



US 20190328537A1

(19) **United States**

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

(10) **Pub. No.: US 2019/0328537 A1**

(43) **Pub. Date: Oct. 31, 2019**

(54) **ADJUSTABLE HEIGHT ARTHROPLASTY PLATE**

Publication Classification

(71) Applicant: **DEPUY SYNTHES PRODUCTS, INC., RAYNHAM, MA (US)**

(51) **Int. Cl.**
A61F 2/40 (2006.01)
A61B 17/88 (2006.01)
A61B 17/80 (2006.01)
A61B 17/68 (2006.01)

(72) Inventors: **JASON M. CHAVARRIA, WARSAW, IN (US); SCOTT A. LUBENSKY, WINONA LAKE, IN (US)**

(52) **U.S. Cl.**
CPC *A61F 2/4014* (2013.01); *A61B 17/88* (2013.01); *A61F 2002/30507* (2013.01); *A61F 2/40* (2013.01); *A61B 17/68* (2013.01); *A61B 17/8061* (2013.01)

(21) Appl. No.: **16/504,446**

(22) Filed: **Jul. 8, 2019**

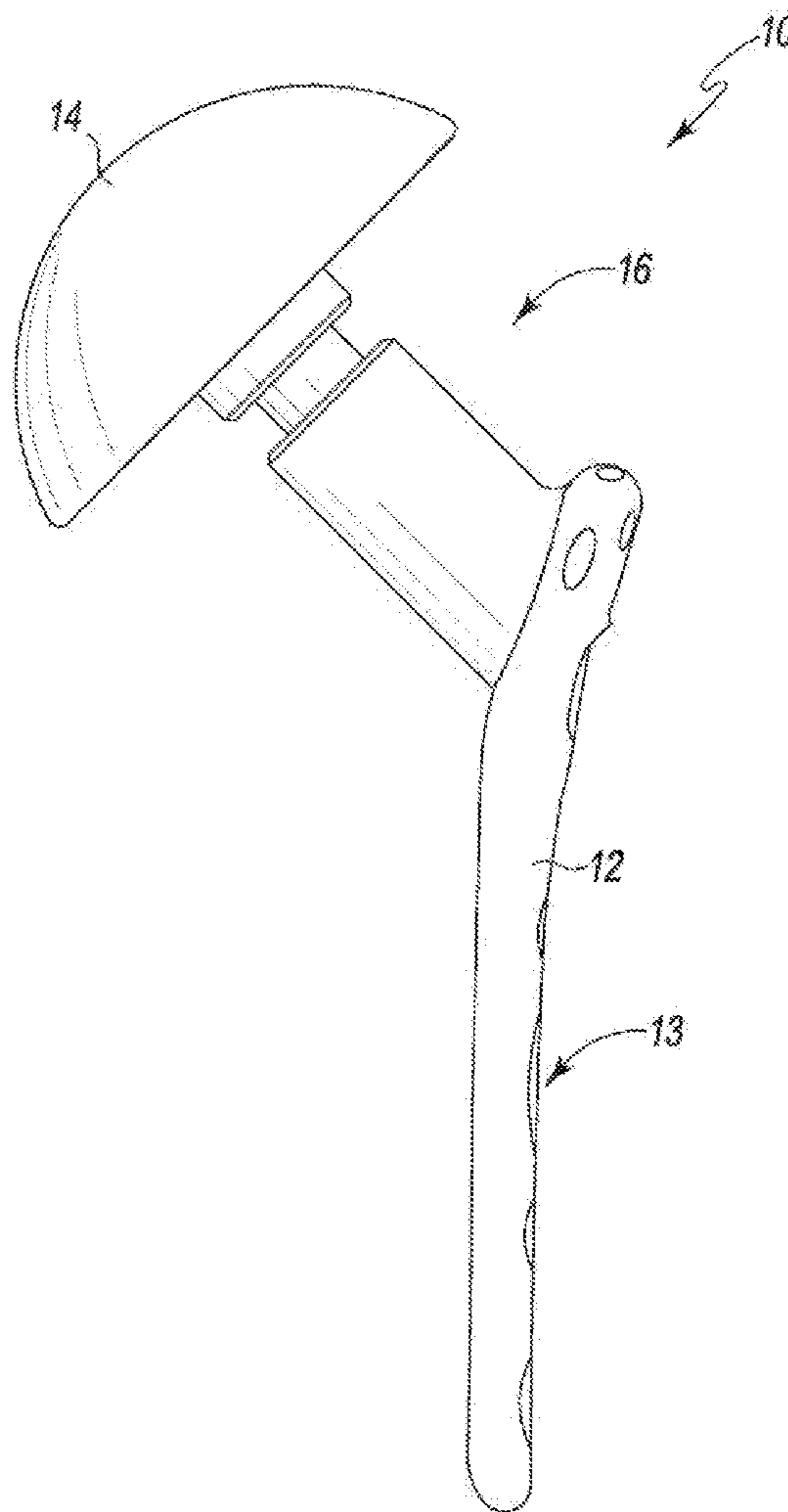
Related U.S. Application Data

(57) **ABSTRACT**

(63) Continuation of application No. 15/888,190, filed on Feb. 5, 2018, now Pat. No. 10,342,668, which is a continuation of application No. 15/193,747, filed on Jun. 27, 2016, now Pat. No. 9,883,948, which is a continuation of application No. 13/834,802, filed on Mar. 15, 2013, now Pat. No. 9,398,928.

(60) Provisional application No. 61/706,916, filed on Sep. 28, 2012.

An orthopaedic assembly. The orthopaedic assembly includes a first member adapted to be coupled to a bone and an adjustable connector assembly including an adjuster having a threaded end and a tapered end. The assembly also includes an articulation component including a bearing surface and adapted to be coupled to the tapered end of the adjustable connector assembly. The threaded end is adapted to engage the fixation plate.



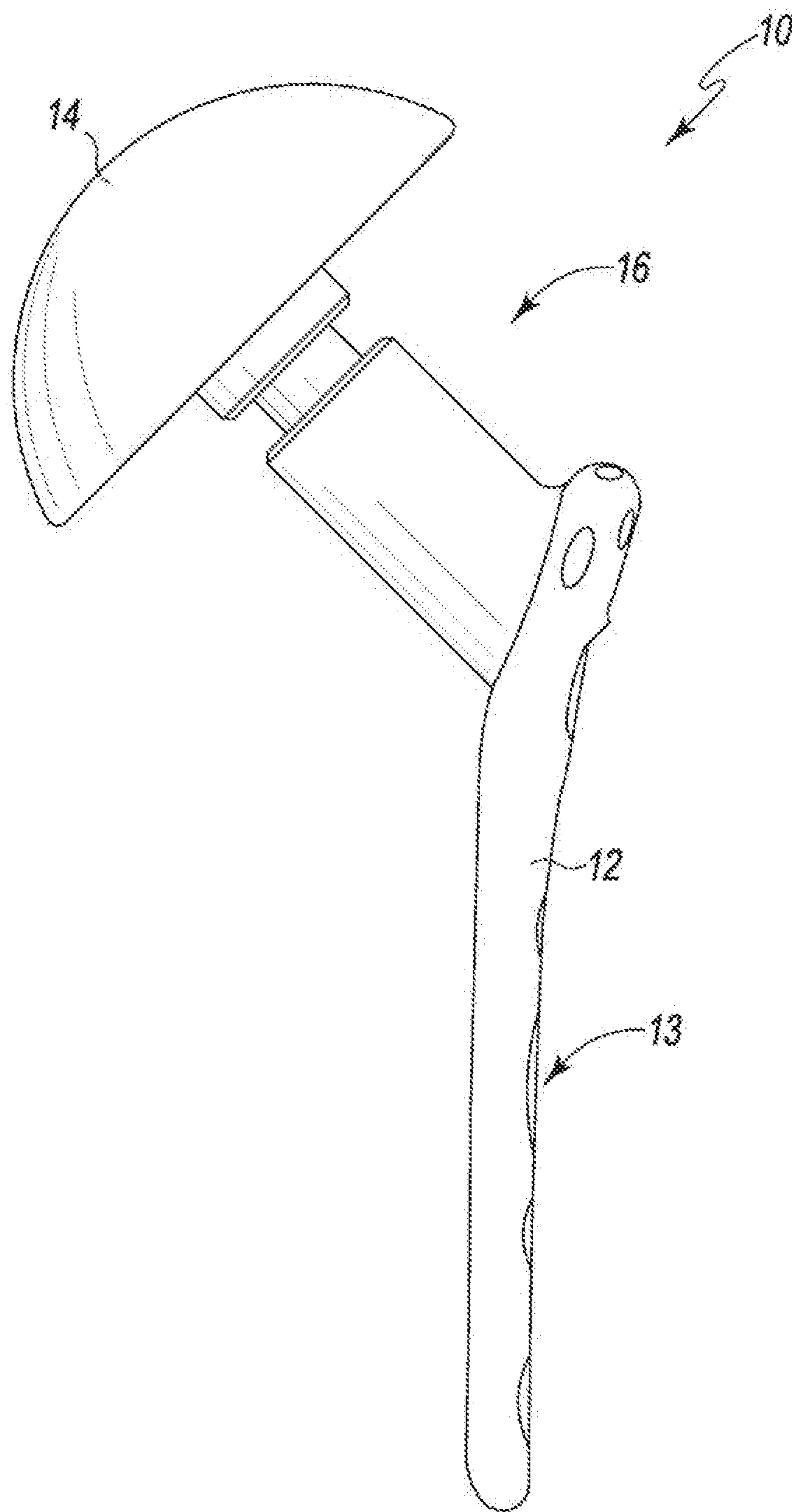


Fig. 1

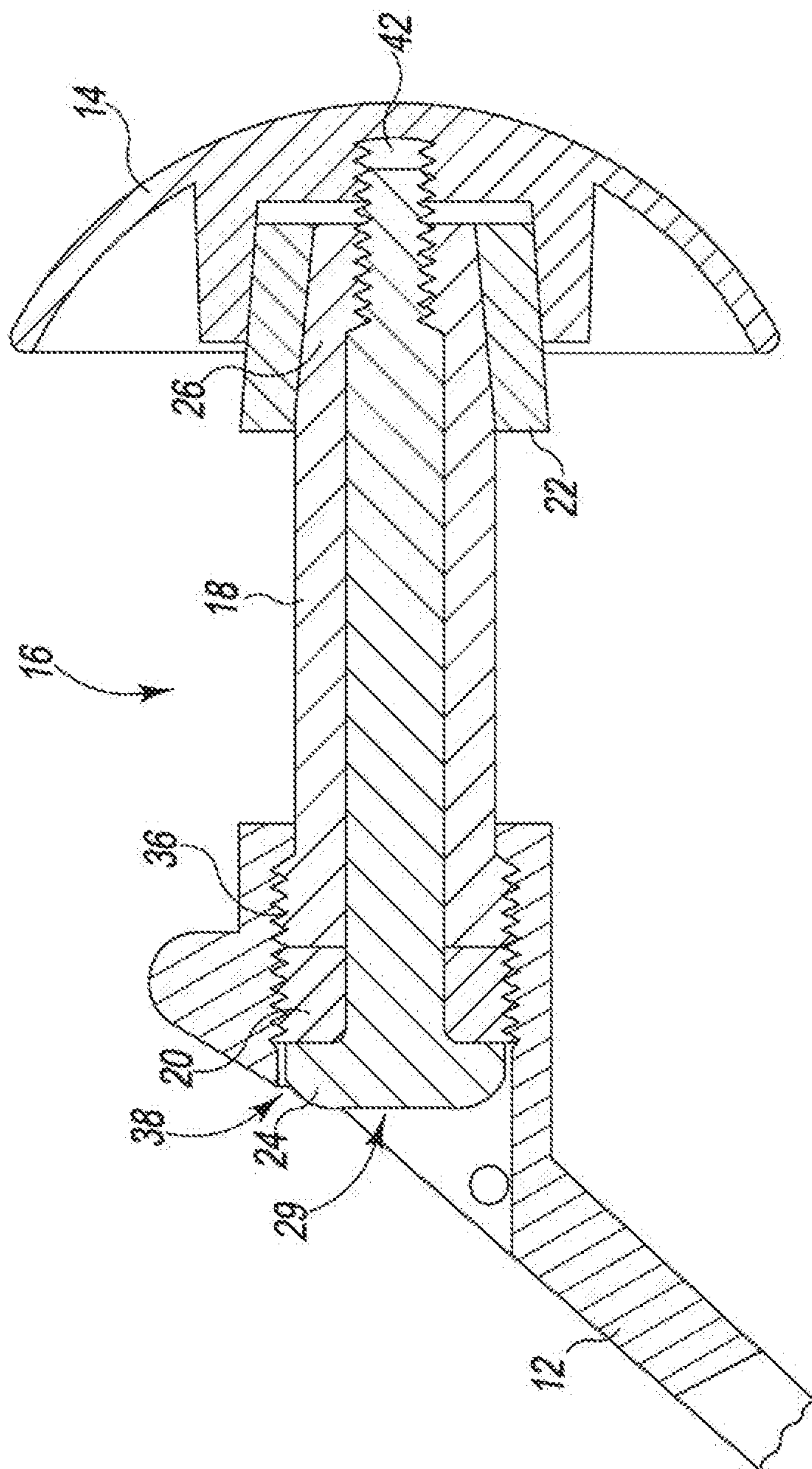


Fig. 2

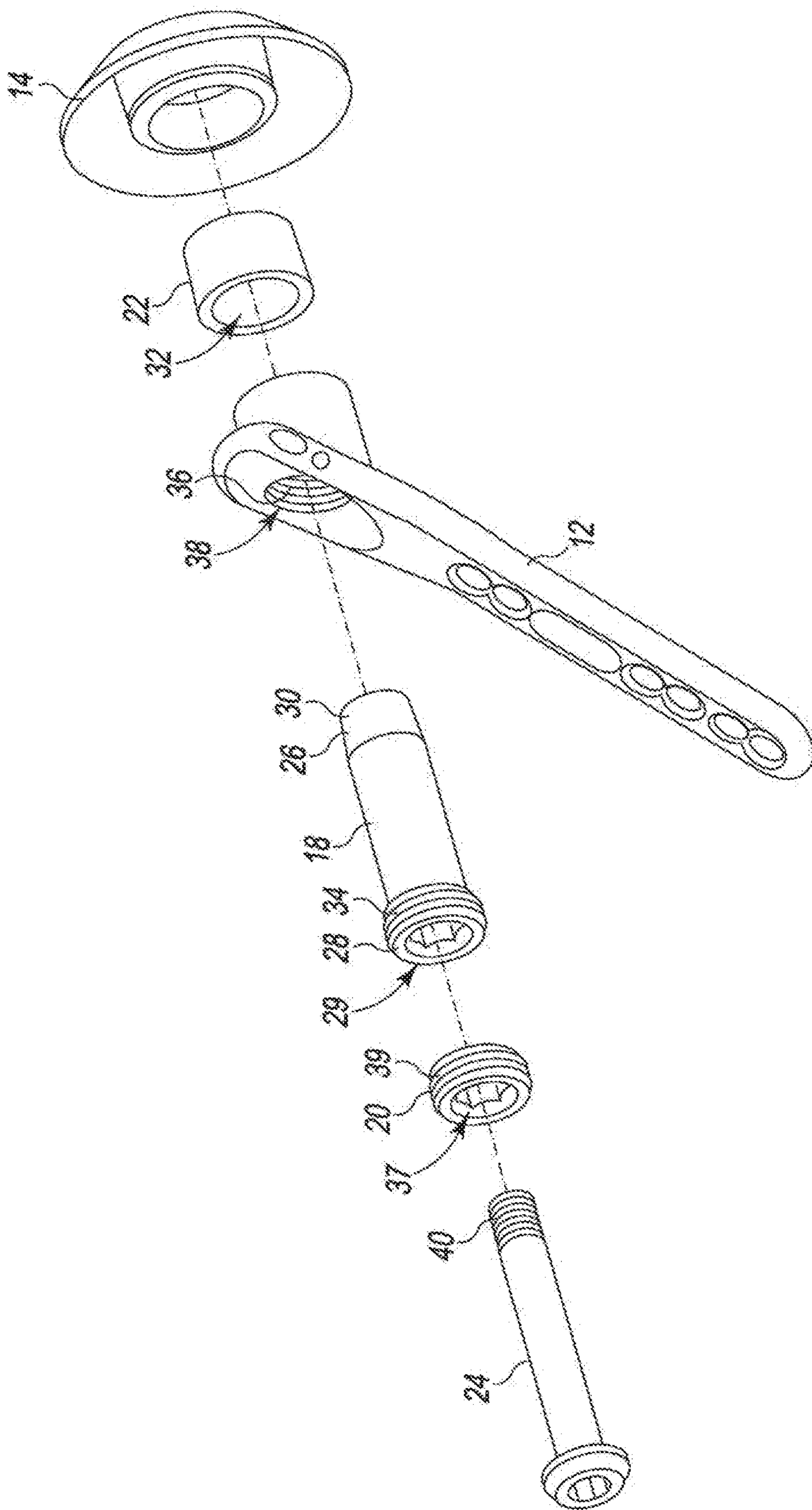


Fig. 3

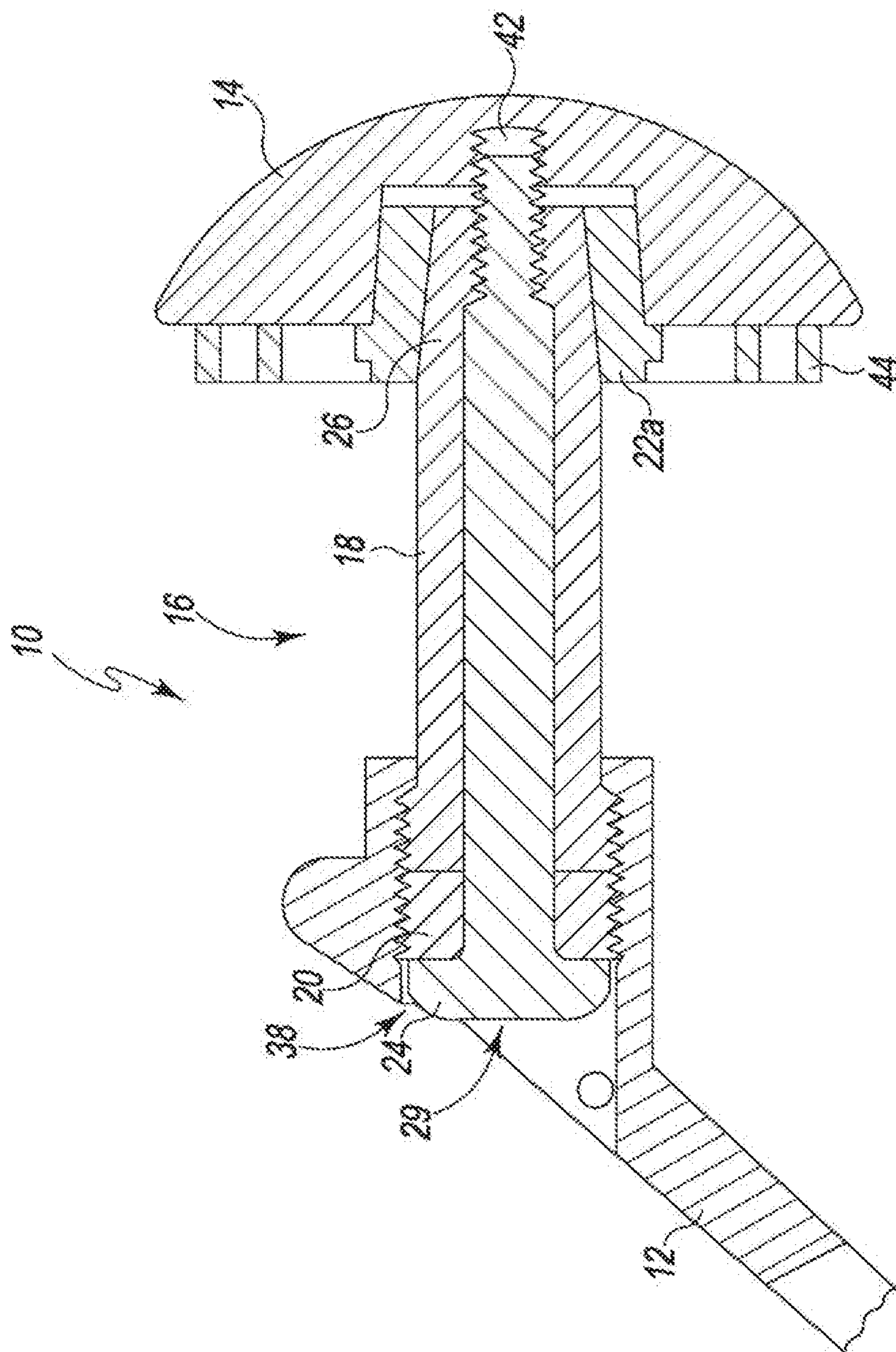


Fig. 4

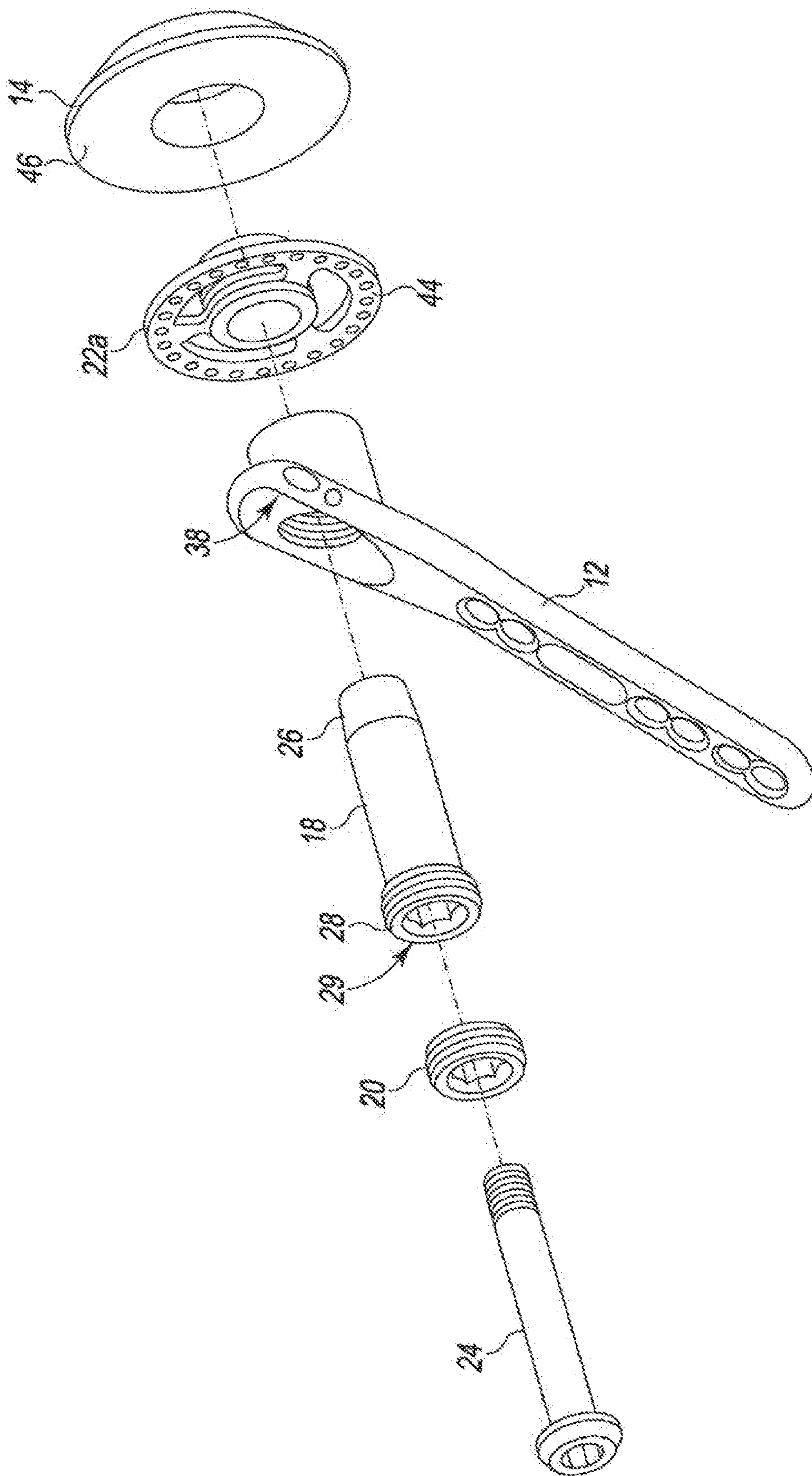


Fig. 5

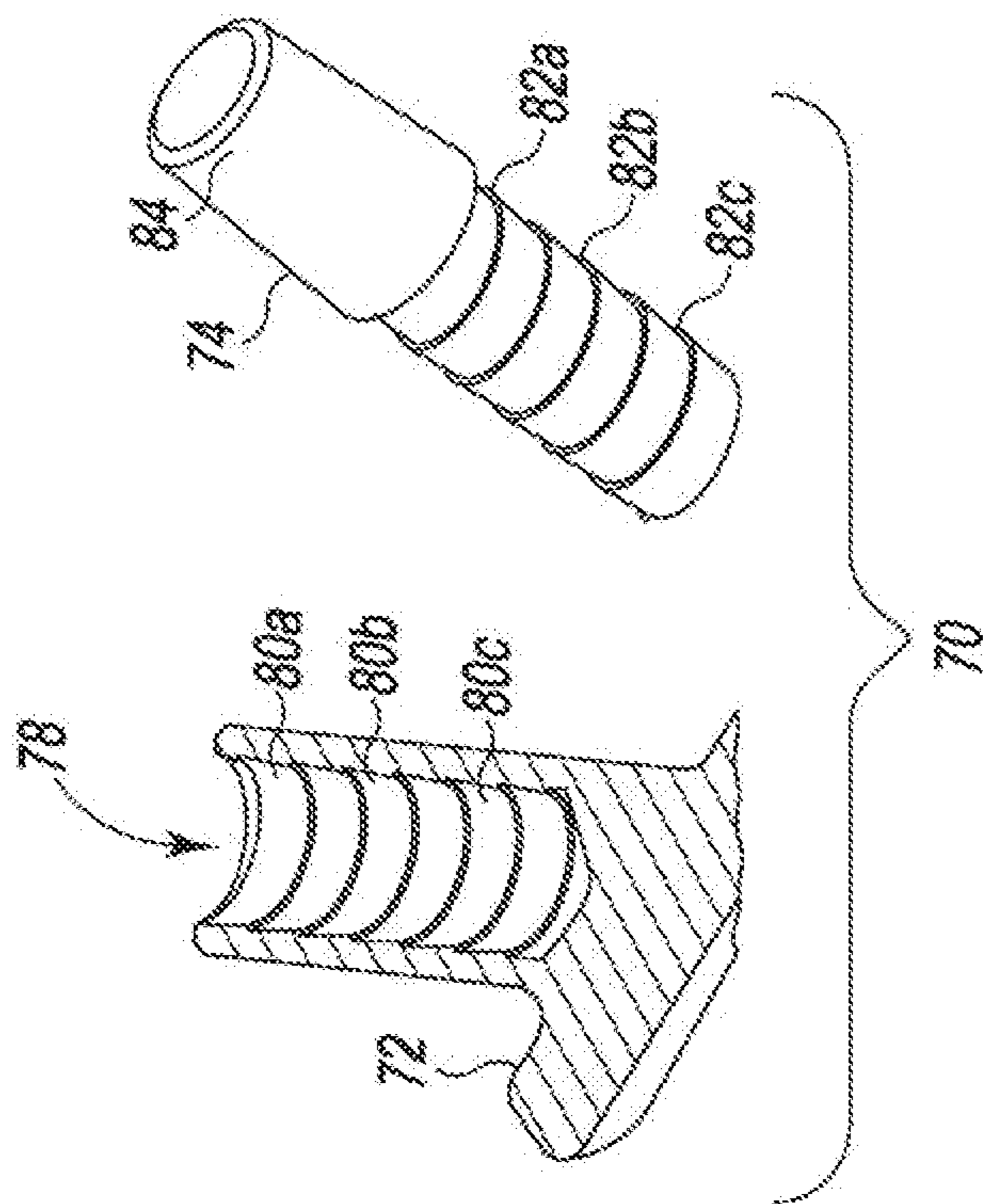


Fig. 7

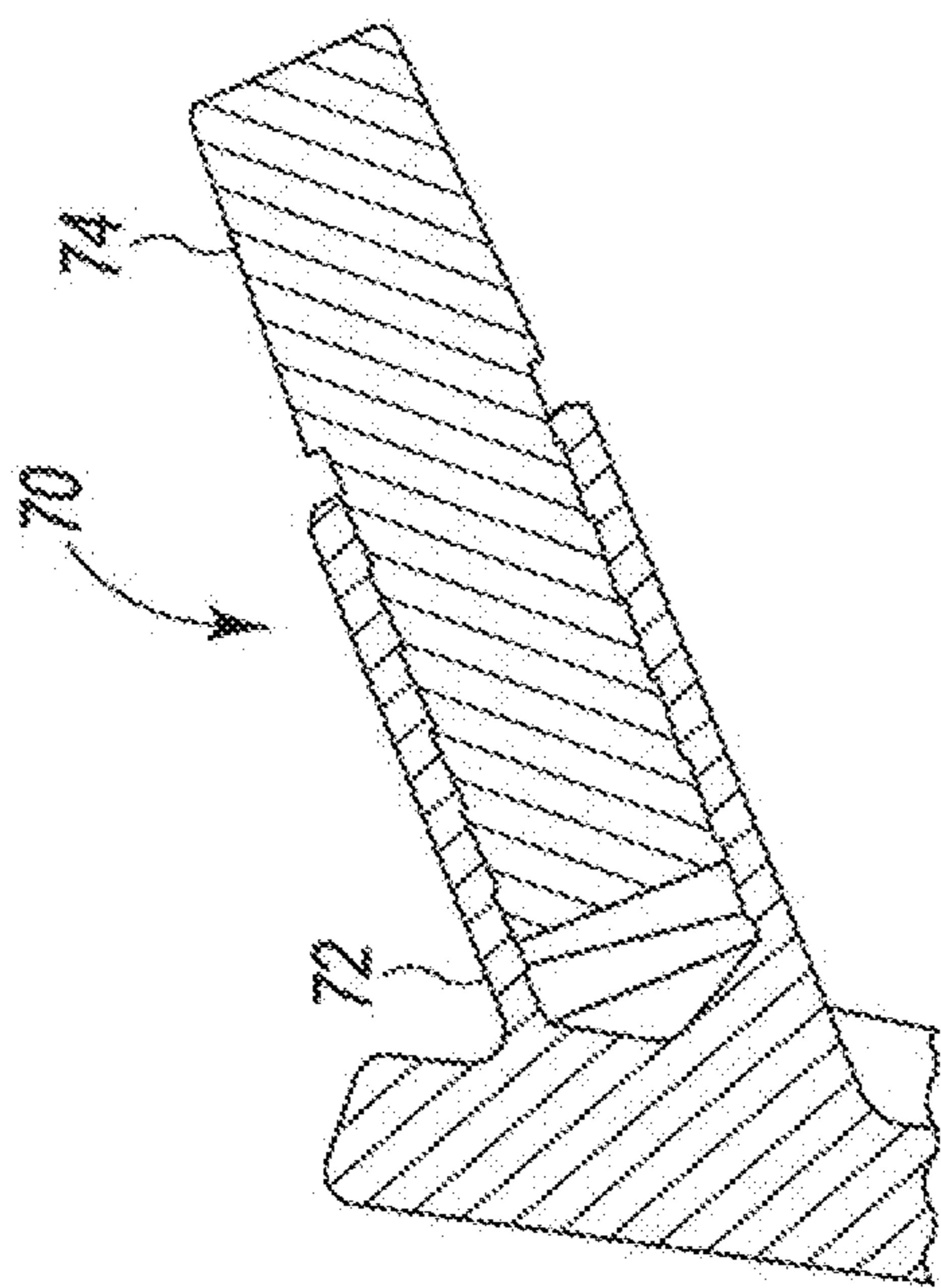


Fig. 6

ADJUSTABLE HEIGHT ARTHROPLASTY PLATE

CROSS-REFERENCE TO RELATED APPLICATION

[0001] This application claims priority to U.S. Prov. App. No. 61/706916 filed Sep. 28, 2012, entitled “Adjustable Height Arthroplasty Plate,” which is incorporated by reference herein in its entirety.

TECHNICAL FIELD OF THE INVENTION

[0002] The present invention relates generally to the field of orthopedics, and, more particularly, to an arthroplasty plate for repairing fractures of an end of a long bone.

BACKGROUND

[0003] A natural shoulder joint may undergo degenerative changes for a variety of reasons, for instance arthritis. Alternatively, a shoulder joint may be fractured or otherwise damaged by an external force. When a shoulder joint is sufficiently degenerated or damaged it may be necessary to replace the natural joint with a prosthetic shoulder joint. Conventional shoulder prostheses comprise a humeral prosthesis, optionally with a glenoid prosthesis. For total or reverse arthroplasty a glenoid component is implanted, however for hemiarthroplasty the humeral component articulates against the natural glenoid cavity.

[0004] When the upper extremity of the humerus is fractured, the humerus generally breaks into several pieces, typically three or four. In particular, for a four part fracture, the humeral head splits off at the level of the anatomical neck, the greater and lesser tuberosities are separated from the humerus shaft below the tuberosities and the tuberosities are separated from one another along the bicipital groove. As there is no longer a blood supply to the humeral head necrosis may begin. For repair of a three-part or four-part fracture, where the blood supply to the humeral head is damaged, the humeral head is replaced, and the greater and lesser tuberosities are reattached to the humeral shaft. Typically, the humeral component of a shoulder prosthesis comprises a stem for insertion into a bore extending into the intramedullary canal of the humerus, generally along the longitudinal axis of the bone, and an articulation component, which may be a convex bearing head in the case of total arthroplasty or a concave cup in the case of reverse arthroplasty. Typically, the articulation component is coupled to a neck portion of the stem, which extends from the intramedullary canal at an inclined angle relative to the longitudinal axis of a distal portion of the stem in order to recreate the anatomy of the natural joint. Commonly, humeral stem prostheses are secured in position within the intramedullary bore using bone cement. Alternatively, the stem may be coated with a material which encourages bone growth to hold the stem in position, such as Porocoat[®] or hydroxyapatite. One such humeral stem prosthesis is commercially available from DePuy Orthopaedics, Inc under the trademark Global FX.

[0005] As an alternative to humeral stem implants, it is known to repair some types of proximal humeral fracture using an intramedullary nail extending along a bore formed within the intramedullary canal. Screws pass from outside of the humerus, through holes formed within the nail and into the humeral head and the tuberosities. This type of fixation

also suffers from vascular damage in the intramedullary canal. Furthermore, a significant drawback is that because the humeral head is not replaced the nail must be inserted through a hole formed in the articular cartilage of the humeral head. A plug must be inserted into the hole to restore the bearing surface.

[0006] For the repair of fractures of the femoral neck it is known to provide a form of prosthesis generally known as a thrust plate prosthesis. One such form of thrust plate prosthesis is disclosed within PCT patent publication WO2007/024980-A2. The prosthesis comprises a plate portion to be attached to the lateral external surface of the proximal femur.

[0007] One issue in dealing with long-limb arthroplasty, is accommodating various anatomic sizes. This becomes even more of a challenge when the desired sizes of linking components that must engage and disengage and be mechanically stable. Also, surgeons desiring the ability to adjust intra-operatively creates additional issues as does the desire by surgeons to have infinite adjustability, not finite size offerings.

[0008] It is an object of embodiments of the present invention to obviate or mitigate one or more of the problems of the prior art, whether identified herein or elsewhere.

SUMMARY OF THE INVENTION

[0009] According to one embodiment of the present invention, an orthopaedic assembly is provided. The assembly includes a first member adapted to be coupled to a bone and an adjustable connector assembly including an adjuster having a threaded end and a tapered end. The assembly further includes an articulation component including a bearing surface and adapted to be coupled to the tapered end of the adjustable connector assembly. The threaded end is adapted to engage the fixation plate.

[0010] According to another embodiment of the present invention, an arthroplasty plate is provided. The arthroplasty plate includes a fixation plate adapted to be fixed to a surface of a bone, having a recess with a plurality of tapers and an adjustable connector assembly including an adjuster having a plurality of tapers on a first end a taper on a second end. An articulation component is also included and has a bearing surface and adapted to be coupled to the tapered second end of the adjustable connector assembly. The plurality of tapers are adapted to engage at least one of the plurality of tapers in the recess of the fixation plate.

[0011] According to yet another embodiment of the present invention, a method of repairing a fractured end of a bone includes removing fractured portions of an articulating bone surface at the end of a bone. The method further includes securing a fixation plate to a lateral surface of the fractured end of the bone. The fixation plate is coupled to adjustable connector assembly, which includes an adjuster having a threaded end and a tapered end. The adjuster is threaded into or out of a threaded bore of the fixation plate to adjust the length of the adjuster extending from the bore of the fixation plate. The position of the adjuster is locked within the threaded bore.

BRIEF DESCRIPTION OF THE DRAWINGS

[0012] FIG. 1 is a view in a frontal plane of an arthroplasty plate according to an embodiment of the present invention.

[0013] FIG. 2 is a cross-sectional view of the arthroplasty plate of FIG. 1.

[0014] FIG. 3 is an exploded view of the arthroplasty plate according to FIG. 1.

[0015] FIG. 4 is a cross-sectional view of an arthroplasty plate according to another embodiment of the present invention.

[0016] FIG. 5 is an exploded view of the arthroplasty plate of FIG. 4.

[0017] FIG. 6 is a cross-sectional view of an arthroplasty plate according to another embodiment of the present invention.

[0018] FIG. 7 is an exploded view of the arthroplasty plate of FIG. 6.

DETAILED DESCRIPTION

[0019] Like reference numerals refer to like parts throughout the following description and the accompanying drawings.

[0020] The below description will be given with respect to a humeral prosthesis, and more particularly, a hemiarthroplasty plate. However, it should be understood that the present invention may be used in other types of prosthesis and that the hemiarthroplasty plate is used as an example only. For example, instead of a fixation plate, a stem or other arthroplasty component could be used. And, any articulation member, not just humeral heads could be used as well.

[0021] Referring first to FIG. 1, this is a view in a frontal plane of an assembled arthroplasty plate 10 for repair of a fractured proximal humerus in accordance with an embodiment of the present invention. The arthroplasty plate comprises a fixation plate (or first member of an orthopaedic assembly) 12 for attachment to a surface of the bone, in this case, a lateral external portion of a proximal humerus of a patient. To aid in the fixation to the bone, the fixation plate 12 may include a plurality of fixation holes 13 adapted to receive bone screws (not shown). An articulation component 14 replaces the natural humeral head. The articulation component 14 comprises a convex bearing surface and is sized and shaped to articulate with either the natural glenoid surface or an implanted glenoid prosthesis to reconstruct the shoulder joint. The articulation component 14 is coupled to the fixation plate 12 via a connector assembly 16 positioned in a bore formed in the proximal humerus.

[0022] Referring now to FIG. 2, this illustrates a cross sectional view of the connector assembly of the arthroplasty plate of FIG. 1 to better illustrate details of the connector assembly 16. In this embodiment, the connector assembly 16 is an adjustable assembly to allow the user to adjust the neck length of the arthroplasty plate 10. The adjustable connector assembly 16 includes an adjuster 18, a locking screw 20, collar 22, and assembly screw 24. The collar 22 is used as a placeholder or substitute for the larger suture collar (FIG. 4).

[0023] Referring to both FIGS. 2 and 3, the various features of the adjustable connector assembly 16 will be further described. The adjuster 18 includes a first end 26 and a second end 28. The first end 26 includes a connection feature 30 adapted to couple to a corresponding connection feature 32 of the collar 22. In the illustrated embodiment, the connection feature 30 of the first end 26 is a male taper and the connection feature 32 of the collar 22 is a female taper. According to some embodiments, the male and female tapers are locking tapers, adapted to lock the adjuster 18 to

the collar 22. In some embodiments, the adjuster 18 may lock directly into the articulation component 14. In those embodiments, the connection feature 30 of the first end 26 of the adjuster 18 is a male taper and is adapted to fit into a corresponding tapered recess in the articulation component 14. The adjuster 18 also includes a through bore 29 extending between the first end 26 and the second end 28.

[0024] The second end 28 of the adjuster 18 includes a connection feature 34 adapted to connect to a corresponding connection feature 36 in the fixation plate 12. In the illustrated embodiments of FIGS. 2 and 3, the connection feature 34 of the second end 28 is a thread and the connection feature 36 in the fixation plate 12 is a corresponding thread form. In use, the adjuster 18 is inserted into through bore 38 in the fixation plate 12. As shown, the through bore 38 is threaded so as to engage the threads of the connection feature 34 of the second end 28. Once the adjuster 18 is at the desired length, the locking screw 20 is screwed into the connection feature 36 of the fixation plate 12, locking the adjuster 18 into place. In order to lock the adjuster 18 into place, the locking screw 20 includes a bore 37 and external threads 39. The external threads 39 of the locking screw 20 engage the threaded through bore 38 of the fixation plate 12, as shown in FIG. 2. The adjuster 18 and the locking screw 20 are locked relative to each other and to the fixation plate 12 on the principal of a locking nut. The adjuster thread 34 is in contact with the top flank of the threads of feature 36 of the plate 12 and the locking screw thread 39 is in contact with the opposite or bottom flank of the threads of the plate 12, making relative movement not possible. The collar 22 is then placed over the first end 26 of the adjuster 18. The tapered connection feature 32 of the collar 22 is then locked into place with the tapered connection feature 30 of the first end 26 of the adjuster 18. The articulation component 14 may then be placed onto the collar 22.

[0025] The assembly screw 24 includes a threaded end 40 and the articulation component 14 includes a corresponding threaded opening 42 (shown in FIG. 2). After the articulation component 14 is placed onto the collar 22, the assembly screw 24 is inserted through the bores 37, 29, 38 of the locking screw 20, adjuster 18, and fixation plate, respectively. The threaded end 40 of the assembly screw 24 then threads into the threads of the threaded opening 42 of the articulation component 14, locking all of the pieces together.

[0026] Turning now to FIGS. 4 and 5, another embodiment of the present invention is illustrated. As shown, the arthroplasty plate assembly 10 includes a fixation plate 12, articulation component 14, an assembly screw 24, a locking screw 20, an adjuster 18, and a collar 22a. In this embodiment, all the features are the same, with the exception that the collar 22a is a plate, having a flange 44 that extends outwardly and abuts an outwardly extending flange 46 (of FIG. 5) of the articulation component 14.

[0027] Turning now to FIGS. 6 and 7, another embodiment of the present invention is illustrated. This embodiment is of an arthroplasty plate 70 having a fixation plate 72, an adjustable connector 74, and an articulation component 14 (of FIG. 1). In this embodiment, the fixation plate 72 includes a through bore 78 that has a thread form that creates series of tapers 80a, 80b, 80c. As illustrated, the thread form is a continuous thread, but in other embodiments in can be discontinuous. In this embodiment, the adjustable connector 74 is the adjuster and includes a plurality of corresponding tapers 82a, 82b, 82c. The user would insert the adjustable

connector **74** into the through bore **78** of the fixation plate **72** and, once at the appropriate length, impacted to engage the tapers **80a**, **82a** to lock the adjuster **74** into place in the fixation plate **72**. In some embodiments, multiple tapers would engage at the appropriate length. At the other end of the adjuster is a male taper **84**. The male taper **84** is adapted to fit inside a female taper (not shown) of the articulation component **14** (of FIG. 1). The articulation component **14** is then impacted onto the adjuster **74**, locking the tapers **84**. In some embodiments, an assembly screw can be used.

[0028] The foregoing description of the invention is illustrative only, and is not intended to limit the scope of the invention to the precise terms set forth. Further, although the invention has been described in detail with reference to certain illustrative embodiments, variations and modifications exist within the scope and spirit of the invention as described and defined in the following claims.

1 to 20. (canceled)

21. A method of shoulder arthroplasty, the method comprising:

- removing an end portion of a humerus;
- securing a fixation plate to a lateral surface of the end portion of the humerus;
- coupling an articulation component to the adjustable connector assembly;

adjusting the length of the adjustable connector assembly wherein to adjust a distance between the end portion of the humerus and the articulation component; and locking the position of the adjustable connector assembly.

22. The method of claim **21**, further comprising coupling an articulation component to the tapered end of the adjuster.

23. The method of claim **22**, wherein the coupling an articulation component to the tapered end of the adjuster includes inserting a collar over the tapered end of the adjuster and inserting the articulation component onto an outer side of the collar.

24. The method of claim **21**, wherein locking the position of the adjuster within the threaded bore includes threading a locking screw into the threaded bore.

25. The method of claim **24**, further comprising inserting an assembly screw through the locking screw and the adjuster.

26. The method of claim **25**, wherein the inserting an assembly screw includes inserting a threaded end of the assembly screw through the locking screw and the adjuster.

27. The method of claim **26**, further comprising threading the threaded end of the assembly screw into a threaded recess of the articulation component.

* * * * *