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(54) Title: INSERTION DEVICES AND METHOD OF USE

(57) Abstract: This invention relates to devices for inserting expandable structures, such as medical balloons, into interior regions of a human or animal body, as well as methods for their use. The insertion devices described herein are capable of directionally guiding and/or inhibiting expansion of an expandable structure within an interior region of an animal or human body to create optimally placed cavities for repair, augmentation and/or treatment of fractured and/or diseased bone.



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INSERTION DEVICES AND METHOD OF USE**Related Applications**

This application is a continuation-in-part of co-pending U.S. Application Serial No. 09/828,470, filed April 6, 2001, entitled "Insertion Devices and Method of Use," which claims the benefit of provisional U.S. Application Serial No. 60/195,207, filed April 7, 2000.

Field of the Invention

This invention relates to an insertion device which is capable of guiding expansion of an expandable structure towards or away from a desired direction and/or interior region of an animal or human body. This invention further relates to an insertion device for inserting expandable structures, such as medical balloons, into an interior region of a human or animal body, wherein the device is capable of expanding at the tip. The present invention also relates to methods of using the disclosed devices in the repair, augmentation and/or treatment of fractured and/or diseased bone.

Background of the Invention

Expandable structures, such as balloon dissectors and catheters, are used in various surgical procedures and for various rehabilitative purposes in the medical arts. In angioplasty, balloon catheters are commonly inserted into veins and arteries to expand blood

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vessels, most commonly to dilate and/or remove obstructions in the blood vessel (e.g. to remove constrictions blocking blood vessels which can cause a heart attack or stroke). Other types of surgical balloons have been used to aid surgeons in accessing specific organs during surgery, usually in lieu of previous insufflation techniques. Such balloons are commonly inserted in a deflated state through an insertion device comprising a cannula, catheter tube, or other similar device, and are positioned under an organ. The balloon may then be inflated to lift and separate a desired organ away from surrounding organs and tissue to make sides of the desired organ easier to access during surgery. The balloon may also be placed and inflated so as to lift and separate other organs and tissues, leaving the desired organ for surgery exposed beneath.

Medical balloons have also been used during procedures for repairing and/or reinforcing fractured and/or diseased bones. Some physicians have used such balloons to create a working space adjacent fractured and/or diseased bone to allow the installation of plates, screws and/or other implantable articles to the bone. In this type of procedure, a cannula is generally inserted through an incision in the skin near the fracture area. A balloon is then inserted through the cannula and inflated between the bone and surrounding tissue around the fracture site to create a working space. A support plate and bone screws, or other similar implements, can then be installed at the fracture site through small incisions in the skin. This type of procedure allows a surgeon to install implantable articles without having to make a long skin incision to isolate and expose the bone.

More recently, balloons have been employed inside fractured and/or diseased bones to repair, reinforce and/or treat the bone. In these procedures,

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balloons can be inserted through a cannula and inflated inside the bone, which can compact cancellous bone, create a cavity and move cortical bone in an effort to restore the natural anatomy. The cavity can be filled
5 with a suitable bone filler, such as bone cement (e.g., polymethylmethacrylate - PMMA), autograft or allograft tissue, or various other bone biocompatible substitutes.

When the bone filler hardens, it essentially creates an internal "cast" which allows the bone to heal properly,
10 but also desirably allows the bone to bear weight immediately.

Summary of the Invention

The present invention provides insertion devices which may be used with an expandable structure
15 such as a balloon catheter to direct the expansion of the structure toward or away from a desired direction. Directed expansion of the expandable structure provides the physician with significant control over the compression of cancellous bone and creation of cavities
20 within the bone, as well as control over the movement of cortical bone. In addition, the controlled expansion of an expandable structure allows the physician to tailor the shape and dimensions of the cavity, and the resulting shape and dimensions of the bolus of filler material
25 contained therein. Moreover, directed expansion of an expandable structure permits a physician to minimize disruption of healthy cancellous and/or cortical bone during a treatment procedure, thereby further enhancing healing of the bone after treatment. Accordingly, the
30 devices and methods disclosed herein permit the physician to optimize the ability of the bone to withstand compressive forces and/or heal as quickly as possible after the procedure is completed.

Expandable structures such as balloon
35 catheters and dissectors are typically formed in

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spherical or elliptical shapes, and normally expand substantially outward. These balloons are desirably fairly low-profile such that they can fit through a cannula. Once inserted through the cannula into the
5 region of treatment, such balloons will generally inflate fairly symmetrically about the axis of the cannula or other insertion device. However, as discussed in U.S. Pat. No. 5,972,015, which is incorporated by reference herein, inflation of a balloon about the cannula's axis
10 can be undesirable in some situations. To account for these situations, various alternative balloon designs and expansion constraint arrangements have been proposed, such as those described in the '015 patent.

The inventions disclosed herein further permit
15 a practitioner to utilize a wide variety of expandable structures in conjunction with the methods and devices disclosed herein. Because the cannula or other insertion device substantially guides the direction of expansion of the expandable structure, there is less need to
20 incorporate expansion constraints in the expandable structure itself. In addition, if the insertion device is comprised of a radiopaque material, the orientation of the device itself can be visualized during the surgical procedure under x-ray fluoroscopy, allowing the
25 practitioner to visually verify the direction of expansion of the structure throughout the entire procedure. Of course, it should be understood that the devices and methods of the present invention could also be used in conjunction with expandable structures
30 incorporating various expansion restraint arrangements.

In a general embodiment of the present invention, an insertion device comprises a hollow member, which is preferably cylindrical, with a distal end and a proximal end, wherein the distal end is the tip, or point
35 of insertion, of the insertion device. The distal end of

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the hollow member desirably comprises a platform which
constrains expansion of the expandable structure in one
or more directions, but permits the expandable structure
to expand in non-constrained directions. In effect, the
5 platform of the hollow member acts as a support or
foundation against which the expandable structure pushes
as it expands. Desirably, the supporting action of the
platform induces the expandable structure to expand away
from the platform, allowing the practitioner to direct
10 expansion towards and/or away from a desired region.

In another general embodiment, the platform
comprises a platform or expansion guide which is inserted
through a hollow member of an insertion device, the guide
desirably extending distally past the tip of the hollow
15 member and into a bone. The guide will desirably act as
a support or foundation against which the expandable
structure expands, inducing the structure to expand away
from the guide. Because this embodiment of the guide
travels through the hollow member, and need not initially
20 penetrate soft tissues and/or the harder cortical bone,
the guide design can be optimized to provide maximum
support for the expandable structure.

In another general embodiment of the present
invention, an insertion device comprises a hollow member
25 with a distal end and a proximal end, wherein the distal
end is the tip, or point of insertion, of the insertion
device. The distal end of the hollow member desirably
comprises a platform which directs the expansion of the
expandable structure in one or more directions. The
30 distal end of the hollow member further comprises one or
more crease or fold lines along which at least a portion
of the platform desirably deforms after insertion into a
bone. By deforming along predetermined lines, sharp
surfaces on the platform are desirably moved away from
35 the expandable structure. In addition, bending of the

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platform can significantly affect the surface area of the platform in contact with the expandable structure as well as the strength and resistance to deformation of the platform. After the expandable structure is contracted,
5 the platform can be withdrawn through the cannula, with the distal end of the insertion device desirably bending the platform towards a lower-profile shape for removal.

In another general embodiment of the present invention, the insertion device comprises a hollow member
10 having a plurality of score lines spaced about the circumference of the distal tip, these score lines desirably form a plurality of adjacent sections oriented in a first, lower profile orientation. After insertion to a desired location within the vertebral body, the
15 adjacent sections can be expanded outward to a second orientation, where the adjacent sections substantially form a funnel, cone or flare at the tip of the insertion device. When removal of the expandable structure is desired, the flared tip desirably guides the expandable
20 structure into the cannula, facilitating passage of the expandable structure into and through the cannula. If desired, the adjacent sections can further incorporate one or more guides or ribs which desirably impinge upon the expandable structure, folding and/or twisting the
25 expandable structure along desired lines and/or in a desired manner, further facilitating removal of the expandable structure through the cannula. When removal of the insertion device from the vertebral body is desired, the withdrawal of the insertion device through
30 the harder cortical bone desirably bends the adjacent sections back towards and/or into their first, lower profile orientation.

The present invention is further related to methods for using the disclosed devices for repair,
35 augmentation and/or treatment of fractured and/or

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diseased bones. One embodiment of an insertion device constructed in accordance with the teachings of the present invention is inserted through cortical bone and into cancellous bone in a vertebral body of a patient.

5 The insertion device is positioned such that the platform directs the expansion of an expandable structure towards a section of cortical bone to be moved to a desired position, such as a depressed upper or lower plate of a vertebral body, such as, for example, an anterior wedge,

10 bi-concave or vertebra plana fracture morphology. The expandable structure is expanded against the platform, which desirably induces the expandable structure to expand substantially away from the platform, compressing cancellous bone to form a cavity and moving the targeted

15 section of cortical body towards a desired position. The expandable structure is contracted, removed, and the cavity is then filled with an appropriate bone filler material. This method, which permits precise manipulation of cortical bone with a minimum of

20 cancellous bone compression, allows a practitioner to move targeted cortical bone while preserving much of the cancellous bone in an uncompressed state. In addition, this method permits the practitioner to maximize the force which the expandable structure exerts on the

25 cortical bone in a targeted or directed manner.

In another embodiment of the present invention, an insertion device constructed in accordance with the teachings of the present invention is inserted into cancellous bone in a vertebral body of a patient.

30 The insertion device is positioned such that the platform directs the expansion of an expandable structure towards a section of cancellous bone to be compressed. The expandable structure is expanded, which desirably compresses some or all of the targeted cancellous bone,

35 creating a cavity within the cancellous bone.

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The expandable structure is then contracted and, if desired, the insertion device is repositioned such that the platform directs the expansion of the expandable structure towards another section of cancellous bone to be compressed. The insertion device
5 may be repositioned by rotation or turning, translationally by pushing or pulling, by angulation by reinserting along either the same or a different access path, or by a combination of methods to create a cavity
10 of a desired size and configuration. The expandable structure is then expanded again, compressing some or all of the targeted cancellous bone and increasing the size and/or altering the shape of the cavity within the bone. If desired, the platform may be repositioned and the
15 procedure can be repeated as necessary to create a cavity of desired dimensions.

The expandable body is removed and the cavity is then filled with an appropriate bone filler material. This method, which facilitates the creation of large
20 cavities within the bone, allows the practitioner to tailor the cavity shape/size to optimize the post-treatment strength and/or healing of the bone. Similarly, the disclosed method can be used to reposition cortical bone towards a desired position, permitting a
25 practitioner to gradually displace small or large sections of the cortical bone, at the practitioner's option.

In another embodiment, the disclosed devices and methods facilitate a practitioner's ability to
30 repair, reinforce and/or treat targeted bone regions in situations where the insertion device is initially positioned near a cortical bone wall of a targeted bone region. Because the disclosed devices and methods provide substantial control over the direction of
35 expansion of the expandable structure, the practitioner

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can position and/or reposition the platform to shield the nearby cortical bone from some or all of the expandable structure during some or all of the surgical procedure. Depending upon the orientation of the platform, the structure can be expanded to differing dimensions, desirably maximizing compression of cancellous bone and/or movement of cortical bone at each orientation. Accordingly, there is no need to reorient the entire insertion device to accomplish the objectives of the procedure, which desirably eliminates a source of additional trauma occurring during the procedure.

In another embodiment, an insertion device is inserted through cortical bone and into cancellous bone in a vertebral body of a patient. A stylet in the insertion device is removed, causing the distal end of the hollow member of the insertion device to expand or flare. An expandable structure is inserted through the insertion device into the vertebra, is expanded to create a cavity, and is contracted and removed through the insertion device. As the expandable structure is withdrawn through the insertion device, the flared distal end of the insertion device desirably guides the structure into the insertion device. The cavity is then filled with an appropriate bone filler.

Another aspect of the invention provides a bone treatment device comprising an expandable structure and a sheath housing the expandable structure. The sheath includes an opening extending along the axis of the sheath and about the axis of the sheath to direct expansion of the expandable structure. The sheath also includes at least one securing element that prevents movement of the expandable structure within the sheath. In one embodiment, the securing element is a tab. In one embodiment, the assembly further comprises a stylet having a distal end and sized and configured to extend

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through the expandable structure. The distal end of the stylet is adapted to extend at least in part beyond the distal end of the sheath. In one embodiment, the distal end of the stylet is blunt.

5 According to another aspect of the invention, a cortical bone wall probe comprises a sheath extending along an axis and having a distal end. A shaft having a distal end is sized and configured for passage through the sheath. A stylet having a blunt distal end is sized
10 and configured for passage through the shaft. The blunt distal end extends at least in part beyond the distal end of the sheath and the distal end of the shaft.

 In one embodiment, the cortical bone wall probe further comprises an expandable structure carried
15 by the shaft. In this embodiment, the sheath includes an opening extending along the axis of the sheath and about the axis of the sheath to direct expansion of the expandable structure.

 In one embodiment, the sheath includes at
20 least one securing element that prevents movement of the shaft within the sheath.

 According to yet another aspect of the invention, a bone treatment device comprises an expandable structure carried by a shaft having a distal
25 end. The shaft includes a lumen extending therethrough. A stylet having a blunt distal tip is sized and configured for passage through the lumen to extend the blunt distal tip, at least in part, beyond the distal end of the shaft to form a balloon catheter assembly. A
30 sheath having a distal end is sized and configured for passage of the catheter assembly. The sheath includes an opening for receiving the expandable structure and at least one securing element for securing the sheath to the distal end of the shaft. In one embodiment, the distal
35 tip of the stylet extends at least in part beyond the

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distal end of the sheath.

Another aspect of the invention provides a method of treating fractured or diseased bone. The method comprises introducing an insertion device through
5 a cortical bone region and into a cancellous bone region of the bone. An expandable structure is introduced through the insertion device in a collapsed or non-expanded condition. The expandable structure is expanded from the collapsed condition to an expanded or enlarged
10 condition extending beyond the radius of the insertion device within the bone to create a void or cavity in the cancellous bone. The expandable structure may take a variety of forms, including, but not limited to, a mechanical jack or retractor device, an expanding spring
15 or coil, or shape-memory device.

In a preferred embodiment, the expandable structure is a balloon, which may be carried by a balloon catheter assembly. The balloon catheter assembly is introduced through the insertion device. The balloon
20 catheter assembly comprises the expandable structure (i.e., an inflatable balloon) and a sheath housing the expandable structure. The sheath includes an opening extending along the axis of the sheath and about the axis of the sheath to direct expansion of the expandable
25 structure. The sheath desirably includes at least one securing element that prevents movement of the expandable structure within the sheath. The expandable structure is expanded outside the opening and beyond the sheath to create a cavity within the bone. The expandable
30 structure (e.g., balloon catheter assembly) is removed, and the cavity can then be filled with a bone filler.

Other objects, advantages and embodiments of the invention are set forth in part in the description which follows, and in part, will be obvious from this
35 description, or may be learned from the practice of the

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invention.

Brief Description of the Drawings

Fig. 1 is a coronal view of a vertebral body, showing a cannula inserted in a vertebral body, with a
5 spherical expandable structure expanded within the vertebral body;

Fig. 2 is a coronal view of a vertebral body, showing one embodiment of an insertion device constructed in accordance with the teachings of the present invention
10 which has been inserted into a vertebral body, with an expandable structure expanded within the vertebral body;

Fig. 3a is a side view of an alternate embodiment of an insertion device constructed in accordance with the teachings of the present invention,
15 wherein the distal end of the hollow member comprises one or more longitudinal score lines at its circumference.

Fig. 3b is a cross-sectional side view of the insertion device of Fig. 3a, showing the adjacent sections in a lower profile orientation;

20 Fig. 4 is a side view of the insertion device of Fig. 3a, showing the adjacent sections in a deployed or flared position;

Fig. 5 is a cross-sectional side view of the insertion device of Fig. 3a, with a stylet positioned
25 within the hollow member of the device;

Fig. 6 is an end of the insertion device of Fig. 4;

Fig. 7 is a side view of another alternate embodiment of an insertion device constructed in
30 accordance with the teachings of the present invention, the device comprising an extension or platform extending from the distal end of the device;

Fig. 8 is a top plan view of another alternate embodiment of an insertion device constructed in
35 accordance with the teachings of the present invention,

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showing a crease line extending along a platform of the device;

Fig. 9a is a side view of another alternate embodiment of an insertion device constructed in accordance with the teachings of the present invention with a stylet positioned within the lumen of the device;

Fig. 9b is a cross sectional side view of the insertion device of Fig. 9a, showing the adjacent sections in a lower profile orientation;

Fig. 9c is a side view of the insertion device of Fig. 9a, showing the adjacent sections moved to a deployed position as the stylet is withdrawn from the device;

Fig. 10 is an end view of another alternate embodiment of an insertion device constructed in accordance with an alternate embodiment of the present invention;

Fig. 11 is a side view of the insertion device of Fig. 10.

Fig. 12 is a side view of the insertion device of Fig. 10, with a stylet positioned within the hollow member of the device;

Fig. 13 is a side view of an alternate embodiment of an insertion device constructed in accordance with the teachings of the present invention, with a stylet positioned within the hollow member of the device;

Fig. 14 is a side view of one embodiment of a stylet constructed in accordance with the teachings of the present invention, for use with various embodiments of the insertion device of the present invention;

Fig. 15 is a side view of an alternate embodiment of a stylet constructed in accordance with the teachings of the present invention, for use with alternate embodiments of the insertion device of the

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present invention;

Fig. 16a is a cross sectional side view of an alternate embodiment of an insertion device and stylet constructed in accordance with the teachings of the present invention, showing one method of assembling the device and stylet;

Fig. 16b is a cross section side view of the insertion device and stylet of Fig. 16a, showing the stylet inserted fully into the hollow member of the device during assembly;

Fig. 17 is a cross sectional side view of the insertion device and stylet of Fig. 16b, with the adjacent sections of the device positioned in a lower profile orientation around the stylet;

Fig. 18 depicts a patient about to undergo a surgical procedure in accordance with the teachings of the present invention;

Fig. 19 depicts an incision point and underlying vertebrae for the patient of Fig. 18;

Fig. 20 is a corona view of a vertebra showing an insertion device approaching the posterior side of the vertebral body;

Fig. 21a depicts an insertion device penetrating the vertebral body of Fig. 20;

Fig. 21b is a coronal view of the vertebra of Fig. 20, with an insertion device positioned within the cancellous bone;

Fig. 22 is a coronal view the vertebra body of Fig. 21a, with the adjacent sections positioned in a deployed orientation.

Fig. 23 is a coronal view of the vertebra body of Fig. 22, wherein an expandable structure is expanded within the vertebra;

Fig. 24 is a coronal view of a vertebra showing an insertion device comprising a platform within

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the vertebra, and an expandable structure expanding away from the platform of the insertion device;

Fig. 25 is a coronal view of the vertebra of Fig. 24, wherein the expandable structure has been contracted, the device rotated, and an expandable structure expanded towards another region of the vertebra;

Fig. 26 is a coronal view of the vertebra of Fig. 24, wherein the created cavity is filled with a bone filler;

Fig. 27 is a coronal view of the vertebra of Fig. 24, wherein an expandable structure and insertion device are used to enlarge a first cavity which has been partially filled with a bone filler;

Fig. 28 are side and perspective views of various embodiments of platforms constructed in accordance with the teachings of the present invention;

Fig. 29 is a side perspective view of one embodiment of an expansion guide constructed in accordance with the teachings of the present invention;

Fig. 30 is a partial side perspective view of the expansion guide of Fig. 29 inserted into an insertion device;

Fig. 31 is a side view of another alternate embodiment of an insertion device constructed in accordance with an alternate embodiment of the present invention;

Fig. 32 is a cross-sectional view of the insertion device of Fig. 31, taken along line 32-32;

Fig. 33 is a cross-sectional view of the insertion device of Fig. 31, taken along line 33-33;

Fig. 34 is a side view of another alternate embodiment of an insertion device constructed in accordance with an alternate embodiment of the present invention;

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Fig. 35 is a cross-sectional view of the insertion device of Fig. 34, taken along line 35-35;

Fig. 36 is a cross-sectional view of the insertion device of Fig. 34, taken along line 36-36;

5 Fig. 37 is a cross-sectional view of a step in one method of manufacturing the insertion device of Fig. 34, taken along line 36-36;

Fig. 38 is a cross-sectional view of the insertion device of Fig. 37, during a subsequent manufacturing step;

Fig. 39. is a cross-sectional view of the insertion device of Fig. 37, during a subsequent manufacturing step.

15 Fig. 40 is a perspective view of a balloon catheter device.

Fig. 41 is an exploded view of the device shown in Fig. 40.

20 Fig. 42 is an enlarged partial view of the sheath shown in Fig. 40 and illustrating an opening within the sheath.

Fig. 43 is an enlarged partial side view of the device shown in Fig. 40, illustrating expansion of an expandable structure outside of the opening and beyond the sheath and crimping of the securing tabs to secure the expandable structure within the sheath.

Fig. 44 is a top view of the device shown in Fig. 43 and illustrating a gap formed between the crimped securing tabs.

30 Fig. 45 is view similar to Fig. 44 and illustrating the crimped securing tabs in contact.

Fig. 46 is a perspective view of an alternative embodiment of a balloon catheter device in which the catheter extends beyond the distal end of the sheath.

35 Fig. 47 is a top partial view of the device

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shown in Fig. 46 and illustrating expansion of the expandable structure through an opening in the sheath.

Fig. 48 is a side view of the device shown in Fig. 47.

5 Fig. 49 is a coronal view of a vertebra showing an insertion device within the vertebra.

Fig. 50 is a coronal view of the vertebra of Fig. 49, wherein a balloon catheter device carrying an expandable structure has been inserted through the
10 insertion device and the expandable structure has been expanded.

Fig. 51 is a side view of a normal human vertebra.

Fig. 52 is a side view of a human vertebra
15 having an anterior wedge fracture.

Fig. 53 is a side view of a human vertebra having a bi-concave fracture.

Fig. 54 is a side view of a human vertebra having a vertebral plana fracture.

20 **Detailed Description of the Preferred Embodiment**

The present invention overcomes the problems and disadvantages associated with current strategies and designs in insertion devices for use with expandable structures, such as medical balloons. In particular, the
25 present invention provides insertion devices which may be used with expandable structures to direct expansion of the structure as well as to assist in insertion and removal of expandable structures from an interior region of a human or animal body. The methods and instruments
30 suitable for such treatment are more fully described in U.S. Patent Nos. 4,969,888, 5,108,404, 5,827,289, 5,972,015, 6,048,346 and 6,066,154, each of which are incorporated herein by reference.

Fig. 1 depicts a vertebra 41 to be treated
35 using an expandable structure 310. An insertion device

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200, such as a cannula or spinal needle, extends through the cortical bone 69 of the vertebra 41, and into the cancellous bone 71. An expandable structure 310 is introduced into the vertebra 41 through the insertion
5 device 200, and desirably expands within the cancellous bone 71, typically expanding outward in a spherical, cylindrical or other manner thereby creating a cavity. To avoid contacting the cortical bone 69 during expansion of the structure 310, a practitioner will typically
10 position the insertion device 200 a sufficient distance away from the cortical bone 69 to allow room for the structure 310 to expand outward. However, if the insertion device 200 is positioned too close to the cortical bone 69, if the structure expands a greater
15 amount towards the cortical bone 69 (such a where the cancellous bone is weaker in that direction), or if the intervening anatomy severely constrains placement of the insertion device to locations near the cortical bone 69, the expansion of the structure and cavity creation may be
20 less than optimal.

Figs. 2 and 7 depict one embodiment of an insertion device constructed in accordance with the teachings of the present invention. The insertion device comprises a hollow member 210 which may be any
25 appropriate shape, but is preferably cylindrical. The hollow member 210 has a distal end 250 and a proximal end 255, wherein the distal end 250 is the tip, or point of insertion, of the insertion device. The hollow member 210 may be any appropriate length to allow the insertion
30 device to provide percutaneous access to an interior region of a body requiring treatment. In one preferred embodiment, the hollow member 210 is approximately 12 cm long.

The hollow member 210 has an appropriate
35 central bore diameter and wall thickness to allow

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surgical instruments and/or medical materials to be passed therethrough, while desirably being strong enough to resist deformation during insertion into an interior region of a body such as a bone. In a preferred
5 embodiment, the hollow member 210 has an internal bore diameter of 0.3 cm and a wall thickness of 0.05 cm. The hollow member 210 may further be made of any material which is appropriate for use within a human or animal
10 body including, but not limited to, stainless surgical steel, teflon, polyethylene, polypropylene, silicates, and liquid crystal polymers (as described in U.S. Pat. No. 6,036,711, which is incorporated herein by reference). In one preferred embodiment, the hollow
15 member 210 is made of stainless steel. If desired, the hollow member 210 may further be coated with any appropriate medical grade coating including, but not limited to, an anti-infective, an anticoagulant, a release coating, and/or a slipping agent.

In one embodiment, an extension or platform
20 220 protrudes from the distal end 250 of the hollow member 210. In this embodiment, the platform 220 comprises a semi-cylindrical section which extends from the walls of the hollow member 210. Of course, the platform could be formed in many different
25 configurations, including one or more of those shown in Fig. 28. In one preferred embodiment, the platform 220 is made of the same material as the hollow member 210. Of course, it should be understood that the platform 220 could be formed of and/or coated with materials different
30 from those incorporated into the hollow member 210. In addition, the platform 220 could be formed integrally with the hollow member 210, such as by cutting away a portion of the hollow member 210 near the distal tip 250 and leaving a cradle shape, or by attaching the platform
35 220 to the distal tip 250 of the hollow member by various

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means well known in the art such as welding, adhesive bonding, etc. In one embodiment, the platform 220 will have sufficient column strength such that it will not buckle and/or significantly deform as the insertion
5 device is introduced through soft tissue and/or the bone.

In the disclosed embodiment, the portion of the distal tip 250 of the hollow member 210 has been longitudinally bisected and removed, with the remaining semi-cylindrical section comprising the platform 220.

10 Desirably, the platform 220 will be positioned near the expandable structure 310 prior to expansion, with the platform 220 located between the expandable structure 310 and a region within the vertebra 41 which is not to be compressed or affected. As the structure
15 310 expands, the platform 220 will act as a support, foundation or barrier to the expandable structure 310, inhibiting the structure 310 from expanding in one or more directions. In effect, the platform 220 will induce the expandable structure 310 to expand away from the
20 platform 220. Because the insertion device and platform 220 can be substantially secured within the cortical bone, the platform 220 will desirably remain substantially rigid and/or immobile within the vertebra as the structure expands. This arrangement allows a
25 practitioner to direct the expansion of the expandable structure 310 towards or away from a specific region of the vertebra.

In another embodiment, best shown in Fig. 8, the platform 220 comprises at least one longitudinal
30 crease line 225, preferably located near the center of the platform, along which at least a portion of the platform 220 desirably deforms prior to, during or after expansion of the structure 310. By causing the platform 220 to deform in a controlled manner, this embodiment
35 facilitates introduction of the platform in a lower

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profile condition, permits the platform to deform to a larger area to best direct the expansion of the structure, and then allows the platform to be withdrawn in a lower profile condition. As an expandable structure
5 310 is expanded, thereby exerting pressure against the platform 220, the crease line 225 facilitates flattening of the platform 220 in a controlled manner, thereby providing a widened and improved support surface for guiding expansion of the expandable structure 310.
10 Flattening of the support 220 can also deform outward the edges of the platform 220, which may be sharp, thereby reducing the risk of damaging or rupturing the expandable structure 310. The crease lines 225 may be created by mechanical cutting, laser etching, welding, brazing, or
15 any other well known means.

In an alternate embodiment, the crease line 225 could reinforce the platform 220, minimizing deformation of the platform 220 during expansion of the structure 220. For example, the rounded underside of the
20 platform 220 could be crimped or bent along the longitudinal axis of the platform to stiffen the platform 220 and resist deformation of this type. If desired, the crimp (not shown) could parallel one or more crease lines 225. In an alternate embodiment, a crimp (not shown)
25 which extends approximately 7mm proximally from the distal tip of the platform 220 results in a significant increase in the resistance of the platform 220 to displacement and/or deformation.

In another alternative embodiment, the distal
30 tip of the platform incorporates one or more serrations or teeth which extend outward from the distal tip and facilitate anchoring of the platform into the opposing cortical wall of the targeted bone region. This arrangement, which allows the platform to be supported at
35 both ends, significantly increases the resistance of the

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platform to displacement and/or deformation during expansion of the expandable structure.

In another embodiment, best shown in Figs. 11, 12 and 13, the platform 220 comprises a relatively flat section extending from the distal end 250 of the hollow member 210. In a preferred embodiment, the platform 220 can incorporate a flattened top surface 212 and a curved outer surface 213, the curved outer surface 213 being formed integrally with the hollow member 210. Although this embodiment slightly constricts the inside bore of the hollow member 210, the shape and increased thickness of the platform 220 greatly increase the amount of force the platform 220 can withstand without deforming. In addition, this embodiment minimizes impingement of sharp edges onto the expandable structure. Moreover, the flat and thickened platform 220, as shown in Fig. 13, can be shaped to have a sharpened tip so that the platform 220 can easily pass through soft tissue and/or bone. In an alternate embodiment, a flexible and/or pliable surface (not shown) may be positioned between the expandable structure and/or the platform, or may be incorporated into the platform or expandable structure, to minimize tearing, cutting and/or other failure of the expandable structure.

Figs. 29 and 30 depict an alternate embodiment of an insertion device and associated component constructed in accordance with the teachings of the present invention. In this embodiment, the insertion device or cannula comprises a hollow member 210 which may be any appropriate shape, but is preferably cylindrical.

The hollow member 210 has a distal end 250, wherein the distal end 250 is the tip, or point of insertion, of the insertion device. An expansion guide 400, best shown in Fig. 29, comprises a handle assembly 405 and a guide shaft 410. The guide shaft 410 is desirably longer than

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hollow member 210, and is also desirably sized to pass through the lumen of the hollow member 210. In the disclosed embodiment, an upper surface 420 of the guide shaft 410 is desirably substantially flattened, and a lower surface 415 of the guide shaft is curved. If desired, the handle assembly 405 and/or guide shaft 410 can incorporate one or more alignment marks 407 to indicate the orientation of the guide shaft, as well as the amount the guide shaft extends from the distal end 250 of the hollow member. In addition, the handle assembly may incorporate mechanical connectors or clips (not shown) to secure the expansion guide 400 to the hollow member 210.

In this embodiment, after an insertion device is positioned within a targeted vertebral body, the expansion guide 400 can be positioned near an expandable structure (not shown) prior to expansion, with the guide shaft 410 located between the expandable structure and an area of the cancellous bone where compression of the cancellous bone is not desired. If desired, the expandable structure can be introduced through the insertion device before the expansion guide 400 is introduced through the insertion device. As the structure expands, the guide shaft 410 will act as a support, foundation or barrier to the expandable structure, desirably inhibiting the structure from expanding in one or more directions. In effect, the guide shaft 410 will act similar to the platform 220 previously described, and will induce the expandable structure 310 to expand away from the guide shaft 410. This arrangement allows a practitioner to direct the expansion of the expandable structure towards or away from a specific region of the vertebral body. In addition, because the expansion guide 400 can be inserted to varying depths within the hollow member, the

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practitioner can choose the desired length of the guide shaft 410 to extend out of the insertion device. In an alternate embodiment, if desired a plurality of platforms (not shown) can be used to shield multiple directions.

5 Because the expansion guide 400 may be introduced after the insertion device is already positioned within the targeted vertebral body, the expansion guide 400 need not have sufficient column strength to penetrate soft tissue and/or cortical bone.
10 This allows the expansion guide 400 to assume a variety of cross sectional forms, including one or more of the forms shown in Fig 28.

 In another alternate embodiment of an insertion device constructed in accordance with the
15 teachings of the present invention, best shown in Figs. 3a, 4 and 6, the insertion device comprises a cylindrical hollow member 210 having a distal end 250 and a proximal end 255, wherein the distal end 250 is the tip, or point of insertion, of the insertion device. The distal end
20 250 of the hollow member 210 is scored longitudinally to form a plurality of score lines 260 around the circumference of the hollow member 210. The plurality of score lines 260 may run parallel or at an angle to one another, and are separated by adjacent sections 265. The
25 score lines 260 may be of any appropriate length and depth to allow the distal end 250 to flare (See Figs. 4 and 6) when an outward pressure is exerted upon the adjacent sections 265.

 In one preferred embodiment, the score lines
30 260 extend approximately 0.5 cm along the longitudinal axis of the hollow member 210, and extend through the wall of the hollow member 210. The score lines 260 are cut into the distal end 250 using any appropriate technique known to those of skill in the art including,
35 but not limited to, laser cutting or etching, chemical

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etching and/or mechanical cutting with carbide or diamond tip saws or high pressure water. The distal end 250 of the hollow member 210 will desirably comprise a sufficient number of longitudinal score lines 260 to
5 allow ease of flaring of the distal end 250. The quantity of score lines 260 required for appropriate flaring is determined by the diameter and wall thickness of the hollow member 210 and the ductility of the material. In one embodiment of the present invention,
10 the hollow member 210 comprises at least three score lines 260 in the distal end 250. In another embodiment, best shown in Figure 6, the hollow member 210 comprises six score lines 260 in the distal end 250.

The flaring of the tip of the hollow member
15 210 may ease insertion and removal of an expandable structure, such as a medical balloon. By flaring the tip, the sharp outer edges of the hollow member 210 are pushed away from the expandable structure and into the surrounding cancellous bone. The expandable structure is
20 thus isolated from these sharp edges, which could contact the expandable structure during expansion, possibly causing the structure to rupture or tear. During withdrawal of the expandable structure, the larger diameter of the flared tip will desirably guide the
25 expandable structure into the smaller diameter hollow member 210, easing withdrawal of the expandable structure into and through the hollow member 210.

If desired, flaring of the tip can be accomplished using an expandable structure to provide a
30 desired outward force, or the tip can be flared mechanically. For example, in the embodiment shown in Fig. 3b, the adjacent sections 265 are thickened on their internal surfaces to form one or more protrusions 266 extending inward from each adjacent section 265. If
35 desired, the protrusions 266 could be formed as a single

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continuous thicker area of the circumference of distal end 250, interrupted by the longitudinal score lines 260.

When a tool, such as a blunt obturator, boring member or stylet 275, which is described below, slides across or
5 presses against the protrusions 266, the adjacent sections 265 are desirably forced outward, flaring the distal end 250 of the hollow member 210 in the desired manner.

The insertion device of the present invention
10 may further comprise a removable blunt obturator or stylet 275. See Figs. 5, 9a, 9b, 12, and 17. The stylet 275 comprises a distal end 279 having a tip 276 which can be blunt or sharpened. If desired, the stylet can be cannulated (not shown) to accommodate the guide wire of a
15 spinal needle assembly, as well known in the art. In one embodiment of the present invention, best shown in Figs. 15, 16a and 17, the tip 276 of the stylet 275 will desirably extend from the distal end 250 of the hollow member 210 when the insertion device is assembled for
20 insertion into an interior body region. The stylet 275 desirably pushes and/or cuts a tunnel or passageway through soft tissue and bone to permit placement of the insertion device into the desired interior body region. If desired, the stylet 275 can further comprise a mating
25 end (not shown) which allows the boring member 275 to be mated to the insertion device during the insertion procedure, in a manner well known in the art. Mating of the hollow member 210 to the stylet 275 desirably prevents slippage and relative movement between these
30 devices during insertion into the patient. The stylet 275 is preferably mated to the hollow member 210 in a manner which allows for easy removal of the stylet 275 from the hollow member 210 after placement of the insertion device in the targeted area.

35 The stylet 275 may be made of any appropriate

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medical grade material and is preferably made of the same material as the hollow member 210. In one preferred embodiment, the stylet 275 is made of stainless steel. The stylet 275 may further be any appropriate shape and size which allows it to slide within and mate with the hollow member 210. In a preferred embodiment, the stylet 275 is approximately the same cylindrical shape as the hollow member 210, is slightly longer than the hollow member 210 so that the tip 276 will protrude from the distal end 250 of the hollow member 210 when assembled for insertion, and is slightly smaller in diameter than the inner bore diameter of the hollow member 210, such that the stylet 275 can freely slide within the hollow member 210 for easy insertion and withdrawal.

In one embodiment of the present invention, shown in Figs. 15, 16a, 16b and 17, the distal end 279 of the stylet 275 desirably comprises one or more grooves or divots 273 located near the tip 276. In a preferred embodiment, the divot 273 is a continuous divot which encircles the circumference of the distal end 279 of the stylet 275. A stylet 275 having at least one divot 273 at the distal end 279 is well suited to mate with a hollow member 210 having a plurality of longitudinal score lines 260 and one or more protrusions 266 on each adjacent section 265 in its distal end 250, as depicted in Figs. 3b, 5, and 17. When such an insertion device is assembled, the stylet 275 can be inserted into the hollow member 210 until the distal end 279 extends out of the hollow member 210. The adjacent sections 265 are then folded or crimped inwards, with the protrusions 266 extending into the divot 273 in the stylet 275, such that the outer wall of hollow member 210 is relatively cylindrical prior to insertion of the insertion device. Once the insertion device is in place within the desired interior region of a body, the stylet 275 is pulled out

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of the hollow member 210, flaring the one or more protrusions 266 and forcing the distal end 279 of the stylet 275 outward. In another embodiment, the collar section 267 adjacent to one or more protrusions 266 is thinner than the rest of the wall of hollow member 210 to make flaring of the distal end 250 easier.

In another embodiment of an insertion device constructed in accordance with the teachings of the present invention, best depicted in Figs. 9a through 9c, an insertion device comprises a hollow member 210 having a platform 220 and one or more adjacent sections separated by a plurality of longitudinal score lines 260 at the distal end 250. In another embodiment, this type of insertion device further comprises a stylet 275 which has at least one divot 273 located near the tip 276. The stylet 275 can be used to exert outward pressure on one or more protrusions 266 on adjacent sections 265 causing the distal end 250 to flare out near the platform 220. Such a preferred embodiment of an insertion device of the present invention allows a user to direct expansion of an expandable structure, such as a medical balloon, while easing insertion and removal of the expandable structure and reducing the risk of damage to the expandable structure.

The present invention further provides methods for using the disclosed insertion devices to direct expansion of the expandable structure and/or to simplify insertion and removal of an expandable structure from an interior region of a human or animal body. For illustrative purposes, a method for osteoporotic vertebral fixation, i.e. insertion and expansion in a vertebral body, will be described. However, a similar method may be used within any appropriate region of a human or animal body.

As shown in Figs. 18-27, in one embodiment of

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the present invention, a patient 10 is placed onto a holder 15, generally U shaped, so that the patient's back is exposed. An x-ray, CAT-scan, MRI, fluoroscope, or other appropriate device 20 which permits a practitioner to visualize the insertion and placement of an insertion device during the surgical procedure may be positioned around the patient. An insertion device 200 comprising a hollow member 210 fitted with a stylet 275, as previously described above, can be introduced through the soft tissues to a vertebral body, which can be located fluoroscopically. The stylet and insertion device will desirably penetrate through the cortical bone 31 of the vertebral body 30, and the stylet 275 can then be removed. In an embodiment of the insertion device wherein the hollow member 210 comprises one or more adjacent sections 265 separated by a plurality of longitudinal score lines 260, the removal of the stylet 275 desirably causes the distal end 250 of the hollow member 210 to flare as depicted in Fig. 22.

An expandable structure 50, such as a medical balloon, can be inserted through the hollow member 210 into the vertebral body 30. Placement of the expandable structure 50 can be monitored by any appropriate means, including x-ray fluoroscopy or real time MRI. The structure is expanded, creating a cavity 55 within the cancellous bone 32 and/or moving cortical bone 31, and then contracted and removed. In an embodiment where the distal end 250 of the hollow member 210 has been flared, the flared end guides the structure 50 into the hollow member 210. The cavity 55 can then be filled with an appropriate bone filler 60.

In another preferred embodiment of the present invention, the hollow member 210 comprises a platform 220 extending from the distal end 250. See Figs. 24 through 27. Once the hollow member 210 is introduced into the

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vertebral body 30, the hollow member can be rotated until the platform 220 shields an area of the vertebral body where expansion of the structure 50 is undesired. When the structure 50 is expanded, the platform 220 induces
5 the structure 50 to expand away from the platform 220. In this way, an appropriate area for a cavity may be formed generally irrespectively of where the insertion device is placed within the vertebral body. Thus, if the
10 insertion device is placed in a position within the vertebral body that is not optimal for cavity formation, instead of torquing, bending, or otherwise adjusting the placement of the entire insertion device, the insertion device 200 may simply be rotated until the platform 220 faces a desired direction of cavity formation. An
15 indicator (not shown) on the handle or proximal portion of the hollow member 210 will desirably indicate to the practitioner the orientation of the platform within the bone. Similarly, if a larger or asymmetrical cavity is desired, after a first cavity is formed by expanding the
20 structure 50, the structure 50 may be contracted, the insertion device 200 may be rotated until the platform 220 faces another direction, and the same or a different structure 50 may be expanded to form a second cavity, etc., as depicted in Fig. 25. Any desired number and/or
25 dimension of cavities may be formed in this way. In another embodiment, different shaped balloons may be inserted to form each different cavity or multiple expandable structures of varying shapes may be used to form each cavity.

30 When the desired cavity or cavities 55 have been formed, the expandable structure 50 may be contracted and removed through the hollow member 210. In an embodiment wherein the distal end 250 of the hollow member 210 is flared, removal of the contracted structure
35 may be easier because the flared tip guides the structure

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into the hollow member 210. A suitable bone substitute, such as polymethylmethacrylate bone cement, a two-part polyurethane material, or any other appropriate biocompatible bone filler 60, is injected into the cavity
5 55 or cavities formed. In one embodiment, a first cavity 55 may be formed and, if desired, at least partially filled with a bone filler 60, then the same or a different expandable structure 50 may be inserted and expanded in the same cavity 55, thereby compacting the
10 hardening bone filler and/or more cancellous bone 32, and the cavity 55 may then be further filled with the same or a different bone filler 60. In another embodiment, a first cavity 55 may be formed, an insertion device 200 with a platform 220 may then be rotated and the same or a
15 different expandable structure 50 may be inserted to create a second cavity or enlarge the first cavity 55, and the cavity(ies) may then be filled with the same or a different bone filler 60. These methods may be followed until all desired cavities have been formed and filled.

20 Once all desired cavities have been filled, the insertion device 200 may be removed from the vertebral body 30. The incision 25 may then be stitched closed and/or covered with bandages.

Figures 31-33 depict an alternate embodiment
25 of an insertion device 600 constructed in accordance with the teachings of the present invention. The insertion device 600 comprises a hollow member 620 and an expandable structure 710. A handle 615 may be provided on the distal end of the hollow member 620 to facilitate
30 manipulation of the tool and/or introduction of a medium to expand the expandable structure 710. The hollow member 620, desirably having a lumen 622 extending therethrough, comprises a shaft 624 and a distal tip 625. The distal end 625 of the shaft 624 can be rounded or
35 beveled to facilitate passage through cortical/cancellous

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bone, or can be flattened to minimize opportunities for penetrating the opposite cortical wall of the targeted bone region. An opening or window 700 is formed in the shaft 624, with an expandable structure 710 desirably positioned within the lumen 622 at a location adjacent the window 700. Upon introduction of the insertion device 600 into a targeted bone region (not shown), the expandable structure 710 can be expanded (See Fig. 33, P1 to P2 to P3), and at least a portion of the expandable structure 710 will desirably expand through the window 700, thereby compressing cancellous bone, creating a cavity and/or displacing cortical bone. Upon contraction of the expandable structure 710, most of the expandable structure 710 will desirably be drawn back into the lumen 622 for removal of the device 600 from the vertebral body. If desired, the handle 615 and/or proximal end 612 of the hollow member 620 can include markings (not shown) which indicate the orientation of the window 700 within the targeted bone region.

The expandable structure 710 may be comprised of a flexible material common in medical device applications, including, but not limited to, plastics, polyethylene, mylar, rubber, nylon, polyurethane, metals or composite materials. Desirably, the shaft 624 will comprise a material that is more resistant to expansion than the material of the expandable structure 710, including, but not limited to, stainless steel, ceramics, composite material and/or rigid plastics. In an alternate embodiment, similar materials for the expandable structure 710 and shaft 624 may be used, but in different thickness and/or amounts, thereby inducing the expandable structure 710 to be more prone to expansion than the shaft 624. The expandable structure 710 may be bonded directly to the shaft 624 by various means well known in the art, including, but not limited

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to, means such as welding, melting, gluing or the like. In alternative embodiments, the expandable structure may be secured inside or outside of the shaft 624, or a combination thereof. In at least one alternative
5 embodiment, at least a portion of the material comprising the expandable structure 710 will plastically deform as it expands.

If desired, the shaft 624 may be sized to pass through the lumen of a cannula or spinal access needle
10 (not shown) already positioned within the targeted bone region. Alternatively, this embodiment of an insertion device 600 can be utilized without an associated insertion device. In such a case, the insertion device 600 will desirably incorporate a sharpened distal tip 625
15 capable of penetrating the soft tissues and cortical/cancellous bone of the vertebral body. The distal tip may be hollow or a solid construct, depending upon the desired penetration strength of the device 600.

Similarly, the window 700 may extend around more or less
20 of the periphery of the shaft 624, depending upon the size and configuration of the expandable structure 710 and the desired penetration strength of the device. For example, where the window 700 extends around approximately 25% of the shaft 624, the penetration
25 strength of the device 600 will be significantly greater than where the window extends around approximately 75% of the shaft 624. If desired, the handle 615 can incorporate an impacting surface (not shown) to facilitate the use of an orthopedic mallet in placing the
30 device 600 in a targeted bone region. In an alternate embodiment, after creation of the cavity, the expandable structure can be removed from the hollow member 600, allowing bone filler to be introduced into the cavity through the hollow member.

35 Figures 34 through 36 depict another alternate

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embodiment of an expansion guide 800 constructed in accordance with an alternate embodiment of the present invention. In this embodiment, the platform 810 comprises a semi-cylindrical section which extends from the walls of the hollow member 820. A notch 825 extends longitudinally along the platform 810. The notch 825 will accommodate a key or projection of an expandable structure (not shown), desirably securing the expandable structure to the platform 810.

Depending on the quality and strength of the surrounding cancellous and/or cortical bone, as an expandable structure expands against the platform 810, the structure can "slide off" the platform 810. In a similar manner, rotation of the platform may displace the expandable structure in an unwanted manner. Desirably, the notch 825 will secure the structure to the platform 810, preventing such occurrences. In addition, the structure may be contracted and the notch 825 used to draw the expanding structure back into proper orientation with the platform 810.

Figures 37 through 39 depict one method of manufacturing the platform 810 of Figure 34. In this embodiment, a distal end of the shaft 820 is cut along a longitudinal line A. Alternatively, the shaft 820 may be cut along longitudinal lines B, depending upon the desired size of the notch and the desired angles of the side walls of the notch. The shaft is placed in a stamping machine 850 and a die 860 stamps the cut walls 821 of the shaft 820 against the opposing walls 822 of the shaft 820. Desirably, the cut walls 821 will contact the opposing walls 822, thereby forming a notch 825 between by the cut walls 821 and the opposing wall 822.

In a similar manner, a notch could be formed in the embodiment of an insertion device described in Figures 29 and 30, and used to guide and secure an

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expandable body to the platform. Once positioned within the targeted bone region, the platform could be manipulated and/or rotated with the expandable structure secured thereto. This embodiment would thereby greatly facilitate proper placement of the expandable structure on the platform in a desired orientation. If desired, the notch could be formed by molding, grinding, stamping or any other machining method known to those in the art.

Figs. 40-50 illustrate another alternative embodiment of a balloon catheter device 900. The device 900 comprises a sheath or hollow member 902. The hollow member 902 is sized and configured to receive a balloon catheter 903. The balloon catheter 903 includes an expandable structure 904 carried by a shaft 906, and a stylet 908 having a blunt, probing ball tip 910. In the illustrated embodiment, the hollow member 902 takes the form of a cannula that forms an outer sheath for the balloon catheter 903. The hollow member 902 may be formed of stainless steel or any other suitable biocompatible material. A handle 912, e.g., a y-adapter, may be provided on the proximal end 914 of the hollow member 902 to facilitate manipulation of the device 900 and/or introduction of a medium to expand the expandable structure 904.

The shaft 906 is desirably sized and configured to permit passage of the stylet 908. The stylet 908 supports the expandable structure 904 throughout the structure's 904 length to provide additional support and stabilization of the expandable structure 904 within the hollow member 902. The stylet 908 may be made of any appropriate medical grade material and is preferably made of the same material as the hollow member 902, e.g., stainless steel. In a preferred embodiment, the stylet 908 is approximately the same cylindrical shape as the shaft 906, is slightly longer

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than the shaft 906 so that the tip 910 will protrude from the distal end 916 of the shaft 906 when assembled for insertion, and is slightly smaller in diameter than the inner bore diameter of the shaft 906, such that the
5 stylet 908 can freely slide within the shaft 906. The stylet 908 can also be fixed with the y-adapter 912 by bonding, such as UV curable adhesive, thermal, ultrasonic or cyanoacrylate adhesive bonding, to a luer lock 909 of the y-adapter 912. If desired, the stylet 908 can be
10 cannulated (not shown) to accommodate the guide wire of a spinal needle assembly, as well known in the art.

The hollow member 902, desirably having a lumen 918 extending therethrough, comprises a shaft 920 and a distal tip 922. The distal tip 922 of the hollow
15 member 902 includes at least one securing element that prevents movement of the expandable structure 904 within the hollow member 902. In the illustrated embodiment, a pair of securing elements are provided and take the form of winged tabs 924.

20 An opening or window platform 926 is formed in the shaft 920 adjacent the winged tabs 924, with the expandable structure 904 desirably positioned within the lumen 918 at a location adjacent the window 926. The window 926 desirably extends along the axis of the hollow
25 member 902 and about the axis of the hollow member 902 for less than 360° to direct expansion of the expandable structure 904. The window 926 desirably extends for less than 180° about the axis of the hollow member 902, and preferably between 90° and 180°. In a representative
30 embodiment, the window extends approximately 135° about the axis of the hollow member 902. The smaller the window 926, the greater the directionality of expansion of the expandable structure 904 and strength of the platform 926.

35 In a representative embodiment, the device 900

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includes the following dimensions:

Window	Length	20 mm.
Expandable Body	Initial length	10 mm.
	Maximum inflation pressure	300 psi.
	Maximum inflation volume	3 cc.
	Maximum diameter	15 mm.
	Maximum length	18 mm.
Hollow Member/Outer Cannula	Outer diameter	4.2 mm (passes through size 3 access tools)
Winged Securing Tab	Width at widest point	2.4 mm.

In the illustrated and preferred embodiment,
 5 the expandable structure 904 is carried tangential to the shaft 906. The directionality of the expandable structure 904 may be indicated by external markers 911 on the handle 912 or by the directionality of the handle 912.

10 The external markers 911 are carried on a strain relief 913 that fits over the y-adapter 912 and hollow member 902. The external markers 911 are winged or flared to show directionality and are aligned with the side port or inflation arm of the y-adapter 912 and the
 15 window 926. The strain relief 913 further acts to prevent kinking of the hollow member 902 at the union of the hollow member 902 and y-adapter 912 and to cover the adhesive securing the union. It is contemplated that various modifications of the markers, as to the location,
 20 type and quantity of markers, could be used which incorporate various directionality arrangements,

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markings, or combinations thereof.

The stylet 908 is passed from the distal end 916 to proximal end 917 of shaft 906 and through the expandable structure 904, and the shaft 906, including
5 expandable structure 904, and stylet 908 are passed through the hollow member 902 from the distal tip 922 to the proximal end 914 for placement of the expandable structure 904 within the window 926. The stylet ball 910 is in contact with the distal tip 922 of the hollow
10 member 902 and may become permanently integrated as part of the assembly. The y-adapter 912 is coupled to the stylet 908 and hollow member 902 as previously described. If desired, the proximal end 914 of the hollow member 902 may be tapered relative to the distal end 922 to permit
15 ease of coupling of the hollow member 902 to the y-adapter 912.

To secure placement of the expandable structure 904 within the window 926, tabs 924 are crimped or folded inwards along the distal end 916 of the shaft
20 906. Crimping or folding of the tabs secures the hollow member 902 to the shaft 906 and thereby prevents lifting and movement of the expandable structure 904 during expansion.

When crimped, the winged tabs 924 also form a
25 taper and reduce the overall distal tip 922 profile, as shown in Fig. 43. The tapered tip 922 allows for easier and less traumatic insertion of the device 900 through bone. The reduced profile of the tapered tip 922 further acts as a landing zone for distended balloon material.
30 After expansion of the expandable structure 904, the distended balloon material plastically deforms and the molded morphology of the expandable structure 904 is altered. The distended balloon material may not completely return to its original shape and a portion can
35 extend outward from the window 926. The tapered tip 922

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allows for distention of the expanded balloon material over the winged tabs 924 preventing bunching or tearing of the expandable structure 904 during removal of the device 900 through the access cannula, thereby reducing withdrawal forces. While two tabs 924 are shown in the illustrated embodiment, it is contemplated that the number and configuration of tabs 924 may vary.

The winged tabs 924 may be adapted to provide a visible gap G in the crimped condition, as shown in Fig. 44. The gap G provides additional landing zone for distended balloon material. In an alternative embodiment, shown in Fig. 45 the winged tabs 924 are adapted to be in contact in the crimped condition, providing no visible gap G.

Figs. 46-48 illustrate an alternative embodiment in which the distal end 916 of the shaft 906 and the stylet tip 910 extend beyond the distal tip 922 of the hollow member 902 to provide additional landing zone for distended balloon material.

The stylet tip 910 allows the device 900 to serve as a cortical wall probe to aid advancement and optimal anterior placement of the device 900 within the vertebral body 30 without breach of the anterior cortical wall. The blunt tip 910 cannot easily pierce the anterior cortical wall of the vertebral body 30. The physician advances the device 900 through an access opening and into cancellous bone 32 until the physician tactilely senses contact between the blunt distal tip 910 and the anterior cortical wall.

Desirably, the stylet 908 is radiopaque, so that its advancement through cancellous bone 32 and its contact with the anterior cortical wall within the vertebral body 30 can be visualized, e.g., by x-ray or real time fluoroscopy or MRI. The stylet 908 therefore allows the physician to gauge the distance between the

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access opening into the vertebral body 30 and the anterior cortical wall, in a manner that avoids penetration of the cortical wall.

In the event of a breach or suspected breach
5 of the anterior cortical wall of the vertebral body 30, the physician can alternatively utilize the stylet tip 910 to safely and easily determine the existence and/or extent of a wall breach. Because the tip 910 of the stylet 908 is blunt, the tip 910 desirably will not
10 easily pass through an intact anterior cortical wall, which allows the physician to "tap" the device along the inner surface of the anterior cortical wall while searching for breaches. Where a wall breach has occurred, and the device 900 could pass through the
15 breach, the blunt tip 910 of the stylet 908 desirably will not pierce or damage soft tissues, such as the aorta or major veins, located forward of the cortical wall. If desired, the blunt tip 910 can alternatively be formed of a soft, deformable material such as rubber or plastic.

20 In use, as shown in Fig. 49, an access opening into a targeted bone region is formed by conventional techniques for insertion of a cannula 928. The physician then passes the device 900 through the lumen of the cannula 928 positioned within the targeted bone region,
25 e.g., a vertebral body 30.

As Fig. 50 illustrates, the expandable structure 904 is then expanded, creating a cavity 55 within the cancellous bone 32 and/or moving cortical bone 31, and then contracted. Fig. 51 illustrates a normal
30 vertebral body 30. The device 900 is positioned such that the window 926 directs the expansion of the expandable structure 904 towards a section of cortical bone 31 to be moved to a desired position, such as a depressed upper or lower plate of a vertebral body 30 such as for example,
35 an anterior wedge (Fig. 52), bi-concave (Fig. 53) or

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vertebra plana (Fig. 54) fracture morphology. The expandable structure 904 may also be rotated or repositioned forward or aft and expanded to create a desired cavity. The expandable structure 904 is then
5 collapsed and removed. The cavity 55 can then be filled with an appropriate bone filler 60.

While the disclosed devices and methods are more specifically described in the context of the treatment of human vertebrae, other human or animal bone
10 types can be treated in the same or equivalent fashion. By way of example, and not by limitation, the present systems and methods could be used in any bone having bone marrow therein, including the radius, the humerus, the vertebrae, the femur, the tibia or the calcaneous.

15 Other embodiments and uses of the invention will be apparent to those skilled in the art from consideration of the specification and practice of the invention disclosed herein. All documents referenced herein are specifically and entirely incorporated by
20 reference. The specification and examples should be considered exemplary only, with the true scope and spirit of the invention being indicated by the following claims.

As will be easily understood by those of ordinary skill in the art, variations and modifications of each of the
25 disclosed embodiments, including combinations thereof, can be easily made within the scope of this invention as defined by the following claims.

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We Claim:

1. A bone treatment device comprising
an expandable structure, and
a sheath housing the expandable structure, the
5 sheath including an opening extending along and about the
axis of the sheath to direct expansion of the expandable
structure, the sheath including at least one securing
element that secures the expandable structure within the
sheath.
- 10 2. A device as in claim 1
wherein the securing element is a tab.
3. A device as in claim 2
wherein the tab is tapered.
4. A device as in claim 1
15 wherein the opening extends along the axis of
the sheath and about the axis of the sheath for less than
360°.
5. A device as in claim 1
wherein the expandable structure is adapted to
20 compact cancellous bone.
6. A device as in claim 1, further
comprising
a stylet having a distal end and sized and
configured to extend through the expandable structure,
25 the distal end of the stylet extending at least in part
beyond the distal end of the sheath.
7. A device as in claim 6
wherein the distal end of the stylet is blunt.
8. A cortical bone wall probe comprising
30 a sheath extending along an axis and having a
distal end,
a shaft having a distal end and being sized
and configured for passage through the sheath, and
a stylet having a blunt distal end and sized
35 and sized and configured for passage through the shaft,

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the blunt distal end extending at least in part beyond the distal end of the sheath and the distal end of the shaft.

9. A cortical bone wall probe as in claim 8,
5 further comprising

an expandable structure carried by the shaft,
and

wherein the sheath includes an opening
extending along the axis of the sheath and about the axis
10 of the sheath to direct expansion of the expandable
structure.

10. A cortical bone wall probe as in claim 9
wherein the expandable structure is adapted to
be expanded outside of the opening and beyond the sheath.

11. A cortical bone wall probe as in claim 9
15 wherein the opening extends along the axis of
the sheath and about the axis of the sheath for less than
360° to direct expansion of the expandable structure.

12. A cortical bone wall probe as in claim 9
20 wherein the expandable structure is adapted to
compact cancellous bone.

13. A cortical bone wall probe as in claim 9
wherein the sheath includes at least one
securing element that prevents movement of the shaft
25 within the sheath.

14. A cortical bone wall probe as in claim 13
wherein the securing element is a tab.

15. A bone treatment device comprising
an expandable structure carried by a shaft,
30 the shaft having a lumen extending therethrough and a
distal end,

a stylet having a blunt distal tip and sized
and configured for passage through the lumen to extend
the blunt distal tip, at least in part, beyond the distal
35 end of the shaft to form a balloon catheter assembly, and

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a sheath having a distal end and sized and configured for passage of the catheter assembly, the sheath including an opening for receiving the expandable structure and at least one securing element for securing
5 the sheath to the distal end of the shaft.

16. A device as in claim 15

wherein the distal tip of the stylet extends at least in part beyond the distal end of the sheath.

17. A method of treating bone comprising the
10 steps of

introducing an insertion device through a cortical bone region and into a cancellous bone region of the bone,

introducing a balloon catheter assembly
15 through the insertion device, the balloon catheter assembly comprising an expandable structure and a sheath housing the expandable structure, the sheath including an opening extending along and about the axis of the sheath to direct expansion of the expandable structure and at
20 least one securing element that secures the expandable structure within the sheath, and

expanding the expandable structure outside the opening and beyond the sheath to create a cavity within the bone.

25 18. A method as in claim 17

wherein expansion of the expandable body compacts cancellous bone.

19. A method as in claim 17, further comprising

30 filling the cavity with a bone filler.

Fig. 1

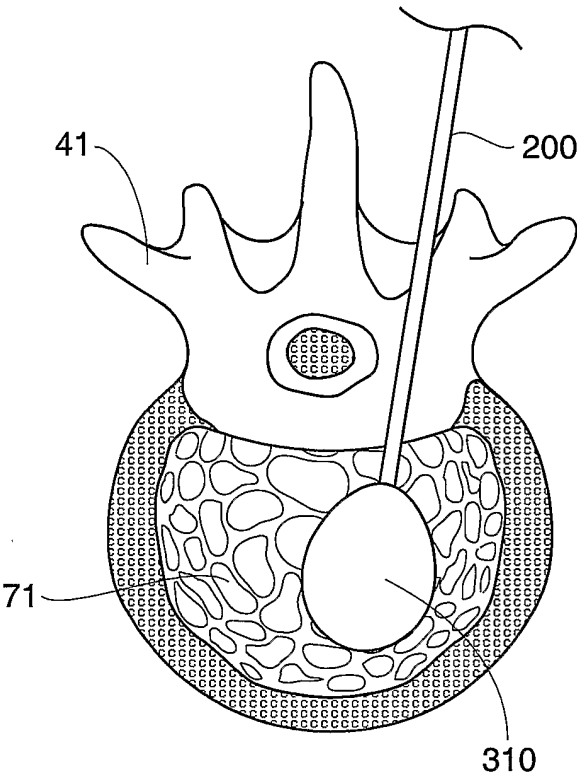
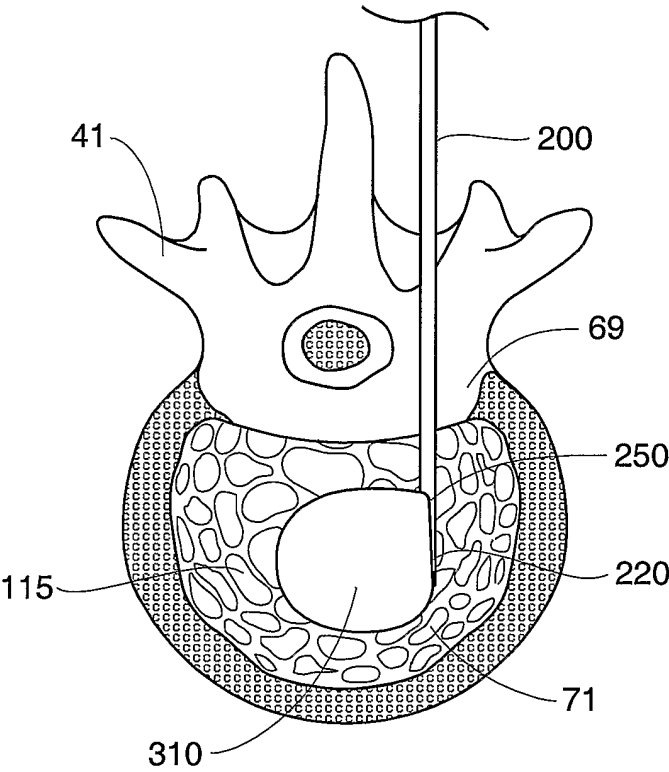
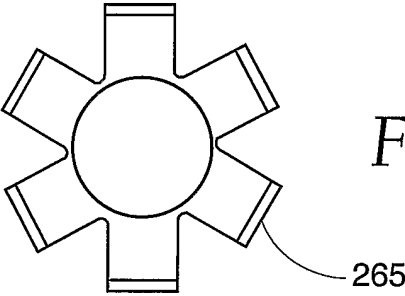
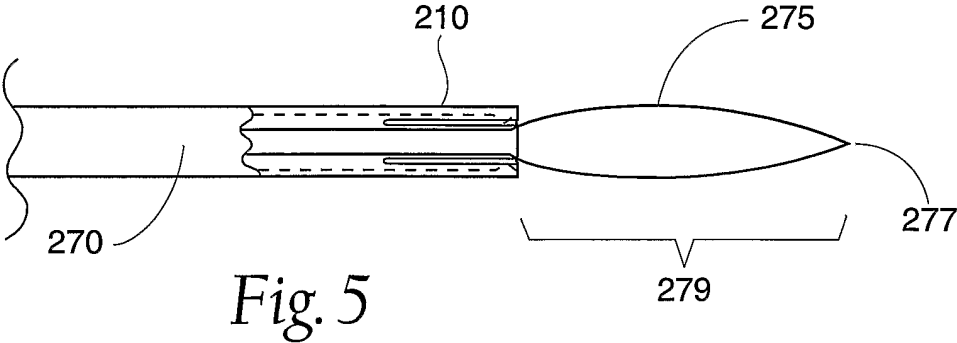
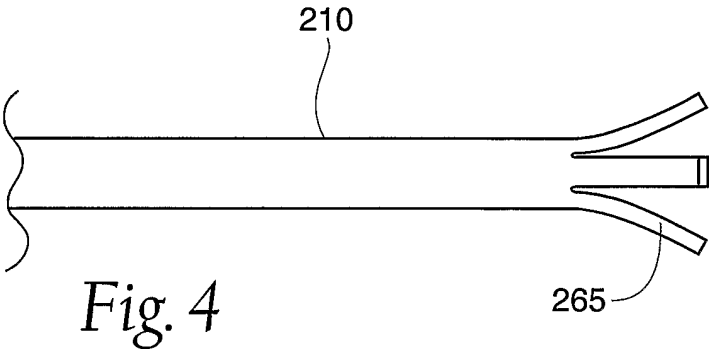
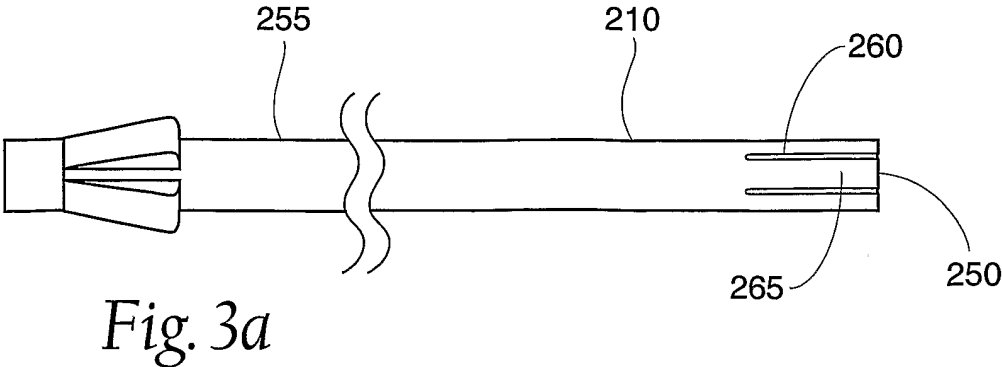


Fig. 2





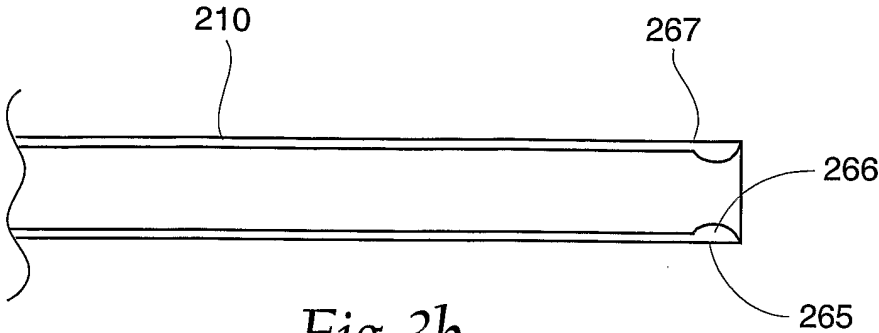


Fig. 3b

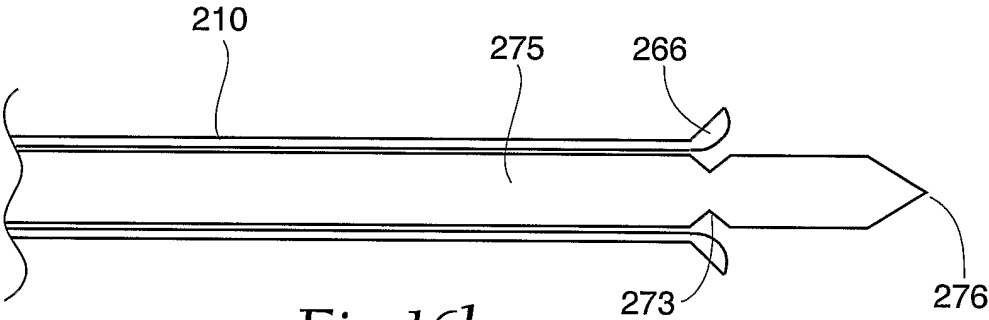


Fig. 16b

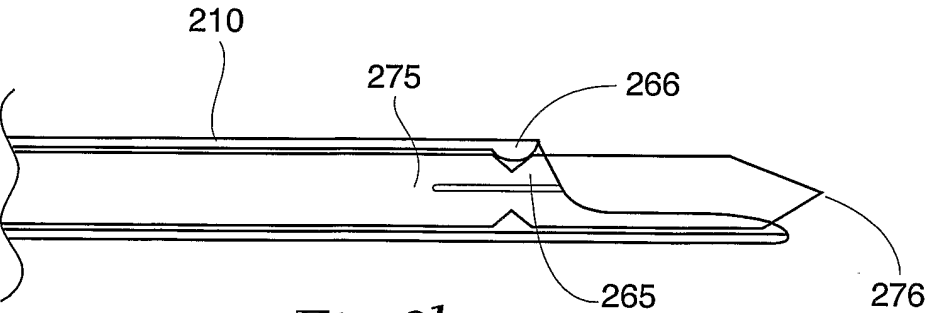


Fig. 9b

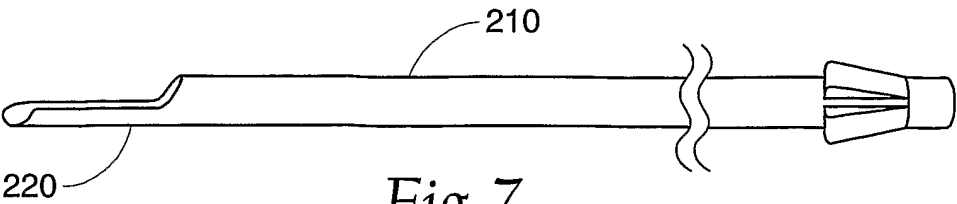


Fig. 7

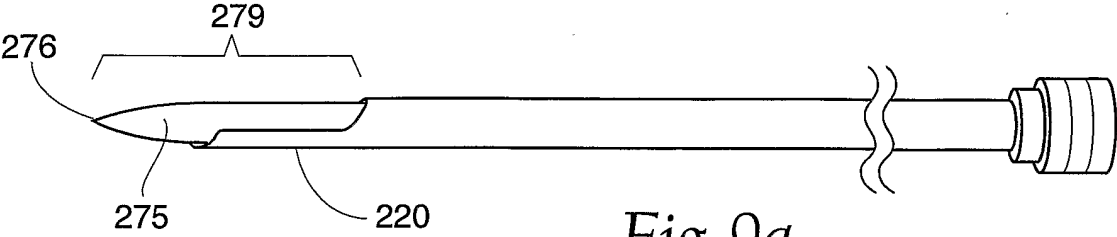


Fig. 9a

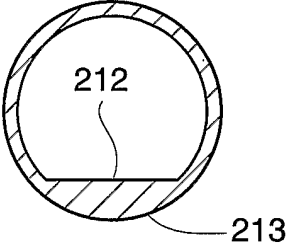


Fig. 10

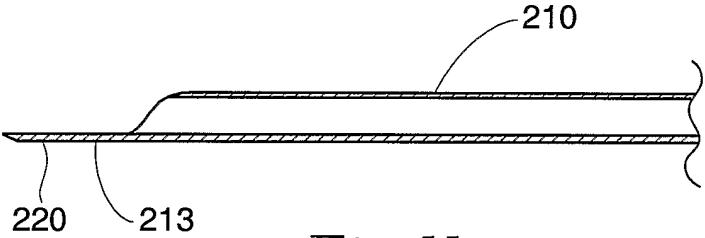


Fig. 11



Fig. 12

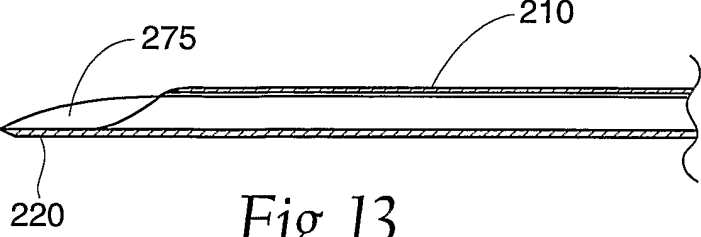


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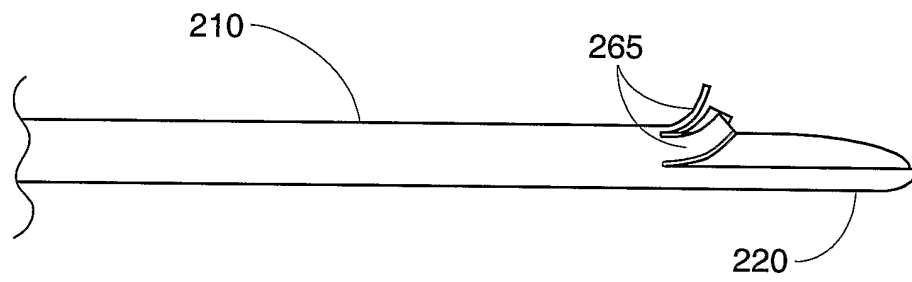


Fig. 9c

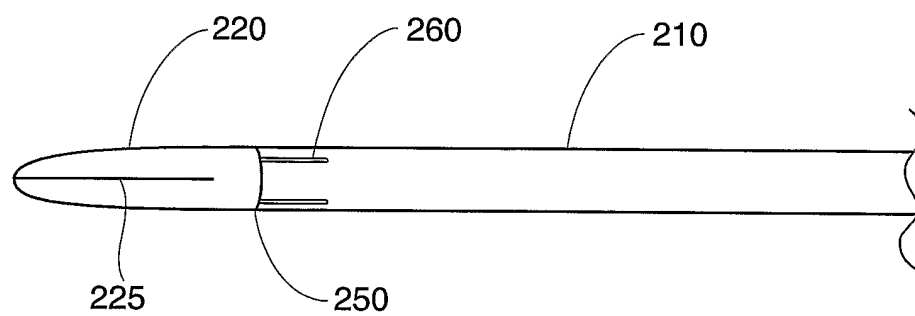


Fig. 8

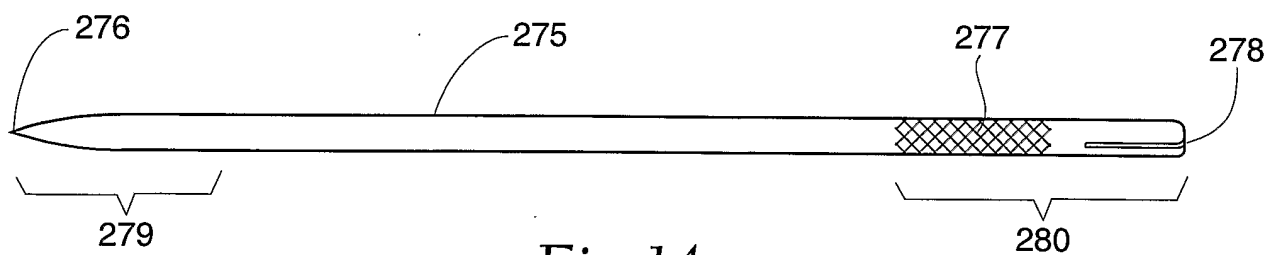


Fig. 14

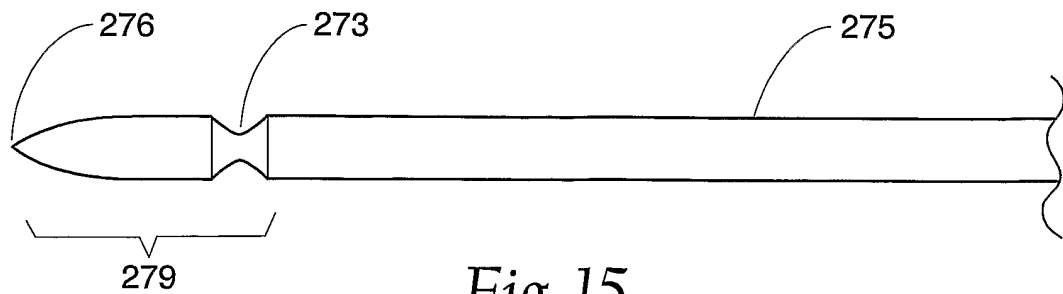


Fig. 15

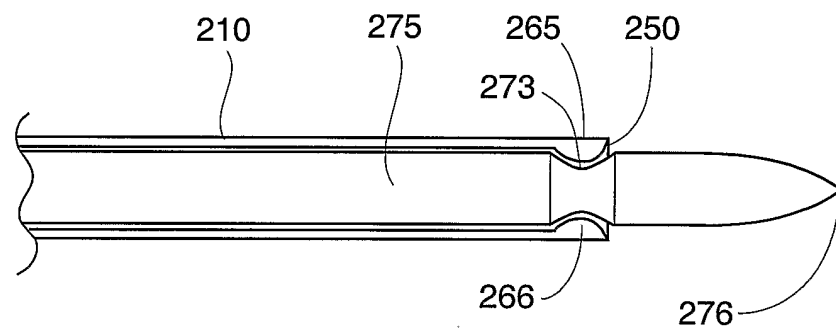


Fig. 17

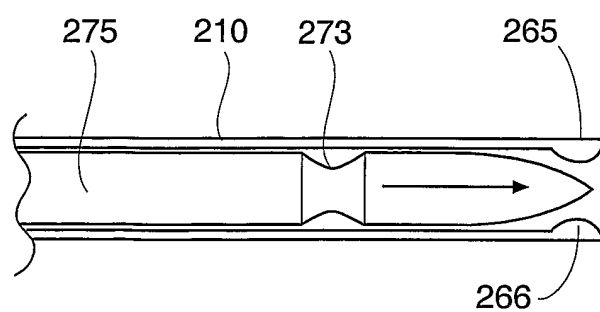


Fig. 16a

Fig. 18

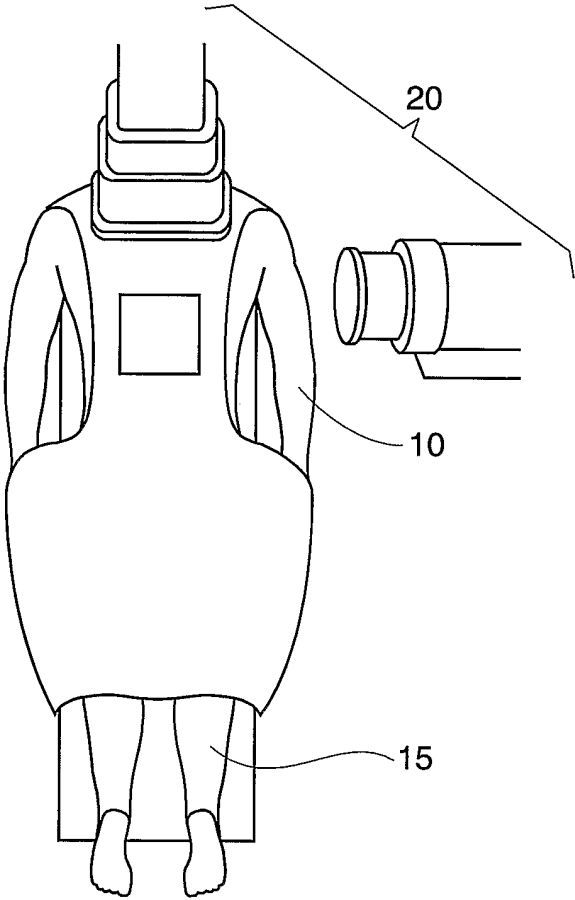
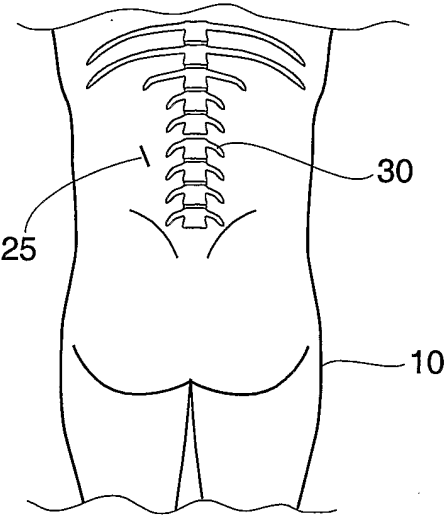


Fig. 19



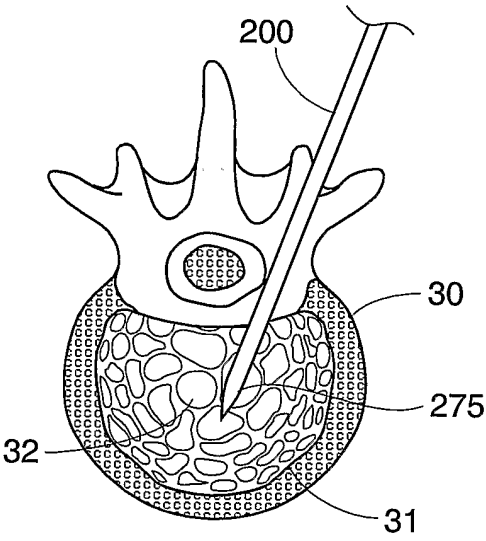
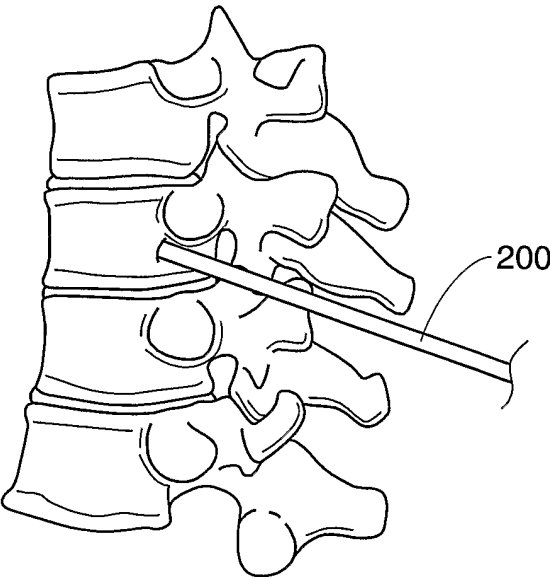
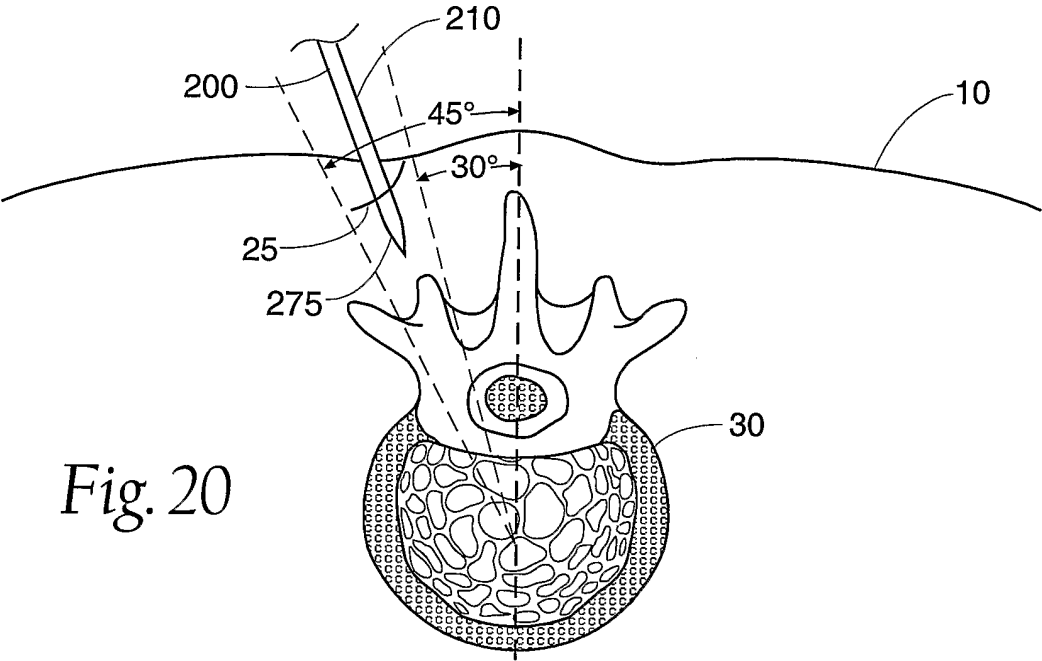


Fig. 22

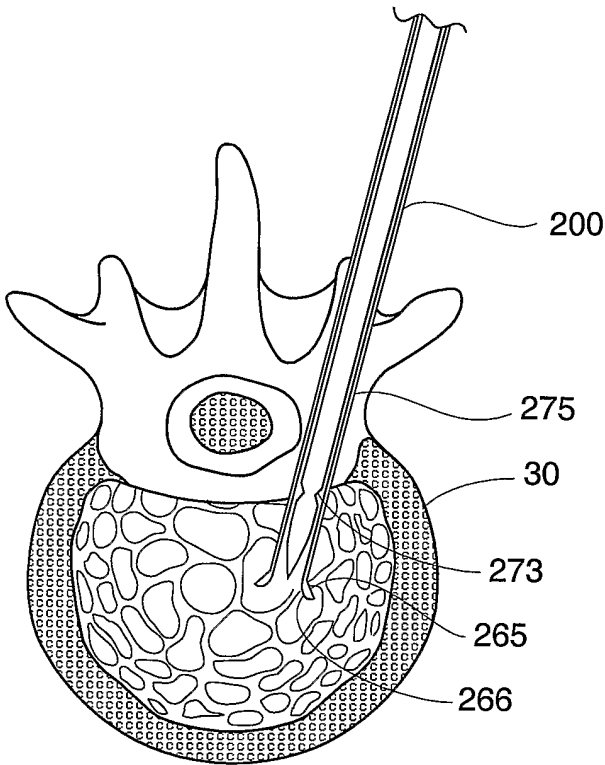


Fig. 23

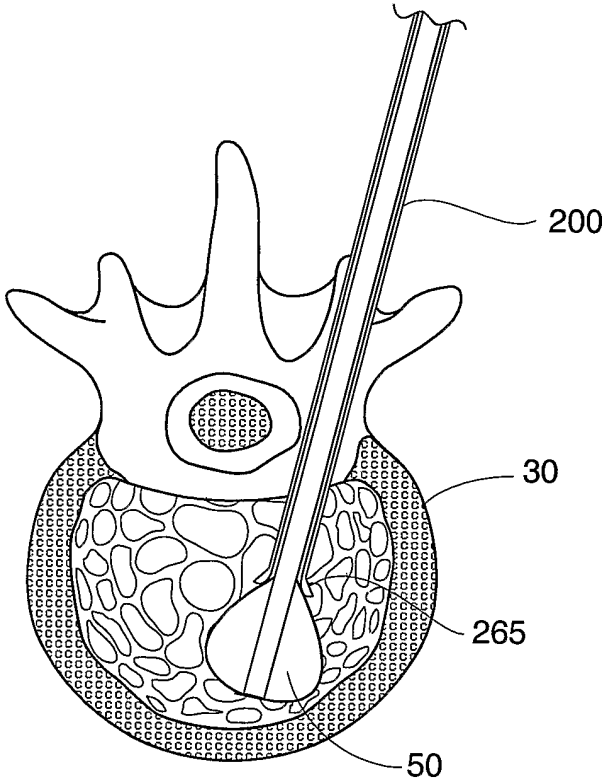


Fig. 24

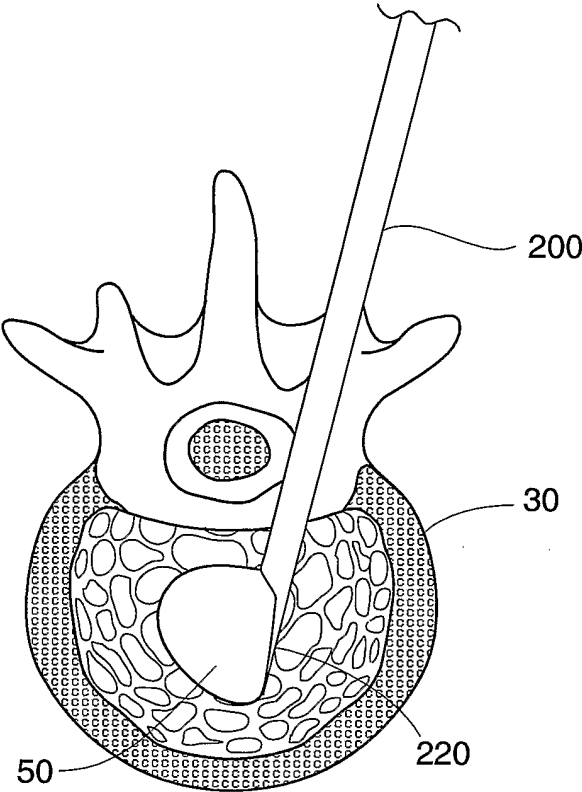


Fig. 25

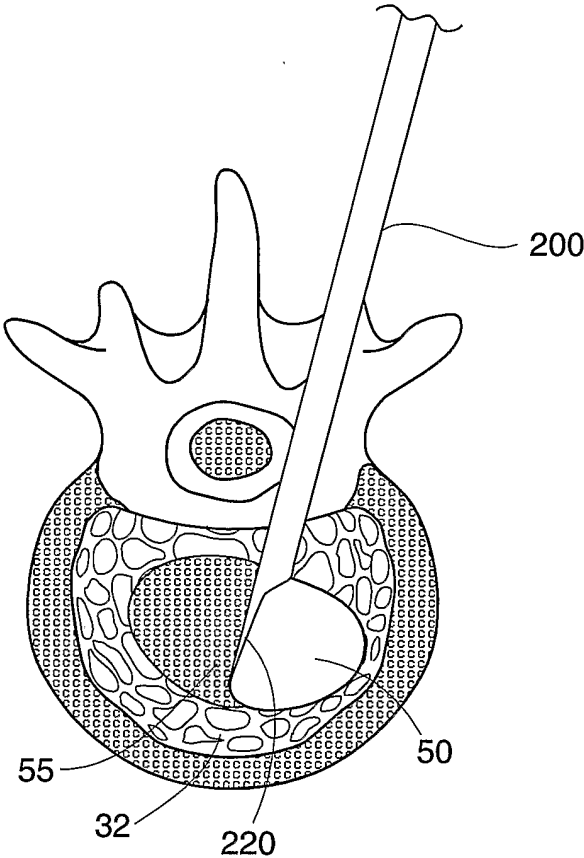


Fig. 26

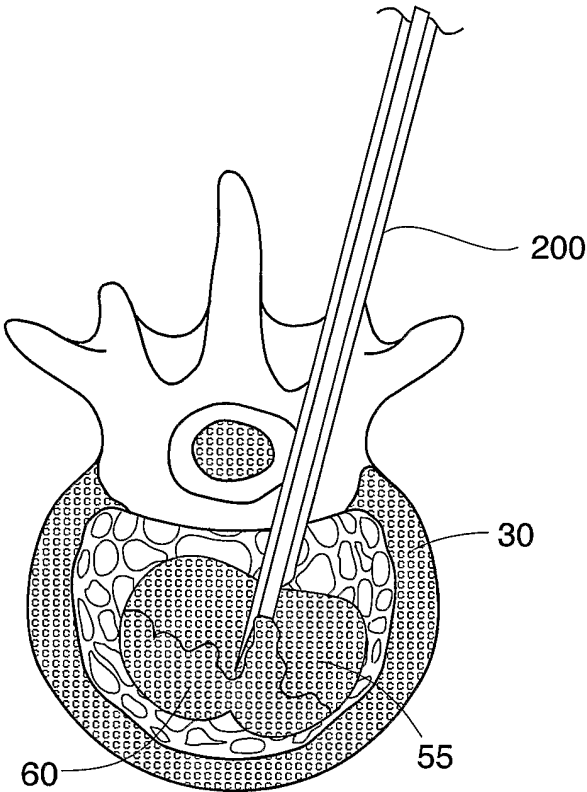
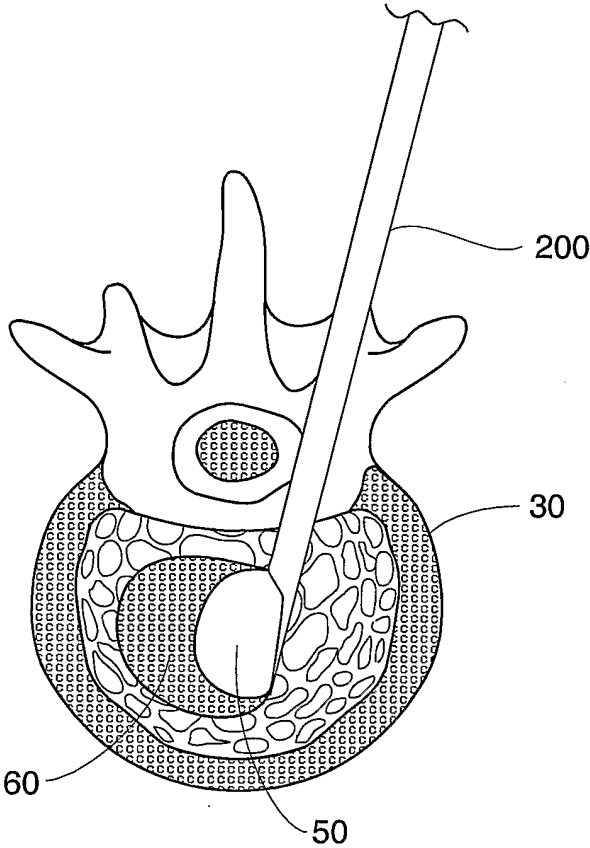
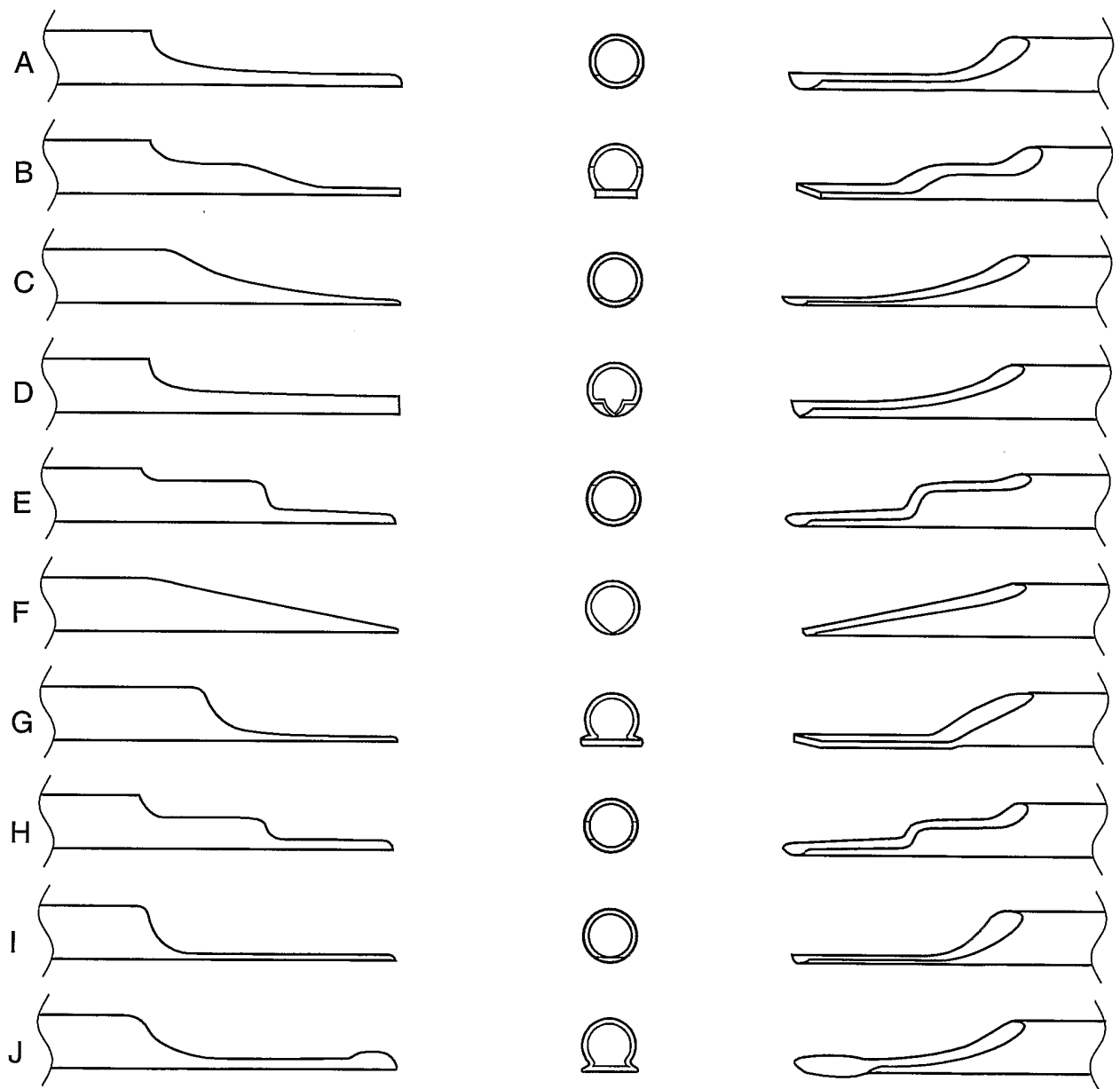
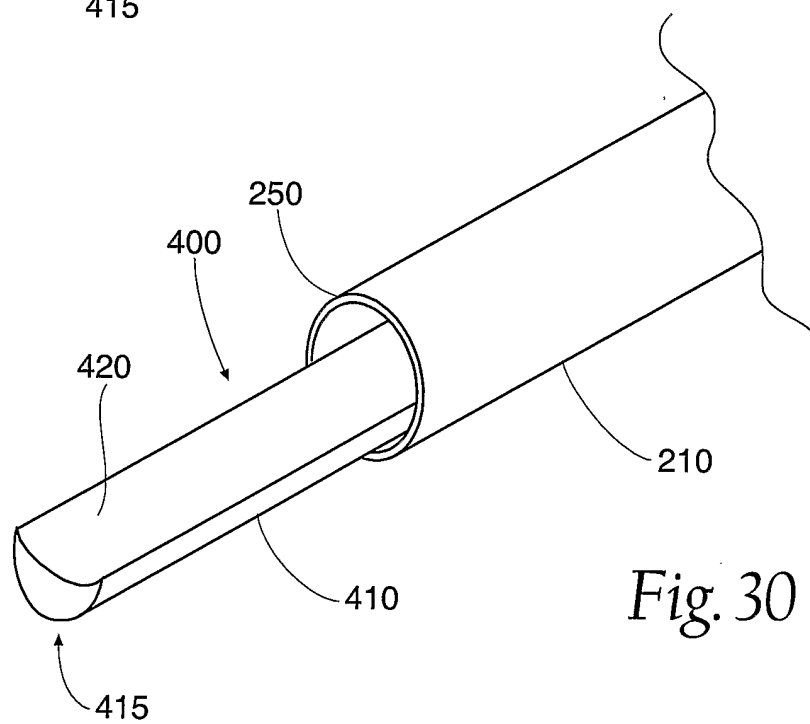
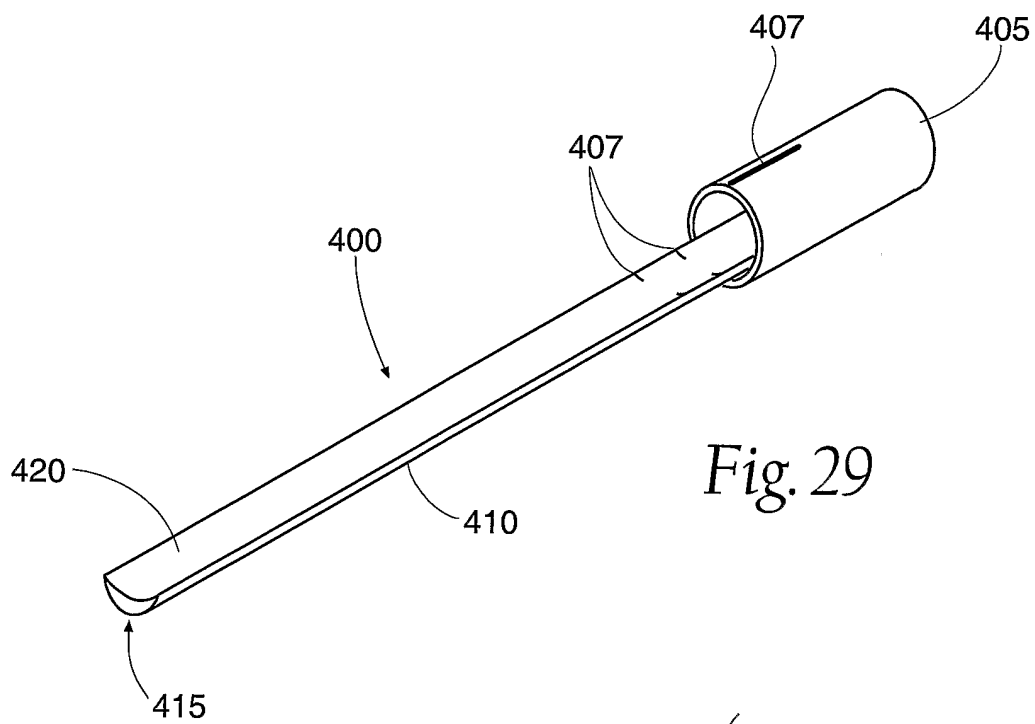


Fig. 27



*Fig. 28*



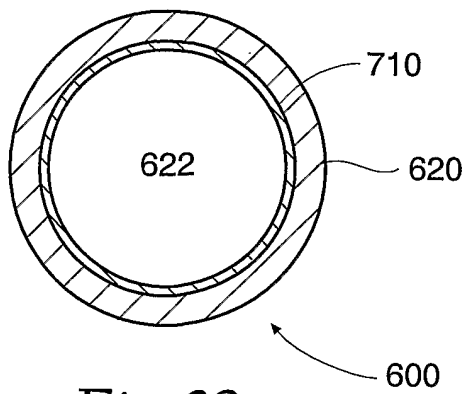


Fig. 32

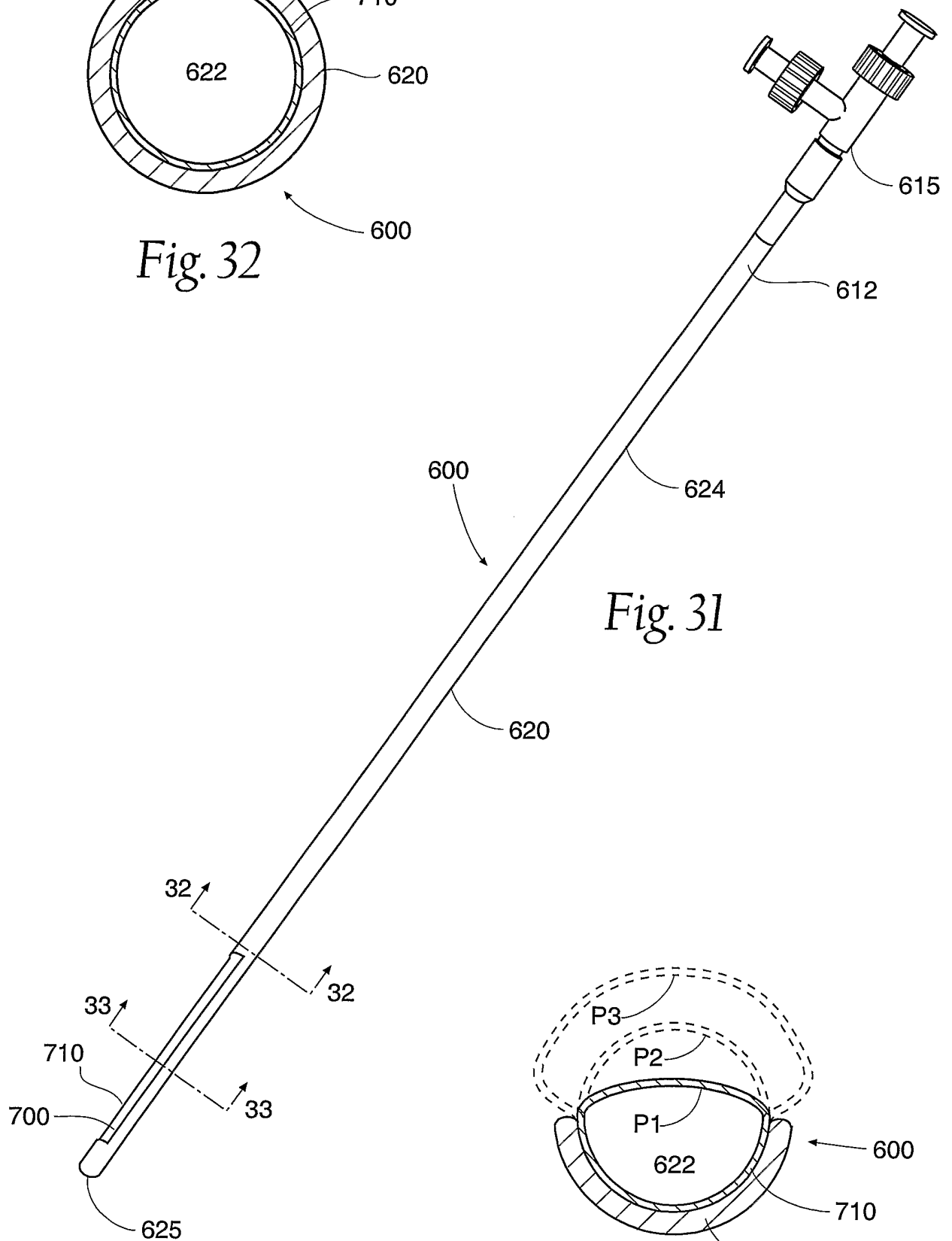


Fig. 31

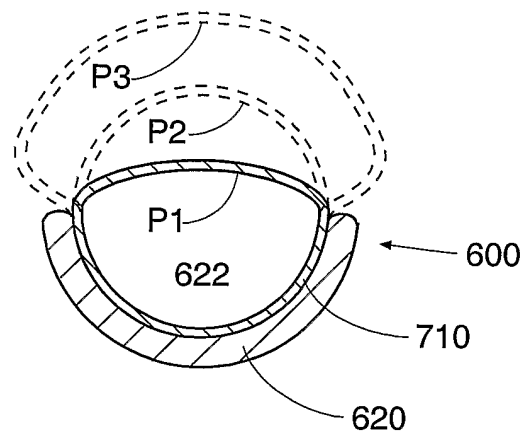


Fig. 33

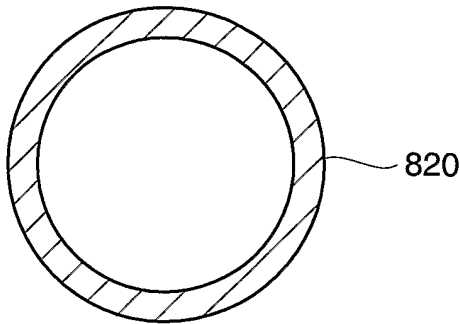


Fig. 35

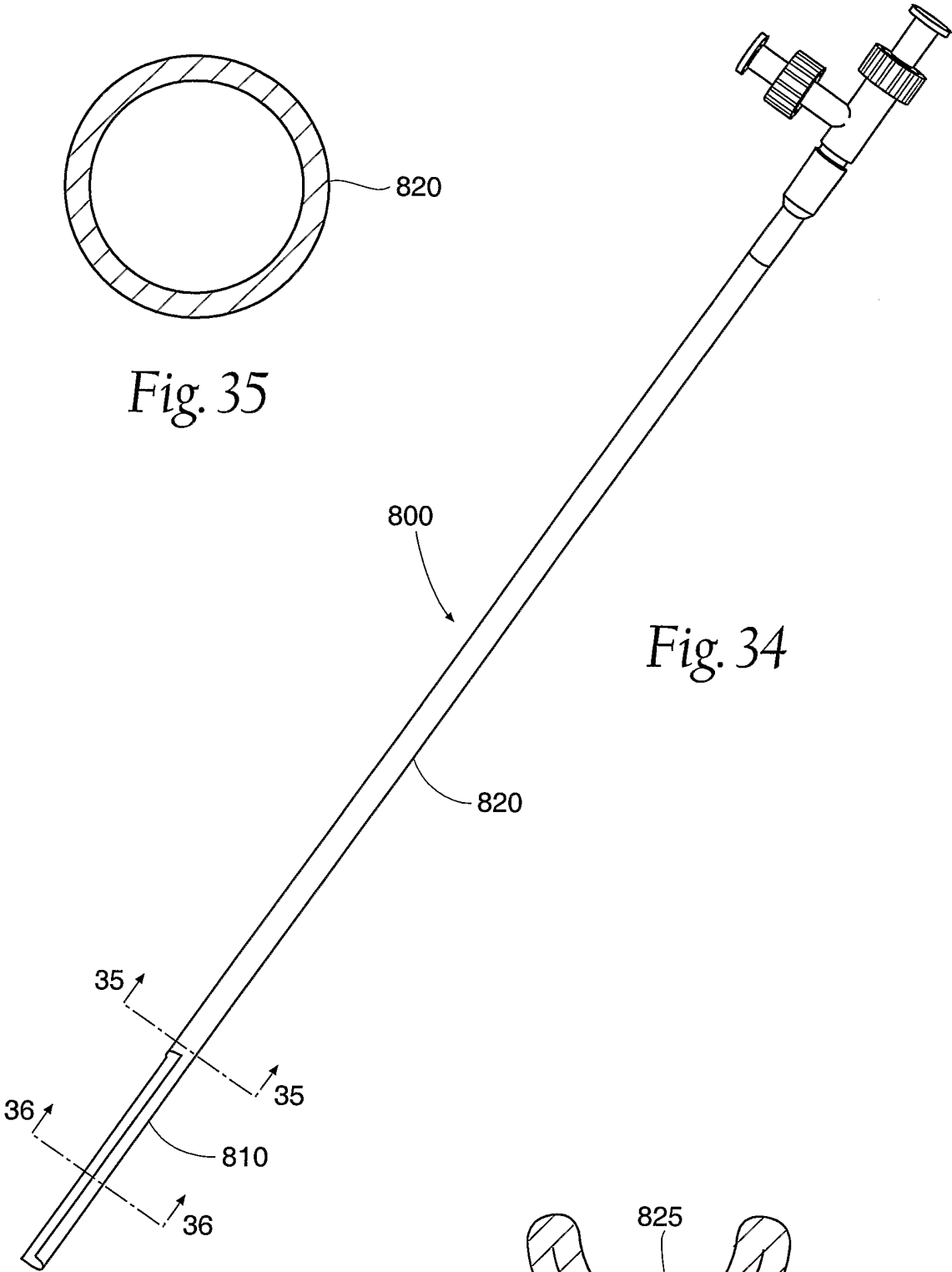


Fig. 34

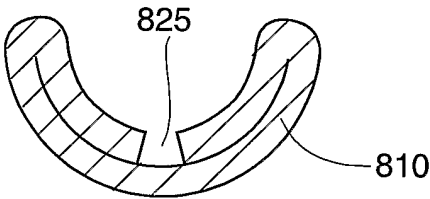


Fig. 36

Fig. 37

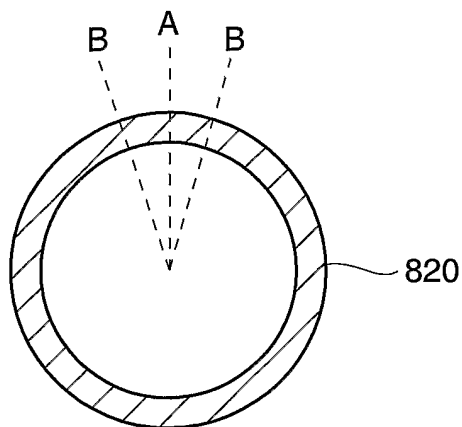


Fig. 38

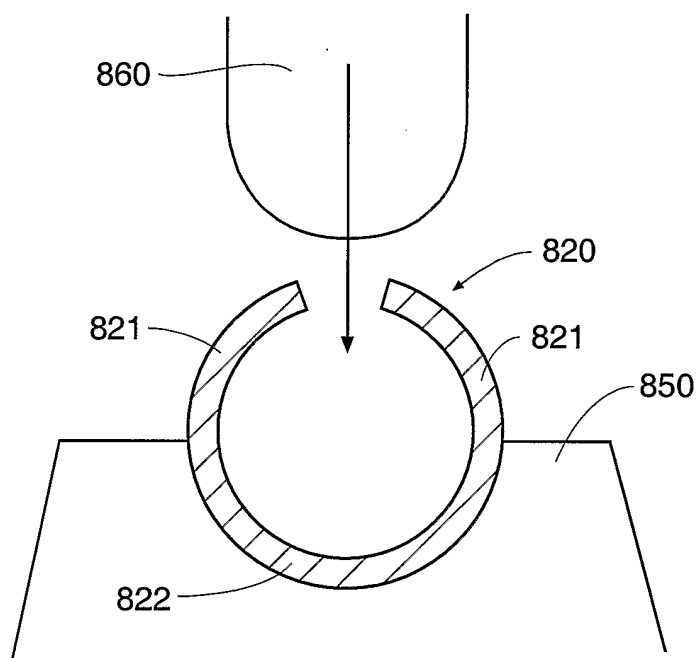
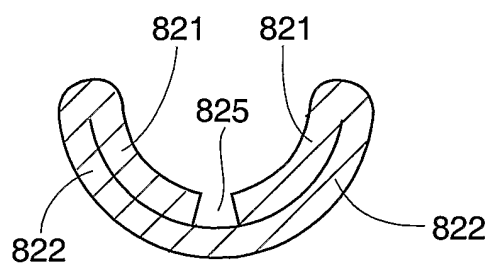
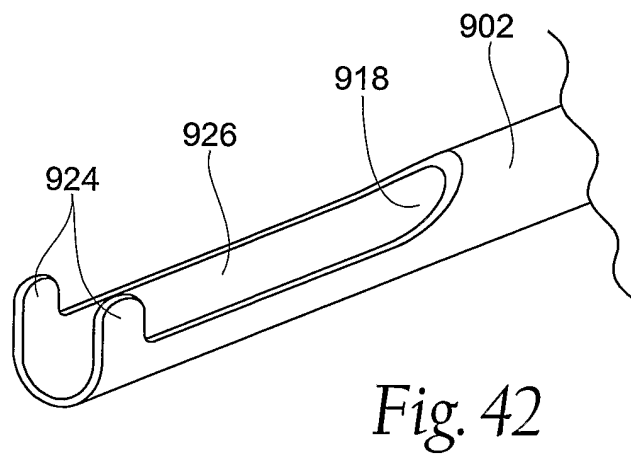
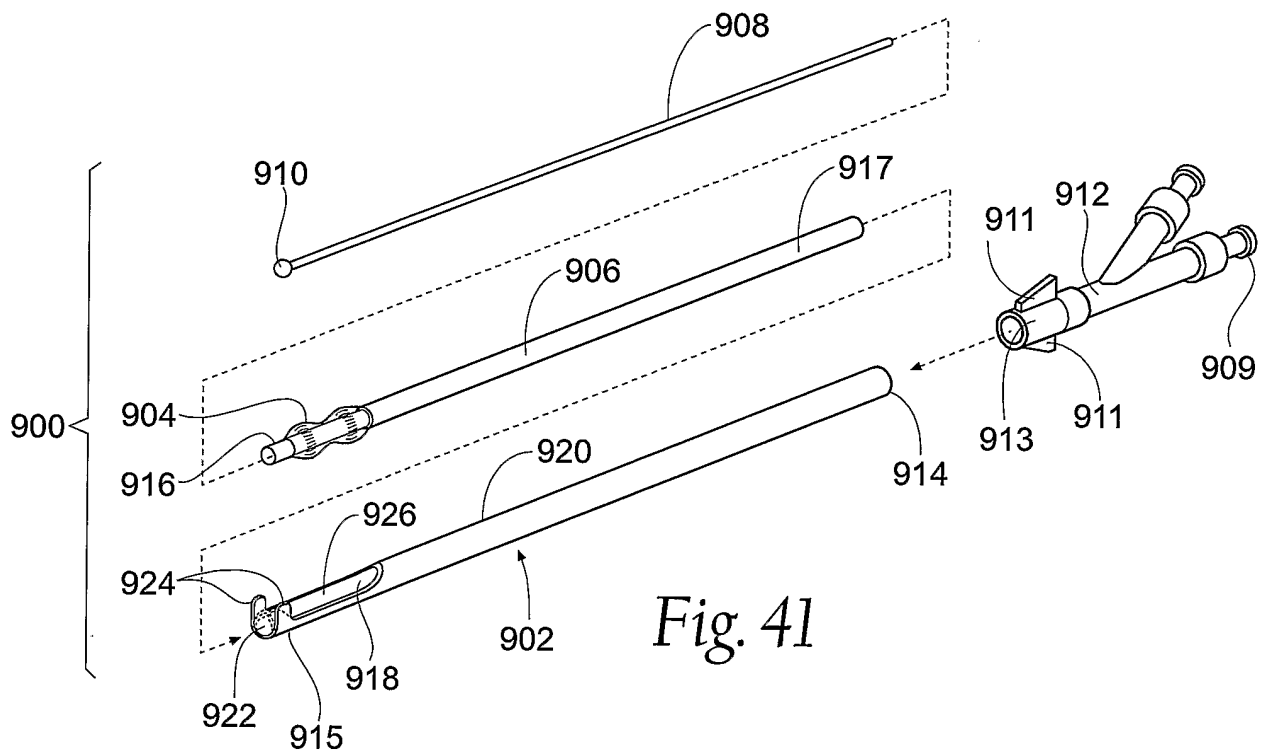
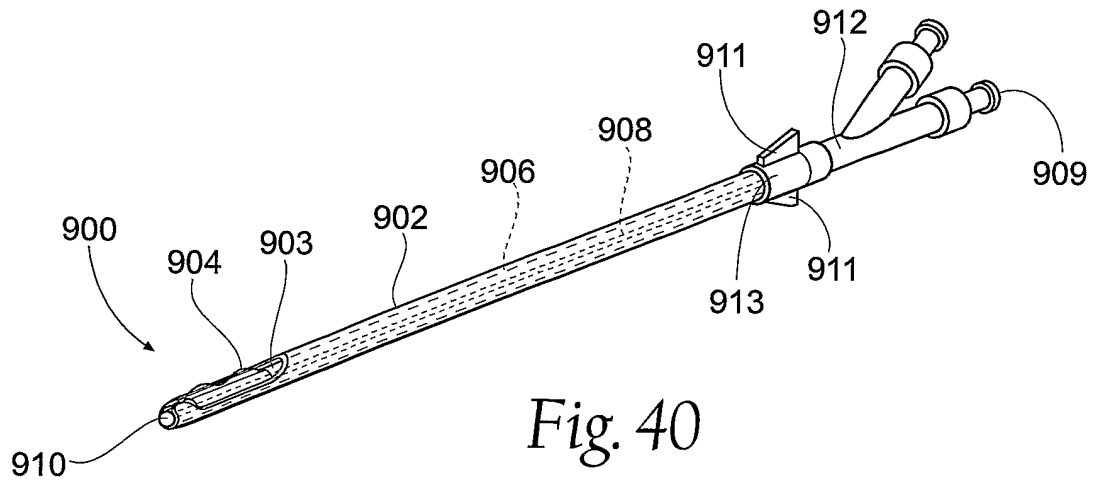
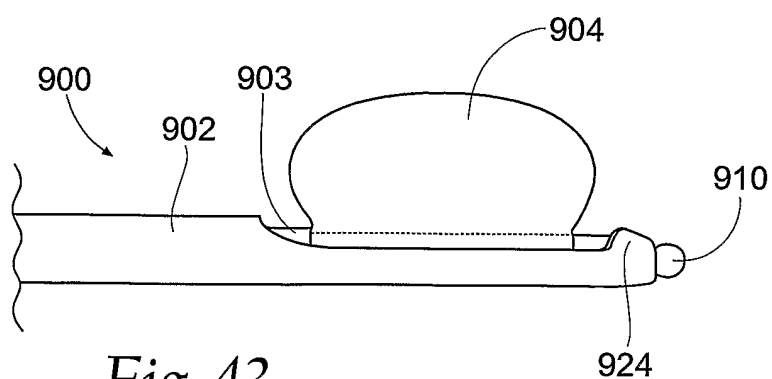
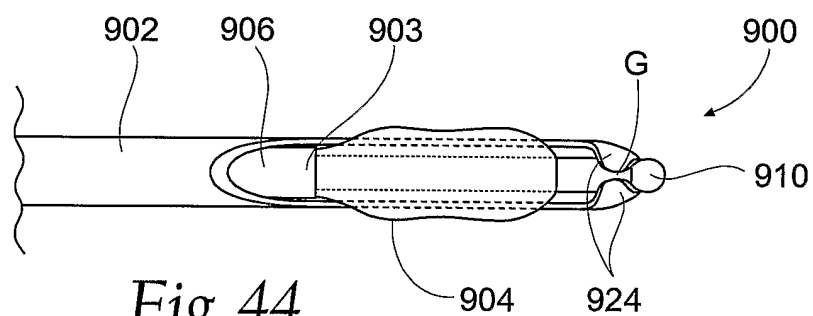
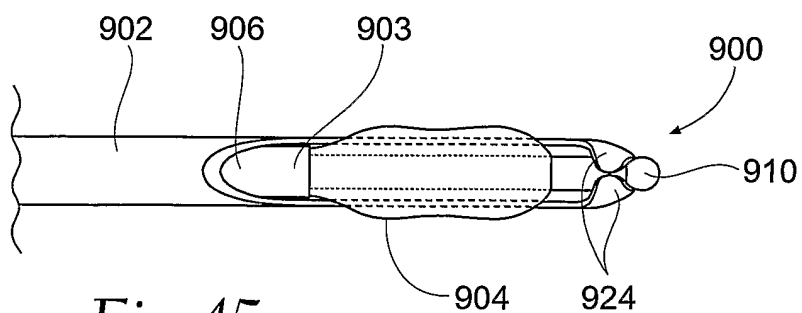


Fig. 39





*Fig. 43**Fig. 44**Fig. 45*

