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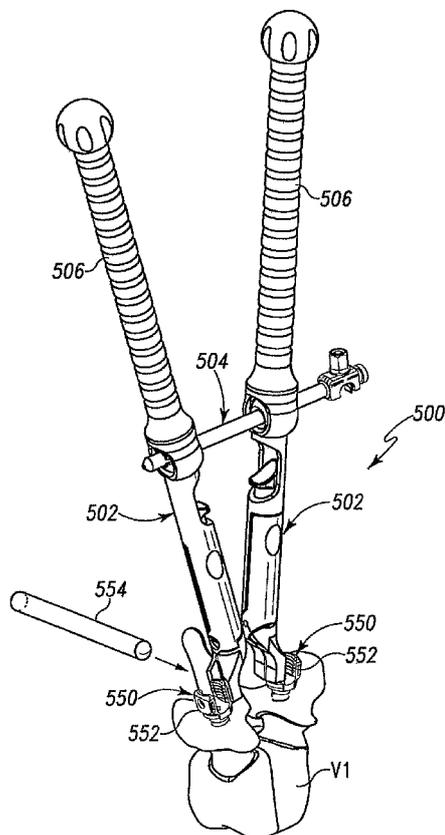
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[Continued on next page]

(54) **Title:** METHODS AND INSTRUMENTS FOR SPINAL DEROTATION



(57) **Abstract:** Derotation instrument assemblies (500, 600, 700) and systems (512, 514, 602) are provided to facilitate positioning one or more vertebrae of a spinal column into a desired alignment. The instrument assemblies (500, 600, 700) and systems (512, 514, 602) include implant holders (10, 502, 610, 710, 710") engageable to respective implants (550) engaged to vertebrae of the spinal column, transverse bridges (100, 504, 750, 770, 1100) to connect implant holders (10, 502, 610, 710, 710) associated with a particular vertebra, and inter-level linking assemblies (510, 1160) to connect instrument assemblies (500, 600, 700) associated with different vertebrae. Derotation handles (506) can be provided on the implant holders (10, 502, 610, 710, 710) to facilitate application of the alignment forces, while the assemblies (500, 600, 700) distribute the corrective forces to the connected implants and vertebrae.

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METHODS AND INSTRUMENTS FOR SPINAL DEROTATION

BACKGROUND

5 Surgical correction of the positioning and alignment of one or more vertebrae in
the spinal column can be desired to address various pathologies and conditions of patients.
However, such repositioning and re-alignment can be time-consuming, cumbersome, and
potentially difficult to achieve during a surgical procedure. For example, the alignment of
multiple vertebral levels can require manipulation of instrumentation at each level to
10 achieve the desired results. Forces applied to the vertebral body need to be controlled to
minimize stresses on the vertebral bodies and implants. Furthermore, the alignment at one
level should be maintained while other levels are aligned. In addition, the instrumentation
employed to achieve the alignment can hinder placement of stabilization constructs that
post-operatively maintain the corrected positioning and alignment achieved during
15 surgery.

 Therefore, instruments, methods and systems that facilitate surgical correction of
the alignment and positioning of a vertebra or vertebrae of the spinal column would be
desirable. Furthermore, instruments, methods and systems that facilitate placement of
stabilization constructs that post-operatively maintain the corrected vertebra or vertebrae
20 are also desirable. In addition, instruments, methods and systems that facilitate control of
the stress exerted on implants and vertebrae to which the implants are attached would be
desirable.

BRIEF DESCRIPTION OF THE DRAWINGS

25 Fig. 1 is a perspective view of a derotation instrument assembly coupled to
implants engaged to a vertebra.

 Fig. 2 is a perspective view of a derotation system for multiple vertebral levels.

 Fig. 3 is a perspective view of another derotation system for multiple levels.

 Fig. 4 is a perspective view of a transverse bridge.

30 Fig. 5 is a longitudinal sectional view of the transverse bridge of Fig. 4.

 Fig. 6 is a perspective view of a connector assembly.

 Fig. 7 is a longitudinal section view of one embodiment implant holder.

Fig. 8 is a longitudinal section view of a locking member useable with the implant holder of Fig. 7.

Fig. 9 is an elevation view of a clamp portion of the implant holder of Fig. 7 in a closed position.

5 Fig. 10 is an elevation view of the clamp portion of Fig. 9 in an open position.

Fig. 11 is an elevation view of the clamp portion of Fig. 9 rotated 90 degrees about its longitudinal axis.

Fig. 12 is a section view along line 12-12 of Fig. 11.

Fig. 13 is a perspective view of a first arm of the implant holder of Fig. 7.

10 Fig. 14 is a perspective view of a second arm of the implant holder of Fig. 7.

Fig. 15 is a sectional view looking proximally at a latch member of the implant holder of Fig. 7.

Fig. 16 is a perspective view of a holding member of the implant holder of Fig. 7.

Fig. 17 is a perspective view of a release button of the implant holder of Fig. 7.

15 Fig. 18 is a perspective of another embodiment derotation instrument assembly engaged to a vertebra.

Fig. 19 is a partially exploded perspective view of a portion of the derotation instrument assembly of Fig. 18.

20 Fig. 20 is a perspective view of a derotation system employing a number of derotation instrument assemblies of Fig. 18 along inter-linked vertebral levels.

Fig. 21 is a perspective view of another embodiment implant holders engaged to implants on respective sides of a vertebral body.

Fig. 22 is a perspective view of a transverse bridge being engaged between the implant holders of Fig. 21.

25 Fig. 23 shows another embodiment transverse bridge engaged between the implant holders of Fig. 21.

DETAILED DESCRIPTION OF THE ILLUSTRATED EMBODIMENTS

30 For the purposes of promoting an understanding of the principles of the invention, reference will now be made to the embodiments illustrated in the drawings and specific language will be used to describe the same. It will nevertheless be understood that no limitation of the scope of the invention is thereby intended. Any such alterations and

further modifications in the illustrated devices, and such further applications of the principles of the invention as illustrated herein are contemplated as would normally occur to one skilled in the art to which the invention relates.

5 Spinal derotation instrumentation is provided to affect one or more derotation maneuvers on a scoliotic spine or on a spine having one or more displaced, misaligned or curved vertebral levels. Specifically, a derotation instrument assembly is attached to at least one vertebral body, with the assembly including at least a pair of bone implants anchored to the vertebral body along the left and right sides of the spinal column; an elongate implant holder removably attached to a head portion of each of the bone
10 implants; a transverse bridge interconnecting the proximal end portions of the implant holders; and a primary handle extending from each of the implant holders proximally of the location in which the transverse bridge is engaged to the respective implant holder. The handles are provided for manipulation by the surgeon, and manipulation forces exerted on one handle are distributed to each implant/vertebra interface by the transverse
15 bridge and other implant holder.

The derotation handles can extend proximally and axially from a distal clamping portion of each of the implants. In one embodiment, only one of the implant holders includes a derotation handle, and the transverse bridge is engaged directly to the proximal end of the other implant holder. The derotation handle or handles may be formed unitarily
20 with the distal clamping portion of the respective implant holder, or may be threadingly or otherwise suitably removably engaged to the distal clamping portion of the respective implant holder to allow for selective attachment and removal.

The bone implants can be configured as pedicle screws, with each screw having a head portion which includes a pair of arms defining a U-shaped channel for receiving a
25 spinal rod, and with the arms defining internal threads for threadingly receiving a set screw for capturing the spinal rod within the U-shaped channel. The screw can be uni-axial, or multi-axial so that the head can pivot relative to the bone engaging portion. . In the illustrated embodiment, the head portions of the screws are configured to receive stabilization element either through a top opening between the pair of arms or to receive
30 an end of the stabilization element as it is passed through the head in an end-wise manner. In another embodiment, the head portion of the screws opens to a side so that the stabilization element can be side-loaded therein. Other embodiments contemplate any

suitable type of implant that can be engaged to a vertebra and coupled to an elongated stabilization element.

The elongate implant holders can each include a distal clamp portion with a distal end portion configured for selective clamping to either arm of the screw head portion. In another embodiment, the implant holder clamps across both arms of the implant. In one specific embodiment, the implant holder includes a clamp portion having a tubular body extending the length of the implant holder and a clamping arm pivotally attached to the tubular body via a pivot pin. A spring may be included for biasing the clamping arm toward an open position along with a releasable latching mechanism to releasably capture the arm of the screw head portion between the distal end portions of the tubular body and the clamping arm. The implant holder may also include a release button to selectively release the tubular body and the clamping arm from the arm of the screw head portion. The implant holder can include a length so that at least its proximal handle is positioned outside the patient through the wound or incision in which the vertebrae are accessed.

The location of the implant holder between the distal clamping end and the proximal handle (or the proximal end of the implant holder if a handle is not provided) can include a joint to facilitate attachment of the transverse bridge at any one of a number of angular orientations relative the implant holder. The joint can be of any suitable configuration, and specific embodiments contemplate a spherical segment unitarily joined with the implant holder, a ball and socket arrangement, or a ball end, for example. The joint of at least one of the implant holders associated with a vertebra can be engaged to the transverse bridge via a clamping mechanism or interference fit that allows at least one implant holder to be engaged to the connecting member at any one of a number of positions along the connecting member. Such engagement between the implant holders and the transverse bridge allows for variable lateral adjustment and variable angular adjustment of the implant holders relative to the transverse bridge. In still another embodiment, the transverse bridge can connect implant holders engaged to respective ones of two or more vertebrae, and extend across the spinal midline to link the implant holders to one another.

In one embodiment, the transverse bridge is configured as a plate defining an elongate slot extending therethrough. One end of the plate is engaged about a spherical joint of one implant holder, and the slot includes a number of recesses or scalloped areas

that can each receive a spherical joint of the other implant holder in any one of a plurality of angular orientations. In another embodiment, the transverse bridge includes a rod-like member positioned through ball-joint mechanisms of the implant holders. In another embodiment, the connecting member is engaged to clamp assemblies that are engaged to the implant holder.

In instances requiring derotation across multiple vertebral levels, a derotation instrument assembly may be attached to respective ones of the multiple vertebral bodies requiring derotation, with the derotation instrument assemblies being interconnected by an inter-level linking assembly coupled between the individual derotation instrument assemblies. As a result, the surgeon may manipulate an integrated frame assembly to affect derotation across multiple vertebral levels, rather than separately manipulating several derotation instrument assemblies to effect derotation at each individual vertebral level. The transverse bridges can be releasably coupled to the implants holders associated with each of the vertebrae such that the spacing and angular orientation between implant holders associated with a particular vertebra can be readily adjusted and maintained by engagement with the respective transverse bridge. The inter-level linking assemblies can be releasably coupled to connecting members of the transverse bridges, for example, such that the spacing and angular orientation between the linked derotation instrument assemblies can be readily adjusted and maintained with clamping and connector assemblies that secure the derotation instrument assemblies to an elongate link member extending between the derotation instrument assemblies.

The inter-level linking assemblies can interconnect the derotation instrument assemblies in a rigid fashion so that the engagement relationship between the components is maintained during derotation of the spinal column. It is further contemplated that at least limited slippage or movement between the inter-level linking assemblies and the derotation instrument assemblies can be provided as the spinal column is straightened to accommodate non-uniform relative displacement among the corrected vertebrae that may be required.

In Fig. 1 there is shown one embodiment of a derotation instrument assembly coupled to implants 550. Implants 550 are engaged to a vertebral body VI. In one specific application, implants 550 are bone anchors secured to respective ones of the pedicles of vertebral body VI. Implants 550 each include a receiver portion 552 for

receiving a respective elongated spinal stabilization element 554 positionable along the spinal column and securable to the implants to maintain a positioning of one or more vertebral bodies. In the illustrated embodiment, the implants are bone screws with a U-shaped head portion providing a receiver to receive a spinal rod. Other embodiments
5 contemplate saddles, posts, clamping members, side-loading receivers or other receiver type members extending from a bone engaging portion in the form of a staple, hook, screw, interbody device, intrabody device or other bone engaging member.

Derotation instrument assembly 500 includes implant holders 502 removably engaged to respective ones of the implants 550 and extending proximally therefrom. The
10 implant holders 502 can be interconnected with one another in a bilateral fashion with a transverse bridge 504 extending therebetween. Each of the implant holders 502 further includes a derotation handle 506 extending proximally from the location along the implant holder 502 to which transverse bridge 504 is engaged. Derotation handles 506 extend in a direction that is generally parallel to and/or forms an axial extension of the clamp portion
15 of the respective implant holder 502, and thus extends in a direction that is generally parallel to the sagittal plane of the spinal column when implants 550 are engaged to the pedicles.

Derotation instrument assembly 500 can be manipulated with one or both of derotation handles 506 to displace, pull, twist or align the vertebra to which implants 550
20 is engaged into the desired alignment with the spinal column. Accordingly, manipulation of multiple anchors engaged to the spinal column can be completed with a single-handle, although the application of such forces through multiple handles is not precluded. The interconnection of the implants 550 results in the corrective forces being distributed to both implants and thus to multiple locations on the vertebral body. This can reduce stress
25 concentrations at any single bone/implant interface as the manipulation forces are applied.

It is further contemplated that a number of derotation instrument assemblies 500 can be coupled to one another by one or more inter-level linking assemblies 510 with a link member 511 extending between and coupled to, for example, transverse bridges 504
30 and 504A of the respective derotation instrument assemblies 500 and 500A shown in Fig. 2. The inter-level, linked instrument assemblies 500, 500A provide a derotation system 512 that facilitates the application of and distribution of derotation, correction, alignment and other forces to various bony structures engaged by the bone implants and

interconnected within the system, such as first and second vertebra V1 and V2.

Accordingly, the resultant stress on any one of the implants and the bone to which the implant is engaged is distributed to multiple locations and/or multiple vertebrae.

It is contemplated that any one, two or three or more vertebral levels with
5 derotation instrument assemblies 500 can be linked. It is further contemplated that any subset of instrumented vertebral levels in a system could be linked. For example, Fig. 3 shows a derotation system 514 for three vertebrae V1, V2 and V3. Derotation instrument assemblies 500 and 500A are linked by inter-level linking assembly 510, and derotation instrument assemblies 500A and 500B are linked by a second inter-level linking assembly
10 510A with a second linking member 511A.

Referring now to Figs. 4-5, one specific example of a transverse bridge 504 is shown in the form of transverse bridge 100. Transverse bridge 100 includes a transverse connecting member 102 and a clamping assembly 140 positioned at one end of connecting member 102. Clamping assembly 140 can be removably engaged to connecting member
15 102 or formed as an integral unitary component therewith.

Transverse connecting member 102 can have a circular shaped cross-section as shown in section view in Fig. 4. Other cross-sectional shapes are also contemplated, including diamond, square, rectangular, polygonal, and non-circular shapes, for example. Connecting member 102 extends between a first end 103 and a second end 104. First end
20 103 can be tapered as shown to facilitate placement of connecting member 102 through an implant holder, as shown in Figs. 1-3. A non-tapered first end 103 is also contemplated. Second end 104 can include clamping assembly 140 either removably or non-removably attached thereto.

Clamping assembly 140 includes a clamping portion 142 and a securing portion
25 144 that is operable to secure and release clamping portion 142 to transverse connecting member 102. Connecting member 102 includes a mounting member 106 adjacent second end 104, and a bore 108 extending through mounting member 106 through which clamping portion 142 extends. Securing portion 144 further includes a bore 148 for receiving a proximal post 150 of clamping portion 142. Securing portion 144 can include
30 internal threads in bore 148 to threadingly engage post 150.

Clamping portion 142 further includes clamping arms 152, 154 at an end of post 150 and a hinge portion 156 between arms 152, 154. Arms 152, 154 are positioned in

recess 110 of mounting member 106 on a side opposite of securing portion 144. Securing portion 144 is rotatable to threadingly displace clamping portion 142 relative thereto. For example, securing portion 144 can be rotated to displace clamping portion 142 in the direction of arrow 160, as shown in Fig. 5. This in turn presses arms 152, 154 against mounting member 106 and moves arms 152, 154 toward one another into engagement with a linking member or other element positioned therebetween, as shown in Figs. 2 and 3.

Mounting member 106 further includes an end flange 112 forming an end of connecting member 102 and a groove 114 adjacent to end flange 112. End flange 112 can receive a connector assembly 160, such as shown in Fig. 6. Connector assembly 160 can be similar to clamping assembly 140 discussed above, but is configured without a connecting member. In contrast, connector assembly 160 includes an end opening defining a receiving slot 161 adjacent one end thereof to receive, for example, end flange 112 of connecting member 102. This allows a connecting member 102 to be extended with one or more additional connector assemblies 160, such as shown in Fig. 3, to receive a linking member of another inter-level linking assembly.

Connector assembly 160 can be provided with a clamping portion 162 and a securing portion 164 mounted to a body 165. The securing portion 164 is operable to move the clamping portion into contact with body 165 and close arms 172 (only one shown in Fig. 6) of clamping portion 162 about, for example, the link member 514 of an inter-level linking assembly 510. Body 165 can further include an end flange 168 and a groove 170 adjacent thereto to receive another connector assembly 160 in end-to-end relation.

Referring now to Fig. 7, one specific example of implant holder 502 will be discussed with respect to an implant holder 10. Implant holder 10 includes a clamp portion 11 and a handle portion 30 extending axially and proximally from clamp portion 11. A joint 80 is provided between clamp portion 11 and handle portion 30 for engaging transverse bridge 504.

Joint 80 can be formed by an end member 82 at the proximal end of clamp portion 11. End member 82 defines a proximally opening receptacle that houses a ball member 84. Ball member 84 includes a passage 86 extending therethrough to receive, for example, connecting member 102 of transverse bridge 100. Ball member 84 is rotatable so that the

angular orientation between implant holder 10 and connecting member 102 can be easily adjusted and accommodated. Ball member 84 further includes relief 88 that allows the ball member to flex and securely engage the connecting member 102 in passage 86 when compressed.

5 Handle portion 30 extends proximally from joint 80, and can include a distal end 32 threadingly engaged to end member 82 to capture ball member 84 in the receptacle. Handle portion 30 further includes an elongated shaft portion 34 defining an internal passage 36 extending axially therethrough. Internal threads 38 are provided adjacent the distal end of passage 36.

10 In Fig. 8 there is shown a locking member 90 that is positionable through passage 36 and into engagement with ball member 84. Locking member 90 includes an elongated shaft 92 extending proximally from a distal engaging end 93. Engaging end 93 can include a threaded portion 94 to threadingly engage internal threads 38 of handle portion 30. Engaging end 93 further includes an end wall 95 that can be concave, inwardly curved
15 or otherwise shaped to engage ball member 84 when shaft 92 is positioned in internal passage 36 of handle portion 30.

 A knob member 96 is provided at the proximal end of shaft 92. Knob member 96 defines an axially extending and distally opening receptacle 97 about shaft 92. The proximal end 35 of shaft portion 34 of handle portion 30 can be received in receptacle 97
20 as locking member is distally, threadingly advanced into handle portion 30. In particular, the length of shaft 92 is sized so that knob member 92 resides proximally of the proximal end 35 of handle portion 30 when threaded portion 94 is threadingly engaged to internal threads 38. Knob member 92 can be grasped by the user to rotate locking member 90 and threadingly advance engaging end 93 into contact with ball member 84. As locking
25 member 90 presses against ball member 84, ball member 84 compresses about relief 88 and into firm engagement with the connecting member 102 in passage 86.

 Referring now to Figs. 9-17, clamp portion 11 of implant holder 10 will be further discussed. Clamp portion 11 includes a first arm 12 in the form of a tubular body and a second arm 14 providing a clamping arm that is pivotally coupled to first arm 12. Each of
30 the first and second arms 12, 14 includes a respective distal end portion 16, 18 of a distal holding end 20 of implant holder 10. Each of the portions 16, 18 forms a space in which to receive a portion of the bone implant, and further includes a projection 17, 19 extending

into the space toward the other portion 16, 18. The projections 17, 19 are received in detents formed in the receiver of the implant to which holder 10 is engaged by clamping arms 12, 14 to the receiver of the implant when implant holder 10 is closed, as shown Fig. 9 for example. To release the implant, implant holder 10 is opened by pivoting second arm 14 about pivotal connection 22 with first arm 12, as shown in Fig. 10.

Arms 12, 14 cross-over one another in a scissors type arrangement, and include inter-fitting recessed portions 30, 32, respectively, at connection 22 so that end portions 16, 18 are aligned with one another. Furthermore, as shown in Fig. 11, arms 12, 14 include a slight bend so that end portions 16, 18 are offset to one side of the longitudinal axis 13 of implant holder 10. Arms 12, 14 are movable relative to one another about connection 22 in a first plane that includes longitudinal axis 13. Distal portions 16, 18 are movable in a relative to one another by pivoting arms 12, 14, and move in a second plane that is generally parallel to the plane including longitudinal axis 13. In addition, the space between end portions 16, 18 opens away from axis 13 to so that the implant to which implant holder 10 is engaged can remain substantially unobstructed for engagement with another implant or system component.

Arms 12, 14 are spring biased toward the open position with a spring 24 positioned in wells 26, 28 formed by respective ones of the arms 12, 14. Wells 26, 28 are oriented toward one another, and located proximally of the pivotal connection 22 between arms 12, 14. In order to secure arms 12, 14 in the closed position in engagement with the implant, a latching mechanism 40 is provided between arms 12, 14. Latching mechanism 40 includes a latch member 42 extending from second arm 14 and a holding member 50 mounted to first arm 12 that is releasably engageable by latch member 42. Latching mechanism 40 also includes a release button 70 coupled to and extending proximally from holding member 50 between arms 12, 14, and a spring 44 biasing holding member 50 into engagement with latch member 42 and further biasing release button 70 proximally.

First arm 12 includes a collar 48 extending therefrom into a receptacle defined between arms 12, 14 in which latching mechanism 40 is located. Holding member 50 extends through collar 48 and is axially movable therein while collar 48 maintains holding member 50 in axial alignment with the remaining portions of latching mechanism 40. In addition, an alignment pin 46 can be press fit in collar 48 and extend therefrom into a slot 52 (Fig. 16) along a portion of the length of holding member 50 to maintain holding

member 50 in rotational alignment with latching mechanism 40. Other embodiments contemplate that collar 48 and/or alignment pin 46 can be eliminated.

Holding member 50 is shown further in Fig. 16. Holding member 50 includes a central body 54 defining axial slot 52 therealong. A connector portion 56 extends from a proximal end of central body 54, and is threadingly received in a distal end opening of release button 70, as shown in Fig. 12. The distal end of central body 54 includes a radially outwardly extending flange 58 that abuttingly engages collar 48 to limit the proximal displacement of release button 70 and holding member 50 under the bias of spring 44.

Holding member 50 also includes a stem 60 extending distally from flange 58 to a latch receiving member 62. Latch receiving member 62 includes a cylindrical body with inclined notched areas 64 and a central projection area 65 between inclined notched areas 64. In the open position, latch member 42 includes sloped portions 43 that reside along inclined notches areas 64, as shown in Fig. 14. When closing arms 12, 14 to engage the implant between portions 16, 18, the sloped portions 43 slide along the respective adjacent inclined notched areas 64 to distally and axially displace holding member 50 until the receptacle 45 (Fig. 15) of latch member 42 aligns with and receives the cylindrical body of latch receiving member 62, as shown in Fig. 12. In the closed position, arm 14 is prevented from pivoting away from arm 12 by engagement of latch member 42 around receiving member 62 of holding member 50. To release latch mechanism 40 and allow arm 14 to pivot away from arm 12, release button 70 is depressed to displace holding member 50 distally sufficiently to align stem 60 with slotted opening 47 (Fig. 15) of latch member 42. This allows receptacle 45 to become disengaged or displaced from about latch receiving member 62, and spring 44 pushes arm 14 away from arm 12 and rotates arm 14 about connection 22 to the open position of Fig. 10.

Release button 70 is further shown in Fig. 17, and includes a body portion 72 extending between a distal end member 74 and a proximal end 76. Distal end member 74 defines the opening which receives connector portion 56 of holding member 50. Proximal end 76 includes a concavely curved surface to facilitate application of manual depression forces with a thumb or finger to proximally displace button 70 and thus latch mechanism 40 between arms 12, 14. Button 70 is accessible through a notched area 13 of first arm 12, as shown in Fig. 11. An outwardly extending lip 78 adjacent proximal end 76 can contact

first arm 12 in notched area 15 to maintain alignment of release button 70 as it is moved therein and to limit distal displacement of button 70.

Other embodiment transverse bridge, implant holder and derotation instrument assemblies and systems are also contemplated. For example, Figs. 18 and 19 show a
5 derotation instrument assembly 600 engaged to implants 550 that are engaged to respective pedicles of vertebra VI. Derotation instrument assembly 600 includes implant holders 610 each having a distal clamp portion 611 releasably engageable to respective ones of the implants 550 and a proximal handle portion 630. A transverse bridge 1100 extends between and is engaged to implant holders 610 at a joint 631 located between
10 handle portion 630 and clamp portion 611 of each of the implant holders 610.

Transverse bridge 1100 includes an elongated transverse connecting member 1102 in the form of a connecting rod with end flanges 1104, 1106 at opposite ends thereof. Transverse bridge 1100 further includes clamping assemblies 1140 at or adjacent to the ends of connecting member 1102 that are movable along connecting member 1102 to
15 allow adjustment in the spacing between the implant holders 610. Clamping assemblies 1140 each include a clamping portion 1142 that includes a split-ring type clamping member that is positioned around implant holder 610 at joint 631, and further includes arms 1144, 1146 that are moveable to selectively release and securely engage clamping assembly 1140 at joint 631 of implant holder 610.

Clamping assembly 1140 further includes a securing portion 1150 having a first end 1152 having a ring-shape defining a receptacle for receiving connecting member 1102 therethrough and a mounting portion 1154 in the form of a post extending from first end 1152. Arms 1144, 1146 define a bore through which mounting portion 1154 extends. A seating washer 1156 is positioned about mounting portion 1154 between first end 1152
25 and arm 1146, and a locking member 1158 is positioned about mounting portion 1154 adjacent arm 1144. Locking member 1158 can be in the form of a wing nut threadingly engaged to the post-like structure of mounting portion 1154, although other engagement relationships and forms for locking member 1158 are contemplated. As locking member 1158 is advanced along mounting portion 1154, first end 1152 of securing portion 1150 is
30 drawn toward locking member 1158, and connecting member 1102 seats against seating washer 1156, and arms 1144, 1146 are compressed between washer 1156 and locking member 1158 to tightly grip joint 631 of the respective implant holder 610. Connecting

member 1102 is clamped between seating washer 1156 and first end 1152 to lock clamping assembly 1140 in position along connecting member 1102.

Derotation instrument assembly 600 can be engaged to multiple vertebrae V1, V2, V3 with an inter-level linking assembly 1160, as shown in Fig. 20, for example. Linking assembly 1160 includes an elongated link member 1162 that extends between and is engaged to the connecting members of each of the transverse bridges 1100, 1100A and 1100B. In the illustrated embodiment, linking assembly 1160 include clamp assemblies like clamp assembly 1140 discussed above that extend between and engage linking member 1162 and to the connecting member of the transverse bridges to provide a derotation system 602, such as shown Fig. 20.

Referring now to Fig. 21, there is shown another embodiment implant holder 710 engaged to one of the implants 550 and a further embodiment implant holder 710' engaged to another of the implants 550. Implant holder 710' includes a distal clamp portion 711' that can be configured similarly to clamp portion 11 of implant holder 10 discussed above. However, implant holder 710' includes a joint 730' at the proximal end of clamp portion 711' and does not include a handle or other structure extending proximally from joint 730'. Implant holder 710, on the other hand, includes a joint 730 at the proximal end of clamp portion 711 and a handle portion 740 extending proximally from joint 730. In both embodiments, joint 730, 730' is configured as a ball-like member or a portion with an outer spherically shaped surface.

In Fig. 22 there is shown a transverse bridge 750. Transverse bridge 750 includes a handle portion 752 at one end thereof, and a connecting member 754 extending from handle portion 752. An end portion 756 of connecting member 754 includes a fork-like shape to receive joint 730 of implant holder 710 therein in pivoting relation. A slotted portion 758 extends between end portion 756 and handle portion 752. Slotted portion 758 defines a central slot 760 with a number of scallops or partially-spherical receiving areas to receive joint 730' therein at any one of the number of locations along the slotted portion 1158.

The partially spherical interface between connecting member 754 and the implant holders 710, 710' allows engagement in any one of a number of angular orientations therebetween. The transverse bridge 750 couples implant holders 710, 710' to one another to provide a derotation instrument assembly 700 that distributes derotation forces applied

with handle portion 730 to each of the implants 550. Furthermore, handle portion 752 of transverse bridge 750 provides a handle that is transversely oriented to the implant holders 710, 710' and when implants 550 are engaged to the pedicles to the sagittal plane of the spinal column. Accordingly, derotation forces and other maneuvers applied through
5 handle portion 752 can likewise be distributed to each of the implants 550.

Fig. 23 shows another embodiment transverse bridge 770 extending between and engaged to implant holders 710, 710'. Transverse bridge 770 includes an elongated connecting member 772 and clamping members 774, 776 at opposite ends of connecting member 772. Clamping members 774, 776 can form a split-ring type arrangement with
10 free ends that can move relative to one another to fit around and clampingly engage the respective joints 730, 730' of implant holders 710, 710'. The free ends of the clamping members can be engaged toward one another to securely clamp the clamping members 774, 776 about the respective joint 730, 730'.

While the invention has been illustrated and described in detail in the drawings and
15 foregoing description, the same is to be considered as illustrative and not restrictive in character. All changes and modifications that come within the spirit of the invention are desired to be protected.

What is claimed is:

1. A system for correcting alignment of one or more vertebrae of a spine, comprising:

5 first and second implants engageable to one of the one or more vertebrae, said first and second implants being configured for engagement with respective ones of first and second stabilization elements positionable along the spine; and

10 a derotation instrument assembly including first and second elongated implant holders releasably engageable to respective ones of said first and second implants, a transverse bridge positionable between and engageable to each of said first and second implant holders at a location spaced proximally from said first and second implants, wherein each of said first and second implant holders includes a derotation handle extending proximally from said location, said derotation handles being associated with said respective implant holders in a manner that permits forces applied to either of said handles to be directed to said transverse bridge, thereby manipulating both said first and second implant holders when engaged to said bridge and both said first and second implants when engaged to said first and second implant holders.

20 2. The system of claim 1, wherein said implant holders each include a distal clamp portion distal of said location, said distal clamp portion including a first arm and a second arm pivotally coupled to said first arm, said location including a joint located adjacent a proximal end of said first arm.

3. The system of claim 2, wherein said joint includes a partially spherical shape.

4. The system of claim 2, wherein said joint includes a ball member pivotally captured in a receptacle formed adjacent said proximal end of said first arm.

25 5. The system of claim 4, wherein said transverse bridge includes an elongated connecting member positionable through a passage formed by each of said ball members of said implant holders.

30 6. The system of claim 5, wherein said handles each includes a locking member movable between a first position wherein said ball member is pivotal in said receptacle and a second position wherein said locking member engages said ball member in said receptacle and secures it in position therein.

7. The system of claim 6, wherein said ball member includes a relief and said locking member is operable to compress said ball member about said elongate member when engaged to said ball member.

5 8. The system of claim 6, wherein said handle includes an elongated shaft portion defining an axial passage, said locking member including an elongated shaft axially received in said passage in threaded engagement with said shaft portion, said locking member including a proximal knob located proximally of a proximal end of said shaft portion of said handle, said knob defining an axially extending and a distally opening receptacle for receiving said proximal end of said shaft portion.

10 9. The system of claim 1, further comprising a second derotation instrument assembly including third and fourth elongated implant holders releasably engageable to respective ones of third and fourth implants engageable to a second one of the one or more vertebrae, a second transverse bridge positionable between and engageable to each of said third and fourth implant holders at a location spaced proximally from said third and fourth implants, wherein each of said third and fourth implant holders includes a derotation handle extending proximally from said location, said derotation handles being associated with said respective implant holders in a manner that permits forces applied to either of said handles to be directed to said second transverse bridge, thereby manipulating both said third and fourth implant holders when engaged to said second transverse bridge and both said third and fourth implants when engaged to said third and fourth implant holders.

15 20 10. The system of claim 9, further comprising an inter-level linking assembly extending between and engaged to each of said transverse bridges.

11. The system of claim 10, wherein said inter-level linking assembly includes an elongate link member extending between said transverse bridges and said transverse bridges each include an elongate connecting member including a clamping assembly removably engageable to said elongate link member.

25 30 12. The system of claim 11, wherein said clamping assemblies each include a clamping portion and a securing portion mounted to said respective connecting member of said transverse bridges, wherein said securing portion is operable to displace said clamping portion relative to said connecting member and move a pair of arms of said clamping portion into contact with said connecting member and relative to one another to clampingly engage said link member between said pair of arms.

13. The system of claim 12, wherein said clamping portion further includes a post extending from said arms through said connecting member, wherein said securing portion threadingly engages said post on a side of said connecting member opposite said pair arms, said securing portion being operable to displace said arms against said connecting member thereby moving said arms toward one another.

14. The system of claim 13, wherein said connecting member includes a mounting portion adjacent one end thereof and said securing portion and clamping portion are mounted to said mounting portion, said connecting member further including a flange extending from said mounting portion and forming an end of said connecting member and further comprising a connector assembly removable engageable to said flange.

15. The system of claim 14, wherein said connector assembly includes:
a body defining a bore and a recess in one side of said body in communication with said recess;

a clamping portion including a pair of arms in said recess and a post extending through said bore; and

a securing portion threadingly engaged with said post and operable to displace said arms against said body in said recess thereby moving said arms toward one another.

16. The system of claim 1, wherein said implant holders each include a distal clamp portion extending along a central longitudinal axis and said clamp portion includes a first arm and a second arm pivotally coupled to said first arm and movable relative to one another in a first plane including said longitudinal axis, said first and second arms each including a distal clamping portion offset to a side of said longitudinal axis and said distal clamping portions move relative to one another in a second plane that is generally parallel to the first plane.

17. A system for correcting alignment of one or more vertebrae of a spine, comprising:

first and second implants engageable to one of the one or more vertebrae, said first and second implants being configured for engagement with respective ones of first and second stabilization elements positionable along the spine; and

a derotation instrument assembly including first and second elongated implant holders releasably engageable to respective ones of said first and second implants, wherein at least one of said implant holders includes a first arm and a second arm extending along

a longitudinal axis, said first and second arms each being pivotally connected to one another with distal portions thereof offset to a first side of said longitudinal axis and movable between a closed position in clamping engagement with a respective one of said first and second implants and an open position for releasing said respective implant, said derotation instrument assembly further including a transverse bridge positionable between and engageable to each of said first and second implant holders at a location spaced proximally from said first and second implants and proximally of said pivotal connection of said arms of said at least one implant holder.

18. The system of claim 17, wherein each of said implant holders further comprises a derotation handle extending proximally from said location on each of said implant holders.

19. The system of claim 17, wherein said distal end portions further define an opening along one side thereof extending along the longitudinal axis that opens away from said longitudinal axis.

20. The system of claim 19, wherein said distal end portions are movable toward one another to grip said implant therebetween by pivoting a proximal end of said second arm toward said first arm.

21. The system of claim 20, wherein said distal end portions each include a projection extending toward the other of said distal end portions, said projection being received in detents in said implant when said implant holder is in said closed position.

22. The system of claim 17, wherein said at least one implant holder includes a latching mechanism for locking said first and second arms in said closed position.

23. The system of claim 22, wherein said latching mechanism includes a holding member between said arms and axially movably therebetween, said latching mechanism further including a latch member removably engageable to said holding member to maintain said first and second arms in said closed position.

24. The system of claim 23, further comprising a release button engaged with said holding member, said release button being operable to release said holding member from said latch member when said first and second arms are in said closed position to permit said arms to be moved to said open position thereby releasing said implant engaged between distal end portions of said first and second arms.

25. The system of claim 24, wherein said first and second arms are spring biased toward said open position.

26. The system of claim 17, wherein said first arm includes a length along said longitudinal axis that is greater than a length of said second arm along said longitudinal axis, wherein said first arm includes a receptacle at said location housing a pivotal ball member for engagement with a portion of said transverse bridge.

27. The system of claim 26, wherein said first arm includes a handle portion extending proximally from said location, and further comprising a locking member in said handle portion selectively engageable with said ball member to lock said ball member in position about said portion of said transverse bridge.

28. A system for correcting alignment of one or more vertebrae of a spine, comprising:

first and second implants engageable to respective first and second vertebrae, said first and second implants being configured for engagement with a stabilization element positionable along the spine between said first and second implants;

a first derotation instrument assembly including a first implant holder releasably engageable to said first implant, a first transverse bridge engaged at a location between distal and proximal ends of said first implant holder, said first transverse bridge extending from said first implant holder to another implant holder associated with the first vertebra for engagement with the another implant holder, wherein said first implant holder includes a derotation handle extending proximally from said location;

a second derotation instrument assembly including a second implant holder releasably engageable to said second implant, a second transverse bridge engaged at a location between distal and proximal ends of said second implant holder, said second transverse bridge extending from said second implant holder to another implant holder associated with the second vertebra for engagement with the another implant holder, wherein said second implant holder includes a derotation handle extending proximally from said location; and

an inter-level linking assembly coupled to said first and second transverse bridges of each of said first and second derotation instrument assemblies.

29. The system of claim 28, wherein each of said first and second implant holders includes a joint at said location, said joint including a ball member pivotally captured in a receptacle of said implant holder.

5 30. The system of claim 29, wherein each of said first and second transverse bridges includes an elongated connecting member positionable through a passage in said ball member of said respective first and second implant holders, said first and second implant holders further each including a locking member operable to lock said ball member in position in said receptacle about said connecting member.

10 31. The system of claim 30, wherein said linking assembly including an elongate linking member and each of said first and second transverse bridges includes a clamping assembly operable to clampingly engage said link member.

32. A method for assembling a system for correcting alignment of a spinal column of a patient, comprising:

engaging first and second implants to a first vertebra;

15 engaging a distal portion of respective first and second implant holders to respective ones of the first and second implants, the first and second implant holders each extending from the distal portion along a longitudinal axis to a proximal end outside the patient;

20 engaging a transverse bridge between each of the first and second implant holders at a location between distal and proximal ends of each of the first and second implant holders with each of the implant holders including a derotation handle extending proximally from the location.

25 33. The method of claim 32, wherein the location on the first and second implant holders is a joint and the transverse bridge includes clamping assemblies to clampingly engage respective ones of the joints.

34. The method of claim 33, wherein the joints include a spherical outer surface and the clamping assemblies includes a split ring configuration clampingly engaged about the outer surface.

30 35. The method of claim 33, wherein the clamping assemblies each include:
a clamping member positioned about the joint of the respective implant holder with a pair of arms extending from the clamping member;

a mounting member with a receptacle at a first end thereof receiving a connecting member of the transverse bridge and a post extending from the first end through the arms of the clamping member; and

a locking member engaged to the post to clamp the arms between the first end and the locking member and secure the clamping member about the joint.

36. The method of claim 35, wherein the clamping assemblies each further include a seating washer between the connecting member and one of the pair of arms, the connecting member being secured between the first end of the mounting member and the seating washer when the clamping member is secured about the joint with the locking member.

37. The method of claim 32, further comprising manipulating the derotation handles to align the spinal column.

38. The method of claim 37, further comprising engaging elongate stabilization elements to each of the first and second anchors after aligning the spinal column to provide post-operative stabilization after manipulating the derotation handles to align the spinal column.

39. The method of claim 32, further comprising:
engaging third and fourth implants to a second vertebra;
engaging a distal portion of respective third and fourth implant holders to respective ones of the third and fourth implants, the third and fourth implant holders each extending from the distal portion thereof along a longitudinal axis to a location outside the patient, wherein each of the third and fourth implant holders includes a handle extending proximally from the location; and

engaging a second transverse bridge to the locations on each of the third and fourth implant holders.

40. The method of claim 39, further comprising engaging an inter-level linking assembly between the transverse bridges.

41. The method of claim 40, wherein each of the transverse bridges includes a clamping assembly and engaging the inter-level linking assembly includes clamping each of the clamping assemblies to an elongated link member of the linking assembly.

42. The method of claim 41, wherein the clamping assemblies are located at an end of the respective transverse bridges and further comprising:

securing at least one connector assembly to an end of at least one of the clamp assemblies; and

clamping the connector assembly to a second elongated link member.

43. The method of claim 32, wherein engaging the transverse bridge includes:
5 positioning an elongate connecting member of the transverse bridge through joints of the implant holders at the locations; and
locking the connecting member in the joints.

44. The method of claim 32, wherein:
the first and second implants each include a receiver defining a proximally opening
10 passage for receiving a respective one of first and second elongate stabilization elements positionable along the spinal column; and

engaging the distal portion of respective first and second implant holders includes clamping the distal portion along one side of the receiver so that the proximally opening passage remains substantially unobstructed for receiving the respective elongate
15 stabilization element.

45. The method of claim 44, further comprising positioning at least one of the stabilization elements in the proximally opening passage of the receiver of at least one of the first and second implants with the respective implant holder engaged to the receiver.

46. The method of claim 44, wherein the first and second implant holders each
20 include a first arm and a second arm pivotally connected to the first arm, the first and second arms being movable in a first plane including a longitudinal axis of the respective implant holder, wherein the distal portions are offset to one side of the longitudinal axis and are movable in a second plane that is generally parallel to the first plane.

47. A method for assembling a system to correct alignment of a spinal column
25 of a patient, comprising:

engaging first and second implants to respective ones of first and second vertebrae;
releasably engaging a first implant holder of a first derotation instrument assembly to the first implant;

engaging a first transverse bridge of the first derotation instrument assembly at a
30 location between a proximal handle of the first implant holder and a distal clamp portion of the first implant holder with the first transverse bridge extending from the first implant

holder to another implant holder associated with the first vertebra and with the handle extending proximally from the first transverse bridge;

releasably engaging a second implant holder of a second derotation instrument assembly to the second implant;

5 engaging a second transverse bridge of the second derotation instrument assembly at a location between a proximal handle of the second implant holder and a distal clamp portion of the second implant holder, the second transverse bridge extending from the second implant holder to another implant holder associated with the second vertebra and with the handle extending proximally from the second transverse bridge; and

10 coupling an inter-level linking assembly to the first and second transverse bridges.

48. The method of claim 47, wherein:

the first and second implants each include a receiver configured for engagement with a stabilization element positionable along the spinal column between the first and second implants; and

15 releasably engaging the respective first and second implant holders includes clamping the distal portion along one side of the receiver so that the passage remains substantially unobstructed for receiving the stabilization element.

49. The method of claim 47, further comprising:

20 pivoting a first ball member at the location of the first implant holder while positioning an elongate connecting member of the first transverse bridge through the ball member;

locking the first ball member to the elongate connecting member of the first transverse bridge;

25 pivoting a second ball member at the location of the second implant holder while positioning an elongate connecting member of the second transverse bridge through the second ball member; and

locking the second ball member to the elongate connecting member of the second transverse bridge.

30 50. The method of claim 47, wherein coupling the inter-level linking assembly includes positioning a link member between the first and second transverse bridges and clamping the first and second transverse bridges to the link member.

51. A method for correcting alignment of a spinal column, comprising:
engaging implants to each of the pedicles of at least one vertebra;
engaging a distal portion of an implant holder to each of the implants, the implant
holders each extending proximally from the respective distal portion to a location outside
the patient, each of the implant holders including a distal clamping portion and a joint at
the location; and

connecting a transverse bridge to the implant holders, the transverse bridge
including a handle portion and a connecting portion extending from the handle portion, the
connecting portion include an end positionable about the joint of one of the implant
holders and a slot between the handle portion and the end that is positionable about the
joint of the other implant holder.

52. The method of claim 51, wherein the joints each include a spherical shape
and the end includes a fork-shape and the slot includes an elongated shape with a series of
partially spherical receiving areas to provide a number of locations to receive the joint of
the other implant holder.

53. The method of claim 52, wherein the one implant holder includes a handle
extending proximally from the joint thereof.

54. The method of claim 53, further comprising applying corrective forces to
the at least one vertebra by manipulating the handle of the one implant holder and the
handle portion of the transverse bridge.

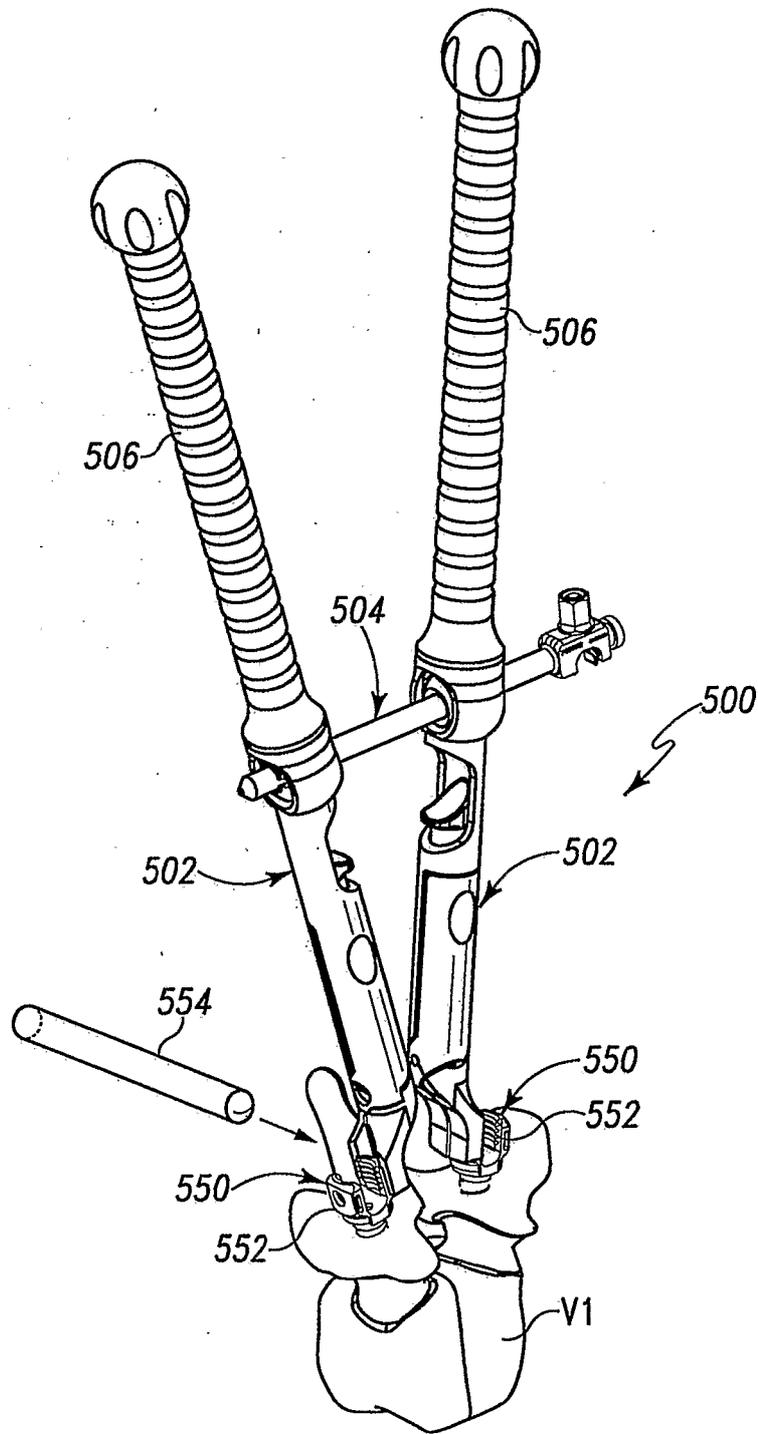


Fig. 1

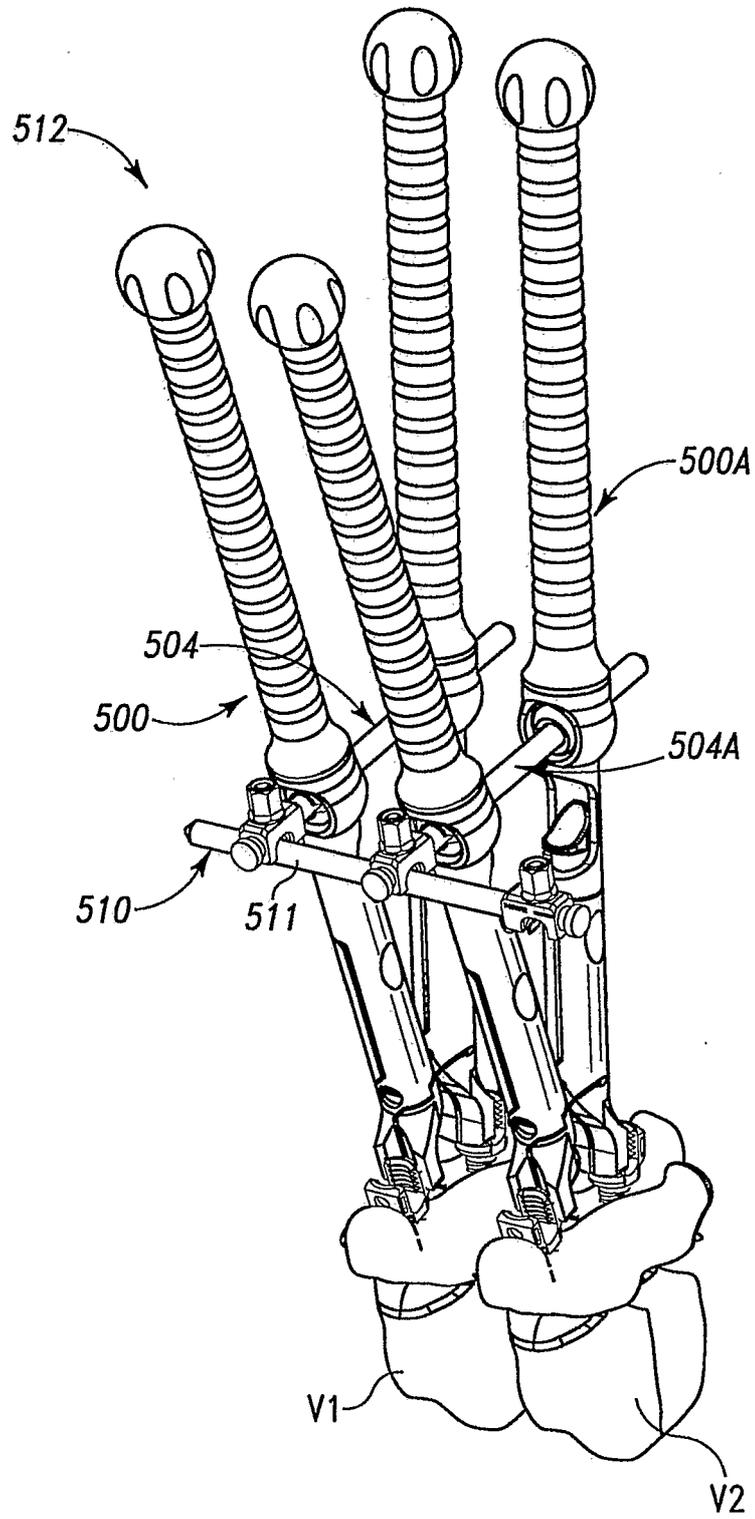


Fig. 2

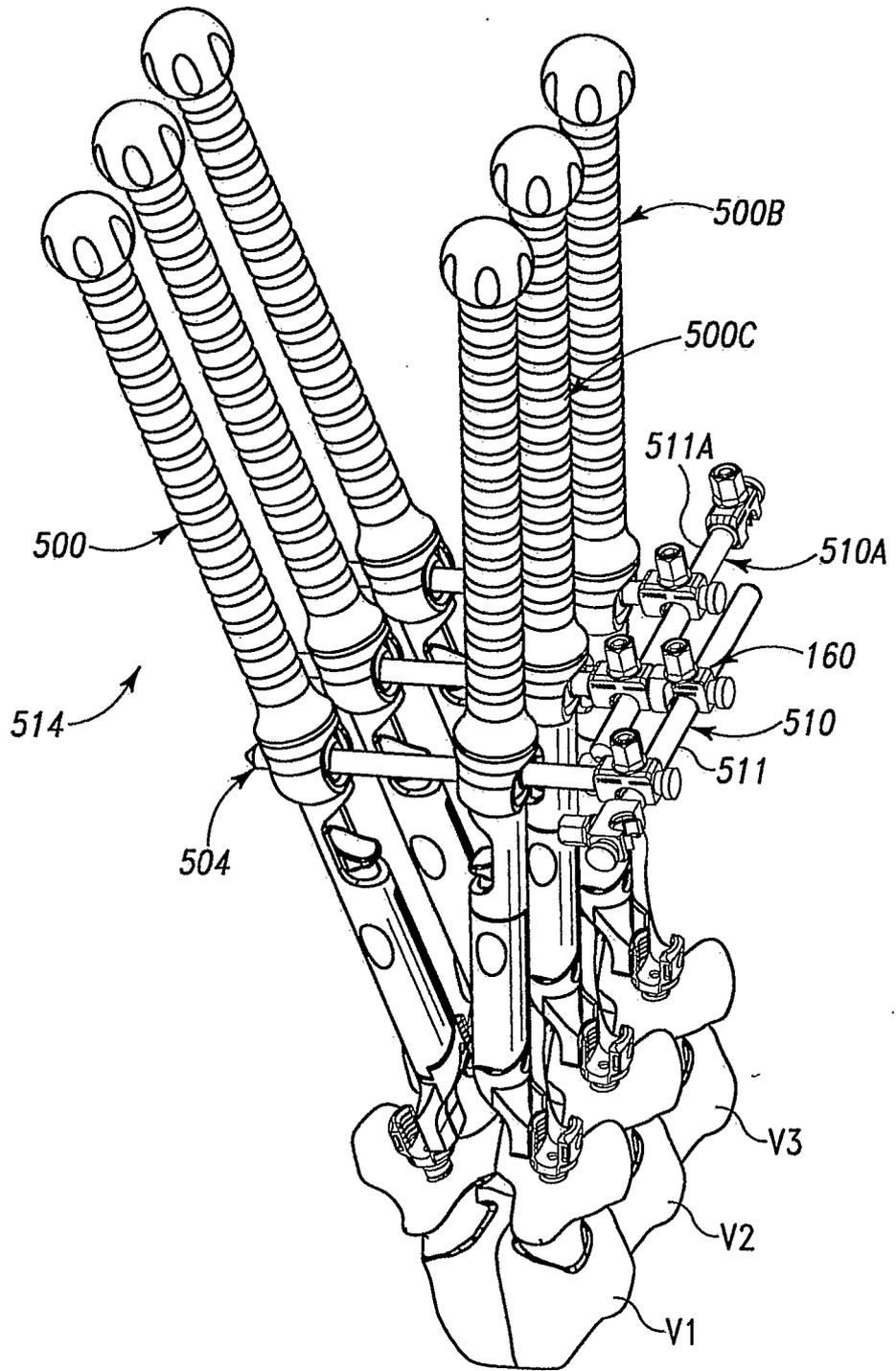


Fig. 3

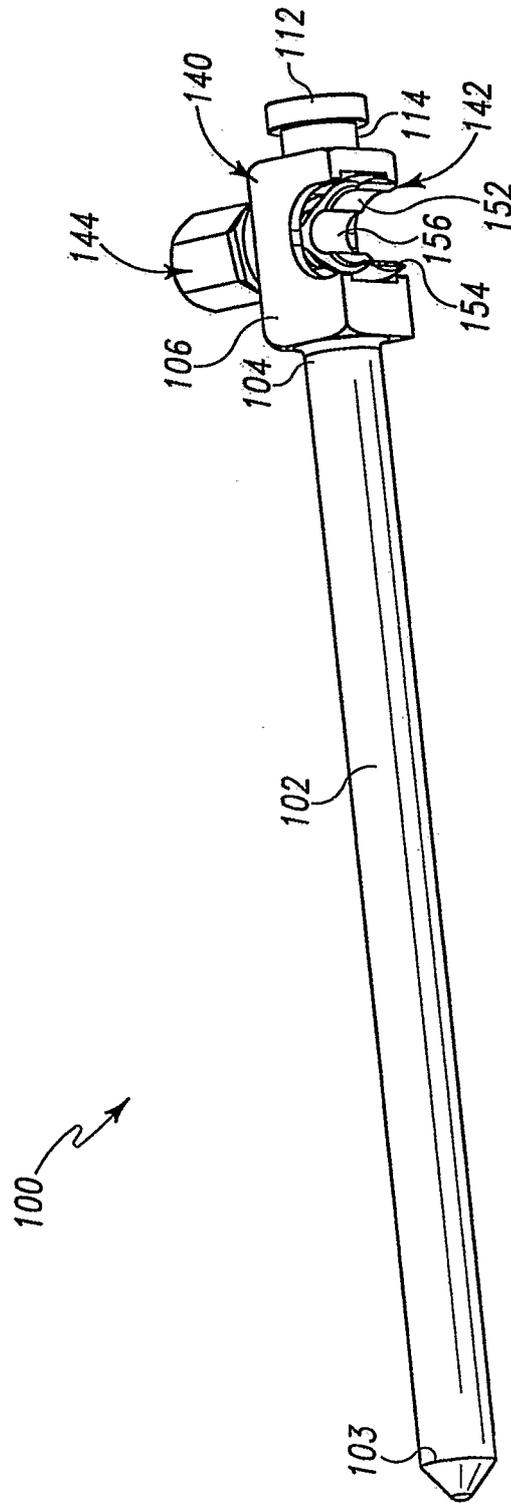


Fig. 4

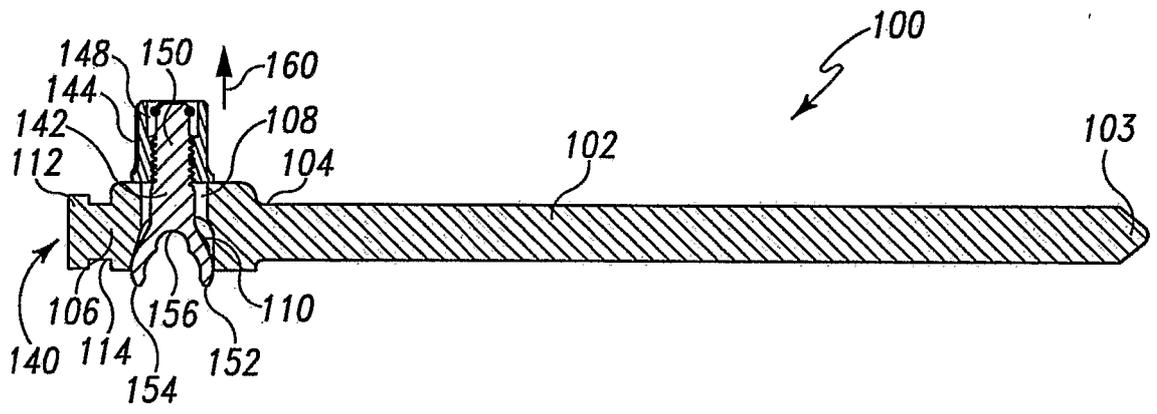


Fig. 5

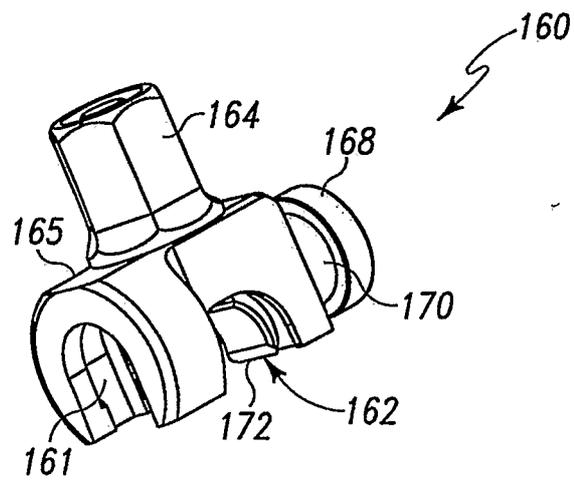


Fig. 6

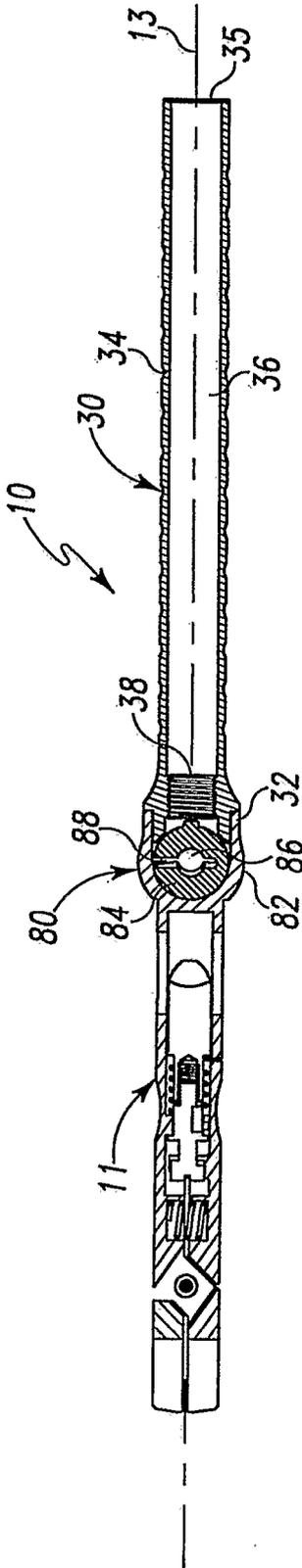


Fig. 7

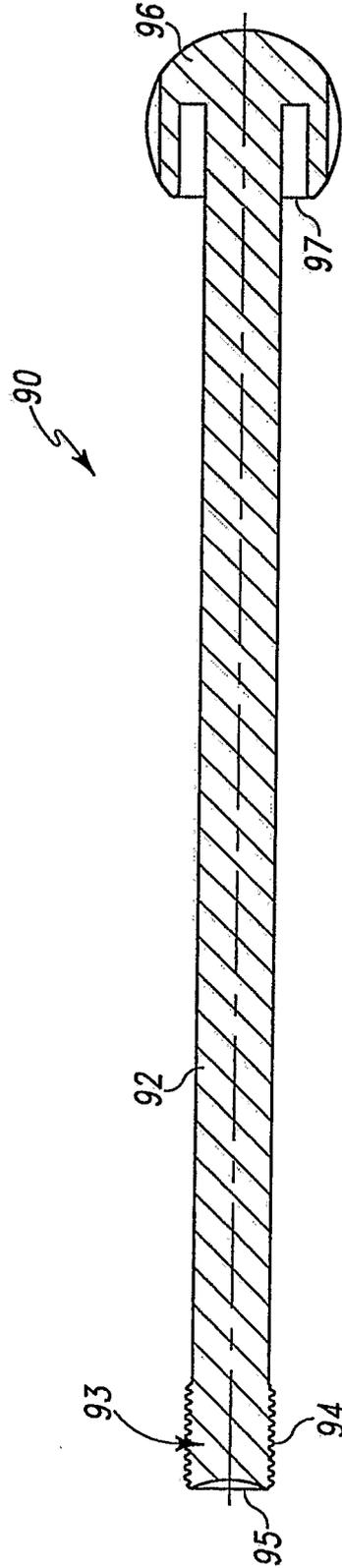


Fig. 8

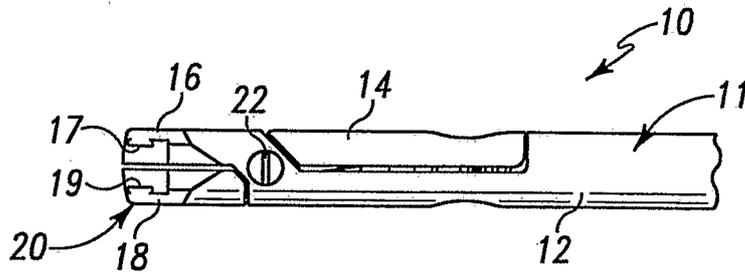


Fig. 9

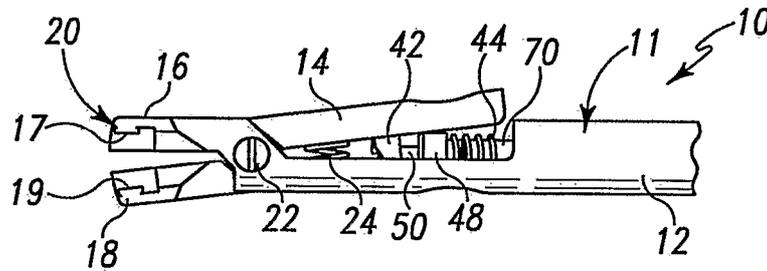


Fig. 10

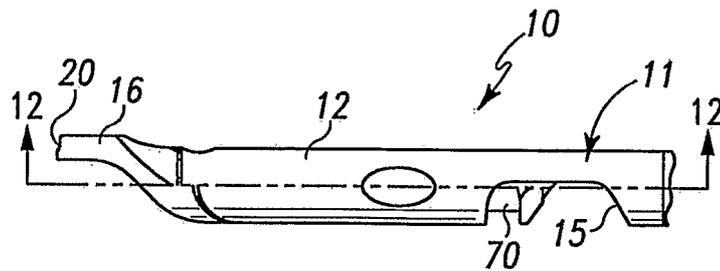


Fig. 11

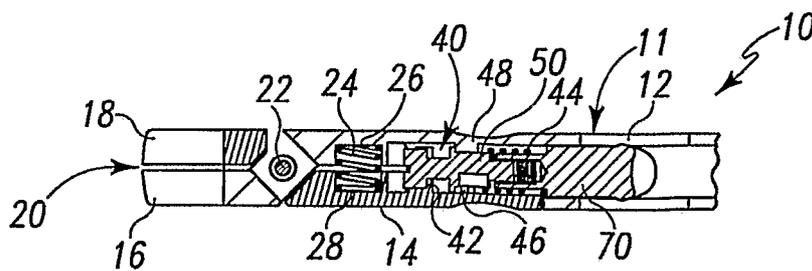


Fig. 12

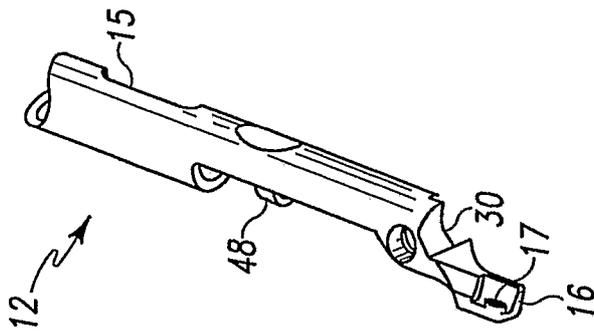


Fig. 12

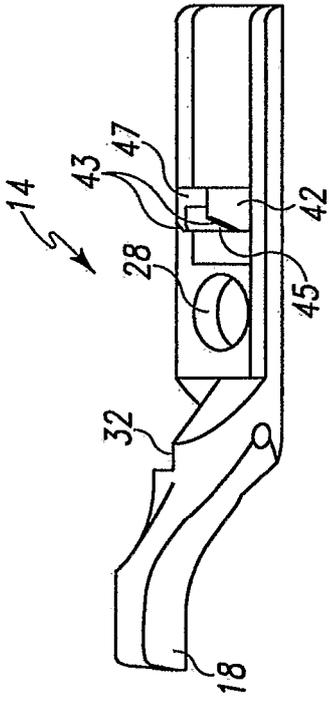


Fig. 14

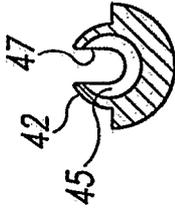


Fig. 15

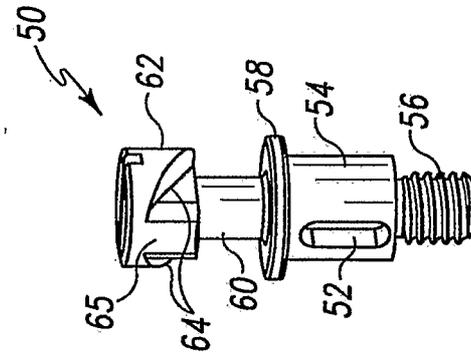


Fig. 16

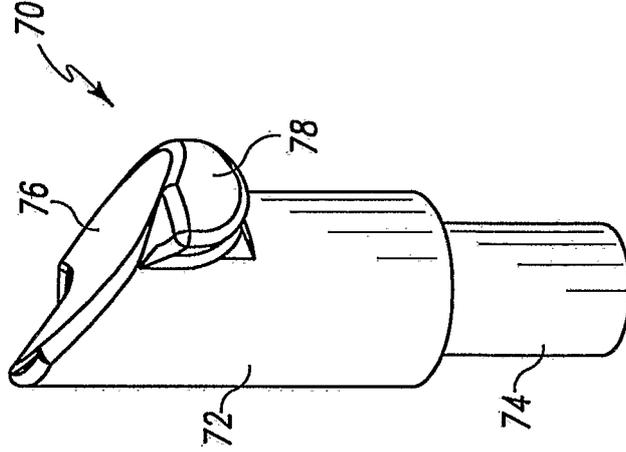


Fig. 17

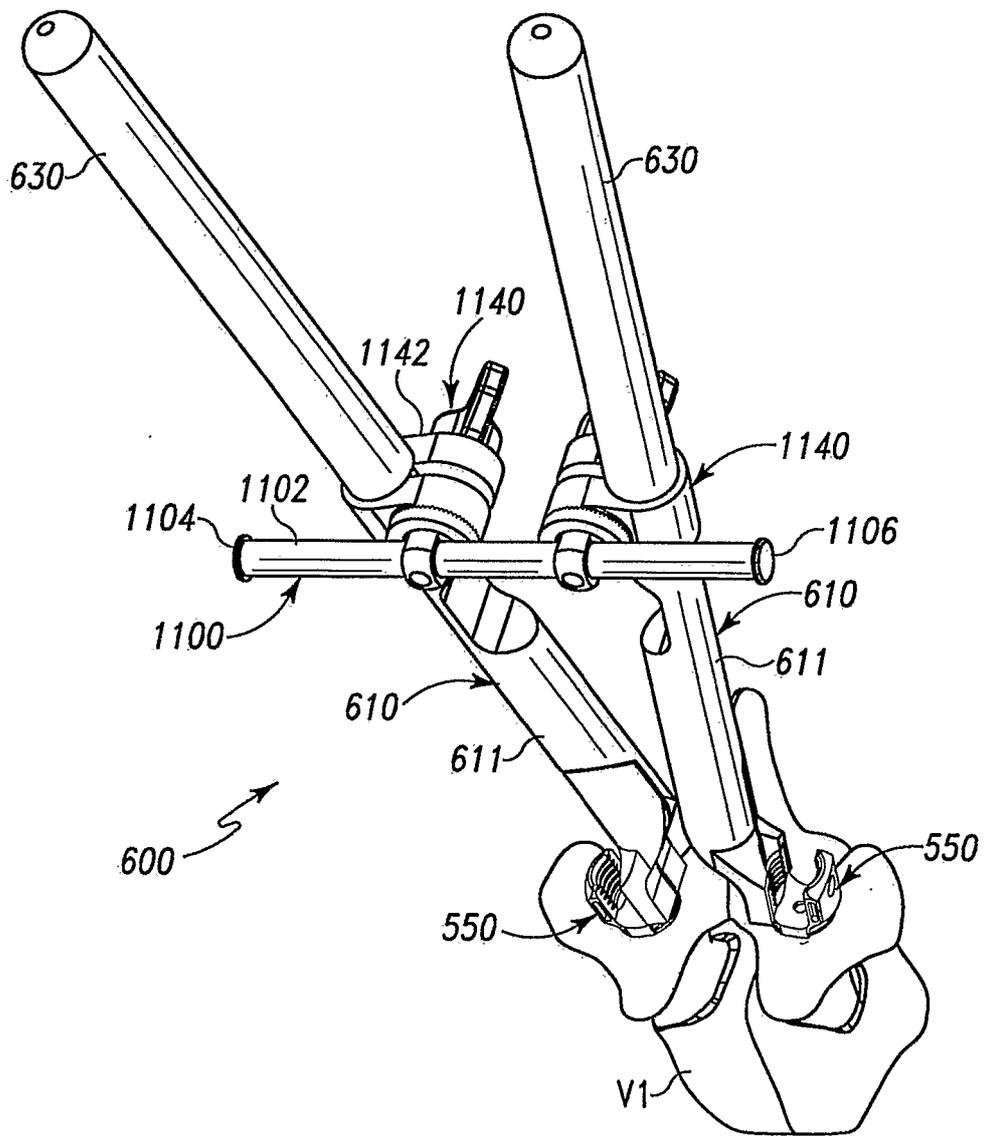


Fig. 18

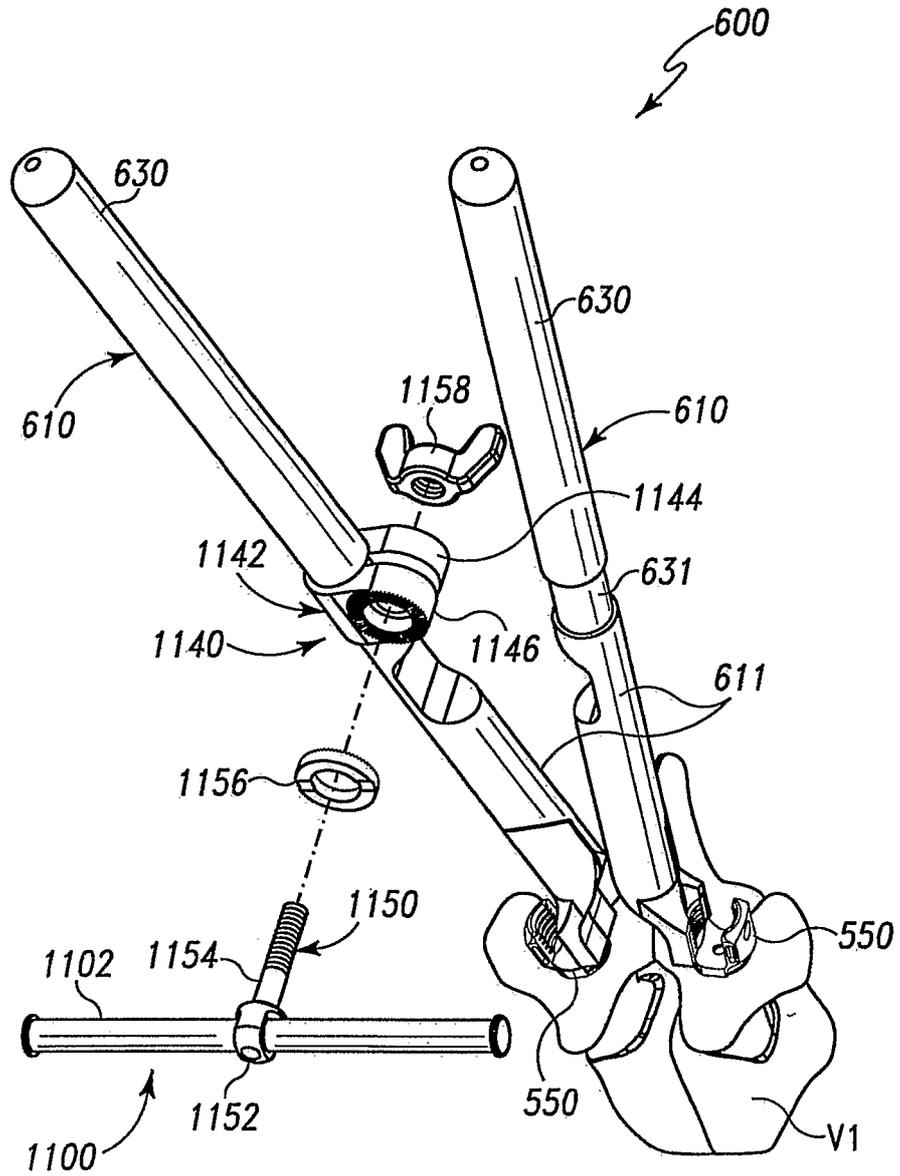


Fig. 19

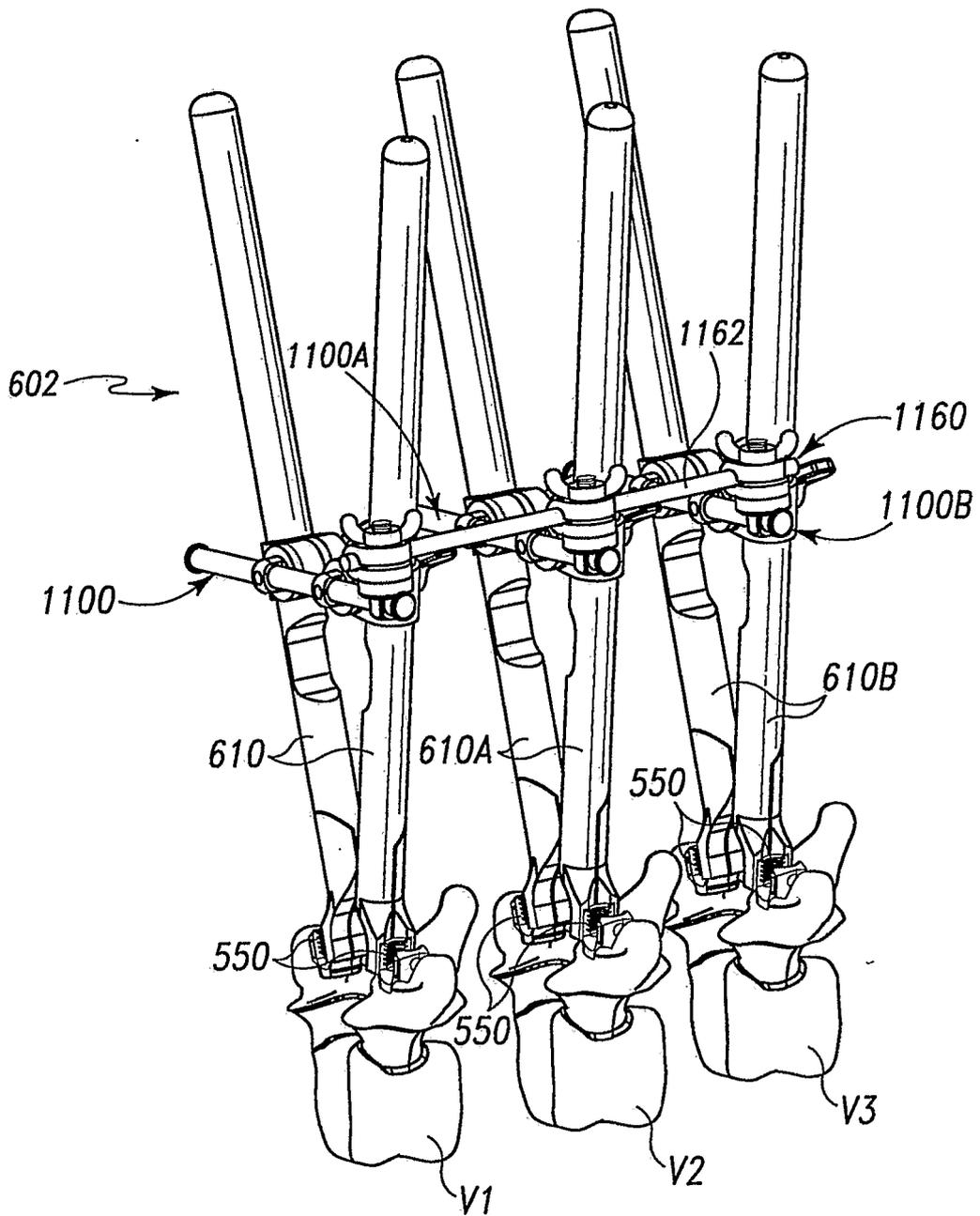


Fig. 20

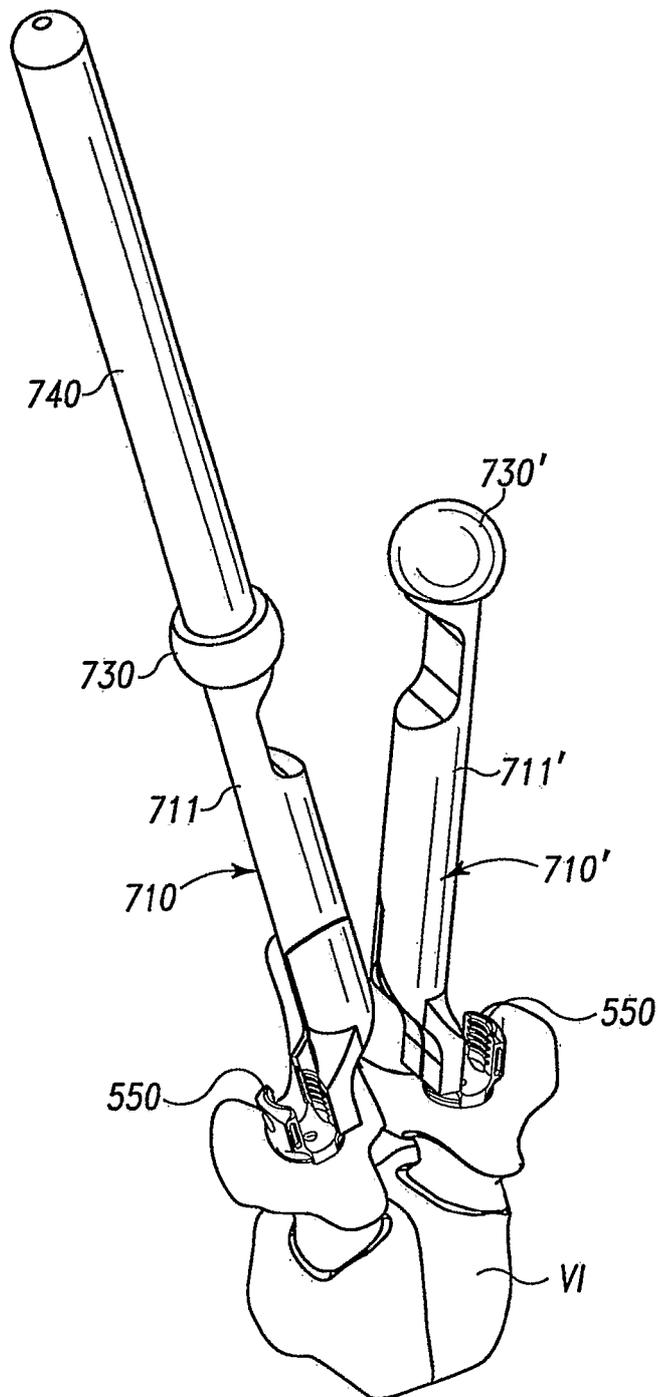


Fig. 21

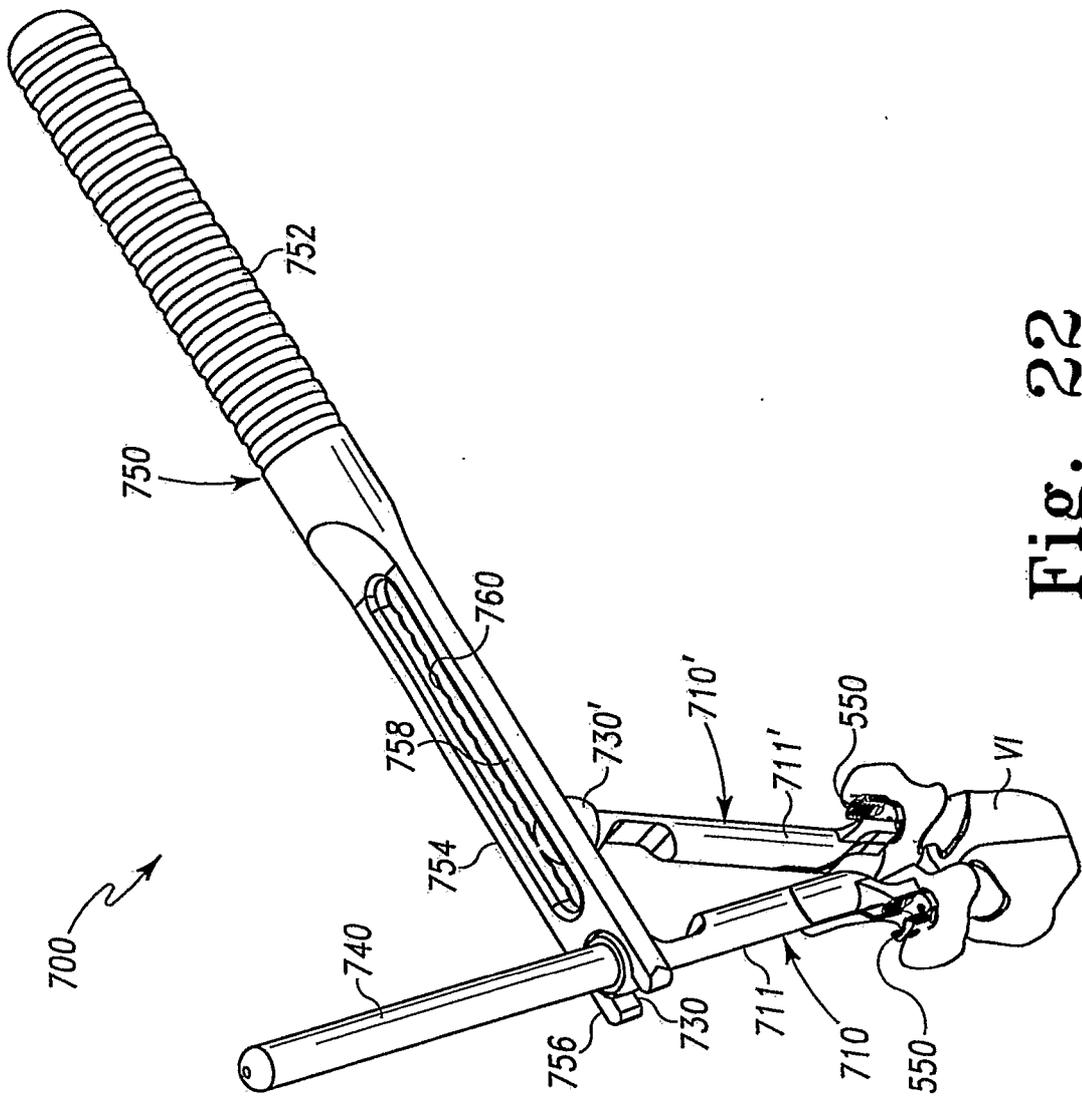


Fig. 22

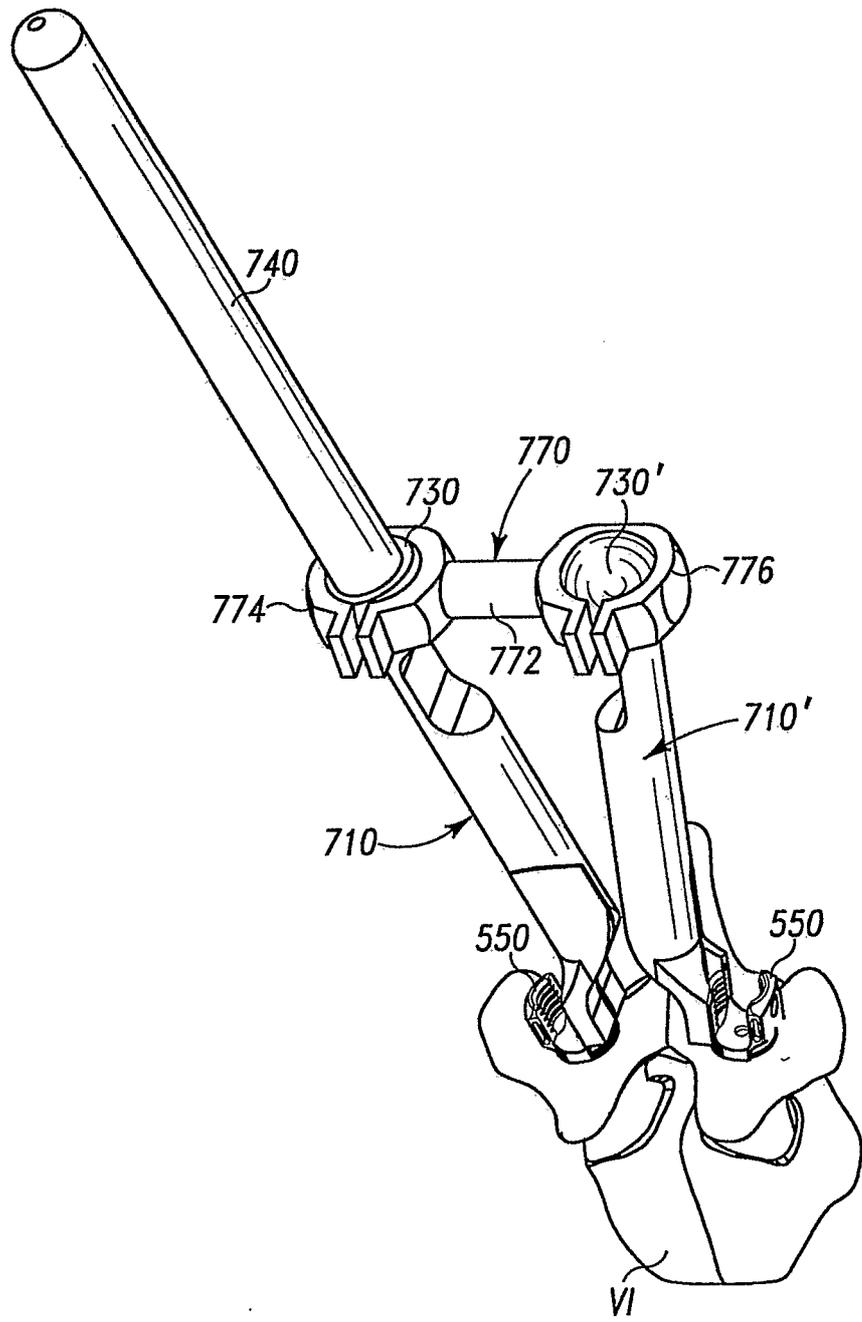


Fig. 23