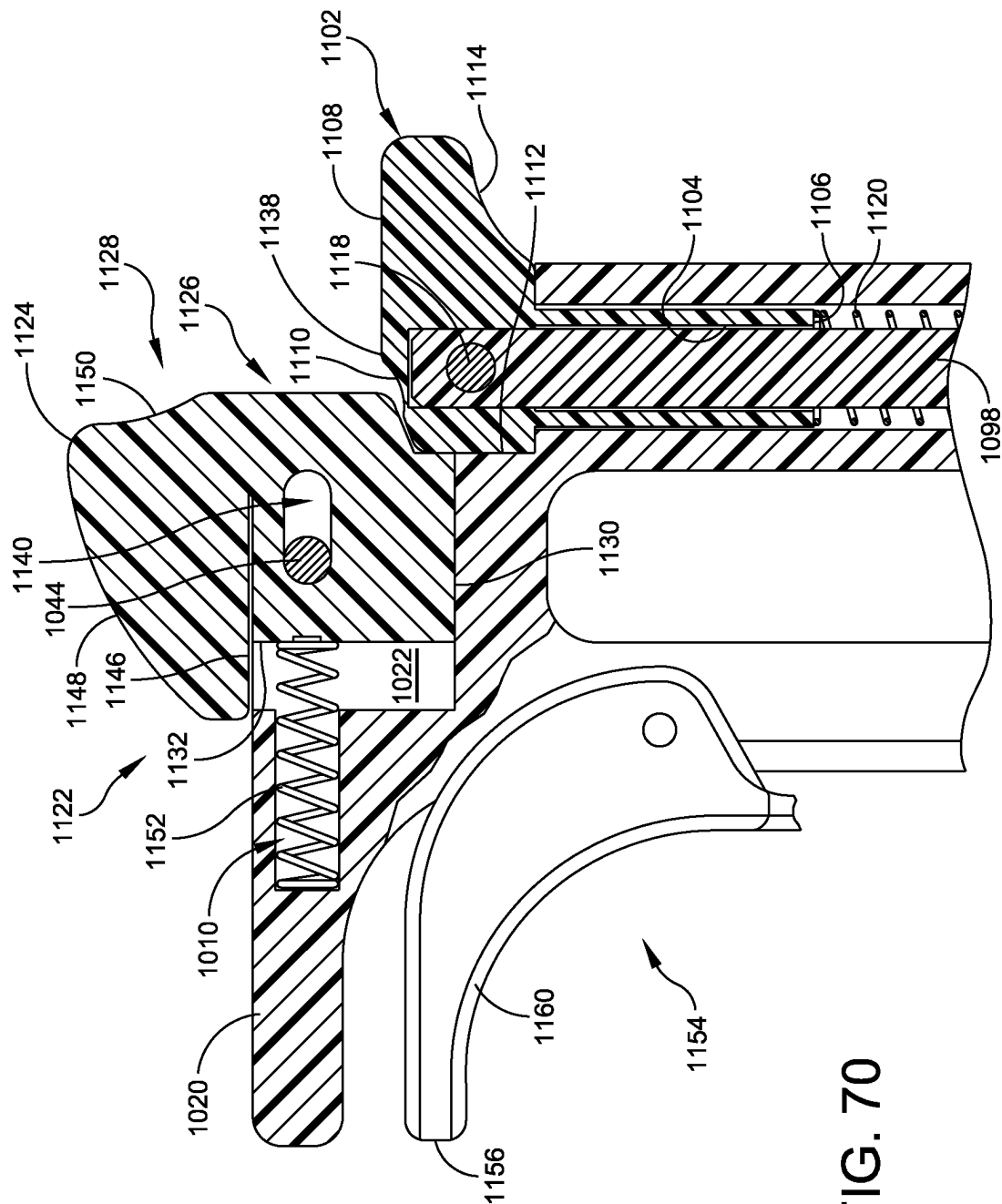


FIG. 70

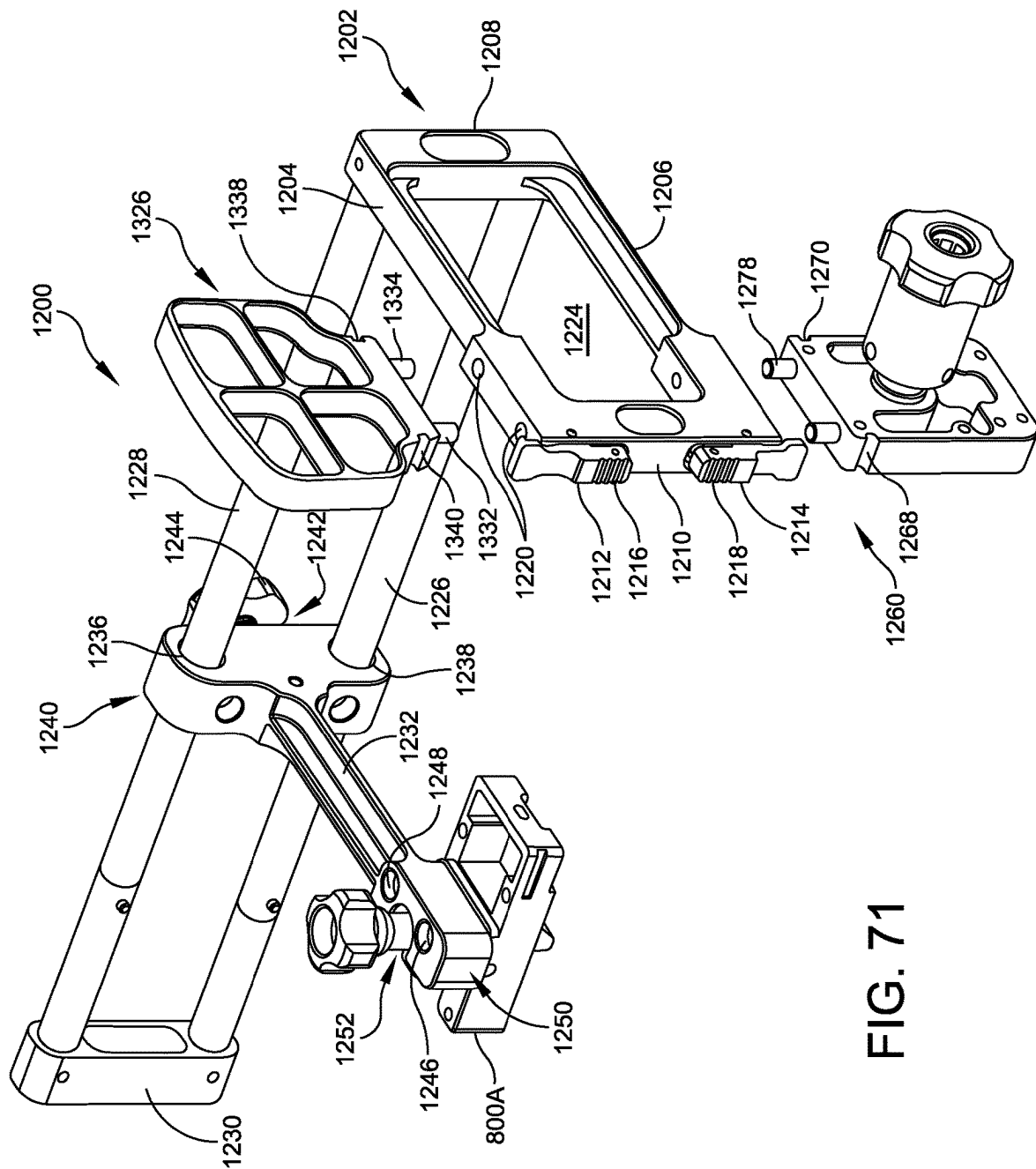
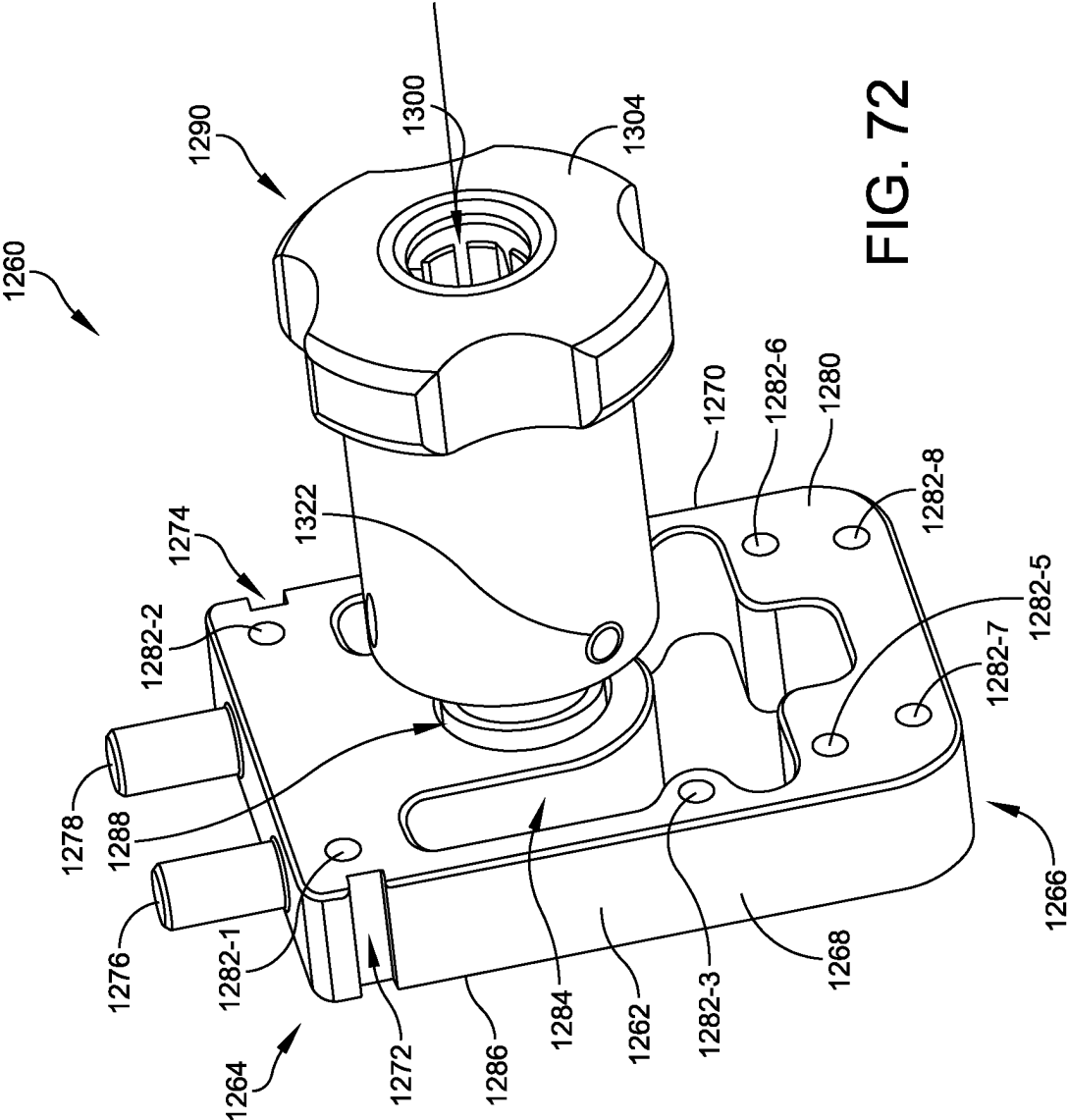
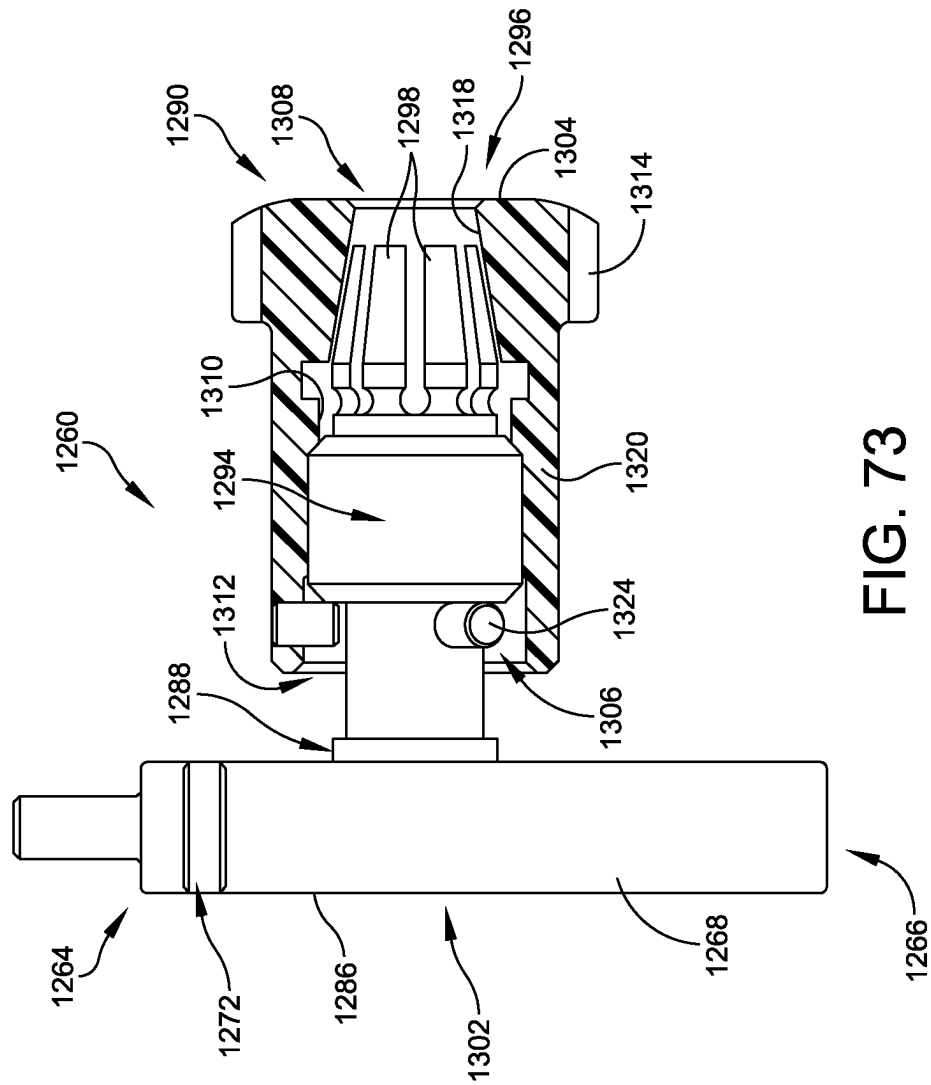


FIG. 71





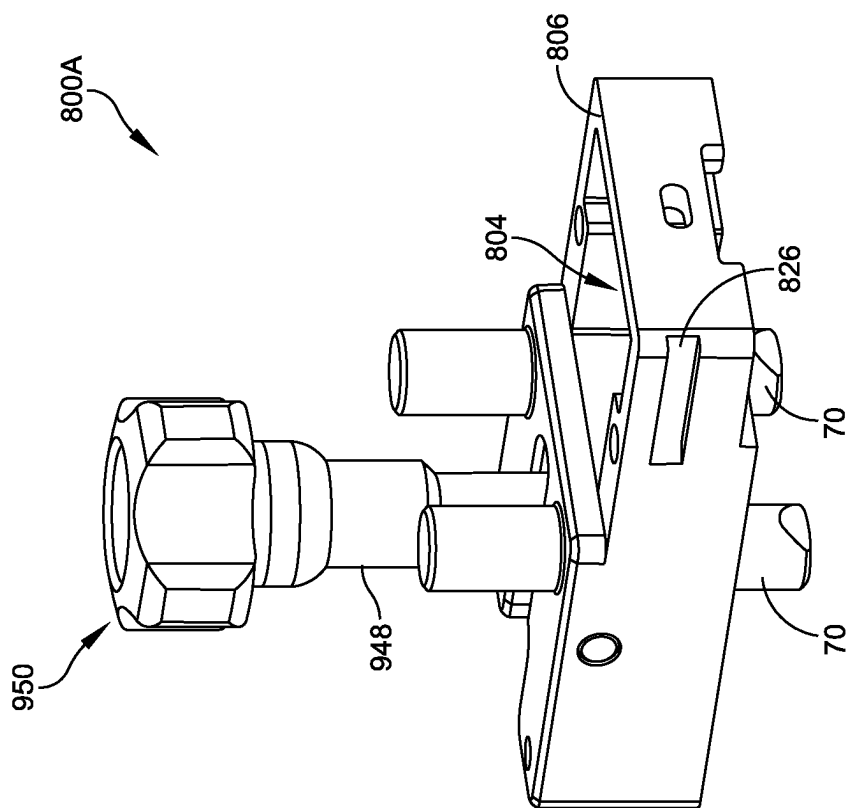


FIG. 74

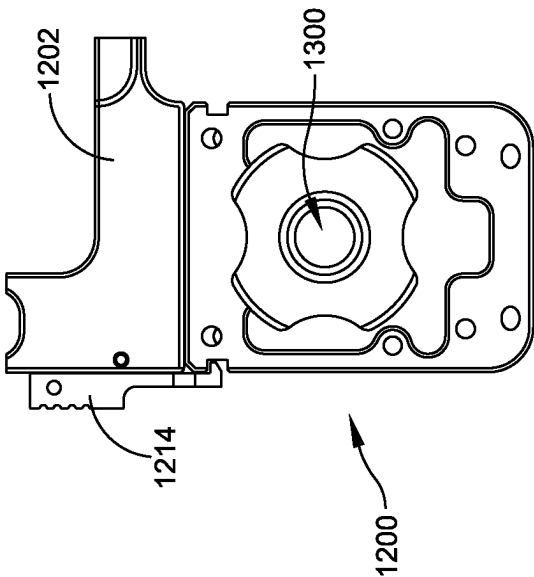


FIG. 76

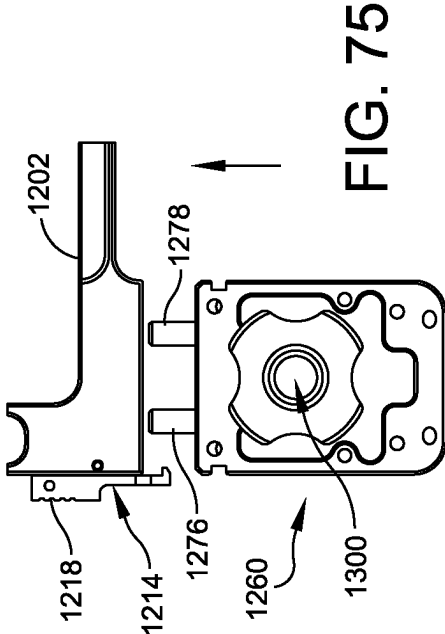


FIG. 75

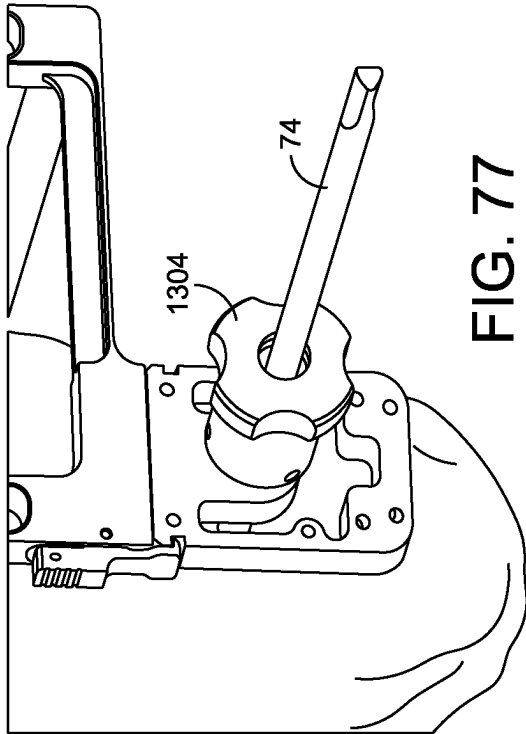


FIG. 77

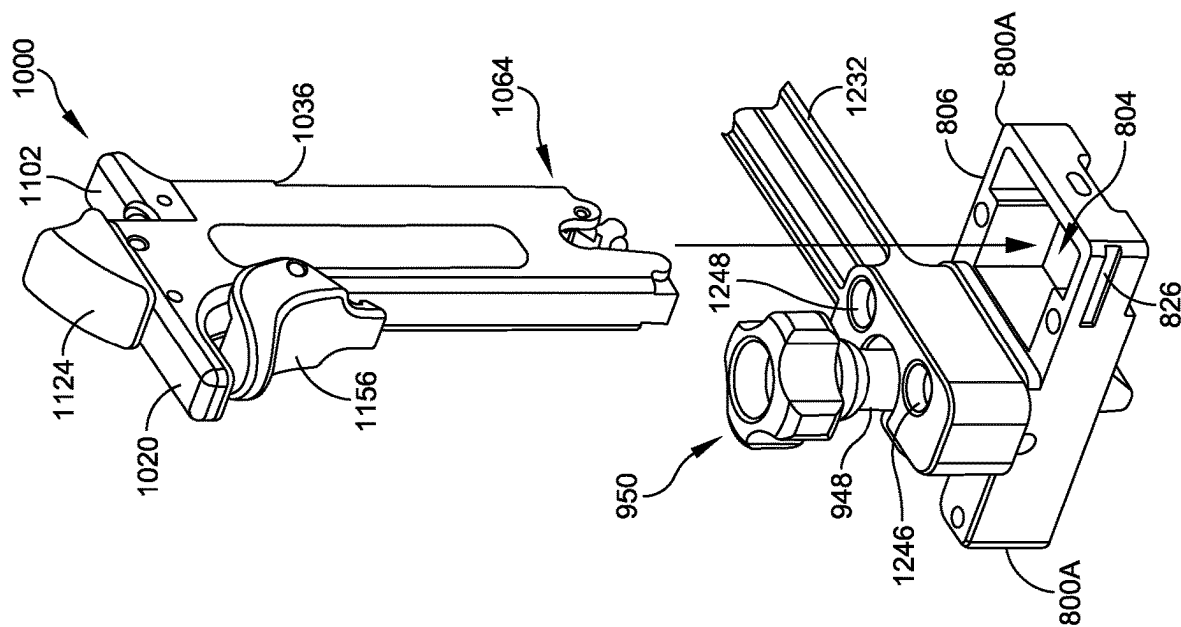


FIG. 78

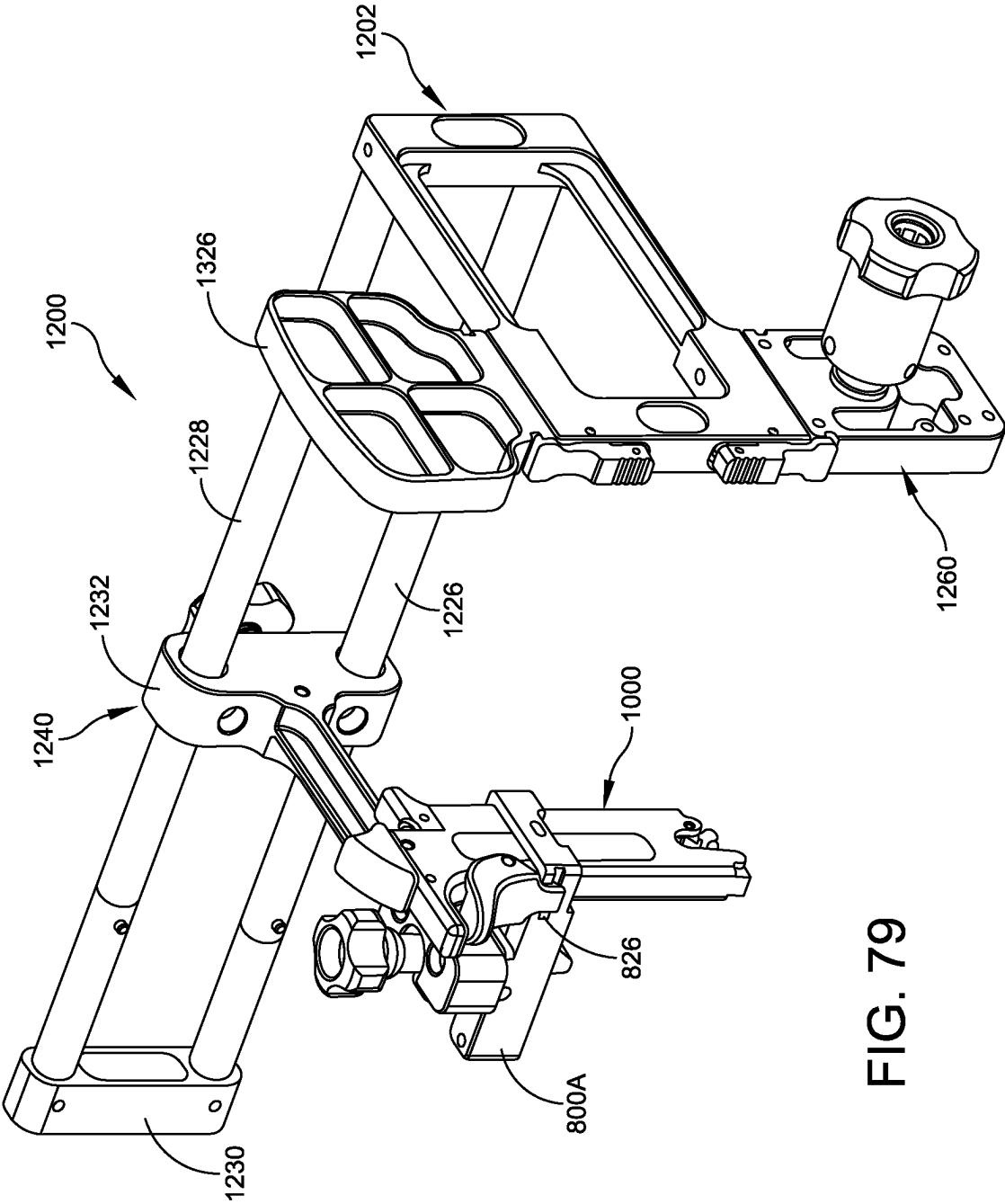


FIG. 79

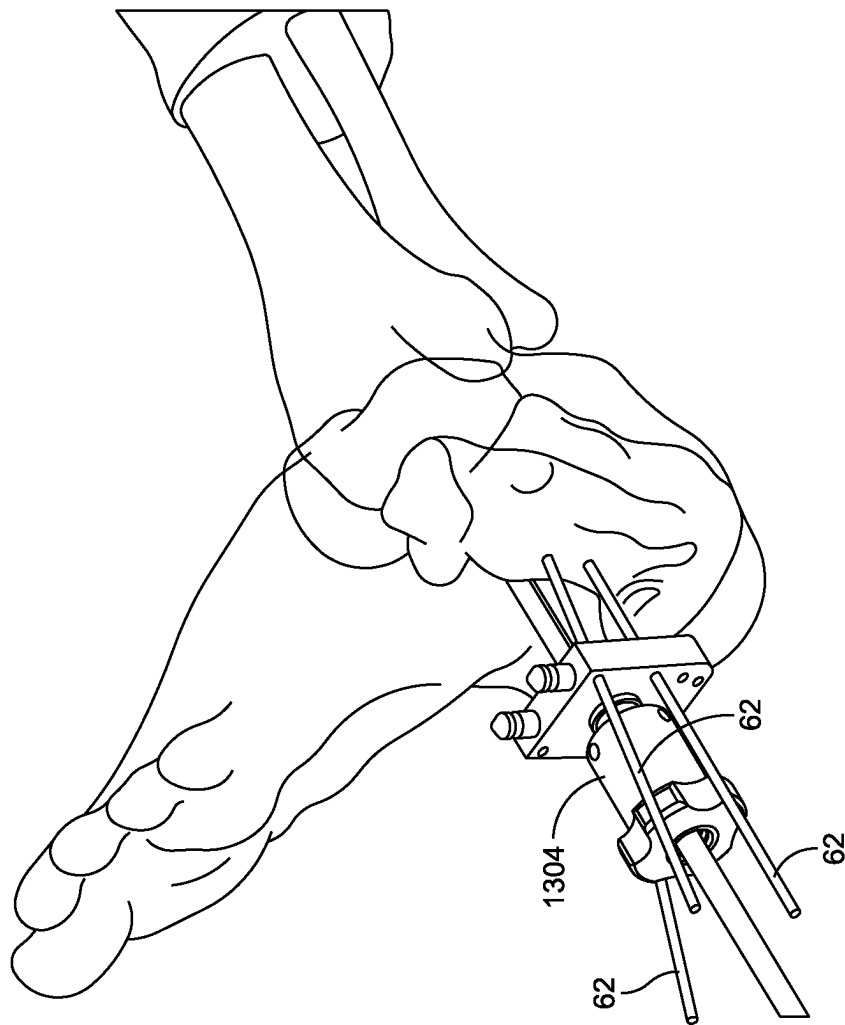


FIG. 80

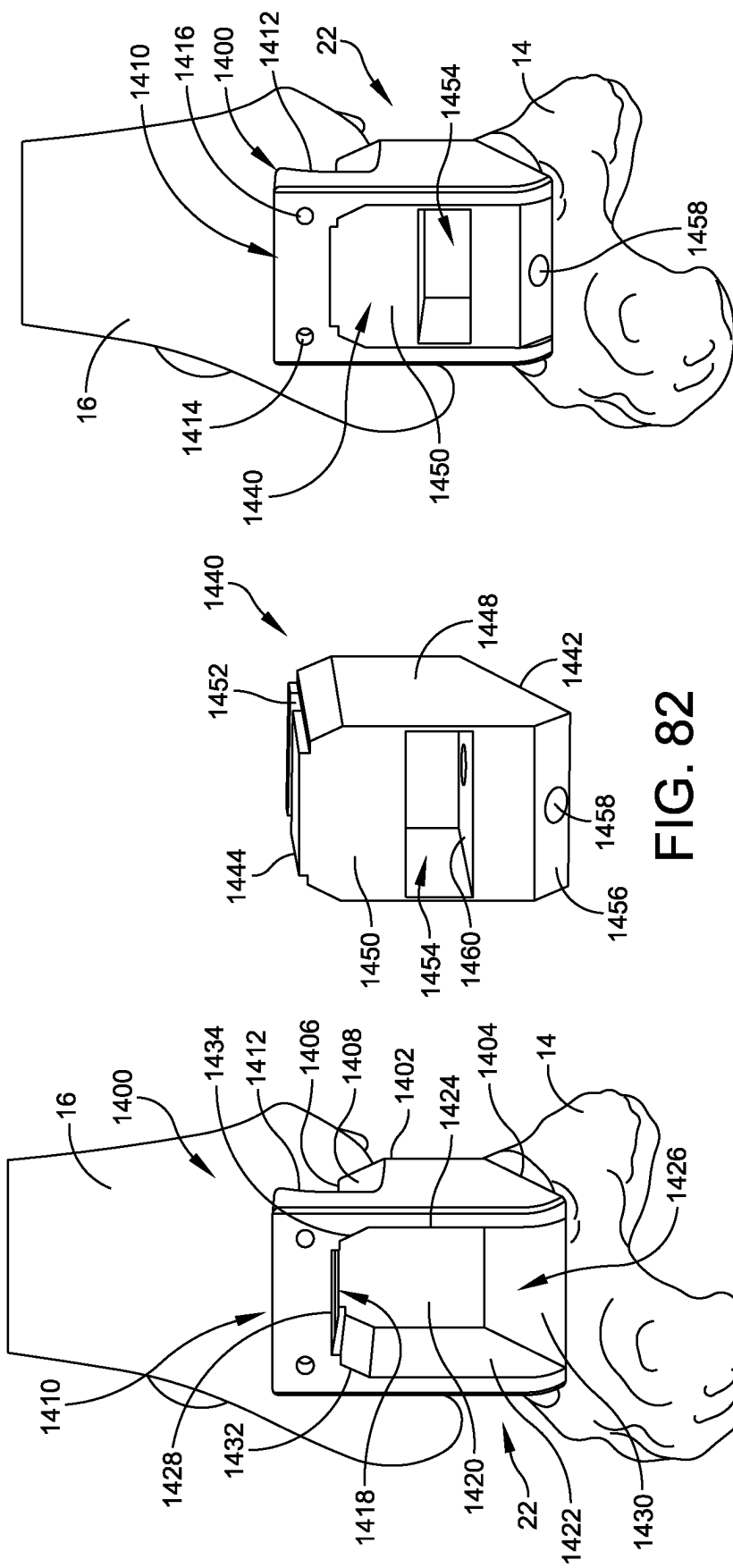


FIG. 83

FIG. 82

FIG. 81

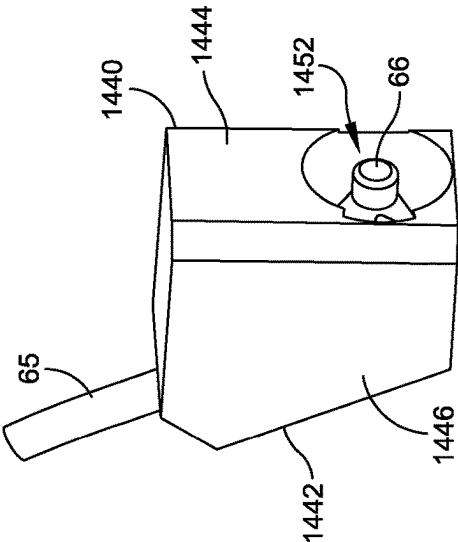


FIG. 85

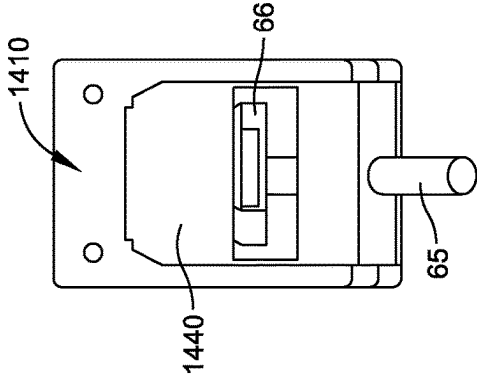


FIG. 86

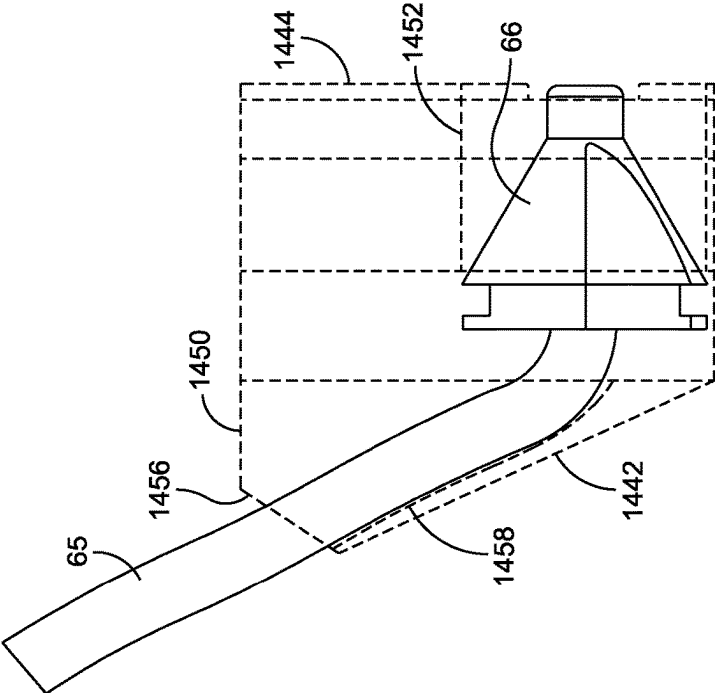


FIG. 84

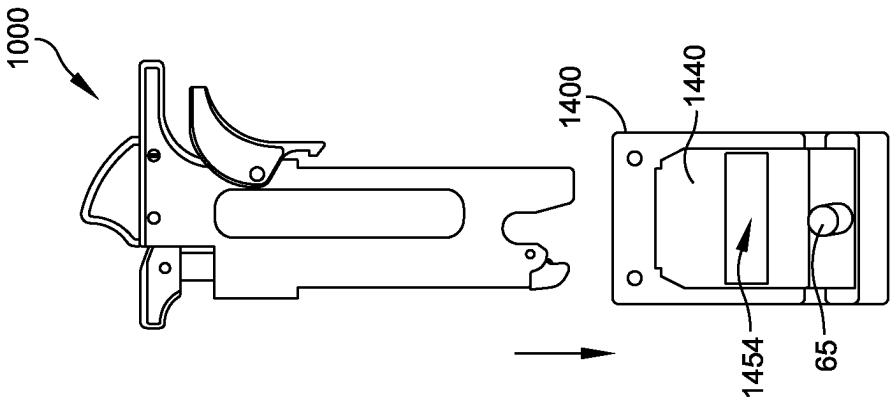


FIG. 87

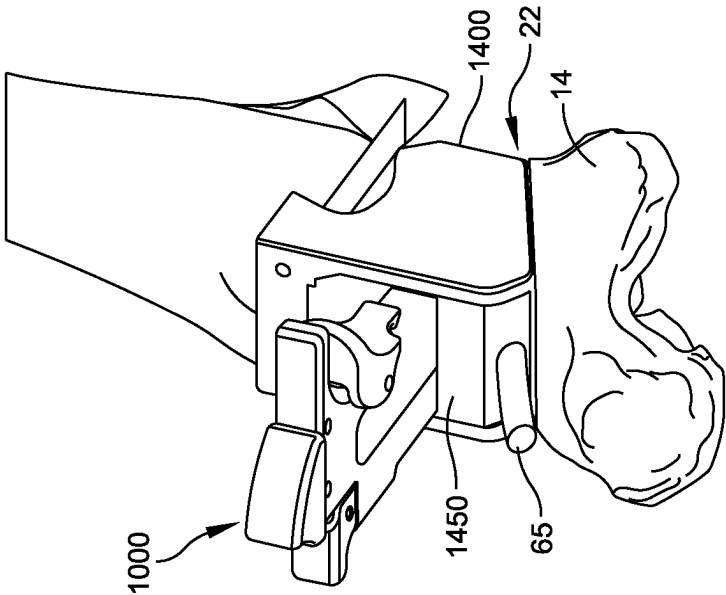


FIG. 88

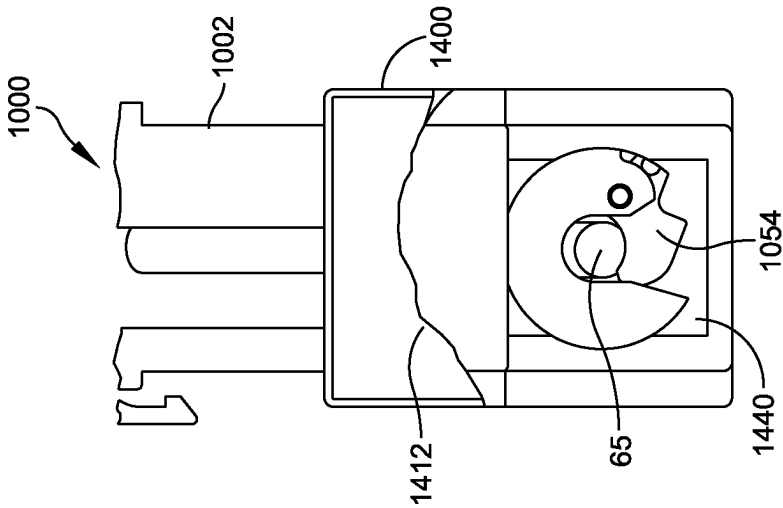


FIG. 89

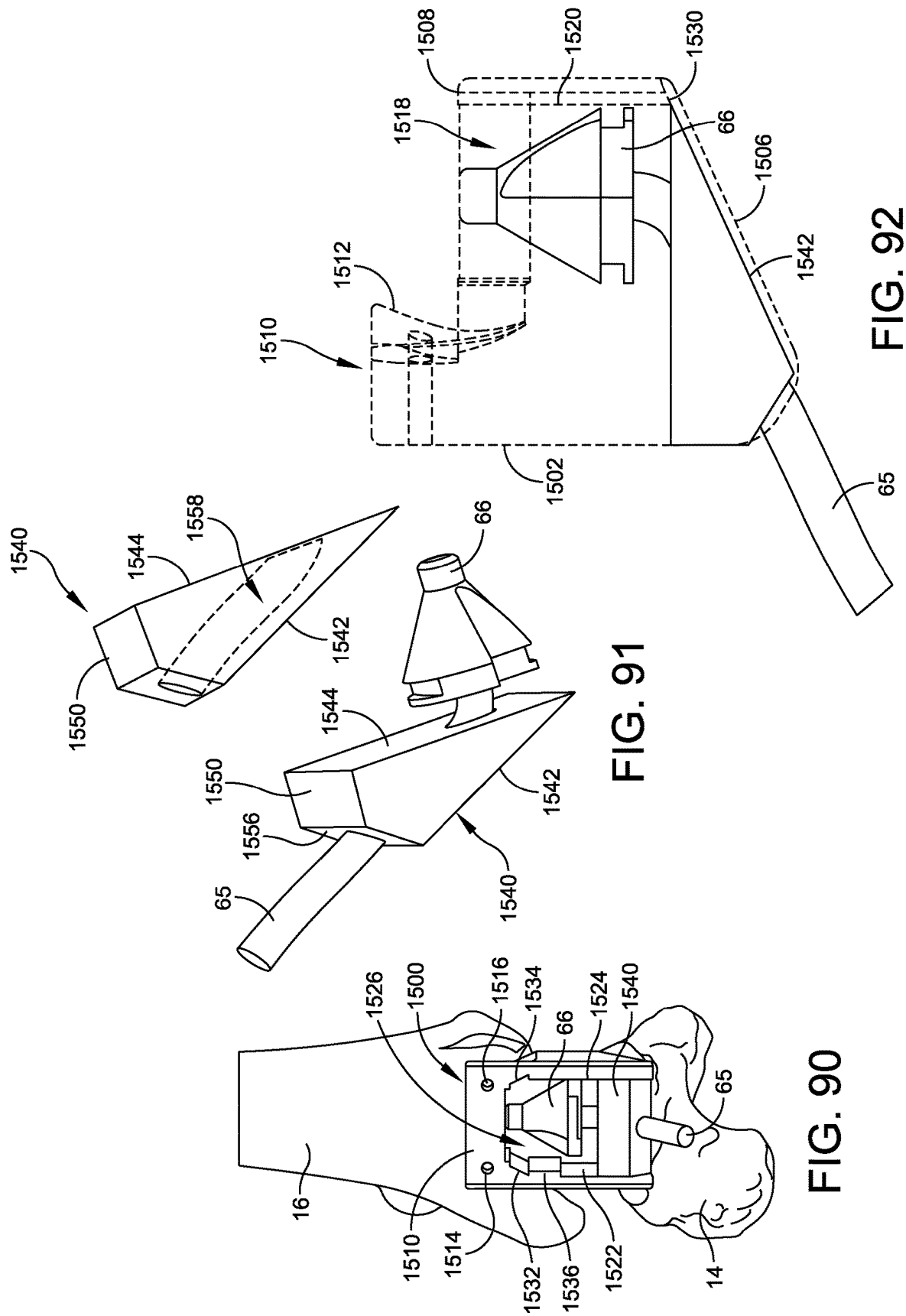
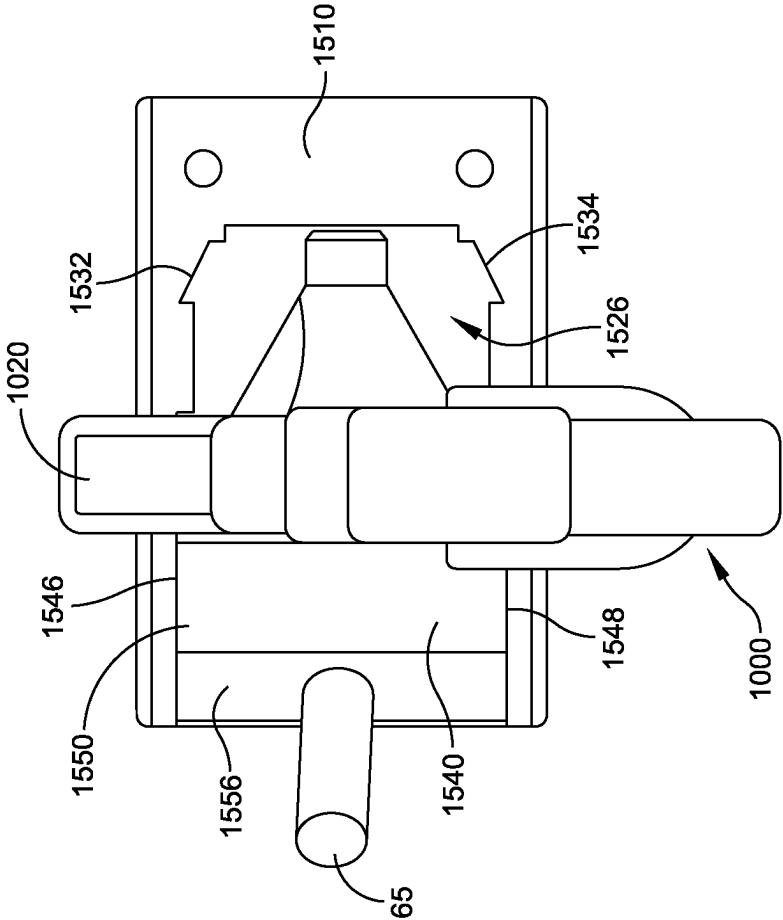
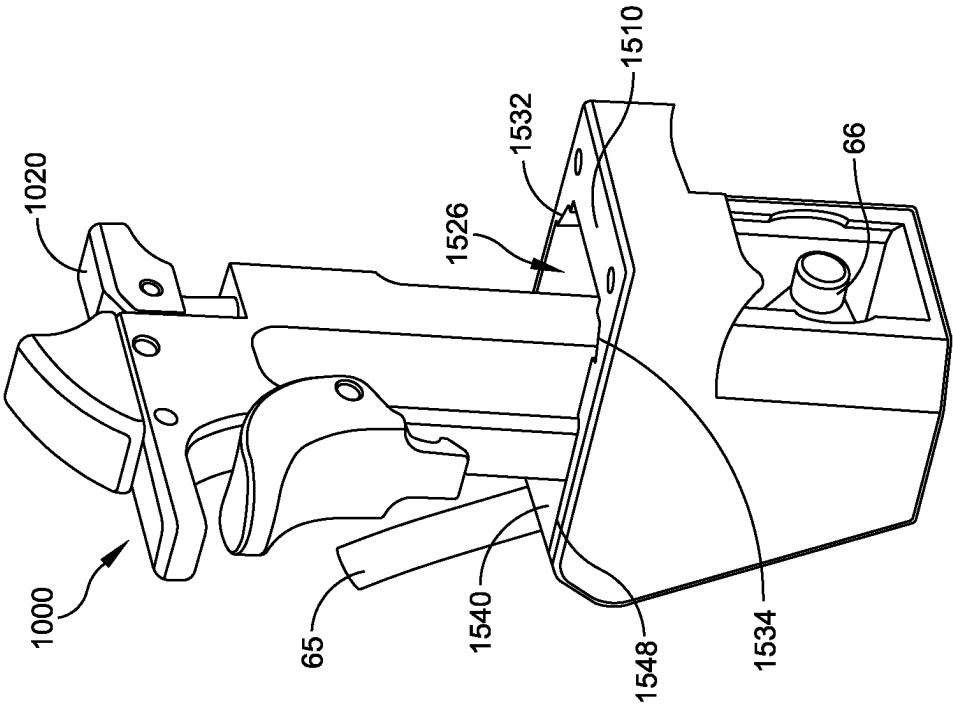
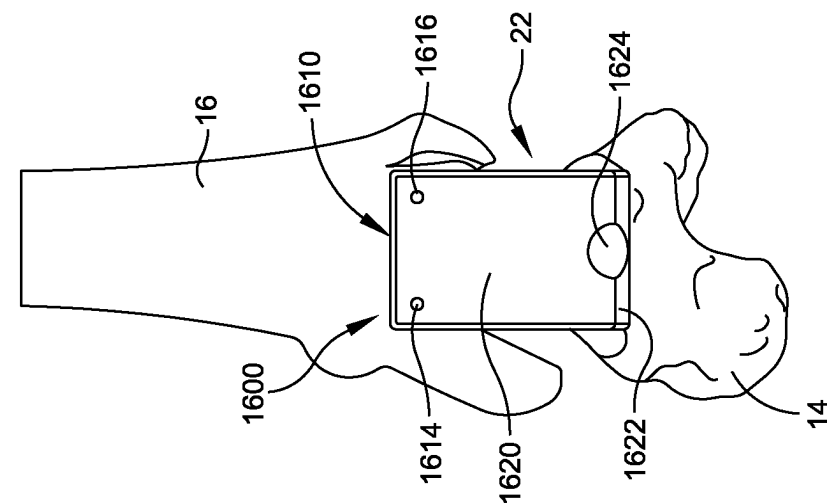
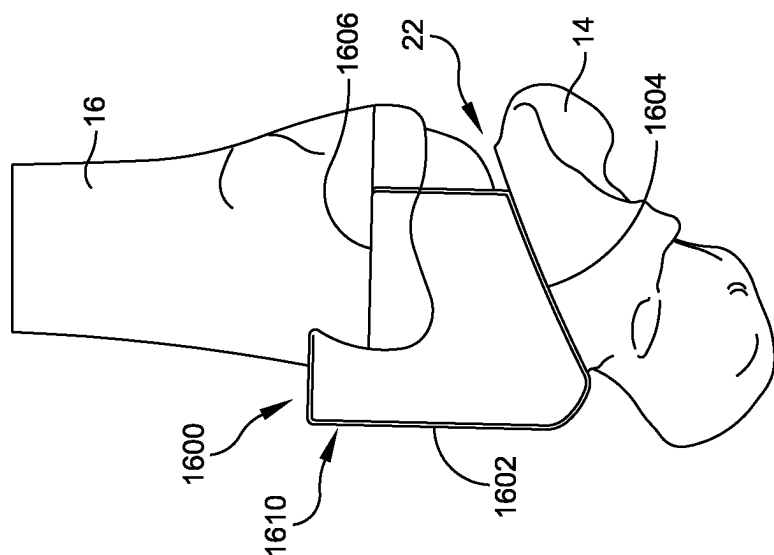
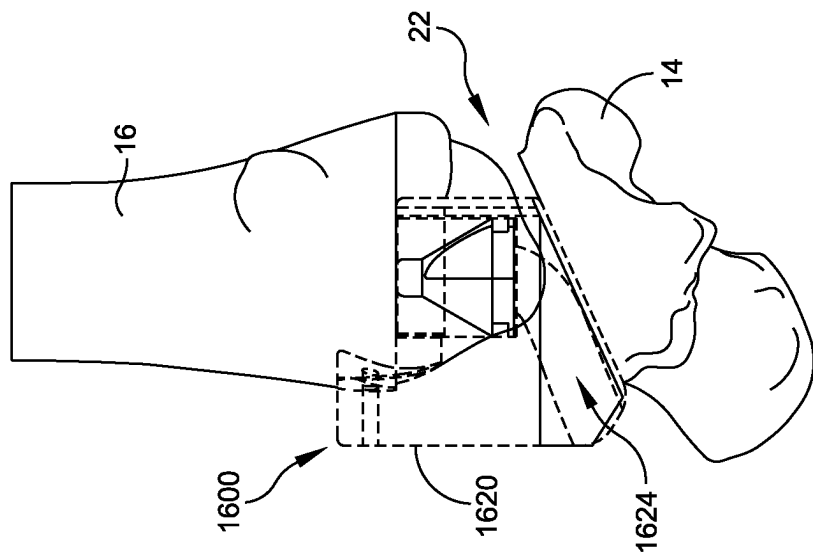


FIG. 92

FIG. 91

FIG. 90





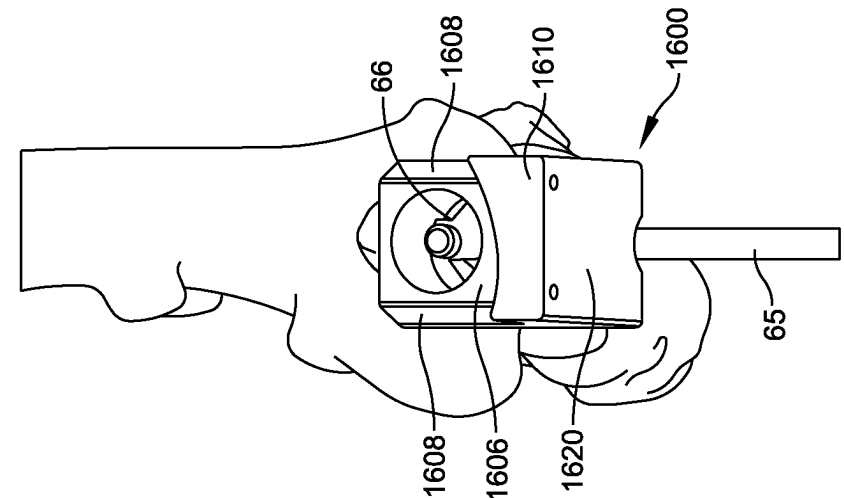


FIG. 98

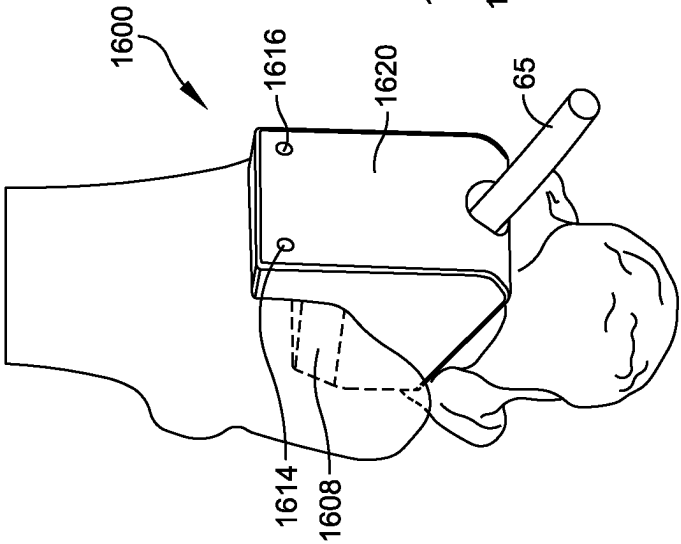


FIG. 99

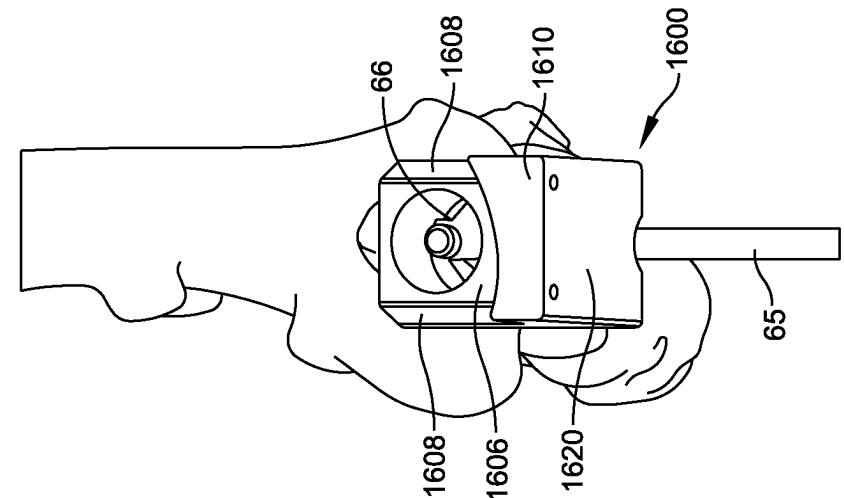


FIG. 100

SYSTEMS AND METHODS FOR INSTALLING AN ORTHOPEDIC IMPLANT

CROSS REFERENCE TO RELATED APPLICATIONS

This application is a continuation of U.S. Ser. No. 16/841,788, filed Apr. 7, 2020, which is a continuation of U.S. patent application Ser. No. 15/933,924, filed Mar. 23, 2018 (now U.S. Pat. No. 10,646,238), which is a division of U.S. patent application Ser. No. 15/440,715, filed Feb. 23, 2017 (now U.S. Pat. No. 9,949,747), which is a continuation of U.S. patent application Ser. No. 14/039,874, filed Sep. 27, 2013 (now U.S. Pat. No. 9,675,365), which is a continuation of U.S. patent application Ser. No. 13/464,175, filed May 4, 2012 (now U.S. Pat. No. 8,808,297), which is a continuation-in-part of U.S. patent application Ser. No. 13/330,091 filed on Dec. 19, 2011 (now U.S. Pat. No. 8,808,303), which claims priority to U.S. Provisional Patent Application No. 61/425,054 filed on Dec. 20, 2010 and to U.S. Provisional Patent Application No. 61/482,657 filed on May 5, 2011, and which is a continuation-in-part of U.S. patent application Ser. No. 12/711,307 filed on Feb. 24, 2010 (now U.S. Pat. No. 9,113,914) claiming priority to U.S. Provisional Patent Application No. 61/154,845 filed on Feb. 24, 2009, the entireties of which are herein incorporated by reference.

FIELD OF DISCLOSURE

The disclosed system and method generally relate to surgical guides. More specifically, the disclosed system and method relate to surgical guides for orthopedic procedures involving an ankle.

BACKGROUND

Total joint replacement prostheses typically include a specially designed jig or fixture to enable a surgeon to make accurate and precise bone resections in and around the joint being prepared to accept the prosthesis. The ultimate goal with any total joint prosthesis is to approximate the function and structure of the natural, healthy structures that the prosthesis is replacing. Should the prosthesis not be properly attached to the joint, i.e., an ankle or knee, the misalignment could result in discomfort to the patient, gait problems, or degradation of the prosthesis.

Many surgical procedures employ the use of intra-operative fluoroscopy to check the alignment of the intramedullary cavities that are prepared to receive the joint replacement prosthesis. However, the use of intra-operative fluoroscopy in the operating room has several drawbacks. One such drawback is that the use of fluoroscopy to check the alignment of intramedullary cavities formed during surgery increases the overall length of the surgical procedure as time is taken to acquire and evaluate the fluoroscopic images. Long surgery times lead to increased tourniquet time for the patient and therefore may increase recovery time.

Another drawback of fluoroscopy is exposing the patient and others in the operating room to the ionized radiation. For example, the U.S. Food and Drug Administration ("FDA") has issued several articles and public health advisories concerning the use of the fluoroscopy during surgical procedures. Consequently, even though steps are taken to protect the patient and other from the ionized radiation, it is virtually impossible to eliminate all risk associated with the ionized radiation.

SUMMARY

In some embodiments, a system includes a cartridge having an elongate body extending from a first end to a second end and having a top side and a bottom side. The cartridge defines a first hole adjacent to the first end that extends through the cartridge from the bottom side to the top side. The top side of the cartridge defines a pair of parallel slots that extend perpendicular with respect to a longitudinal axis of the cartridge. Each slot of the pair of parallel slots is equidistant from a central axis defined by the first hole.

In some embodiments, a system includes a first component, a second component, and a third component. The first component has a body defining a first opening. The second component has a body defining a hollow interior and a first side that defines a second opening. The second component is sized and configured to engage the first component such that the second opening aligns with the first opening. The third component has an elongate body extending from a first end to a second end and having a top side and a bottom side. The third component defines a first hole adjacent to the first end that extends through the cartridge from the bottom side to the top side. The third component is sized and configured to be received within the first opening defined by the first component, the second opening defined by the second component, and the hollow interior of the second component when the first and second components are engaged.

In some embodiments, a method includes inserting a second component into a resected joint space located between a first bone and a second bone; coupling a first component to the second component such that a first opening defined by the first component aligns with a second opening defined by the second component; and inserting a third component into the first opening defined by the first component, the second opening defined by the second component, and a hollow interior of the second component.

BRIEF DESCRIPTION OF THE DRAWINGS

These and other features and advantages of the present invention will be more fully disclosed in, or rendered obvious by, the following detailed description of the preferred embodiment of the invention, which is to be considered together with the accompanying drawings wherein like numbers refer to like parts and further wherein:

FIG. 1 illustrates the bones of a human foot and ankle;

FIGS. 2A and 2B are schematic representations of a scanned image of a human foot and ankle joint;

FIG. 3 is a perspective view of tibial and talar resection guides located upon portions of a tibia and a talus;

FIG. 4 is an exploded perspective view of a tibial cutting guide mount and tibial resection guide;

FIG. 5 is a perspective view of a tibial cutting guide disposed within a tibial cutting guide mount located on an inferior portion of a tibia;

FIG. 6 is a front elevational view of a tibial cutting guide disposed within a tibial cutting guide mount located on an inferior portion of a tibia;

FIG. 7 is a side elevational view of a tibial cutting guide disposed within a tibial cutting guide mount located on an inferior portion of a tibia during resection of the tibia;

FIG. 8 is a schematic representation of a resected tibia following application and use of the tibial cutting guide and tibial cutting guide mount;

FIG. 9 is a perspective view of a talar cutting guide disposed within a talar cutting guide mount;

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FIG. 10 is an exploded perspective view of the talar cutting guide mount and the talar cutting guide illustrated in FIG. 9;

FIG. 11 is a perspective view of the talar cutting guide disposed within the talar cutting guide mount located on a superior portion of a talus;

FIG. 12 is a front elevational view of the talar cutting guide disposed within the talar cutting guide mount located on a superior portion of a talus;

FIG. 13 is a side perspective view of the talar cutting guide disposed within the talar cutting guide mount located on a superior portion of a talus during resection of the talus;

FIG. 14 is a schematic representation of a resected talus following application and use of the talar cutting guide and talar cutting guide mount;

FIG. 15 is a schematic representation of a resected joint space following application and use of the talar and tibial cutting guide mounts and cutting guides;

FIG. 16 is a perspective view of one example of a custom tibial drill guide mount;

FIG. 17 is a front elevational view of the tibial drill guide mount illustrated in FIG. 16;

FIG. 18 is a rear elevation view of the tibial drill guide mount illustrated in FIG. 16;

FIG. 19 is a bottom elevational view of the tibial drill guide mount illustrated in FIG. 16;

FIG. 20 is a top elevational view of the tibial drill guide mount illustrated in FIG. 16;

FIG. 21 is a perspective view of one example of a tibial drill guide;

FIG. 22 is a side elevational view of the tibial drill guide illustrated in FIG. 21;

FIG. 23 is a top elevational view of the tibial drill guide illustrated in FIG. 21;

FIG. 24 is an exploded perspective view of the tibial drill guide mount and the tibial drill guide;

FIG. 25A is a side elevational view of the tibial drill guide disposed within the tibial drill guide mount being inserted into resected joint space;

FIG. 25B is a perspective view of the assemblage of the tibial drill guide mount and tibial drill guide disposed within the resected joint space;

FIG. 25C is a perspective view of the assembly of the tibial drill guide mount and tibial drill guide disposed and pinned within the resected joint space;

FIG. 26 is a perspective view of one example of an alignment tool;

FIG. 27 is an exploded perspective view of the alignment tool illustrated in FIG. 26;

FIGS. 28A and 28B illustrate the relative movement permitted between each of the components of the alignment tool illustrated in FIG. 26;

FIG. 29 is a perspective view of one example of an adapter bar for coupling the assemblage of the tibial drill guide mount and tibial drill guide to the alignment tool;

FIG. 30 is a perspective view of the adapter bar coupled to the assemblage of the tibial drill guide mount and tibial drill guide and to the alignment tool;

FIG. 31 is a top isometric view of another example of an alignment tool/foot holder assembly for use with a tibial drill guide mount and tibial drill guide;

FIG. 32 is a bottom isometric view of the alignment tool/foot holder assembly illustrated in FIG. 31;

FIG. 33 is an elevational front view of the alignment tool/foot holder assembly illustrated in FIG. 31;

FIG. 34 is an elevational side view of the alignment tool/foot holder assembly illustrated in FIG. 31;

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FIG. 35 is a top isometric view of another example of an alignment tool/foot holder assembly for use with the tibial drill guide mount and tibial drill guide;

FIG. 36 is a top elevational view of the alignment tool/foot holder assembly illustrated in FIG. 35;

FIG. 37 is an elevational front view of the alignment tool/foot holder assembly illustrated in FIG. 35;

FIG. 38 is an elevational side view of the alignment tool/foot holder assembly illustrated in FIG. 35;

FIG. 39 is a perspective view of another example of a tibial cutting guide mount;

FIG. 40 is a front side elevational view of the tibial cutting guide mount illustrated in FIG. 39;

FIG. 41 is a side elevational view of the tibial cutting guide mount illustrated in FIG. 39;

FIG. 42 is a top side view of the tibial cutting guide mount illustrated in FIG. 39;

FIG. 43 is a bottom side view of the tibial cutting guide mount illustrated in FIG. 39;

FIG. 44 is a perspective view of a tibial drill guide cartridge for use with the tibial drill guide mount illustrated in FIG. 39;

FIG. 45 is a front end view of the tibial drill guide cartridge illustrated in FIG. 44;

FIG. 46 is a bottom side plan view of the tibial drill guide cartridge illustrated in FIG. 44;

FIG. 47 is a side view of the tibial drill guide cartridge illustrated in FIG. 44;

FIG. 48 is an exploded perspective view of a mounting plate and dowel pins configured to for use with the tibial drill guide mount illustrated in FIG. 39;

FIG. 49 is a partially exploded perspective view of a mounting plate and dowel pins configured to for use with the tibial drill guide mount illustrated in FIG. 39;

FIG. 50 is a partially exploded perspective view of a mounting plate, dowel pins, and tibial drill guide mount configured to receive a tibial drill guide cartridge in accordance with FIG. 44;

FIG. 51 is a perspective view of the tibial drill guide mount, tibial drill guide cartridge, dowel pins, and mounting plate assembled together;

FIG. 52 is a side view of the assembly illustrated in FIG. 51;

FIG. 53 is a top side plan view of the assembly illustrated in FIG. 51;

FIG. 54 is a bottom side plan view of the assembly illustrated in FIG. 51;

FIG. 55 is a perspective view of a foot holder assembly for use with the assembly illustrated in FIG. 51;

FIG. 56 is a perspective view of a pivoting arrangement used to secure the assembly illustrated in FIG. 51 to the foot holder assembly;

FIG. 57 is a top side plan view of the foot holder assembly illustrated in FIG. 55;

FIG. 58 is a side view of the foot holder assembly illustrated in FIG. 55;

FIG. 59 is an opposite side view of the foot holder assembly illustrated in FIG. 55;

FIG. 60 is a rear end view of the foot holder assembly illustrated in FIG. 55;

FIG. 61 is a front end view of the foot holder assembly illustrated in FIG. 55;

FIG. 62 is a perspective view of a drill being extended through the foot holder assembly and tibial drill guide;

FIG. 63 is an isometric view of one example of a reamer stabilizer in accordance with some embodiments;

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FIGS. 64 and 65 illustrate the reamer stabilizer illustrated in FIG. 63 during various stages of operation;

FIG. 66 is an exploded isometric view of the reamer stabilizer illustrated in FIG. 63;

FIGS. 67 and 68 are cross-sectional detailed view of the coupling assembly of the reamer stabilizer illustrated in FIG. 63 during various stages of operation;

FIG. 69 is a cross-sectional detail view of the coupling between a plunger rod, pivot rod, and reamer guide body in accordance with the reamer stabilizer illustrated in FIG. 63;

FIG. 70 is a cross-sectional detail view of the locking assembly of the reamer stabilizer illustrated in FIG. 63;

FIG. 71 is an isometric view of an embodiment of a foot holder assembly;

FIG. 72 is an isometric view of one example of a drill guide assembly that is configured to be releasably coupled to the foot holder assembly illustrated in FIG. 71;

FIG. 73 is a partial cross-sectional view of the drill guide assembly illustrated in FIG. 72;

FIG. 74 is an isometric view of one example of a modified mounting member in accordance with the foot holder assembly illustrated in FIG. 71;

FIGS. 75 and 76 illustrate the coupling of the drill guide assembly illustrated in FIG. 72 to the foot holder assembly illustrated in FIG. 71;

FIG. 77 illustrates a trocar being received within the drill guide assembly;

FIGS. 78 and 79 illustrate a reamer stabilizer in accordance with FIG. 63 being coupled to the foot holder assembly illustrated in FIG. 71;

FIG. 80 illustrates the drill guide assembly coupled to the foot of a patient during an operation;

FIG. 81 illustrates one example of an anterior reaming guide mount disposed within a resected joint space in accordance with some embodiments;

FIG. 82 is an isometric view of one example of an insert for use with the anterior reaming guide mount illustrated in FIG. 81;

FIG. 83 illustrates the insert illustrated in FIG. 82 disposed within the anterior reaming guide mount, which is received within a resected joint space;

FIG. 84 is a side view of a flexible reaming rod and reaming head disposed within the insert illustrated in FIG. 82;

FIG. 85 is an isometric side view of the flexible reaming rod and reaming head disposed within the insert;

FIG. 86 is a front elevation view of the flexible reaming rod and reaming head disposed within the insert;

FIGS. 87-89 illustrate the reamer stabilizer, anterior reaming guide mount, and insert during various stages of an operation;

FIG. 90 illustrates another example of an anterior reaming guide mount and insert disposed within a resected joint space during an operation;

FIG. 91 are isometric side view of the insert illustrated in FIG. 90;

FIG. 92 is a side view of the insert disposed within the anterior reaming guide mount in accordance with FIG. 90;

FIGS. 93 and 94 illustrate the reamer stabilizer in use with the anterior reaming guide mount and insert illustrated in FIG. 90;

FIGS. 95-100 illustrate another example of an anterior reaming guide mount.

DETAILED DESCRIPTION

This description of preferred embodiments is intended to be read in connection with the accompanying drawings,

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which are to be considered part of the entire written description. The drawing figures are not necessarily to scale and certain features may be shown exaggerated in scale or in somewhat schematic form in the interest of clarity and conciseness. In the description, relative terms such as "horizontal," "vertical," "up," "down," "top" and "bottom" as well as derivatives thereof (e.g., "horizontally," "downwardly," "upwardly," etc.) should be construed to refer to the orientation as then described or as shown in the drawing figure under discussion. These relative terms are for convenience of description and normally are not intended to require a particular orientation. Terms including "inwardly" versus "outwardly," "longitudinal" versus "lateral" and the like are to be interpreted relative to one another or relative to an axis of elongation, or an axis or center of rotation, as appropriate. Terms concerning attachments, coupling and the like, such as "connected" and "interconnected," refer to a relationship wherein structures are secured or attached to one another either directly or indirectly through intervening structures, as well as both movable or rigid attachments or relationships, unless expressly described otherwise. When only a single machine is illustrated, the term "machine" shall also be taken to include any collection of machines that individually or jointly execute a set (or multiple sets) of instructions to perform any one or more of the methodologies discussed herein. The term "operatively connected" is such an attachment, coupling or connection that allows the pertinent structures to operate as intended by virtue of that relationship. In the claims, means-plus-function clauses, if used, are intended to cover the structures described, suggested, or rendered obvious by the written description or drawings for performing the recited function, including not only structural equivalents but also equivalent structures.

The disclosed systems and methods advantageously utilize custom manufactured surgical instruments, guides, and/or fixtures that are based upon a patient's anatomy to reduce the use of fluoroscopy during a surgical procedure. In some instances, the use of fluoroscopy during a surgical procedure may be eliminated altogether. The custom instruments, guides, and/or fixtures are created by imaging a patient's anatomy with a computer tomography scanner ("CT"), a magnetic resonance imaging machine ("MRI"), or like medical imaging technology prior to surgery and utilizing these images to create patient-specific instruments, guides, and/or fixtures.

Although the following description of the custom patient-specific instruments are described with respect to a foot 10 and ankle 12 (FIG. 1), one skilled in the art will understand that the systems and methods may be utilized in connection with other joints including, but not limited to, knees, hips, shoulders, and the like. As shown in FIG. 1, a typical human foot 10 includes an ankle joint 12 formed between a talus 14, which is disposed on a calcaneus 20, and a tibia 16 and fibula 18.

A CT or MRI scanned image or series of images may be taken of a patient's ankle 12 (or other joint) and then converted from, e.g., a DICOM image format, to a solid computer model of the ankle including the calcaneus, talus, tibia, navicular, and fibula to determine implant alignment, type, and sizing using specialized modeling methods that are often embodied in computer software. Computer generated solid models that are derived from the data of the CT or MRI scan image will often include precise and accurate information regarding the surface contours surrounding the structures that have been imaged, e.g., the surface topography of the bones or contour of fascia that have been imaged. It will be understood that by surface topography it is meant the

location, shape, size and distribution of surface features such as concavities and prominences or the like.

The methods disclosed in U.S. Pat. No. 5,768,134, issued to Swaelens et al., which is incorporated by reference herein in its entirety, have been found to yield adequate conversions of data of CT or MRI scan images to solid computer models. In some embodiments, images are made of a foot **10**, i.e., the calcaneus **20**, talus **14**, tibia **16**, and fibula **18** of a patient using a CT or MRI machine, or other digital image capturing and processing unit as is understood by one skilled in the art. The image data is processed in a processing unit, after which a model **50** is generated using the processed digitized image data as illustrated in FIGS. **2A** and **2B**.

Interactive processing and preparation of the digitized image data is performed, which includes the manipulation and introduction of additional extrinsic digital information, such as, predefined reference locations **52** for component positioning and alignment so that adjustments to the surgical site **54**, that will require resection during surgery, may be planned and mapped onto computer model **50** (FIGS. **2A** and **2B**). After the interactive processing of the digitized image data, it is possible to go back to original CAD data to obtain a higher resolution digital representation of the patient specific surgical instruments, prostheses, guides, or fixtures so as to add that digital representation to the patient's image data model.

FIG. **3** illustrates a pair of custom cutting guides for an ankle replacement surgery including a tibial resection guide mount **100** and a talar resection guide mount **102**, which are formed and mounted to the patient's lower tibia **16a** and upper talus **14a**. A custom tibial drill guide mount **200** (FIGS. **16-20**) is also formed and configured to be received within ankle space created by using the custom tibial and talar resection guide mounts **100**, **102**. Although custom cutting guides are described for preparing a patient's talus, tibia, and femur, one skilled in the art will understand that other cutting guides may be implemented and that custom guides may be created for other joints including, but not limited to, the knee, hip, shoulder, or other joint.

Tibial resection guide mount **100** illustrated in FIG. **3** is formed from a resilient polymer material of the type that is suitable for use in connection with stereo lithography, selective laser sintering, or like manufacturing equipment. Resection guide mount **100** includes a unitary body including a cruciform tibial yolk **104** projecting upwardly from a base **106** that further defines a guide receptacle recess **108** as best seen in FIG. **4**. Cruciform yolk **104** includes a pair of spaced apart arms **110**, **112** that project outwardly from a central post **114**. Arms **110**, **112** and central post **114** each have a conformal bone engaging surface **116** that is complementary to the contours of a corresponding portion of the patient's lower tibia **16a** as illustrated in FIG. **7**. Through the previously discussed imaging operations, conformal bone engaging surfaces **116** of arms **110**, **112** and central post **114** are configured for complementary matching with anatomical surface features of a selected region of the patient's natural bone. For tibial resection guide mount **100**, the selected bone region comprises the lower surfaces of the patient's tibia **16a**.

As best seen in FIGS. **3-5**, a pilot block **118** projects outwardly from central post **114**, adjacent to the intersection of arms **110**, **112**. A support block **120** (FIG. **4**) is located on base **106** in spaced relation to pilot block **118**. Guide receptacle recess **108** is defined by a pair of wings **122**, **124** that extend outwardly from either side of central post **114** in opposite directions on base **106**, with support block **120** located between them. Each wing **122**, **124** includes a

respective pylon **126** projecting outwardly from base **106** so as to provide lateral support for tibial resection guide **132** (FIGS. **4** and **5**). An elongate slot **128** is defined transversely in a central portion of base **106** below pilot block **118**, but above support block **120**. Each wing **122**, **124** also defines a respective slot **130** that is oriented at an angle relative to central post **114**. In some embodiments, slots **130** are disposed at a non-perpendicular angle relative to central post **114**, although one skilled in the art will understand that slots **130** may be disposed at perpendicular angles with respect to the direction in which central post **114** extends. Slots **128** and **130** are sized and shaped to allow a typical surgical saw **60** (FIG. **7**) of the type often used for bone resection, to pass through from a correspondingly positioned and sized slot in resection guide **132** without contact, or with only incidental contact with resection guide mount **100**.

Referring again to FIG. **4**, tibial resection guide **132** includes a pair of arms **134** that project downwardly and outwardly in diverging angular relation from the ends of a bridge beam **136**. The shape of tibial resection guide **132** is complementary to the shape of guide receptacle recess **108** as defined by the inwardly facing surfaces of pilot block **118**, support block **120**, and pylons **126**. Bridge beam **136** defines an elongate slot **138** that aligns with slot **128** when tibial resection guide is coupled to and supported by resection guide mount **100**. Arms **134** each define a respective slot **140** that align with a respective slot **130**.

The inwardly facing surfaces **142** of pilot block **118**, support block **120**, and pylons **126**, that together define guide receptacle recess **108**, have a shape that is complementary to the outer profile of tibial resection guide **132**. Guide receptacle recess **108** is sized so as to accept tibial resection guide **132** with a "press-fit". By press-fit it should be understood that the inwardly facing surfaces **142** of pilot block **118**, support block **120**, and pylons **126** are sufficiently resilient to deflect or compress elastically so as to store elastic energy when tibial resection guide **132** is pushed into guide receptacle recess **108**. Of course, it will also be understood that tibial resection guide **132** will have an outer peripheral shape that is complementary to the circumferential shape of guide receptacle recess **108**, but slightly larger in size, for press-fit embodiments. Also, tibial resection guide **132** may be retained within guide receptacle recess **108** by only frictional engagement with the inwardly facing surfaces of pilot block **118**, support block **120**, and pylons **126**. In some embodiments, tibial resection guide **132** can simply slide into guide receptacle recess **108** without operative contact or only incidental engagement with the inwardly facing surfaces of pilot block **118**, support block **120**, and pylons **126**.

Referring now to FIGS. **9** and **10**, a talar resection guide mount **102** is formed from a resilient polymer material of the type that is suitable for use in connection with stereo lithography, selective laser sintering, or the like manufacturing equipment, e.g., a polyamide powder rapid prototype material is suitable for use in connection with selective laser sintering. Talar resection guide mount **102** also includes a conformal bone engaging surface **144** that is complementary to the contours of a corresponding portion of the patient's upper talus **14a** (FIGS. **11** and **13**). Through the previously discussed imaging operations, conformal bone engaging surface **144** of talar resection guide mount **102** is configured for complementary matching with anatomical surface features of a selected region of the patient's natural bone. For talar resection guide mount **102**, the selected bone region comprises the outer, upper surfaces of the patient's talus.

Talar resection guide mount **102** comprises a unitary block that defines a central guide receptacle recess **146** and a pair of through-bores **148** (FIG. **10**). Guide receptacle recess **146** is defined by the inwardly facing surfaces **150** of a pair of wings **152**, **154** that project outwardly, in opposite directions from a base **156**. Each wing **152**, **154** includes a pylon **158** projecting upwardly to support guide housing **160** such that an elongate slot **162** is defined within base **156** and below guide housing **160** (FIGS. **10** and **11**). Slot **162** is sized and shaped to allow a typical surgical saw **60**, of the type often used for bone resection, to pass through from a correspondingly positioned and sized slot **164** in talar resection guide **166** without contact, or with only incidental contact with talar resection guide locator **102** (FIGS. **11** and **13**). An annular wall **168**, having a shape that is complementary to the outer profile of talar resection guide **166**, projects outwardly in substantially perpendicular relation to a back wall and so as to further defines guide receptacle recess **146**.

Still referring to FIGS. **9** and **10**, talar resection guide **166** includes a pair of confronting, parallel plates **170**, **172** that define elongate slot **164** between them, and are joined to one another at their ends by wings **174**. In this way, the shape of talar resection guide **166** is complementary to the shape of guide receptacle recess **146** as defined by the inwardly facing surfaces **150** of wings **152**, **154**, base **156**, and pylons **158**. Guide receptacle recess **146** is sized so as to accept talar resection guide **166** with a press-fit. Of course, it will also be understood that talar resection guide **166** will have an outer peripheral shape that is complementary to the circumferential shape of guide receptacle recess **146**, but slightly larger in size, for press-fit embodiments. Also, talar resection guide **166** may be retained within guide receptacle recess **146** by only frictional engagement with the inwardly facing surfaces **150** of wings **152**, **154**, base **156**, and pylons **158**. In some embodiments, talar resection guide **166** can simply slide into guide receptacle recess **146** without operative contact or only incidental engagement with the inwardly facing surfaces **150** of wings **152**, **154**, base **156**, and pylons **158**.

Tibial drill guide mount **200** illustrated in FIGS. **16-20** also may be fabricated from a resilient polymer material of the type that is suitable for use in connection with stereo lithography, selective laser sintering, or the like manufacturing equipment, e.g., a polyamide powder repaid prototype material is suitable for use in connection with selective laser sintering. As shown in FIGS. **16-20**, tibial drill guide mount **200** includes a somewhat rectangular body **204** that defines an aperture **206** that extends from a top surface **208** of body **204** to a bottom surface **210** of body **204**. Top surface **208** of body **204** may include a pair of chamfers **212** that are sized and configured to be mate against the resected surfaces of the lower tibia **16a** (FIG. **8**). Put another way, the top or upper surface **208** of body **204**, including chamfers **212**, is complementary to the geometry and locations of slots **138** and **140** of tibial resection guide **132**.

Front side **214** of body **204** defines one or more blind holes **216**. As illustrated in the embodiment shown in FIG. **17**, body **204** may define three blind holes **216-1**, **216-2**, and **216-3**. In some embodiments, blind holes **216-1** and **216-2** may be reamed holes that are sized and configured to receive a dowel pin, and blind hole **216-3** may also be a reamed hole for receiving a dowel pin or blind hole **216-3** may be threaded for engaging a screw as described below.

Aperture **206** may have a circular cross sectional area and include a shoulder **218** having a reduced diameter compared to aperture **206** and includes an anti-rotational feature **220** as best seen in FIG. **20**. Anti-rotational feature **220** of shoulder

218 may include one or more flats or other geometric structure(s) to prevent tibial drill guide **202** from rotating with respect to tibial drill guide mount **200** when tibial drill guide **202** is disposed within aperture **206**.

Extending from body **204** of tibial drill guide mount **200** are tibial engagement structure **222** and talar engagement structure **224**. The outer surface **226** of tibial engagement structure **222** may have a rectangular shape that is substantially planar, and the internal and substantially conformal engagement surface **228** of tibial engagement structure **222** may be somewhat convex for engaging the tibia **16** of the patient. Tibial engagement structure **222** may define one or more holes **230** for receiving a k-wire or pin as described below.

Talar engagement structure **224** may also include a substantially planar and rectangular outer surface **232**. The lower portion **234** of talar engagement structure **224** may be a conformal surface having a geometry that matches the geometry of the talar bone **14** (FIG. **14**). Talar engagement structure **224** may also define one or more holes **236** sized and configured to receive a k-wire as described below.

Tibial drill guide **202** illustrated in FIGS. **21-23** is preferably fabricated from a material having more structural integrity than tibial drill guide mount **200** to enable drill guide **202** to guide a drill bit without being damaged. Examples of materials include, but are not limited to, metals, ceramics, or the like. Drill guide **202** has a cylindrically shaped first portion **238** that is sized and configured to be received within the portion of aperture **206** that extends through the shoulder or reduced diameter area **218**. A second portion **240** of drill guide **202** has a larger cross-sectional diameter than first portion **238** and is sized and configured to be received within aperture **206** of tibial drill guide mount **200**. A flat **242**, which is best seen in FIGS. **21** and **23**, is formed along an exterior surface **244** of first portion **238** of drill guide **202**. The internal surface **248** of second portion **240** of tibial drill guide **202** has a conical shape that intersects and communicates with aperture **246** such that a drill or reamer may be received through drill guide **202**.

As with the digital image models **50** disclosed above, and considering a generalized digital model of a tibial resection guide mount **100** added to the patient's lower tibia image data, the anatomic surface features of the patient's lower tibia, e.g., the surface topography, may be complementarily mapped onto each of conformal bone engaging surfaces **116** of arms **110**, **112**, and central post **114**, i.e., the surfaces that will engage the bones unique surface topography, of tibial resection guide mount **100**. It will be understood that complementary mapping of the digital images results in localized prominences on the surface of a bone becoming localized concavities on conformal bone engaging surfaces **116** of arms **110**, **112**, and central post **114** of tibial resection guide mount **100**, while localized concavities on the surface of a bone become localized prominences on conformal bone engaging surfaces **116** of arms **110**, **112**, and central post **114**.

Each of conformal bone engaging surfaces **116** of arms **110**, **112**, and central post **114** of resection guide mount **100** is redefined with a complementary, substantially mirror image of the anatomic surface features of a selected region of the patient's lower tibia **16a**. As a consequence of this complementary bone surface mapping, tibial resection guide mount **100** releasably "locks" on to the complementary topography of the corresponding portion of the patient's natural tibia without the need for other external or internal guidance fixtures. In other words, the mating of bone surface asperities in their corresponding concavities formed in con-

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formal bone engaging surfaces **116** of tibial resection guide mount **100** ensures that little or no relative movement, e.g., slipping sideways, occurs between tibial resection guide mount **100** and the tibial surface.

A substantially identical mapping is carried out in connection with the design of a patient specific talar resection guide mount **102** and tibial drill guide mount **200**. Notably, the mapping for the design of tibial drill guide mount **200** is performed by extrapolating where the resections to the tibia **16** and talus **14** will be made using tibial and talar resection guide mounts **100** and **102** and mapping the tibial drill guide mount **200** onto the extrapolated geometry of the tibia and talus.

A visual presentation of the virtual alignment results between the patient's lower tibia **16a** and resection guide mount **100**, the patient's upper talus **14a** and resection guide mount **102**, and the proposed resected area that is to be created by resecting the talus **14** and tibia utilizing the tibial resection guide mount **100** and the talar resection guide mount **102** are created and forwarded to the surgeon to obtain approval of the results prior to manufacturing. Additionally, the surgeon may be provided with a visual representation of the virtual alignment results between the proposed resected joint space and tibial drill guide mount **200** are created and forwarded to the surgeon to obtain approval of the results prior to manufacturing. Upon receipt of the surgeon's approval, resection guide mount **100**, resection guide mount **102**, and tibial drill guide mount **200** are manufactured and returned to the surgeon for use in the surgery.

During a total ankle replacement, for example, the surgeon makes an anterior incision to gain initial access to the ankle joint. The surgeon orients tibia resection guide mount **100** on lower tibia **16a** until the conformal bone engaging surfaces **116** of arms **110**, **112** and central post **114** of tibial resection guide mount **100** securely engage one another so as to releasably "interlock" with the topography of the exposed surface of lower tibia **16a** as best seen in FIGS. 5-7. With tibial resection guide mount **100** locked onto the patient's lower tibia **16a**, a surgeon press-fits an appropriately configured distal resection guide **132** in guide receptacle recess **108** of tibial resection guide mount **100**. This results in the resection guide mount **100** being sandwiched between the resection guide **132** and the patient's bone tibia **16a** (FIGS. 5 and 6). With the resection guide mount **100** accurately positioned with respect to the selected bone region and resection guide mount **100** construct appropriately secured to the patient's bone by virtue of the mating of bone surface asperities in their corresponding concavities formed in conformal bone engaging surfaces **116**, the surgeon uses a conventional surgical blade **60** and the resection slots **128** and **130** of resection guide **132** to resect the patient's bone **16** (FIGS. 7 and 8).

In a similar fashion, when talar resection guide mount **102** is added to the patient's talar image data, the anatomic surface features of the patient's upper talus, e.g., the surface topography, may be complementarily mapped onto conformal bone engaging surface **144**. It will again be understood that complementary mapping of the digital images results in localized prominences on the surface of a bone becoming localized concavities on conformal bone engaging surface **144**, while localized concavities on the surface of a bone become localized prominences on conformal bone engaging surface **144**. In this way, conformal bone engaging surface **144** is redefined with a complementary, substantially mirror image of the anatomic surface features of a selected region of the patient's lower tibia. As a consequence of this

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complementary bone surface mapping, talar resection guide mount **102** releasably "locks" on to the complementary topography of the corresponding portion of the patient's natural talus without the need for other external or internal guidance fixtures.

To continue the total ankle replacement the surgeon orients resection guide mount **102** on upper talus **14a** until conformal bone engaging surface **144** of resection guide mount **102** "locks" to the topography of the exposed surface of upper talus **14a** (FIG. 11). With resection guide mount **102** locked onto the patient's upper talus, a surgeon press-fits an appropriately configured distal resection guide **166** in guide receptacle recess **146** of talar resection guide mount **102**. This results in resection guide mount **102** being sandwiched between resection guide **166** and the patient's bone **14** (FIGS. 12 and 13). With the resection guide mount **102** accurately positioned with respect to the selected bone region and resection guide **166** and guide mount **102** appropriately constructed and secured to the patient's bone, by virtue of the mating of bone surface asperities in their corresponding concavities formed in conformal bone engaging surfaces **144**, the surgeon uses a conventional surgical blade **60** and the resection slot **164** of resection guide **166** to resect the patient's bone **14** (FIGS. 13 and 14).

Once the tibia **16** and talus **14** have been resected, tibial drill guide mount **200** and tibial drill guide **202** are coupled together and installed into resected joint space **22** (FIG. 15). Tibial drill guide mount **200** and tibial drill guide **202** are coupled together by inserting first portion **238** of tibial drill guide **202** into aperture **206** defined by body **204** of tibial drill guide mount **200** (FIG. 24). Flat **242** formed on the first portion **238** of tibial drill guide **202** is aligned with anti-rotation feature **220** of shoulder **218** such that tibial drill guide **202** slides into aperture **206** until a lower surface **250** of second portion **240** of drill guide **202** contacts and abuts shoulder **218** of tibial drill guide mount **200**.

Body **204** of tibial drill guide mount **200**, in which tibial drill guide **202** is disposed, is inserted into resected joint space **22** in an anterior posterior direction with chamfers **212** sliding along resected areas of tibia **16** formed by utilizing slots **140** of tibial resection guide **132** as best seen in FIGS. 25A and 25B. The assemblage of tibial drill guide mount **200** and tibial drill guide **202** are slid into resected joint space **22** until talar engagement structure contacts talus **14**. A surgeon may move tibial guide mount **200** within resected joint space until conformal surface **228** is appropriately secured to the patient's bone by virtue of the mating of bone surface asperities in their corresponding concavities formed in conformal bone engaging surface **228**. Once properly located, k-wires **62** may be inserted into holes **230** and/or holes **236**, respectively defined by tibial engagement structure **222** and talar engagement structure **224**, to secure the assemblage of the tibial drill guide mount **200** and tibial drill guide **202** to the patient's tibia **16** and talus **14** as illustrated in FIG. 25C.

With tibial drill guide mount **200** and tibial drill guide **202** secured within resected joint space **22**, the patient's leg is inserted into a foot holder and alignment tool **300**. FIGS. 26-28B illustrate one example of an alignment tool **300**, which serves the task of supporting the ankle joint during a prosthesis installation procedure. Alignment tool **300** includes a foot holder assembly **302** and a leg rest **304**. Foot holder assembly **302** includes a foot rest **306**, to which the foot is secured by a foot clamp **310** and heel clamps **308** during a prosthesis installation procedure. The calf of the leg is suitably secured to the leg rest **304** once the ankle joint has been resected and tibial drill guide mount **200** and tibial drill guide **200** have been installed. Together, foot holder

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assembly 302 and leg rest 304 hold the foot and ankle relative to the leg during an installation procedure.

As shown in FIG. 26, foot holder assembly 302 is sized and configured for pivoting, under control of the physician, from a vertical or upright condition (shown in solid lines in FIG. 26) toward a more horizontal or tilted condition (shown in phantom lines in FIG. 26). In the upright condition, assembly 302 serves to hold the ankle joint in a desired orientation with respect to the natural anterior-to-posterior and medial-to-lateral axes.

As best seen in FIG. 27, foot holder assembly 302 includes a back plate 312 and a mid-plate 314, which is sandwiched between foot rest 306 and back plate 312. Mid-plate 314 is coupled to the foot rest 306 by sliding dovetail couplings 316 for up-and-down (i.e., vertical) movement relative to foot rest 306. A pair of oppositely spaced alignment rods 318 is carried by the mid-plate 314.

Alignment rods 318 are disposed in the same horizontal plane and extend from mid-plate 314 through vertically elongated slots 320 defined by foot rest 306 such that rods 318 are disposed on opposite sides of the tibia in the medial-to-lateral plane when a foot is supported by foot holder assembly 302. Vertical movement of mid-plate 314 moves alignment rods 318 up-and-down in unison within slots 320 on opposite sides of the foot rest 306 (FIG. 28A).

Back plate 312 is coupled to mid-plate 314 by sliding dovetail couplings 322 for side-to-side (i.e., horizontal) movement relative to foot rest 306 as illustrated in FIG. 28B. Back plate 312 also carries a bushing 324, which extends through openings 326 defined by mid-plate 314 and foot rest 306 and terminates at or near the plane of the foot rest 306 against which the bottom of the foot contacts. The center of the bushing 324 coincides with the intersection of the horizontal plane of the rods 318.

An adapter bar 400 for coupling tibial drill guide mount 200 to alignment tool 300 is illustrated in FIG. 29. Adapter bar 400 includes an elongate body 402 linearly extending from a first end 404 to a second end 406. Each of the ends 404, 406 includes a respective extension 408, 410 that extends from elongate body 402 at an angle. In some embodiments, extensions 408 and 410 orthogonally extend from elongate body 402, although one skilled in the art will understand that extensions 408 and 410 may diverge from elongate body 402 at other angles. In some embodiments, elongate body 402 may not have a linear shape, but may have a curved or arced shape as will be understood by one skilled in the art.

Each extension 408 and 410 defines a respective hole 412, 414 that is sized and configured to slidably receive alignment rods 318 that extend from alignment tool 300. Elongate body 402 defines one or more holes 416-1, 416-2, and 416-3 (collectively referred to as "holes 416") for coupling to adapter bar 400 to tibial drill guide mount 200. In some embodiments, the one or more holes 416 align with one or more holes 216 defined by body 204 of tibial drill guide mount 200 such that a pin or other device for maintaining the alignment and engagement of adapter bar 400 and tibial drill guide mount 200. For example, holes 216-1 and 216-2 of tibial drill guide mount 200 align with holes 416-1 and 416-2 of adapter bar 400, and hole 216-3 of drill guide mount 200 aligns with hole 416-3 of adapter bar 400. Dowel pins 70 (shown in FIG. 25C) may be inserted into holes 216-1 and 416-1 as well as into holes 216-2 and 416-2 to align tibial drill guide mount 200 with adapter bar 400 in both the horizontal and vertical directions (e.g., in the x- and y-directions), and a screw (not shown) may be inserted through

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hole 416-3 into threaded hole 216-3 to secure tibial drill guide mount 200 to adapter bar at the proper height or depth (e.g., in the z-direction).

With tibial drill guide mount 200 and tibial drill guide 202 disposed within the resected ankle space 22, the foot and lower leg are placed in foot rest 306 and leg rest 304 (FIG. 30). The physician estimates the ankle's axis of dorsi-plantar rotation and visually aligns the ankle to the axis of rotation of the alignment tool 300. Foot rest 306 is adjusted to rotate the foot so that the big toe is essentially pointing in a vertical direction with respect to the leg that extends in a horizontal direction. The forefoot and heel are secured to foot rest 306 with clamps 308 and 310. Leg rest 304 is adjusted to the calf so that the tibia 16 is approximately parallel to the floor. The foot and calf are desirably aligned so that the anterior-posterior ("A-P") line of the talus's trochlea is essentially vertical.

Adapter bar 400 is coupled to alignment tool 300 by aligning holes 412 and 414 that are respectively defined by extensions 408 and 410 with alignment rods 318 of alignment tool 300. Adapter bar 400 is then slid along alignment rods 318 until holes 416 of adapter bar align with holes 216 defined by body 204 of tibial drill guide mount 200 (FIG. 30). As described above, dowel pins 70 are inserted into holes 416-1 and 416-2 of adapter bar 400 and holes 216-1 and 216-2 of tibial drill guide mount 200. With dowels 70 disposed within holes 216-1, 216-2, 416-1, and 416-2, tibial drill guide mount 200 is properly aligned with alignment tool 300 in the medial lateral (e.g., x-direction) and superior-inferior (e.g., y-direction) directions. A screw is inserted through hole 416-3 into threaded hole 216-3, which secures tibial drill guide mount 200 to adapter bar 400 and provides proper alignment in the anterior-posterior direction (e.g., the z-direction).

With the patient's foot disposed within alignment tool 300, bushing 324 on back plate 312 establishes alignment with the mechanical axis of tibia 16 and alignment of rods 318. Thus, after using adapter bar 400 to align tibial drill guide mount 200 with alignment tool 300 as described above, in line drilling of the center of the ankle and tibia for introduction of a bottom foot cannula is made possible without the use of fluoroscopy since aperture 246 of tibial drill guide 202 disposed within tibial drill guide mount 200 is aligned with an axis defined by bushing 324. Such arrangement enables an intramedullary channel to be formed that is substantially collinear with a mechanical axis defined by the tibia.

Various minimally invasive surgical techniques may be used to introduce a bottom foot cannula into the calcaneus 20, talus 14, and tibia 16. In one representative embodiment, bushing 324 is temporarily separated from the back plate 312 (e.g., by unscrewing) to provide access to the bottom of the foot. The physician uses a scalpel to make an initial incision in the bottom of the foot and replaces bushing 324. A cannulated trocar loaded with a k-wire (not shown) can be inserted through bushing 324, into the bottom of the foot, until the calcaneus 20 is contacted and the k-wire is firmly set within the calcaneus 20. The trocar can then be removed, and the k-wire lightly tapped further into the calcaneus 20. In a representative embodiment, the bushing 324 measures 6 mm in diameter, and the cannulated trocar can be 6 mm loaded with a 2.4 mm k-wire. The physician can now operate a cannulated first reamer (e.g., 6 mm) (not shown) over the k-wire up into the calcaneus 20 and talus 14. The first reamer opens an access path for insertion of a bottom foot cannula.

After withdrawing the first reamer and bushing 324, the physician then inserts a bottom foot cannula 64 as shown in

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FIG. 30. With the bottom foot cannula **64** in place, a second reamer **66** (e.g., 5 mm) can be operated through the cannula **64** to drill approximately another 100 mm through the talus **14** and up into the tibia **16** to establish an intramedullary guide path through the calcaneus **20** and talus **14** leading into the tibia **16** (FIG. 30). As second reamer **66** is advanced towards tibia **16**, the tip **68** of reamer **66** is guided by the conical interior surface **248** of tibial drill guide **204**, which is aligned with bushing **324** of alignment tool **300**.

Once an intramedullary channel through the calcaneus **20**, talus **14**, and tibia **16** has been established, adapter bar **400** is decoupled from drill guide mount **200** and alignment rods **318**. Drill guide mount **200** is removed from resected joint space **22** to expose the resected joint space to the surgeon.

With the resected ankle joint space **22** exposed to the surgeon, an ankle prosthesis is then installed. In one example, the ankle prosthesis includes a stem that may extend from the bottom of the calcaneus up to the top of the talus (i.e., a talo-calcaneal stem), although in some embodiment the stem is completely disposed within the talus (i.e., a talar stem). A convex dome is coupled to the stem and provides an articulating joint surface. A tibial stem may be monolithic or include a plurality of segments that may be coupled together in situ. A tibial platform couples to the tibial stem and either includes or is coupled to a convex joint surface for articulating with the articulating joint surface coupled to the talar/talo-calcaneal stem. Examples of such ankle prosthesis and methods of installing such prosthesis are disclosed in U.S. Pat. No. 7,534,246 issued to Reiley et al., the entirety of which is herein incorporated by reference.

The disclosed tibial drill guide mount **200** and drill guide **202** may be used with a variety of alternative alignment tools. For example, FIGS. 31-34 illustrate another example of an alignment tool in the form of a foot holder assembly **500** to which tibial drill guide mount **200** may be directly coupled. As shown in FIGS. 31 and 32, foot holder assembly **500** includes a base plate **502** defining a plurality of slots **504** and **506** and an aperture **503**.

Slots **504** are sized and configured to slidably receive a pair of heel clamps **508**, and slots **506** are sized and configured to slidably receive a pair of forefoot clamps or guides **510**. Heel clamps **508** and forefoot clamps **510** cooperate to maintain a foot of a patient in a desired position with respect to base plate **502** by utilizing a locking mechanism such as, for example, a set screw or other locking device, to fix the position of heel clamps **508** and forefoot clamps **510** to base plate **502**. The respective foot engaging surfaces **512** and **514** of heel clamps **508** and forefoot clamps **510** may have a shape that complements the medial and lateral shape of a human foot.

Extending from base plate **502** are a pair of alignment rods **516** that are arranged on base plate **502** such that one alignment rod is disposed on a medial side of a patient's foot and the other alignment rod is disposed on a lateral side of a patient's foot. A coupling bar **518** is sized and configured to slidably engage alignment rods **516** as best seen in FIGS. 32 and 34. Coupling bar **518** includes a pair of spaced apart legs **520** that define channels **522** (FIG. 32) in which alignment rods **516** are slidably received. One or both of legs **520** include a clamp or other locking mechanism **524** for increasing the friction between coupling bar **518** and alignment rods **516** in order to releasably lock coupling bar **518** at a certain position along the length of alignment rods **516**.

Medial-lateral cross bar **526** couples together legs **520** of coupling bar **518**. Extending from medial-lateral cross bar **526** is mount coupling member **528**. Mount coupling member **528** includes one or more holes **530-1**, **530-2**, and **530-3**

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(collectively referred to as "holes **530**") that are sized and configured to align with holes **216** defined by tibial drill guide mount **200**.

A peg **532** (FIG. 33) extends from medial-lateral cross bar **526** for coupling shin engaging member **534** via slot **536** defined by shin engaging member **534**. Shin engaging member **534** includes a shelf **538** having a concave surface **540** for abutting a shin of a patient. A nut or other locking mechanism (not shown) for engaging peg **532**, which may be threaded, may be used to fix the position of shelf **538** relative to medial-lateral cross bar **526**.

The use of foot holder assembly **500** in connection with the assemblage of tibial drill guide mount **200** and tibial drill guide **202** is similar to the use of alignment tool **300** described above. For example, once the assembly of tibial drill guide mount **200** and tibial drill guide **202** are disposed within resected joint space **22**, the heel of the patient's foot is placed between heel clamps **508** and the patient's forefoot is placed between forefoot clamps **510**. The locking mechanisms of heel and forefoot clamps **508** and **510** may be engaged to initially set positions of heel and forefoot clamps **508** and **510** relative to base plate **502**.

Holes **530** of coupling member **528** are aligned with holes **216** defined by tibial drill guide mount **200** by sliding legs **520** of coupling bar **518** along alignment rods **516**. Dowel pins **70** and/or a threaded screw (not shown) may be used to couple holes **530** of coupling member **528** to holes **216** of tibial drill guide mount **200**. The surgeon may check to ensure that the patient's foot is firmly against base plate **502** and then engage clamps **524** such that coupling bar **518** is fixed to alignment rods **516**.

Shin engaging member **534** is adjusted until concave surface **540** contacts the patient's shin. The adjustment of shin engaging member **534** is guided by the engagement between slot **536** and peg **532**. With shin engaging member **534** in the desired position, the nut or other locking mechanism (not shown) locks shin engagement member **534** in place. The surgeon may make final adjustments to the heel and forefoot clamps **508** and **510** and then create the intramedullary channel as described above.

Another example of an alignment tool **600** for use with tibial drill guide mount **200** and tibial drill guide **202** is illustrated in FIGS. 35-38. As shown in FIG. 35, alignment tool **600** includes a base plate **602** comprising a plurality of bars **602a**, **602b**, and **602c**. Although three bars **602a**, **602b**, and **602c** are illustrated, one skilled in the art will understand that fewer or more bars may be implemented. Bar **602b** defines a hole **603** sized and configured to receive a surgical tool, such as, for example, a cannulated drill. Additional elements including, but not limited to, heel clamps and/or forefoot clamps (not shown) may be coupled to the bars **602a**, **602b**, and **602c** of base plate **602** for aiding in the positioning of a patient's foot with respect to hole **603**.

Extending from base plate **602** is a pair of spaced apart alignment rods **604**. One of alignment rods **604** may be disposed on a medial side of a patient's leg, and the other alignment rod **604** disposed on a lateral side of the patient's leg. Alignment rods **604**, like alignment rods **318** of alignment tool **300**, may be slidably receiving within holes **412**, **414** of adapter bar **400**.

The use of alignment tool **600** in connection with the assemblage of tibial drill guide mount **200** and tibial drill guide **202** and the adapter bar **400** is similar to the use of alignment tool **300** described above. For example, once the assembly of tibial drill guide mount **200** and tibial drill guide **202** are disposed within resected joint space **22**, adapter bar **400** is coupled to alignment tool **600** by aligning holes **412**

and **414** that are respectively defined by extensions **408** and **410** with alignment rods **604** of alignment tool **600**. Adapter bar **400** is slid along alignment rods **604** until holes **416** of adapter bar align with holes **216** defined by body **204** of tibial drill guide **200**. As described above, dowel pins are inserted into holes **416-1** and **416-2** of adapter bar **400** and **216-1** and **216-2** of tibial drill guide mount **200**. With dowels disposed within holes **216-1**, **216-2**, **416-1**, and **416-2**, tibial drill guide mount **200** is properly aligned with alignment tool **600** in the medial lateral (e.g., x-direction) and superior-inferior (e.g., y-direction) directions. A screw is inserted through hole **416-3** into threaded hole **216-3**, which secures tibial drill guide mount **200** to adapter bar **400** and provides proper alignment in the anterior-posterior direction (e.g., the z-direction). The surgeon may make final adjustments to the heel and forefoot clamps **508** and **510** and then create the intramedullary channel as described above.

FIGS. **39-63** illustrate another embodiment of a system for performing a surgical procedure. Specifically, FIGS. **39-43** illustrate a tibial drill guide mount **700** sized and configured to receive the tibial drill guide cartridge **702** illustrated in FIGS. **44-47**. Tibial drill guide mount **700** may also receive other drill guide cartridges for use during other stages of the surgical procedures. Like tibial drill guide mount **200**, tibial drill guide **700** may be fabricated from a resilient polymer material of the type that is suitable for use in connection with stereo lithography, selective laser sintering, or the like manufacturing equipment, e.g., a polyamide powder repaid prototype material is suitable for use in connection with selective laser sintering.

As shown in FIG. **39-43**, tibial drill guide mount **700** has a somewhat rectangular body **704** having a front side **706**, a rear side **708**, top side **710**, bottom side **712**, and a pair of opposed sides **714** and **716**. Front side **706** defines a recess **718** sized and configured to slidably receive tibial drill guide **702** therein. Recess **718** communicates with a recess **720** (FIGS. **39** and **43**) defined by bottom side **712** and a recess **722** (FIGS. **39**, **42**, and **43**) defined by top side **710** such that body **704** is substantially hollow.

The respective inner surfaces **724**, **726** of sides **714**, **716** have different geometries that correspond with the cross-sectional geometry of tibial drill guide cartridge **702** to ensure that tibial drill guide cartridge **702** is properly inserted into recess **718**. In the embodiment illustrated in FIGS. **39-43**, side **716** includes first and second ledges **728**, **730** that inwardly extend into recess **718**, and side **714** has an inwardly tapered upper region **732** and an inwardly extending ledge **734**. One skilled in the art will understand that sides **714**, **716** may include other features for ensuring proper insertion of tibial drill cartridge **702** into recess **718**. In some embodiments, sides **714**, **716** may have the identical geometry and tibial drill guide cartridge may be reversibly inserted into recess **718**.

Front side **706** defines one or more dowel holes **736-1**, **736-2** (collectively referred to as "dowel holes **736**") sized and configured to receive a dowel pin **70** therein. One or more through holes **738-1**, **738-2**, **738-3** (collectively referred to as "through holes **738**") extend through front side **706**, which also defines a blind hole **740**. Through holes **738** are sized and configured to receive k-wires for pinning tibial drill guide mount to a patient's bone as described below.

Top side **710** of tibial drill guide mount **700** includes a pair of chamfers **742** that are sized and configured to be mate against and reference the resected surfaces of the lower tibia **16a** (FIG. **8**). Tibial drill guide mount **700** also includes a tibial engagement structure **744** and a talar engagement structure **746**. Tibial engagement structure **744** extends from

top side **710** and includes a substantially conformal engagement surface **748**. Talar engagement structure **746** extends from bottom side **712** and also includes a substantially conformal engagement surface **750**.

Tibial drill guide cartridge **702** has a substantially rectangular elongate body **754** that may be formed from a more substantial material than tibial drill guide mount **700** such as, for example, metals, ceramics, or the like. As best seen in FIGS. **44** and **45**, the geometry of sides **756**, **758** is respectively complementary to the sides **714**, **716** of tibial drill guide mount **700**. For example, side **758** includes ledges **760** and **762** that respectively correspond to ledges **728** and **730**, and side **756** includes a ledge **764** and an angled section **766**, which respectively correspond to ledge **734** and upper region **732** of tibial drill guide mount **700**.

Front side **768** of tibial drill guide cartridge **702** defines a blind hole **770**, which may be threaded for reasons described below. Tibial drill guide cartridge **702** defines a pair of holes **772** and **774** that extend from bottom surface **776** to top surface **778**. Hole **772** may be a reamed hole that is sized and configured to receive a ball detent therein, and hole **774** has an internal surface **780** that tapers from a larger diameter at bottom surface **776** to a smaller surface that is sized and configured to receive a surgical tool, such as a drill and/or reamer. Top surface **778** defines a pair of parallel slots **782-1**, **782-2** (collectively referred to as "slots **782**") that extend from side **756** to side **758**. As best seen in FIGS. **44** and **47**, slots **782** are disposed equidistant from a central axis defined by hole **774** to provide a visual key for a physician that wants check the alignment of hole **774** with a mechanical axis of a patient's tibia using fluoroscopy.

As illustrated in FIGS. **48**, a mounting plate **800** has a substantially rectangular body **802** that is fabricated from a material including, but not limited to, metals, ceramics, or the like. Body **802** defines an aperture **804** that extends from front side **806** to back side **808** and has a similar geometry of recess **718** of tibial drill guide mount **700** such that tibial drill guide cartridge **702** may be received therein. Body **802** also defines a pair of through holes **810-1**, **810-2** (collectively referred to as "holes **810**") that are arranged on body **802** such that they correspond to holes **738** of tibial drill guide mount **700** and are sized and configured to receive a k-wire or pin therein.

A mounting base **812** extends from front side **806** of mounting plate **800** and defines a hole **814** that extends from a first side **816** to a second side **818**. Mounting base **812** defines a notch **820** and one or more dowel pin holes **822-1**, **822-2** (collectively referred to as "holes **822**") that are aligned with holes **736** of tibial drill guide mount **700**. Notch **820** bisects hole **814**. Mounting base **812** may also define one or more recesses **824** that correspond to one or more protrusions **784** that extends from front side **706** of tibial drill guide mount **700**. Recesses **824** and protrusions **784** cooperate to ensure that mounting base **812** and tibial drill guide mount **700** are properly aligned. One skilled in the art will understand that other geometric features may be implemented to ensure proper alignment between mounting base **812** and tibial drill guide mount **700**.

As illustrated in FIGS. **49-54**, mounting plate **800** may be coupled to tibial drill guide mount **700** using dowel pins **70**, which are received through holes **822** and **734**. Tibial drill guide cartridge **702** is received through aperture **804** and recess **718** as best seen in FIG. **51**. FIGS. **53** and **54** illustrate that when tibial drill guide cartridge **702** is properly inserted into the assemblage of tibial drill guide mount **700** and mounting plate **800**, hole **772** aligns with hole **828** defined by mounting plate **800**, which may include a ball detent (not

shown) disposed therein. Consequently, the ball detent is received within hole 772 to retain tibial drill guide cartridge 702 disposed within aperture 804 and recess 718 such that hole 774 is disposed within recesses 754 and 756. A screw or other threaded object (not shown) can be inserted into threaded hole 770 and then pulled to remove tibial drill guide cartridge 702 from aperture 804 and recess 718 as illustrated in FIGS. 53 and 54.

Tibial drill guide mount 700, tibial drill guide 702, and mounting plate 800 may be used in connection with alignment tool 300, adapter bar 400, foot holder assembly 500, and alignment tool 600 as described above. Additionally, tibial drill guide mount 700, tibial drill guide 702, and mounting plate 800 may also be used in conjunction with foot holder assembly 900 illustrated in FIGS. 55-60 as can

As shown in FIG. 55, foot holder assembly 900 includes a base plate 902 that extends from a first end 904 to a second end 906. First and second ends 904, 906 each define a pocket 908 and a hole 910. Pocket 908 is sized and configured to receive a drill bushing 912 having a cylindrical body defining hole 914 that aligns with through hole 910. Accordingly, both first end 904 and second end 906 may support an ankle or forefoot of a patient. Each pocket 908 includes a spring loaded detent 916 communicatively coupled to it that include a finger receiving surface 918 and is configured to slide relative to base plate 902 and secure drill bushing 912 within pocket 908. In some embodiments, drill bushing may be threaded and configured to be coupled to base plate 902 with complementary threads disposed on an inner surface of holes 910.

Base plate 902 also includes a medial/lateral extension 920 that extends in a substantially perpendicular direction from an approximate mid-point between first end 904 and second end 906. Base plate 902 may also define a viewing opening 922 such that a surgeon may be able to view the bottom of a patient's foot when the foot is secured to foot holder assembly 900.

One or more rods 924 extend from base plate 902 in a substantially perpendicular direction with respect to an upper foot holding surface 926 (FIG. 56). Rods 924 may be secured to base plate 902 using screws or through other securing means as will be understood by one skilled in the art. A cap 928 is secured to an upper end of rods 924 and be secured to rods 924 using screws or other fixation means.

A mounting member 930 has an elongate body 932 that defines a pair of holes 934, 936 at one end 938 that slidably receive rods 924 such that mounting member 930 may be slid along rods 924 in order to position tibial drill guide mount 700 with respect to base plate 902. A spring loaded button 940 is disposed at first end 938 of mounting member 930 and is coupled to a locking mechanism (not shown) disposed within mounting member 930 for locking mounting member 930 at a position along rods 924.

One or more holes 942 are defined at the second end 944 of mounting member 930 and correspond to holes 716 of drill guide mount 700 for coupling drill guide mount 700 to foot holder assembly 900. Second end 942 also defines a slot 946, as best seen in FIGS. 56 and 60, that is sized and configured to receive an internally threaded rod 948 of a pivoting arrangement 950, which includes a lower portion 952 that is received within slot 820 of mounting plate 800 and is cross-pinned through hole 814. The cross-pinning of pivoting arrangement 950 may pivot about an axis defined by hole 814 and is configured to receive an support tightening knob 954. Bottom surface 956 (FIG. 60) of knob 954 has an outer dimension that is greater than slot 946 and is

configured to engage mounting member 930 in order to secure the assemblage of mounting plate 800 and tibial drill guide mount 700, which may include tibial drill cartridge 702.

In operation, tibial drill guide mount 700 is inserted into resected joint space 22. Mounting plate 800 is connected to tibial drill guide mount 700 using dowel pins 70 as best seen in FIGS. 49 and 50. With pivoting arrangement 950 cross-pinned to mounting plate 800, the assemblage of mounting plate 800 and pivoting arrangement 948 is coupled to tibial drill guide mount with dowel pins 70, which may be press fit into holes 822 of mounting plate 800 and holes 716 of tibial drill guide mount 700 as will be understood by one skilled in the art. Tibial drill guide mount 700 and mounting plate may be secured within resected joint space 22 by inserting k-wires (not shown) into holes 736, 790 defined by tibial drill guide mount 700 and holes 830-1, 830-2 (corresponding to holes 736-1, 736-2) and 832-1, 832-2 defined by mounting plate 800.

With mounting plate 800 coupled to tibial drill guide mount 700 that is disposed within resected joint space 22, pivoting arrangement 948 is rotated such that it extends in a direction approximately parallel to a longitudinal axis defined by a patient's leg and the cartridge-style tibial drill guide 702 is inserted into aperture 804 of mounting plate 800 and recess 718 of tibial drill guide mount 700. Tibial drill guide cartridge 702 is inserted until leading end 786 of tibial drill cartridge 702 abuts rear wall 788 of tibial drill guide mount 700 at which point the ball detent disposed within hole 772 engages hole 828 defined by mounting plate 800 and the front side 768 of tibial drill guide cartridge 702 is flush with front side 806 of mounting plate 800.

Holes 940 of mounting member 930 are aligned with, and received over, dowel pins 70 that extend from front side 806 of mounting plate to couple mounting member 930 of foot holder assembly 900 to the assemblage of mounting plate 800, tibial drill guide mount 700, and tibial drill guide cartridge 702. With mounting member 903 coupled to dowel pins 70 and mounting plate 800, pivoting arrangement 948 is rotated with respect to mounting plate 800 such that rod 946 of pivoting arrangement 948 is received within slot 944 of mounting member 930. Knob 952 is then rotated about its axis (clockwise or counterclockwise) such that the bottom surface 954 of knob 952 contacts mounting member 930 to maintain engagement between mounting member 930 and the assemblage of tibial drill guide mount 700 and mounting plate 800.

Drill bushing 912 is coupled to hole 910 that is aligned with the heel of a patient's foot. As described above, drill bushing 912 may be slid into pocket 908 defined by bottom plate 902 until spring loaded detents 916 releasably lock drill bushing 912 in place. In some embodiments, drill bushing 912 may be screwed into base plate 902 by way of corresponding threads disposed on an outer surface of drill bushing 912 that engage threads defined by an inner surface of pocket 908 and/or hole 910. With drill bushing 912 in place and the patient's leg secured to foot holder assembly 900, various minimally invasive surgical techniques may be used to introduce a bottom foot cannula into the calcaneus 20, talus 14, and tibia 16 as described above.

Once access to the patient's calcaneus has been achieved, a bottom foot cannula 64 is inserted through the patient's calcaneus 20. A reamer 66 is operated through the cannula 64 to drill approximately another through the talus 14 and up into the tibia 16 to establish an intramedullary guide path through the calcaneus 20 and talus 14 leading into the tibia 16. As reamer 66 exits talus 14, the conically shaped internal

surface **748** guides the tip **68** into hole **788**. An axis defined by hole **788** is substantially axially aligned with a mechanical axis of tibia **16** such that as reamer **66** is extended through hole **788**, it bores an intramedullary canal within tibia **16**.

Reamer Stabilizer

FIGS. **63-70** illustrate one example of a reamer stabilizer **1000** that may be used to stabilize the reamer as it is advanced into the tibia of a patient. Referring first to FIG. **63**, reamer stabilizer **1000** includes an elongate body **1002** extending from a distal end **1004** to a proximal end **1006**. As best seen in FIGS. **64** and **65**, body **1002** defines a longitudinal channel **1008** extending along the length of body **1002**. Body **1002** also defines a pair of cavities **1010**, **1012** for receiving buttons and biasing members as described in greater detail below.

Distal end **1004** of body **1002** includes a pair of spaced apart prongs **1014**, **1016**. In some embodiments, prong **1014** has a length that is longer than a length of prong **1016**. As shown in FIGS. **64** and **65**, longitudinal channel **1008** extends along prong **1016**. A notch **1018** is defined between prongs **1014** and **1016**.

Proximal end **1004** includes a handle **1020** that extends at from body **1002** at an angle relative to the longitudinal axis defined by body **1002** as best seen in FIG. **63**. Opposite handle **1020** proximal end **1006** includes a cutout region **1022** defined by a pair of perpendicular walls **1024**, **1024** as illustrated in FIG. **66**. Although walls **1024** are illustrated and described as perpendicular to one another, one of ordinary skill in the art will understand that cutout region **1022** may be defined by walls having other configurations.

In some embodiments, body **1002** has a rectangular cross-sectional geometry defined by four sides **1026**, **1028**, **1030**, **1032**. Opposed sides **1026**, **1028** each include a respective step **1034**, **1036** along their respective lengths. Steps **1034**, **1036** are positioned at a same distance from notch **1018**.

Opposed sides **1030**, **1032** defines holes **1038**, **1040**, **1042** each configured to receive a respective pin **1044**, **1046**, **1048** in a press-fit engagement as described below. Hole **1038** is positioned near proximal end **1006**. Hole **1040** is disposed adjacent to wall **1026** and step **1034**. Hole **1042** is formed in prong **1016**. In some embodiments, opposed sides **1030**, **1032** define an opening **1050**, which reduces the overall weight of reamer stabilizer **1000** and provides a surgeon or user with additional surfaces to manipulate reamer stabilizer **1000**.

As best seen in FIGS. **64** and **65**, a slidable guiding assembly **1052** is disposed within longitudinal channel **1008**. Guiding assembly **1052** includes a reamer guide body **1054** that is pivotably coupled to stabilizer body **1002** by pin **1048**, which is received within hole **1056**. Reamer guide body **1054** includes a concave guiding surface **1058** disposed adjacent to hole **1056**. Opposite concave guiding surface **1058** guide body **1054** includes a step **1060**, which is disposed adjacent to a hole **1062**, which is defined in a forked end **1064** of guide body **1054**. Formed end **1064** is formed by a pair of spaced apart tabs **1066**, **1068** that together define a recess **1070** therebetween.

A pin **1072** (FIGS. **66** and **67**) is received in hole **1062** of reamer guide body **1054** for pivotably coupling reamer guide body **1054** to pivot rod **1074**, which includes a corresponding hole **1076** at its distal end **1078**. Pivot rod **1074** defines another hole **1080** at its proximal end **1084**. Hole **1080** is sized and configured to receive a pin **1082** for coupling pivot rod **1074** to plunger rod **1086**.

As best seen in FIGS. **66-69**, plunger rod **1086** defines a hole **1088** at its distal end **1090**, which has a flared geometry relative to the remainder of plunger rod **1086**. In some embodiments, distal end **1090** includes a pair of opposed flats **1092**, **1094** and defines a slot **1096** as best seen in FIG. **69** in which pivot rod **1074** is received. Distal end **1090** also forms a shoulder **1091** configured to maintain plunger rod **1086** within longitudinal channel **1008** as described in greater detail below.

Proximal end **1098** of plunger rod **1086** defines a hole **1100** that is size and configured to receive a pin **1118** for coupling plunger rod **1086** to head **1102**. Head **1102** defines a blind hole **1104** that inwardly extend from distal end **1106** and is sized and configured to receive proximal end **1098** of plunger rod **1086** therein. In some embodiments, top side **1108** of head **1102** includes an angled surface **1110** that terminates at side **1112**. Head **1102** also includes an arced surface **1114** for providing an ergonomic contour to a user's finger (FIG. **70**). A hole **1116** extends through head **1102** in a direction that is perpendicular to the direction in which blind hole **1104** extends and is configured to align with hole **1100** of plunger rod **1086** for coupling plunger rod **1086** to head **1102** using pin **1118**. Although a cross-pin arrangement is described, one of ordinary skill in the art will understand that other coupling means may be used to couple head **1102** to plunger rod **1086** including, but not limited to, a taper fit, ultrasonic welding, a snap fit arrangement, or the use of adhesive to list but only a few possibilities.

A biasing member **1120** is configured to be disposed over plunger rod **1086** and abut the distal end **1106** of head **1102**. In some embodiments, biasing member **1120** is a compression spring that applies a biasing force to head **1102** in a proximal direction as biasing member **1120** is disposed between distal end **1106** and a reduced diameter area **1009** of longitudinal channel **1008**.

Turning now to FIG. **70**, locking assembly **1122** is disposed at the proximal end **1006** of reamer guide body **1002** and is configured to lock guiding assembly **1052** in a position in which guiding assembly **1052** engages a reamer body **65** as described in greater detail below. Locking assembly **1122** includes a locking button **1124** slidably coupled to reamer stabilizer body **1002**. Locking button **1124** includes a lower portion **1126** that is configured to be received within cavity **1022** defined by stabilizer body **1002** and an upper portion **1128** extending above stabilizer body **1002** for facilitating engagement by a surgeon or other user.

In some embodiments, lower portion **1126** has a substantially rectangular geometry comprising a bottom surface **1130**, an internal side surface **1132**, and an outer side surface **1134**. Bottom surface **1130** is flat and configured to slide along a surface of cavity **1010**. The interface between bottom surface **1130** and outer side surface **1134** includes an angled surface **1138** that is complementary to angled surface **1110** of head **1102**. A slot **1140** is defined by sides **1142**, **1144**. Slot **1140** extends parallel to bottom surface **1130** and is sized and configured to receive pin **1044**, which is received through hole **1038** defined by stabilizer body **1002**.

In some embodiments, upper portion **1128** has a triangular shape although one of ordinary skill in the art will understand that upper portion **1128** can take on other geometric shapes. Upper portion **1128** includes a substantially flat bottom surface **1146** configured to slide along an upper or proximal-most surface of handle **1020**. Upper sides **1148**, **1150** form the other two sides of upper portion **1128**. Side **1150** is curved to facilitate ergonomic engagement with a finger of a surgeon or user.

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A biasing member 1152 is disposed within cavity 1010 and is configured to urge locking button 1124 away from handle 1020 and towards guiding assembly 1052. In some embodiments, such as the embodiment illustrated in FIGS. 66-70, biasing member 1152 is a compression spring that is disposed in cavity 1010 in an abutting relationship with inner side surface 1132 of lower portion 1126 of locking button 1124. Biasing member 1152 is positioned such in cavity 1010 such that biasing member 1152 is substantially collinear with slot 1140 to prevent rotation and jamming of locking button 1124 as will be understood by one of ordinary skill in the art.

Coupling assembly 1154 is coupled to side 1026 of stabilizer body 1002 and is configured to couple reamer stabilizer 1000 to other surgical devices as described in greater detail below. Coupling assembly 1154 includes a pivoting button 1156 and a biasing member 1158. As best seen in FIG. 66, pivoting button 1156 has an arcuate body 1160 extending from a lower end 1162 to an upper end 1164. A pair of ears 1166, 1168 extend from an approximate middle of body 1160 that together define depression 1170. Each ear 1166, 1168 defines a respective hole 1172, 1174 for receiving pin 1046.

Lower end 1162 includes a detent 1176 extending from inner surface 1178 adjacent to depression 1170. A recess 1180, which is illustrated in FIGS. 64 and 65, is defined within depression 1170 at a location that is disposed proximally of holes 1172, 1174 defined by ears 1166, 1168. Detent 1176 is disposed distally of step 1034. The relative locations of detent 1176 with respect to step 1034 and recess 1180 with respect to holes 1172, 1174 are provided for coupling reamer stabilizer 1000 to other surgical device as described in greater detail below. Concave outer surface 1182 provides an ergonomic surface for the finger of a surgeon or user of reamer stabilizer 1000 when the reamer stabilizer 1000 is to be decoupled from other surgical devices.

To assemble reamer stabilizer 1000, guiding assembly 1052 is assembled by placing pivot rod 1074 within slot 1096 at the distal end 1090 of plunger rod 1086. Pivot rod 1074 is coupled to plunger rod 1088 by inserting pin 1082 through holes 1080 and 1100. Reamer guide body 1054 is coupled to the distal end 1078 of pivot rod 1074 by inserting pin 1072 into holes 1062 and 1076.

Proximal end 1098 of plunger rod 1086 is inserted into longitudinal channel 1008 at the opening at defined by 1016 at the distal end 1004 of stabilizer body 1002. When shoulder 1091 defined by distal end 1090 contacts reduced diameter area 1009 of longitudinal channel 1008, proximal end 1098 of plunger rod 1086 outwardly extends from longitudinal channel 1008. Biasing member 1116 is inserted into longitudinal channel 1008 over plunger rod 1086 as is head 1102. Hole 1116 of head 1102 is aligned with hole 1100 of plunger rod 1086 and the two pieces are coupled together by inserting pin 1118 through holes 1116 and 1100. Reamer guide body 1054 is coupled to stabilizer body 1002 by inserting pin 1048 through holes 1042 and 1056.

With guiding assembly 1052 coupled to stabilizer body 1002, locking assembly 1122 is coupled to stabilizer body 1002. Locking assembly 1122 is coupled to stabilizer body 1002 by inserting biasing member 1152 into cavity 1010 defined by body 1002. Lower portion 1126 of locking button 1124 is inserted into cavity 1010 until slot 1140 defined by lower portion 1126 aligns with hole 1038 defined by body 1002. With slot 1140 aligned with hole 1038, pin 1044 is inserted into hole 1038 and slot 1140 to cross-pin locking button 1124 to body 1002.

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Coupling assembly 1154 is installed by inserting biasing member 1158 into cavity 1012, and pivoting button 1156 is placed over biasing member 1158 such that holes 1172, 1174 defined by ears 1166, 1168 aligns with hole 1040 defined by body 1002. With holes 1166, 1168 aligned with hole 1040, pin 1046 is inserted into the holes 1166, 1168, and 1040 to secure pivoting button 1156 to body 1002.

Foot Holder Assembly

Reamer stabilizer 1000 is configured to be used in connection with a foot holder assembly such as foot holder assembly 1200 illustrated in FIG. 71. Foot holder assembly 1200 includes a base plate 1202 having a generally rectangular shape extending from a first side 1204 to a second side 1206 and from a third side 1208 to a fourth side 1210.

A pair of biased detents 1212, 1214 are disposed at opposite ends of side 1210 and are configured to couple foot plate 1326 and drill guide assembly 1260 to one of sides 1204, 1206 of base plate 1202 as described in greater detail below. Foot plate 1326 and drill guide assembly 1260 can advantageously be coupled to either of sides 1204, 1206 such that foot holder assembly 1200 is reversible and can be used for an operation on a patient's left and/or right foot and ankle. Detents 1212, 1214 each include a respective finger-engaging surface 1216, 1218 that are manipulated by a surgeon or other user to disengage foot plate 1326 and/or drill guide assembly 1260 from base plate 1202.

Sides 1204, 1206 of base plate 1202 each define a pair of holes 1222, 1224 that are sized and configured to receive pegs 1332, 1334 of foot plate 1326 and pegs 1276, 1278 of drill guide assembly 1260 as described in greater detail below. Sides 1204, 1206, 1208, 1210 collectively define a viewing opening 1224 such that a surgeon may be able to view the bottom of a patient's foot when the foot is secured to foot holder assembly 1200.

One or more rods 1226, 1228 extend from side 1208 of base plate 1202 in a perpendicular direction with respect to the direction in which sides 1204 and 1206 extend from side 1208. In some embodiments, rods 1226, 1228 are secured to base plate 1202 using screws although one of ordinary skill in the art will understand that other securing means for securing rods 1226, 1228 to base plate 1202 can be used. A cap 1230 is coupled to the ends of rods 1226, 1228 opposite the ends to which base plate 1202 is coupled. Cap 1230 can also be coupled to rods 1226, 1228 using screws or other securement means.

A mounting member 1232 having an elongate body 1234 that defines a pair of holes 1236, 1238 at one end 1240 for slidably receiving rods 1226, 1228. A locking screw 1242 comprising a knob 1244 provides a locking mechanism for locking mounting member 1232 at a certain position along rods 1226, 1228. One or more holes 1246, 1248 are defined at the second end 1250 of mounting member 1232 and correspond to holes 736 of drill guide mount 700 and holes 822 of modified mounting plate 800A, which is described in greater detail below. Second end 1250 also defines a slot 1252 that is sized and configured to receive an internally threaded rod 948 of pivoting arrangement 950.

Drill guide assembly 1260 is now described with reference to FIGS. 72-73. Referring first to FIG. 72, drill guide assembly 1260 includes a rectangular base 1262 extending from a coupling end 1264 to an opposite end 1266. Sides 1268, 1270 extend between ends 1264, 1266 and each define a respective recess 1272, 1274 adjacent to coupling end 1264. Pegs 1276, 1278 extend from coupling end 1264 and are sized and configured to be received within holes 1220, 1222 defined by sides 1204, 1206 of foot holder assembly

1200. Although two pegs 1276, 1278 are illustrated, one of ordinary skill in the art will understand that fewer or more pegs may be implemented.

Top side 1280 defines one or more holes 1282-1, 1282-2, 1282-3, 1282-4, 1282-5, 1282-6, 1282-7, 1282-8 ("holes 1282") for receiving k-wires. An opening 1284 is defined by top side 1280 and extends through base 1262 to patient-contact side 1286, which is disposed opposite top side 1280. Opening 1284 enables a surgeon or other professional to view the bottom of a patient's foot. A passageway 1288 also extends through base 1262 and is sized and configured to receive a locking bushing assembly 1290.

As best seen in FIG. 73, locking bushing assembly 1290 includes a central member 1292 that is coupled within passageway 1288. Central member 1292 includes a threaded flared region 1294 that is disposed adjacent to end 1296. A plurality of flexible prongs 1298 are disposed at end 1296 and have a tapered configuration that narrows from flared region 1294 to end 1296. Central member 1292 defines a bore 1300 extending from end 1296 to end 1302. Bore 1300 is sized and configured to receive a drill bushing as described in greater detail below.

A knob 1304 defines an internal space 1306 and a hole 1308 that aligns with bore 1300 of central member 1292. Inner surface 1310 adjacent to open end 1312 of knob 1304 includes threads for engaging the threads of threaded flared region 1294 of central member 1292. Opposite open end 1312, knob 1304 includes a plurality of outwardly extending gripping surfaces 1314 at end 1316. Internally, end 1316 includes a taper 1318. Side wall 1320 of knob 1304 defines one or more holes 1322 for receiving a respective pin 1324 for preventing knob 1304 from being separated from central member 1292.

Referring again to FIG. 71, foot plate 1326 has a rectangular base portion 1328 and a coupling portion 1330. Coupling portion 1330 includes a pair of pegs 1332, 1334 that are sized and configured to be received within holes 1222, 1223 defined by sides 1204, 1206 of base plate 1202. Sides 1336, 1338 each define a respective slot 1340, 1342 that are sized and configured to receive biased detents 1212, 1214 of base plate 1202 of foot holder assembly 1200.

Operation

The use of reamer stabilizer 1000, foot holder assembly 1200, drill guide assembly 1260, and foot plate 1326 is now described. As described above, a surgeon uses tibial resection guide mount 100 and tibial resection guide 132 to resect the inferior end of a patient's tibia 16 and uses talar resection guide mount 102 and talar resection guide 166 to resect the superior surface of a patient's talus 14 to create resected joint space 22 as illustrated in FIG. 15.

Tibial drill guide mount 700 is inserted into resected joint space 22, and mounting plate 800A is connected to tibial drill guide mount 700 using dowel pins in the same way mounting plate 800 is connected to tibial drill guide mount 700 as described above with reference to FIGS. 49 and 50. Cartridge-style tibial drill guide 702 is inserted into aperture 804 of mounting plate 800A and recess 718 of tibial drill guide mount 700. Tibial drill guide cartridge 702 is inserted until leading end 786 of tibial drill cartridge 702 abuts rear wall 788 of tibial drill guide mount 700 at which point the ball detent disposed within hole 772 engages hole 828 defined by mounting plate 800 and the front side 768 of tibial drill guide cartridge 702 is flush with front side 806 of mounting plate 800A, which is illustrated in FIG. 74.

Mounting member 1232 of foot holder assembly 1200 is coupled to tibial drill guide mount 700 and mounting plate 800A using dowel pins 70. For example, holes 1246, 1248

defined by second end 1250 are aligned with and receive dowel pins 70 that extend from mounting plate 800A. Pivoting arrangement 948 of mounting member 800A is pivoted from a horizontal position in which lower portion 952 is not received within slot 1252 defined by mounting member 1232 to a vertical position in which lower portion 952 is received within slot 1252. Knob 952 is rotated about its axis (clockwise or counterclockwise) such that the bottom surface 954 of knob 952 contacts mounting member 1232 to maintain engagement between mounting member 1232 and the assemblage of tibial drill guide mount 700 and mounting plate 800A.

As illustrated in FIGS. 75 and 76, drill guide assembly 1260 is coupled to the appropriate side 1204, 1206 of base plate 1202 such that drill guide assembly 1260 will be disposed directly adjacent to the heel of a patient's foot. The coupling of drill guide assembly 1260 to base plate 1202 includes inserting pegs 1276, 1278 into holes 1220 defined by side 1204 or into holes 1222 defined by side 1206. As pegs 1276, 1278 are inserted into holes 1220 or 1222, biased detent 1212 or 1214 outwardly flexes in response to contacting base 1262 of drill guide assembly 1260 and is then urged into a locking engagement within one of slots 1272 or 1274 defined by sides 1268, 1270 by a biasing member when a detent 1212 or 1214 is aligned with a slot 1272 or 1274.

Foot plate 1326 is coupled to the side 1204, 1206 of base plate 1202 that is opposite the side 1204, 1206 to which drill guide assembly 1260 is coupled such that foot plate 1326 is disposed adjacent to the forefoot of the patient. The coupling of foot plate 1326 to base plate 1202 includes inserting pegs 1332, 1334 into holes 1220 defined by side 1204 or into holes 1222 defined by side 1206 of base plate 1202. As pegs 1332, 1334 are inserted into holes 1220 or 1222, biased detent 1212 or 1214 outwardly flexes in response to contacting coupling portion 1330 of foot plate 1326. Detent 1212 or 1214 is urged into a slot 1340 or 1342 defined by a side 1336 or 1338 of coupling portion 1330 when detent 1212 or 1214 is aligned with slot 1340 or 1342.

The distance between base plate 1202 and mounting member 1232 can be adjusted by unscrewing locking screw 1242 such that mounting member 1232 can be slid along rods 1226, 1228. When the desired positioning of mounting member 1232 relative to base plate 1202 has been achieved, locking screw 1242 is rotated to lock mounting member 1232 at its position along rods 1226, 1228.

A trocar 74, which is illustrated in FIG. 77, having ink applied to its tip is inserted into bore 1300 defined by central member 1292 of locking bushing assembly 1290 and touched the skin of the patient's foot to create a mark. Drill guide assembly 1260 is removed from its engagement with base plate 1202 by pressing the biased detent 1212, 1214 that is engaged with a slot 1268, 1270 defined by base 1262 such that detent 1212, 1214 is urged out of its engagement with the slot 1268, 1270. With detent 1212, 1214 disengaged from slot 1268, 1270, base 1262 is pulled away from base 1202 until pegs 1278, 1280 are removed from holes 1220 or 1222.

With drill guide assembly 1260 removed, access to the calcaneus 20 of the patient is made by making a small incision at the marked location using a scalpel or other surgical cutting tool. Drill guide assembly 1260 is then re-coupled to base plate 1202 as described above.

A drill bushing or cannula (not shown) is inserted into bore 1300 and then locked in place by rotating knob 1304 of locking bushing assembly 1294. Rotating knob 1304 causes the threads formed on inner surface 1310 of knob 1304 to engage the threads of threaded flared region 1294. As knob

1304 is rotated in one direction, e.g., a clockwise direction, the rotation of knob 1304 relative to central member 1292 causes knob 1304 to be advanced along central member 1292 towards base 1262, which results in taper 1318 contacting flexible prongs 1298. Flexible prongs 1298 are urged inwardly towards one another as knob 1304 moves towards base 1262 thereby providing a frictional lock between locking bushing assembly 1290 and drill bushing or cannula.

With drill bushing or cannula locked to locking bushing assembly 1290, a drill is used to create a pilot hole through the calcaneus 20, talus 14, and into tibia 16. As the drill exits talus 14, the conically shaped internal surface 748 of tibial drill cartridge 702 guides the tip of the drill into tibia 14. Once the pilot hole has been drilled to a desired depth into tibia 14, the drill is backed out and tibial drill cartridge 702 is removed from tibial drill guide mount 700. Removal of cartridge 702 includes inserting a threaded dowel or rod into threaded blind hole 770 and pulling on threaded dowel or rod to remove cartridge 702 from tibial guide mount 700.

A reamer head 66 is inserted into the space vacated by cartridge 702 and is coupled to a driving rod 65 of a reamer that is received within the vacated space having been inserted through the drill bushing or cannula locked in locking bushing assembly 1290.

Once reamer head 66 is coupled to reamer rod 65, reamer stabilizer 1000 is secured to mounting plate 800A as described with reference to FIGS. 78 and 79. Distal end 1004 of stabilizer body 1002 is inserted into aperture 804 of mounting plate 800A and into body 704 via recess 718 defined by front side 706 of tibial drill guide mount 700. Reamer stabilizer 1000 continues to be advanced until steps 1034, 1036 contact top surface 806 in an abutting relationship. With steps 1034, 1036 contacting top surface 806, detent 1176 disposed on the inner surface 1178 of pivoting button 1156 is received within slot 826 of mounting plate 800A.

When detent 1176 is disposed within slot 826 and reamer stabilizer 1000 is coupled to mounting plate 800A, reamer driving rod 65 is received within notch 1018 defined at the distal end 1004 of stabilizer body 1002. Guiding assembly 1052 is actuated such that reamer guide body 1054 in combination with notch 1018 encloses and surrounds the reamer driving rod 65 as best seen by comparing FIGS. 64 and 65. Guiding assembly 1052 is actuated by applying a downward pressure (i.e., pressure in a distal direction) to head 1102, which urges plunger rod 1086 and pivot rod 1074 in a distal direction. The movement of plunger rod 1086 and pivot rod 1074 in the distal direction forces reamer guide body 1054 to pivot about hole 1056, which is pinned to stabilizer body 1002 by pin 1042. The distal end 1078 of pivot rod 1074 may outwardly flex with respect to hole 1080 at the proximal end 1084 as reamer guide body 1054 pivots about hole 1056. Although reamer guide body 1054 is illustrated as entirely extending across notch 1018, one of ordinary skill in the art will understand that reamer guide body 1054 will extend only partially across notch 1018 in some embodiments.

Still referring to FIGS. 64 and 65, locking assembly 1122 is configured to automatically lock guiding assembly 1052 in its engaged position with the reamer driving rod 65. Locking button 1124 is urged by biasing member 1152 towards in the direction towards head 1102 such that, when angled surface 1110 of head 1102 is disposed below angled surface 1138 of locking button 1124, locking button 1124 slides over the head 1102 to maintain the engagement of the reamer 65 and concave guiding surface 1058 of reamer guide body 1054. The reamer 65, 66 is advanced into the

pilot intramedullary channel previously formed by the drill while being supported by reamer stabilizer 1000, which maintains the direction in which reamer 65, 66 is advanced into tibia 16 and prevents the reamer 65, 66 from wandering.

Once the intramedullary channel has been reamed to a desired depth, the reamer 65, 66 is retracted through the intramedullary channel until the reamer head 66 is received within the resected joint space 22. Reamer stabilizer 1000 is then removed from its engagement with reamer rod 65 and mounting plate 800A. To disengage reamer stabilizer 1000 from its engagement with the reamer 65, locking button 1124 is pushed in a direction away from head 1102 until locking button 1124 is received within cavity 1010 defined by stabilizer body 1002.

Biasing member 1120 of guiding assembly 1052, which is disposed in abutting contact with distal end 1106 of head 1102, causes head 1102, plunger rod 1086, and pivot rod 1074 to move in a proximal direction when locking button 1124 does not contact head 1102 or otherwise impede head 1102 from moving in the proximal direction. The proximal movement of head 1102, plunger rod 1086, and pivot rod 1074 causes reamer guide body 1054 to pivot about pin 1048 due to the cross-pinned engagement between pivot rod 1074 and reamer guide body 1054.

With guiding assembly 1054 disengaged from the reamer, reamer stabilizer 1000 is disengaged from mounting plate 800A by pressing pivoting button 1156 such that button 1156 pivots about pin 1076 and detent 1176 is removed from its engagement with slot 826. Reamer stabilizer 1000 is then pulled from aperture 804. The reamer head 66 is then removed from resected joint space 22.

Knob 952 is rotated in a direction opposite to the direction in which knob 952 was rotated to tighten pivoting arrangement to mounting member 800A such that the bottom surface 954 loosens its frictional engagement with mounting member 1232. Pivoting arrangement 948 is pivoted back to a horizontal position, and locking screw 1242 of mounting member 1232 is loosened by rotating knob 1244 in a direction that is opposite the direction in which knob 1244 was rotated to tighten locking screw 1242. Mounting member 1232 slides along rods 1226, 1228 as base plate 1202 is moved away from the patient's foot.

With the drill bushing or cannula still disposed within the calcaneus 20 and talus 14, drill guide assembly 1260 is decoupled from its engagement with base plate 1202 in the same manner as described above. Foot holder assembly 1200 is then removed such that drill guide assembly 1260, tibial drill guide mount 700, and mounting plate 800A are still engaged with the patient's foot. K-wires 62 used to maintain the position of tibial drill guide mount 700 and mounting plate 800A are removed, and then tibial drill guide mount 700 and mounting plate 800A are removed.

With drill bushing or cannula still disposed within the calcaneus 20 and talus 14, k-wires 62 are inserted through one or more holes 1282 to secure drill guide assembly 1260 to the foot of the patient as illustrated in FIG. 80. Once drill guide assembly 1260 is secured, a tool for driving components of a modular ankle prosthesis into the intramedullary canal formed by the reamer is inserted through drill guide or cannula held by drill guide assembly 1260. The remainder of the installation prosthesis is described in U.S. Pat. No. 7,534,246 issued to Reiley et al.

Anterior Approaches

The disclosed systems and methods described above can also be adapted to enable an intramedullary cavity to be formed in the tibia of a patient via an anterior approach once resected joint space 22 has been formed using tibial resec-

tion guide mount **100** and tibial resection guide **132** to resect the inferior end of a patient's tibial and uses talar resection guide mount **102** and talar resection guide **166** to resect the superior surface of a patient's talus **14** to create resected joint space **22** as illustrated in FIG. **15**. The anterior approach of forming an intramedullary channel in a patient's tibia avoids drilling through the calcaneus and talus of the patient.

Referring now to FIGS. **81-89**, a custom anterior reaming guide mount **1400** is illustrated as being disposed within resected joint space **22**. Reaming guide mount **1400** is formed from a resilient polymer material of the type that is suitable for use in connection with stereo lithography, selective laser sintering, or like manufacturing equipment.

Reaming guide mount **1400** includes a body **1402** having an inferior surface **1404** configured to mate against the flat formed on the superior surface of the resected talus. The superior surface **1406** includes a pair of opposed angled surfaces **1408** that are configured to correspond to the cuts made using tibial resection guide **166**.

A mating portion **1410** extends from superior surface **1406** and includes a conformal bone engaging surface **1412**, which is complementary to a surface of the patient's tibia **16**. Mating portion **1410** defines holes **1414**, **1416** that are sized and configured to receive k-wires **62** for securing reaming guide mount **1400** to talus **16**. Superior surface **1406** also defines an opening **1418** through which a reamer head **66** can be received.

Body **1402** also includes a rear wall **1420** and a pair of opposed side walls **1422**, **1424** that define a cavity **1426** with superior wall **1428** and inferior wall **1430**. In some embodiments, the respective interfaces between superior wall **1428** and side walls **1422**, **1424** include chamfers **1432**, **1434** or other geometric features used for properly locating insert **1440**.

As best seen in FIGS. **82,84**, and **85**, insert **1440** has an overall shape that is complementary to the internal geometry of cavity **1426** defined by reaming guide mount **1400** and can be fabricated from a more durable material than reaming guide mount **1400** such as, for example, a metal material. In particular, insert **1440** includes an inferior surface **1442**, a superior surface **1444**, side surfaces **1446**, **1448**, and a front surface **1450**. Superior surface **1444** defines an opening **1452** (FIG. **85**) through which a reamer head **66** can be received as described in greater detail below.

Front surface **1450** also defines an opening **1454** that is sized such that a reamer head **66** can be received within opening **1454**. Openings **1452** and **1454** communicate with each other such that the reamer head inserted within opening **1454** can be received within opening **1452** via internal communication between the openings **1452**, **1454**. In some embodiments, opening **1454** is smaller than the size of a reamer head **66**, but provides a surgeon access a reamer head **66** disposed within opening **1454** such that reamer head **66** can be coupled to a reamer driving rod **65**.

An angled front face **1456** is disposed between front face **1450** and inferior surface **1442**. Angled front face **1456** defines a passageway **1458** that extends from angled front face **1456** to bottom surface **1460** of internal chamber **1562**. Passageway **1562** is sized and configured to receive a flexible reamer.

In operation, reaming guide mount **1400** is inserted into resected joint space **22**. Angled surfaces **1408**, **1410** of superior surface **1406** and conformal bone engaging surface **1414** precisely locate reaming guide mount **1400** within the resected joint space **22**.

A reamer head **66** is inserted into opening **1452** defined by superior surface **1452** of insert **1440**. Insert **1440** is inserted into cavity **1428** until opening **1452** defined by superior surface **1444** of insert **1440** aligns with opening **1420** defined by superior surface **1420** of reaming guide mount **1400**. A reamer rod **65** is inserted into passageway **1458** defined by angled front face **1456** and coupled to reamer head **66** disposed within opening **1452**. A surgeon may insert one or more tools in opening **1454** to secure reamer head **66** to reamer rod **65**. Reamer head **66** can then be advanced into the patient's tibia **16**.

In some embodiments, reamer stabilizer **1000** is used in connection with reaming guide mount **1400** and insert **1440**. For example and as illustrated in FIGS. **87-89**, with a reamer head **66** and reamer body **65** assembled together within the construct of reaming guide mount **1400** and insert **1440**, which are disposed within resected joint space **22**, reamer stabilizer **1000** is coupled to stabilizer driving rod **65**. To couple reamer stabilizer **1000** to stabilizer driving rod **65**, distal end **1004** of stabilizer body **1002** is inserted into opening **1454** defined by front surface **1450** of insert **1440** until driving rod **65** is received within notch **1018** defined at the distal end **1004** of stabilizer body **1002**.

Guiding assembly **1052** is actuated such that reamer guide body **1054** and notch **1018** encloses and surrounds the reamer driving rod **65** as best seen in FIG. **89**. Guiding assembly **1052** is actuated by applying a downward pressure (i.e., pressure in a distal direction) to head **1102**, which urges plunger rod **1086** and pivot rod **1074** in a distal direction. The movement of plunger rod **1086** and pivot rod **1074** in the distal direction forces reamer guide body **1054** to pivot about hole **1056**, which is pinned to stabilizer body **1002** by pin **1042**. The distal end **1078** of pivot rod **1074** may outwardly flex with respect to hole **1080** at the proximal end **1084** as reamer guide body **1054** pivots about hole **1056**.

Locking assembly **1122** is configured to automatically lock guiding assembly **1052** in its engaged position with the reamer driving rod **65**. As described above, locking button **1124** is urged by biasing member **1152** towards in the direction towards head **1102** such that, when angled surface **1110** of head **1102** is disposed below angled surface **1138** of locking button **1124**, locking button **1124** slides over the head **1102** to maintain the engagement of the reamer rod **65** and concave guiding surface **1058** of reamer guide body **1054**.

The reamer **65**, **66** is advanced into tibia **16** to form a reamed intramedullary channel while being supported by reamer stabilizer **1000**, which maintains the direction in which reamer **65**, **66** is advanced into tibia **16** and prevents the reamer **65**, **66** from wandering within tibia **16**.

Once the intramedullary channel has been reamed to a desired depth, the reamer **65**, **66** is retracted through the intramedullary channel until the reamer head **66** is received within opening **1420** defined by superior surface **1406** of reaming guide mount **1400** and/or within opening **1452** defined by superior surface **1444** defined by insert **1440**. Reamer stabilizer **1000** is then removed from its engagement with reamer rod **65**.

To disengage reamer stabilizer **1000** from its engagement with the reamer rod **65**, locking button **1124** is pushed in a direction away from head **1102** until locking button **1124** is received within cavity **1010** defined by stabilizer body **1002**. Biasing member **1120** of guiding assembly **1052**, which is disposed in abutting contact with distal end **1106** of head **1102**, causes head **1102**, plunger rod **1086**, and pivot rod **1074** to move in a proximal direction when locking button **1124** does not contact head **1102** or otherwise impede head

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1102 from moving in the proximal direction. The proximal movement of head 1102, plunger rod 1086, and pivot rod 1074 causes reamer guide body 1054 to pivot about pin 1048 due to the cross-pinned engagement between pivot rod 1074 and reamer guide body 1054. With guiding assembly 1054 disengaged from the reamer, reamer stabilizer 1000 is pulled out of opening 1454 defined by front surface 1450 of insert 1440.

As will be understood by one of ordinary skill in the art, the size and shape of reaming guide mount and insert may be varied. For example, FIGS. 90-94 illustrate another embodiment of reaming guide mount 1500 and insert 1540. As shown in FIGS. 90 and 92, body 1502 of reaming guide mount 1500 has an inferior surface 1504 configured to mate against the flat formed on the superior surface of the resected talus. The superior surface 1506 includes a pair of opposed angled surfaces 1508 that are configured to correspond to the cuts made using tibial resection guide 166.

Mating portion 1510 extends from superior surface 1506 and includes a conformal bone engaging surface 1512 (FIG. 92), which is complementary to a surface of the patient's tibia 16. Holes 1514, 1516 are defined by mating portion 1512 and are sized and configured to receive k-wires 62 for securing reaming guide mount 1500 to talus 16. Superior surface 1508 also defines an opening 1518 through which a reamer head 66 can be received.

Body 1502 also includes a rear wall 1520 (FIG. 92) and a pair of opposed side walls 1524, 1526 that define a cavity 1526 with superior wall 1528 and inferior wall 1530 (FIGS. 90 and 92). The respective interfaces between superior wall 1528 and side walls 1524, 1526 include chamfers 1432, 1434 or other geometric features used for properly locating insert 1540. An inwardly projecting structure 1538 extends from side wall 1524.

As best seen in FIG. 91, insert 1540 has a triangular wedge shape such that it is able to be received between inwardly projecting structure 1538 and inferior wall 1530 of reaming guide mount 1500. Insert 1540 includes an inferior surface 1542, a superior surface 1544, side surfaces 1546, 1548 (FIG. 94), and a front surface 1550. Angled front face 1556 is disposed between front face 1550 and inferior surface 1542 and defines a passageway 1558 that extends from angled front face 1556 to superior surface 1544. Passageway 1558 is sized and configured to receive a flexible reamer rod 65.

As shown in FIGS. 93 and 94, reamer stabilizer 1000 is received within cavity 1526 adjacent to insert 1540 such that reamer stabilizer 1000 abuts both inwardly projecting structure 1536 and insert 1540. Reamer stabilizer 1000 stabilizes reamer 65, 66 as reamer 65, 66 is advanced into the tibia 16 of a patient as described above.

Another embodiment of an anterior reaming guide mount is illustrated in FIGS. 95-100. Reaming guide mount 1600 includes a body 1062 having an inferior surface 1604 (FIG. 96) configured to mate against the flat formed on the superior surface of the resected talus 14. The superior surface 1606 includes a pair of opposed angled surfaces 1608 (FIGS. 99 and 100) that are configured to correspond to the cuts made using tibial resection guide 166.

Mating portion 1610 extends from superior surface 1606 and includes a conformal bone engaging surface 1614, which is complementary to a surface of the patient's tibia 16. Holes 1614, 1616 are defined by mating portion 1612 and are sized and configured to receive k-wires 62 for securing reaming guide mount 1600 to talus 16. Superior surface 1606 also defines an opening 1618 through which a reamer head 66 can be received.

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Body 1602 also includes a front surface 1622 and an angled front surface 1624 that defines a passageway 1626 that communicates with opening 1620. Passageway 1626 is configured to receive a flexible reamer driving rod 65 that is to be coupled to a reamer head 66 disposed within opening 66.

The disclosed systems and methods advantageously utilize custom manufactured surgical instruments, guides, and/or fixtures that are based upon a patient's anatomy to reduce the use of fluoroscopy during a surgical procedure. In some instances, the use of fluoroscopy during a surgical procedure may be eliminated altogether. The custom instruments, guides, and/or fixtures are created by imaging a patient's anatomy with a computer tomography scanner ("CT"), a magnetic resonance imaging machine ("MRI"), or like medical imaging technology prior to surgery and utilizing these images to create patient-specific instruments, guides, and/or fixtures.

Although the invention has been described in terms of exemplary embodiments, it is not limited thereto. Rather, the appended claims should be construed broadly, to include other variants and embodiments of the invention, which may be made by those skilled in the art without departing from the scope and range of equivalents of the invention.

What is claimed is:

1. A surgical device, comprising:

- a elongate body having a proximal end and a distal end with a step formed along at least one side of the body, the distal end of the elongate body defining a notch sized and configured to receive a reamer;
- a coupling assembly supported by the elongate body including (i) a reamer guide body disposed at the distal end of the elongate body and configured to move between a first position and a second position, and (ii) first and second rods disposed within a channel defined by the elongate body, the first rod coupled to a head extending outwardly from the longitudinal channel, and the second rod pivotably coupled to the first rod and to an end of the reamer guide body; and
- a locking assembly supported by the elongate body, the locking assembly configured to releasably engage the coupling assembly to maintain the reamer guide body in the second position, wherein the reamer guide body extends at least partially across the notch in the second position.

2. The surgical device of claim 1, wherein the coupling assembly includes a biasing member disposed within the channel, the biasing member configured to bias the head in a proximal direction.

3. The surgical device of claim 1, wherein the locking assembly includes a button slidably positioned within a cavity defined at the proximal end of the elongate body, and a biasing member configured to urge the button in a direction towards the coupling assembly.

4. The surgical device of claim 3, wherein a lower portion of the button includes an angled surface that complements an angled surface of the head of the coupling assembly.

5. The surgical device of claim 1, further comprising a second coupling assembly coupled to the elongate body, the second coupling assembly configured to releasably secure the surgical device to another instrument in combination with the step.

6. The surgical device of claim 5, wherein the second coupling assembly includes a button pivotably coupled to the at least one side of the elongate body, and a biasing member disposed within a cavity formed along the at least

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one side of the elongate body, the biasing member configured to exert a force on the button.

7. The surgical device of claim 6, wherein the button includes a detent at a first end that is disposed opposed an end on which the biasing member exerts the force.

8. The surgical device of claim 1, wherein the reamer guide body includes an arcuate surface configured to support the reamer, the reamer guide body disposed adjacent to a first hole configured to receive a first pin for pivotably coupling the reamer guide body to the distal end of the elongate body, and a second hole is defined adjacent to the first hole, the second hole configured to receive a second pin for pivotably coupling the reamer guide body to the pivotable second rod of the coupling assembly.

9. The surgical device of claim 1, wherein the coupling assembly includes a plunger rod disposed within a channel defined by the elongate body, a first end of the plunger rod is coupled to the pivotable second rod, and a second end of the plunger rod is coupled to the head including an upper portion that is disposed outwardly of the channel; and a first biasing member disposed within the channel, the biasing member disposed between a lower portion of the head and a reduced diameter area of the channel defined by the elongate body, the biasing member configured to urge the head in a proximal direction.

10. The surgical device of claim 9, wherein the locking assembly includes a button slidably positioned within a cavity defined at the proximal end of the elongate body, and a second biasing member configured to urge the button in a direction towards the head of the coupling assembly.

11. The surgical device of claim 10, wherein a lower portion of the button includes an angled surface that complements an angled surface of the head of the coupling assembly.

12. A reamer stabilizer, comprising: an elongate body extending from a proximal end to a distal end, the distal end of the elongate body defining a notch for receiving a reamer; a slidable guide assembly supported by the elongate body, the slidable guide assembly including a reamer guide body pivotably coupled to the distal end of the elongate body, the reamer guide body configured to move between a first

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position and a second position and including an arcuate surface for supporting the reamer wherein the reamer guide body defines first and second holes, the first hole is configured to receive a first pin for pivotably coupling the reamer guide body to the distal end of the elongate body, and the second hole is configured to receive a second pin for pivotably coupling the reamer guide body to a pivoting rod of the slidable guide assembly; and a locking assembly slidably supported by the elongate body, the locking assembly configured to move between a third position and a fourth position in which the locking assembly releasably engages the slidable guide assembly to maintain the reamer guide body in the second position, wherein the reamer guide body extends at least partially across the notch in the second position.

13. The reamer stabilizer of claim 12, wherein the slidable guide assembly includes a plunger rod disposed within a longitudinal channel defined by the elongate body, a first end of the plunger rod is coupled to the pivoting rod, and a second end of the plunger rod is coupled to a head including an upper portion that is disposed outwardly of the elongate channel; and a first biasing member disposed within the elongate channel, the biasing member disposed between a lower portion of the head and a reduced diameter area of the longitudinal channel defined by the elongate body, the biasing member configured to urge the head in a proximal direction.

14. The reamer stabilizer of claim 12, wherein the locking assembly includes a button slidably positioned within a cavity defined at the proximal end of the elongate body, and a second biasing member configured to urge the button in a direction towards the slidable guide assembly.

15. The reamer stabilizer of claim 12, further comprising a coupling assembly includes a button pivotably coupled to a side of the elongate body, and a biasing member disposed within a cavity formed along the side of the elongate body, the biasing member configured to exert a force on the button.

16. The reamer stabilizer of claim 15, wherein the button includes a detent at a first end that is disposed opposed an end on which the biasing member exerts the force.

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