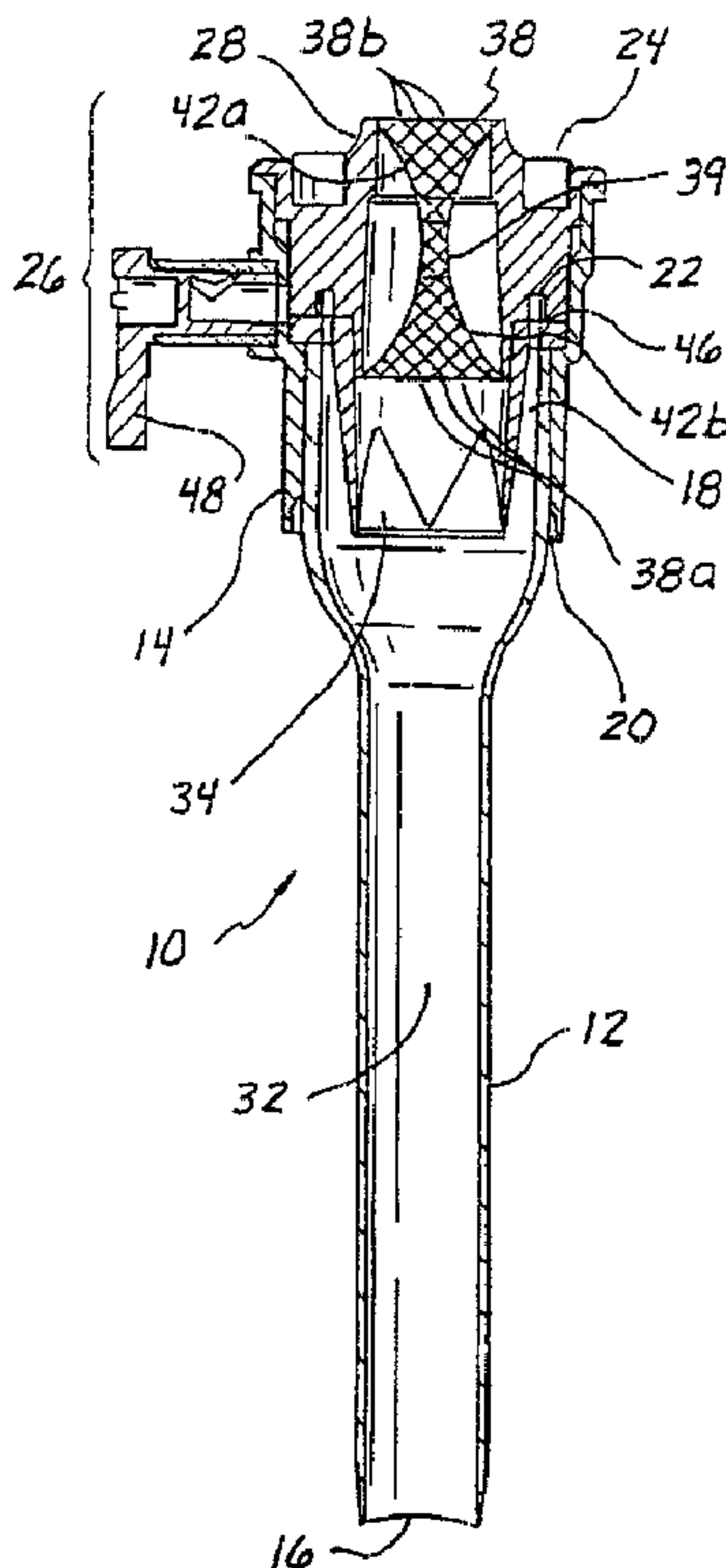




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(54) **Titre : JOINT D'ACCES UNIVERSEL**  
 (54) **Title: UNIVERSAL ACCESS SEAL**



(57) **Abrégé/Abstract:**  
 A surgical access device (10) including a valve housing (14) and a receiving element (38) mounted in the valve housing (14) having an aperture (39) for flexibly receiving and directing instruments having a wide range of diameters. The instrument receiving element

**(57) Abrégé(suite)/Abstract(continued):**

includes a braid or mesh tube (38) generally shaped like an hourglass. The surgical access device may include an access septum seal (36) molded from a gel material. The surgical access device (10) can flexibly engage instruments having diameters ranging from about 3.5 mm to about 12.9 mm.

Abstract of the Disclosure

A surgical access device (10) including a valve housing (14) and a receiving element (38) mounted in the valve housing (14) having an aperture (39) for flexibly receiving and directing instruments having a wide range of diameters.

- 5 The instrument receiving element includes a braid or mesh tube (38) generally shaped like an hourglass. The surgical access device may include an access septum seal (36) molded from a gel material. The surgical access device (10) can flexibly engage instruments having diameters ranging from about 3.5 mm to about 12.9 mm.

## UNIVERSAL ACCESS SEAL

### Background of the Invention

#### 5 Field of the Invention

The present invention generally relates to surgical access devices and, more particularly, to an access seal providing passage of instrumentation and for maintaining pneumoperitoneum during laparoscopic surgeries.

#### Description of Related Art

10 Surgical access devices, such as a trocar 10 illustrated in FIG. 1, typically include a cannula 2 and a valve housing 4 that define a working channel 6 across a body wall 7, such as an abdominal wall, and into a body cavity 8, such as an abdominal cavity. The cannula is typically formed as an elongate rigid cylinder that is inserted, with the help of an obturator, into the body cavity to  
15 provide access across the body wall.

In a typical abdominal laparoscopic surgery, the abdomen is insufflated to pressurize and thereby enlarge the cavity within which a surgical procedure is to be performed. Various instruments used in the procedure are inserted, previously one at a time, through the working channel of the trocar to perform the  
20 surgery. In order to maintain the insufflation pressure when the instrument is inserted through the trocar, a valve has been provided in the housing to form a seal around the instrument. These instrument valves have typically been

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provided in the form of septum valves. When the instrument is removed, a zero-closure valve has typically been provided to seal the trocar in order to maintain the insufflation pressure. A zero-closure valve such as a double duckbill valve disclosed in U.S. Patent No. 6,162,196, may be used.

5           Surgical instruments, however, vary in size and diameter. While the zero-closure valves can accommodate a relatively wide range of diameters, the septum valves are generally capable of stretching only a nominal amount to accommodate larger diameters. Specifically, the septum valves are generally formed in valve sets that are limited to the range of instruments that they can  
10 accommodate. When an instrument was required that had a diameter outside the range of a valve set, the entire trocar or at least the housing supporting the valve set had to be replaced with one that could accommodate the new instrument. In some cases, septum valves having universal seals were provided to accommodate different ranges of instrument diameters. These universal  
15 seals were typically made of elastic and tearable materials that often tear or puncture causing loss of insufflation gases. Attempts have also been made to include multiple septum seals to accommodate instruments having various diameters. For example, a septum valve may include one septum seal to engage large diameter instruments and another septum seal to engage smaller  
20 diameter instruments. These septum valves with multiple septum seals are typically more expensive to manufacture. Moreover, the seals are still limited to the specific range of instruments they can support.

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Accordingly, there is a need in the art for a universal access seal capable of accommodating a wide range of instrument sizes. In particular, the universal access seal should be able to sealingly engage instruments of various diameters ranging from about 3.5 mm to about 12.9 mm. An access seal covering this  
5 range of instruments would reduce adjustments and, thus, time and costs required during surgery. It is also desirable for the universal access seal to perform when a sharp instrument is inserted off-center or when an instrument is moved radially after insertion. It is further desirable that the universal access seal facilitates the insertion and removal of instruments including tissue removal.

10

#### Summary of the Invention

A surgical access device including a universal access seal is capable of accommodating instruments of various diameters. The universal access seal comprises a braid or mesh tube that is preferably shaped like an hourglass. The  
15 universal access seal can sealingly engage instruments of various diameters ranging from about 3.5 mm to about 12.9 mm. The braid facilitates insertion and manipulation of surgical instruments by directing the instruments along an axis of the surgical access device. In another embodiment of the invention, the surgical access device further includes a septum seal that is preferably molded from a  
20 gel material. The gel septum seal further facilitates the insertion and removal of instruments. These and other features and advantages of the invention will

become more apparent with the description of preferred embodiments and reference to the associated drawings.

#### Description of the Drawings

5           FIG. 1 illustrates a common surgical access device such as a trocar of the prior art;

          FIG. 2 illustrates a surgical access device in accordance with a first embodiment of the invention;

          FIG. 3 is an axial cross-section view taken along line A-A of FIG. 2;

10          FIG. 4 illustrates a surgical access device in accordance with a second embodiment of the invention;

          FIG. 5 illustrates a valve housing of the surgical access device of the invention;

          FIG. 6 is an axial cross-section view taken along line B-B of FIG. 5;

15          FIG. 7 illustrates a valve septum seal of a surgical access device in accordance with another embodiment of the invention;

          FIG. 8 is an axial cross-section view taken along line C-C of FIG. 7;

          FIG. 9 illustrates a valve septum seal of a surgical access device in accordance with another embodiment of the invention; and

20          FIG. 10 is an axial cross-section view taken along line D-D of FIG. 9.

### Description of Preferred Embodiments

The following detailed description refers to the accompanying drawings that illustrate the embodiments of the present invention. The scope of the claims should not be limited by the preferred embodiments set forth in the examples, but should be given the broadest interpretation consistent with the description as a whole. Rather the scope of the invention is defined by the appended claims.

Referring to FIGS. 2 and 3, there is shown a first embodiment of a trocar 10 including a cannula 12 and a valve housing 14. Cannula 12 defines an interior lumen having an open distal end portion 16 and an open proximal end portion 18. Proximal end portion 18 extends into and is mounted in a distal end portion 20 of valve housing 14. Valve housing 14 has an open proximal end portion 22 that defines an opening 24. An obturator can be inserted into valve housing 14 and cannula 12 through opening 24 as further described below.

Valve housing 14 includes an access port 26, which comprises a braid or mesh tube 38 having an aperture or central sealing orifice 39 adapted to receive a wide range of instrument sizes. Access port 26 further includes an end cap 28 for mounting braid 38. End cap 28 is typically disposed in a radial plane generally perpendicular to a trocar axis 30. Braid 38 includes braid elements 38b and is preferably made of polyester, which provides a low-friction, expandable lead-in to aperture 39. Braid 38 can sealingly engage instruments inserted therethrough having diameters ranging from about 3.5 mm to about 12.9

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mm. As an instrument such as an obturator is inserted into access port 26, braid 38 expands to the size of the instrument so that it forms a tight seal with the outer surface of the instrument and directs the instrument through aperture 39. By directing the instrument through aperture 39, braid 38 minimizes the possibility of tearing even if the instrument is inserted off-center or off-axis.

Braid 38 is generally shaped like an hourglass having converging and diverging sidewalls 42a and 42b, respectively, that facilitate the insertion and removal of instruments through access port 26. Braid 38 can be made from a variety of natural and synthetic monofilament thread materials including polyester, Kevlar™, carbon fiber, Gore-Tex™ (expanded PTFE), Nomex™, nylon, fiber glass, cotton, polypropylene and ceramic. Braid elements 38b, which are preferably woven from a polyester monofilament having a diameter of about 0.005", may stretch, flex, slide and/or expand in response to the direction and movement of the inserted instrument. Braid elements 38b can be made from various metal wire materials including music wire, stainless steel and Nitinol™. These materials allow greater interstitial spacings within braid elements 38b that result in less contact between the inserted instrument and braid 38 to produce a more effective seal between an elastomer and the instrument as further described below.

Braid 38 can be permanently coated or treated with a variety of materials and/or processes designed to reduce friction between inserted instruments and braid 38. The coatings may be applied on each individual braid element 38b or

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as layers over the braid elements. The layers may be external, internal or may encapsulate braid 38. The friction reducing materials include any soft or low-durometer elastomeric material. The elastomeric material could be at least one of a thermoplastic and a thermoset. Examples of the elastomeric materials include silicone, polyurethane, polyisoprene and Kraton™. Examples of other coatings and treatments include hydrophilic polymer coatings, Teflon™ (PTFE) coatings, cyanoacrylate coatings, Parylene™ coatings, plasma surface treatments and chlorination treatments.

In a second embodiment of the invention as illustrated in FIG. 4, access port 26 further includes a septum seal 36 having an aperture 40. Aperture 40 measures about 0.115" in diameter and is in line with trocar axis 30. Braid 38 is configured to line aperture 40 of septum seal 36. Septum seal 36 may also stretch, flex, swivel and/or slide to receive the inserted instrument. Septum seal 36 may be configured to float within valve housing 14 to minimize the cat-eye effect around the inserted instrument, which can result in seal leakage during manipulation of the instrument. In another embodiment of the invention, valve housing 14 further includes finger tabs 11a and 11b providing means for engaging trocar 10 and manipulating cannula 12 into a preferred operative position. In another embodiment of the invention, access port 26 is configured as a hand-access port to allow passage of a surgeon's hand or finger into the peritoneal cavity of a patient. In another embodiment of the invention, access port 26 is utilized as a hemostasis valve for vascular or cardiovascular surgeries

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to prevent loss of blood yet allows passage of guidewires, catheters and other devices into the arterial or venous system of a patient. In yet another embodiment of the invention, access port 26 is utilized as an endoscopic valve for urological procedures to prevent loss of fluids yet allows passage of  
5 guidewires, catheters and other devices into the urethra or ureter.

Septum seal 36 is preferably molded from a gel material and is preferably encased in a seal housing to affect a radial compressive force about the outside diameter of septum seal 36. As an instrument is inserted into braid 38, braid 38 expands while septum seal 36 resists expansion due to the outer compressive  
10 force provided by the seal housing. This forces the gel material to extrude through interstitial spaces 38a to sealingly engage the outside diameter of any instrument inserted through access port 26 while minimizing the frictional contact between the inserted instrument and septum seal 36. The gel material has a low durometer that enables it to extrude through interstitial spaces 38a. The gel  
15 material is preferably a composite material comprising mineral oil and a thermoplastic elastomer such as a Kraton™ material.

Septum seal 36 could also be manufactured from a closed cell foam material or an open cell foam material sealed with a film coating. Examples of the foamed materials include silicone, urethane, Kraton™, polyethylene,  
20 polyisoprene, polyvinylchloride (PVC), polyurethane, ethylene propylene diene monomer (EPDM), Neoprene™ and styrene butadiene (SBR). Septum seal 36 may be coated or treated with a variety of materials and/or processes designed

to reduce friction between the inserted instruments and the gel material.

Examples include hydrophilic polymer coatings, Teflon™ (PTFE) coatings, thermoplastic coatings, cyanoacrylate coatings, Parylene coatings, plasma surface treatments, cornstarch powder coatings and chlorination treatments.

5 Septum seal 36 may also be lubricated with a variety of materials to facilitate the insertion and withdrawal of instruments. Examples of these materials include silicone oil, silicone grease, liquid soaps, Astroglide™ lubricants, mineral oil, glycerin, alcohol, saline, Teflon™ (PTFE) lubricants, Krytox™ lubricants, molybdenum disulfide lubricants and graphite.

10 Another aspect of the invention is braid elements 38b also serve to reduce the force required to insert and advance an instrument through septum seal 36. The coefficient of kinetic friction ( $f$ ) for polyester braid 38/septum seal 36 verses a metal or polymer instrument shaft is significantly less than that of an elastomeric septum seal verses a metal or polymer instrument shaft. Typical  
15 coefficient of kinetic friction values range from about 0.15 to about 0.5 for polymers such as polyester verses steel, whereas the typical coefficient of kinetic friction values for elastomers verses steel range from about 1.6 to about 10. As a result, braid elements 38b minimize the contact between the shaft of the inserted instrument and septum seal 36 and minimize the frictional forces  
20 required to insert and advance the instrument through septum seal 36.

Braid elements 38b also serve to capture lubricants such as oils and greases within interstitial spaces 38a. In particular, interstitial spaces 38a

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capture lubricants to prevent inserting instruments from wiping all of the lubricants from braid 38 and septum seal 36 during instrument exchanges. That is, some lubricant will always be present within braid elements 38b to facilitate manipulation and exchange of instruments throughout the surgical procedure.

5 The presence of lubricants also improves the sealing properties of the present invention. As observed with prior art trocar seals, lubricants such as oils and greases are typically completely transferred from the lubricated septum seals to the inserted instruments after a few instrument exchanges resulting in a non-lubricated septum seal for the remainder of the surgical procedure. A drawback  
10 of the prior art seals is subsequent instrument manipulations and exchanges become increasingly difficult for the operating surgeon or user.

With the flexibility of the braid and septum seal of the invention, an instrument having a sharp, irregular, forked or otherwise potentially damaging distal features may be directed through the access port in a minimally  
15 threatening position. The braid, either alone or in combination with the septum seal, can stretch, flex, slide and/or expand so as to easily receive an approaching instrument. The flexibility of the braid and septum seal thus provides a very durable and relatively friction-free insertion and removal of instrumentation.

20 Access port 26 may further comprise a zero-closure valve 34 such as a double duckbill valve, which maintains pneumoperitoneum in the absence of inserted instrumentation as described in U.S. Patent No.

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6,162,196. With this embodiment, valve housing 14 and cannula 12 extend along trocar axis 30 and define a working channel 32 for receipt of a surgical instrument. In the absence of an instrument, zero-closure valve 34 closes on itself forming a gas-tight seal at very low retrograde pressure and preventing loss of insufflation gas. When an instrument is present in working channel 32, braid 38 and/or septum seal 36 forms a seal with the instrument in order to seal working channel 32. In particular, access port 26 provides a positive seal with respect to instruments inserted therethrough.

Access port 26 may further comprise a manifold 46 and a stopcock 48, both of which are preferably molded from polycarbonate. Manifold 46 is positioned within valve housing 14 and serves to locate zero-closure valve 34 and septum seal 36 relative to valve housing 14. Manifold 46 also facilitates the flow of insufflation gasses from an insufflator, through cannula 12 and into the surgical site. End cap 28 is preferably ultrasonically welded to valve housing 14 and serves to fix manifold 46, zero-closure valve 34, septum seal 36 and braid 38 within valve housing 14. The proximal end of braid 38 is preferably bonded to end cap 28 and the distal end of braid 38 is not attached to any component and is free to float within access port 26. Braid 38 is preferably flared at both the proximal end and the distal end in an hourglass shape. In another embodiment of the invention, the proximal end of braid 38 is bonded or fixed to manifold 46 to prevent migration during instrument insertion and removal. Zero-closure valve 34 is preferably molded from polyisoprene and is located distal to septum seal

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36. As illustrated in FIGS. 5 and 6, valve housing 14 including access port 26 may be removably attached to a disposable or reusable cannula 12.

The following describes the preferred method of manufacturing an access port in accordance with an embodiment of the invention. Zero-closure valve 34, which is preferably transfer molded, is first placed into valve housing 14, which is preferably injection molded. Manifold 46, which is preferably injection molded, is then mounted on top of zero-closure valve 34. Septum seal 36, which is preferably injection molded, is then placed on top of manifold 46. Braid 38, which is preferably heat set to form flared sections at its proximal and distal ends, is then bonded to end cap 28, which is preferably injection molded. The distal end of braid 38 is then threaded through aperture 40 of septum seal 36, and end cap 28 is positioned on top of valve housing 14 to effectively capture all of the access port components. End cap 28 is then ultrasonically welded to valve housing 14. Stopcock 48 is then bonded to valve housing 14. The housing assembly is then removably attached, via bayonet locks, to cannula 12, which is preferably injection molded from polycarbonate.

In another embodiment of the invention, FIGS. 7 and 8 illustrate a septum seal 60 comprising a plurality of cored sections 60a-60d spaced about a central aperture 62 that enable septum seal 60 to deform in response to insertion of an instrument. Cored sections 60a-60d may be formed in either or both of the top and bottom surfaces of septum seal 60 as illustrated in cross-section view C-C in FIG. 8. In yet another embodiment of the invention, FIGS. 9 and 10 illustrate a

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septum seal 80 comprising a plurality of small apertures 82a and 82b connected to pockets 88a and 88b, respectively. Small apertures 82a and 82b are in fluid communication with and are oriented perpendicular to instrument insertion aperture 86. Pockets 88a and 88b are located within septum seal 80 and serve to store lubricants. When an instrument is inserted through aperture 86, septum seal 80 deforms forcing the lubricants to ooze from storage pockets 88a and 88b through apertures 82a and 82b, respectively, thereby lubricating the instrument and facilitating its insertion and manipulation.

It is appreciated that various fillers and additives could be incorporated into the various elastomeric septum seal materials to reduce the tackiness and to increase the lubricity of the material thereby facilitating the insertion and removal of instruments. Examples of the additives include waxes, soaps, paraffin wax, beeswax, calcium stearate, stearic acid, silicone oil, silicone grease, mineral oil, glycerin, graphite, silica, glass spheres, Teflon™ (PTFE), Parylene, talc and molybdenum disulfide.

The scope of the claims should not be limited by particular embodiments set forth herein, but should be construed in a manner consistent with the description as a whole.

20

**CLAIMS:**

1. A surgical access device operatively connected to a cannula having an axis extending between a proximal end and a distal end, comprising:  
a valve housing disposed at the proximal end of the cannula; and  
an instrument receiving element mounted in the valve housing having an aperture for flexibly receiving and directing instruments having a wide range of diameters; and  
an access septum seal disposed in the valve housing having a septum seal aperture, the instrument receiving element extending through the septum seal aperture;  
wherein the instrument receiving element includes a tubular braid.
2. The surgical access device recited in Claim 1 wherein the braid is coated with a material for reducing friction.
3. The surgical access device recited in Claim 1 wherein the braid is treated with at least one of plasma surface treatment and chlorination treatment for reducing friction during passage of instrumentation.
4. The surgical access device recited in Claim 2 wherein the coated braid expands to form a seal around an outer surface of an instrument.
5. The surgical access device recited in claim 1 wherein the tubular braid contacts the septum seal aperture.
6. The surgical access device recited in Claim 1 wherein the access septum seal moves along one or more axes in response to movement of an inserted instrument.
7. A surgical access device operatively connected to a cannula having an axis extending between a proximal end and a distal end, comprising:  
a valve housing disposed at the proximal end of the cannula; and

an instrument receiving element mounted in the valve housing having an aperture for flexibly receiving and directing instruments having a wide range of diameters; and  
an access septum seal disposed in the valve housing having a septum seal aperture, the instrument receiving element extending through the septum seal aperture;  
wherein the instrument receiving element includes a mesh tube.

8. The surgical access device recited in Claim 7 wherein the mesh tube contacts the septum seal aperture.

9. The surgical access device recited in Claims 1 or 7 wherein the instrument receiving element is generally shaped like an hourglass.

10. The surgical access device recited in Claims 1 or 7 wherein the instrument receiving element includes a converging sidewall facilitating insertion of an instrument and a diverging sidewall facilitating removal of the instrument.

11. The surgical access device recited in Claims 1 or 7 further comprising a zero-closure valve disposed in the valve housing.

12. The surgical access device recited in Claims 1 or 7 further comprising an end cap for mounting the instrument receiving element.

13. The surgical access device recited in Claims 1 or 7 wherein the access septum seal further includes additives including at least one of wax, soap, paraffin wax, beeswax, calcium stearate, stearic acid, silicone oil, silicone grease, mineral oil, glycerin, graphite, silica, glass spheres, PTFE (Teflon™), elastomer poly(*p*-xylylene) (Parylene™), talc, and molybdenum disulfide to reduce tackiness and increase lubricity.

14. The surgical access device recited in Claims 1 or 7 wherein the access septum seal is molded from a gel material extruding through interstitial spaces of the

instrument receiving element and sealingly engaging an instrument while minimizing frictional contact between the instrument and the access septum seal.

15. A surgical access device operatively connected to a cannula having an axis extending between a proximal end and a distal end, comprising:  
a valve housing disposed at the proximal end of the cannula; and  
an instrument receiving element mounted in the valve housing having an aperture for flexibly receiving and directing instruments having a wide range of diameters; and  
an access septum seal disposed in the valve housing having a septum seal aperture, the instrument receiving element extending through the septum seal aperture;  
wherein the access septum seal is molded from a gel material extruding through interstitial spaces of the instrument receiving element and sealingly engaging an instrument while minimizing frictional contact between the instrument and the access septum seal.

16. The surgical access device recited in Claims 14 or 15 wherein the gel material is a composite material comprising mineral oil and a thermoplastic elastomer.

17. The surgical access device recited in Claims 14 or 15 wherein the gel material comprises a low-durometer elastomeric material that is at least one of a thermoplastic and a thermoset.

18. The surgical access device recited in Claim 17 wherein the elastomeric material includes at least one of silicone, polyurethane, polyisoprene and styrenic block copolymer (Kraton™).

19. The surgical access device recited in Claims 14 or 15 wherein the gel material comprises a closed cell foam material or an open cell foam material.

20. The surgical access device recited in Claim 19 wherein the foam material includes at least one of silicone, urethane, styrenic block copolymer (Kraton™),

polyethylene, polyisoprene, polyvinylchloride (PVC), polyurethane, ethylene propylene diene monomer (EPDM), polychloroprene (Neoprene™), and styrene butadiene (SBR).

21. The surgical access device recited in Claims 1, 7, 14 or 15 wherein the access septum seal comprises cored sections spaced about the septum seal aperture enabling the access septum seal to deform during insertion and removal of an instrument.

22. A surgical access device operatively connected to a cannula having an axis extending between a proximal end and a distal end, comprising:

a valve housing disposed at the proximal end of the cannula; and

an instrument receiving element mounted in the valve housing having an aperture for flexibly receiving and directing instruments having a wide range of diameters; and

an access septum seal disposed in the valve housing having a septum seal aperture, the instrument receiving element extending through the septum seal aperture;

wherein the access septum seal comprises cored sections spaced about the septum seal aperture enabling the access septum seal to deform during insertion and removal of an instrument.

23. The surgical access device recited in Claims 21 or 22 wherein the access septum seal comprises pockets storing lubricants and lubricant releasing apertures connecting the pockets to the septum seal aperture.

24. The surgical access device recited in Claim 23 wherein the lubricant releasing apertures are oriented perpendicular to the septum seal aperture.

25. The surgical access device recited in Claim 23 wherein the access septum seal deforms forcing the lubricants to ooze from the pockets through the lubricant releasing apertures thereby lubricating the instrument and facilitating instrument insertion and manipulation.

26. The surgical access device recited in Claims 1, 7, 14, 15, 21 or 22 wherein a distal most portion of the instrument receiving element is in a non-contacting relationship with the access septum seal.

27. The surgical access device recited in Claims 1, 7, 14, 15, 21 or 22 wherein a distal most portion of the instrument receiving element is positioned axially away from the septum seal aperture.

28. The surgical access device recited in Claims 1, 7, 14, 15, 21 or 22 wherein the septum seal aperture is increased by the instrument receiving element engaging the septum seal.

29. A surgical access device operatively connected to a cannula having an axis extending between a proximal end and a distal end, comprising:

a valve housing disposed at the proximal end of the cannula; and

an instrument receiving element mounted in the valve housing having an aperture for flexibly receiving and directing instruments having a wide range of diameters; and

an access septum seal disposed in the valve housing having a septum seal aperture, the instrument receiving element extending through the septum seal aperture;

wherein the septum seal aperture is increased by the instrument receiving element engaging the septum seal.

30. The surgical access device recited in Claims 1, 7, 14, 15, 21 or 22 wherein the septum seal aperture is deflected in an axial direction by the instrument receiving element engaging the septum seal.

31. A surgical access device operatively connected to a cannula having an axis extending between a proximal end and a distal end, comprising:

a valve housing disposed at the proximal end of the cannula; and

an instrument receiving element mounted in the valve housing having an aperture for flexibly receiving and directing instruments having a wide range of diameters; and

an access septum seal disposed in the valve housing having a septum seal aperture, the instrument receiving element extending through the septum seal aperture;  
wherein the septum seal aperture is deflected in an axial direction by the instrument receiving element engaging the septum seal.

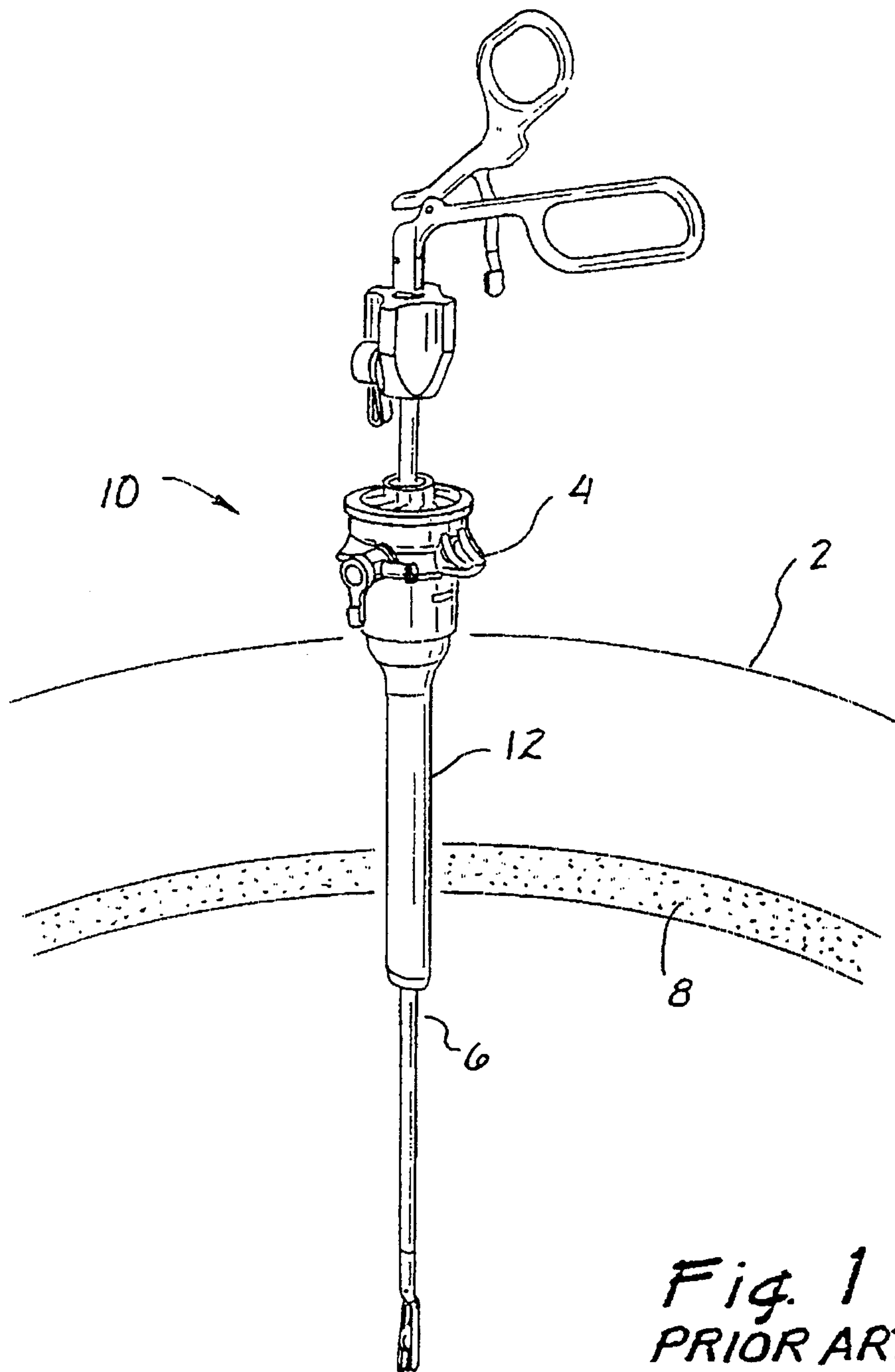


Fig. 1  
PRIOR ART

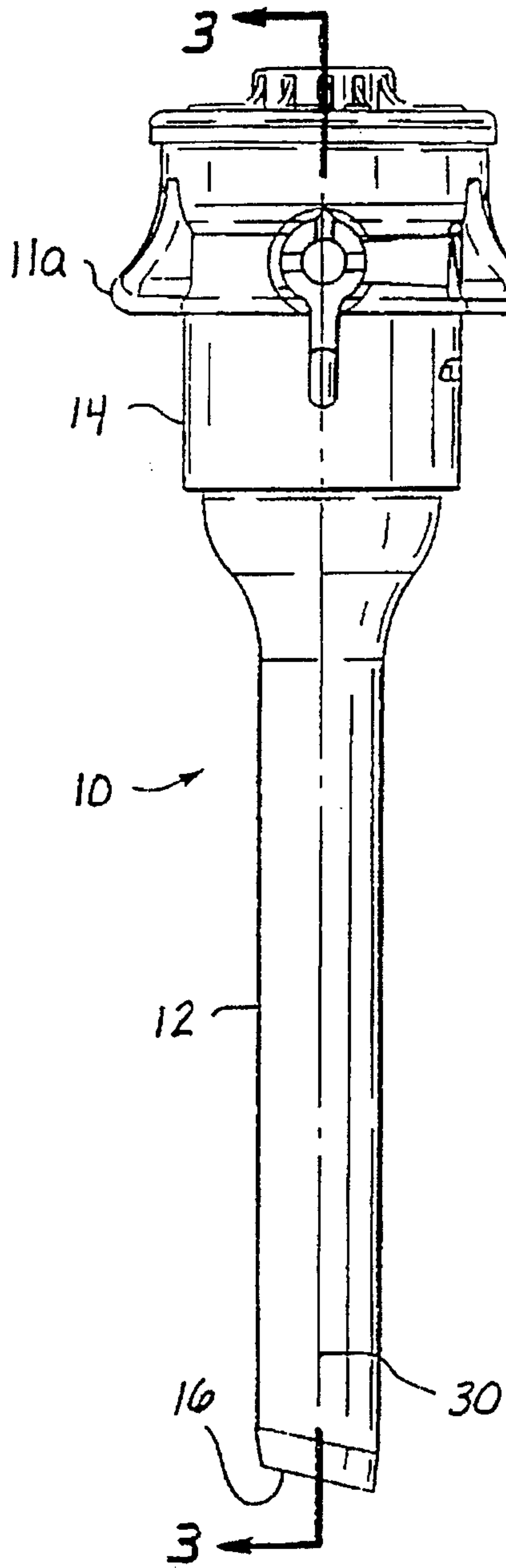


Fig. 2

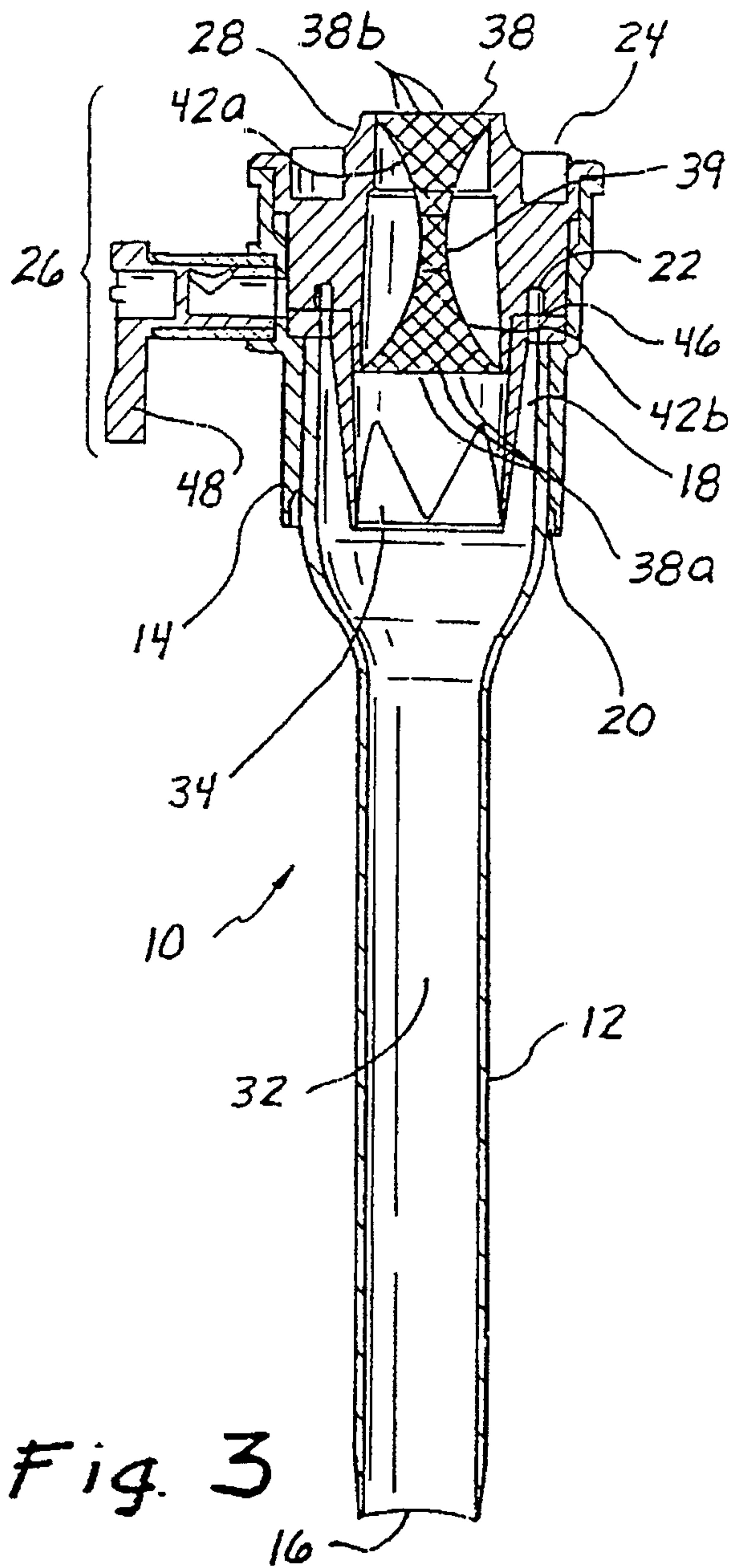


Fig. 3

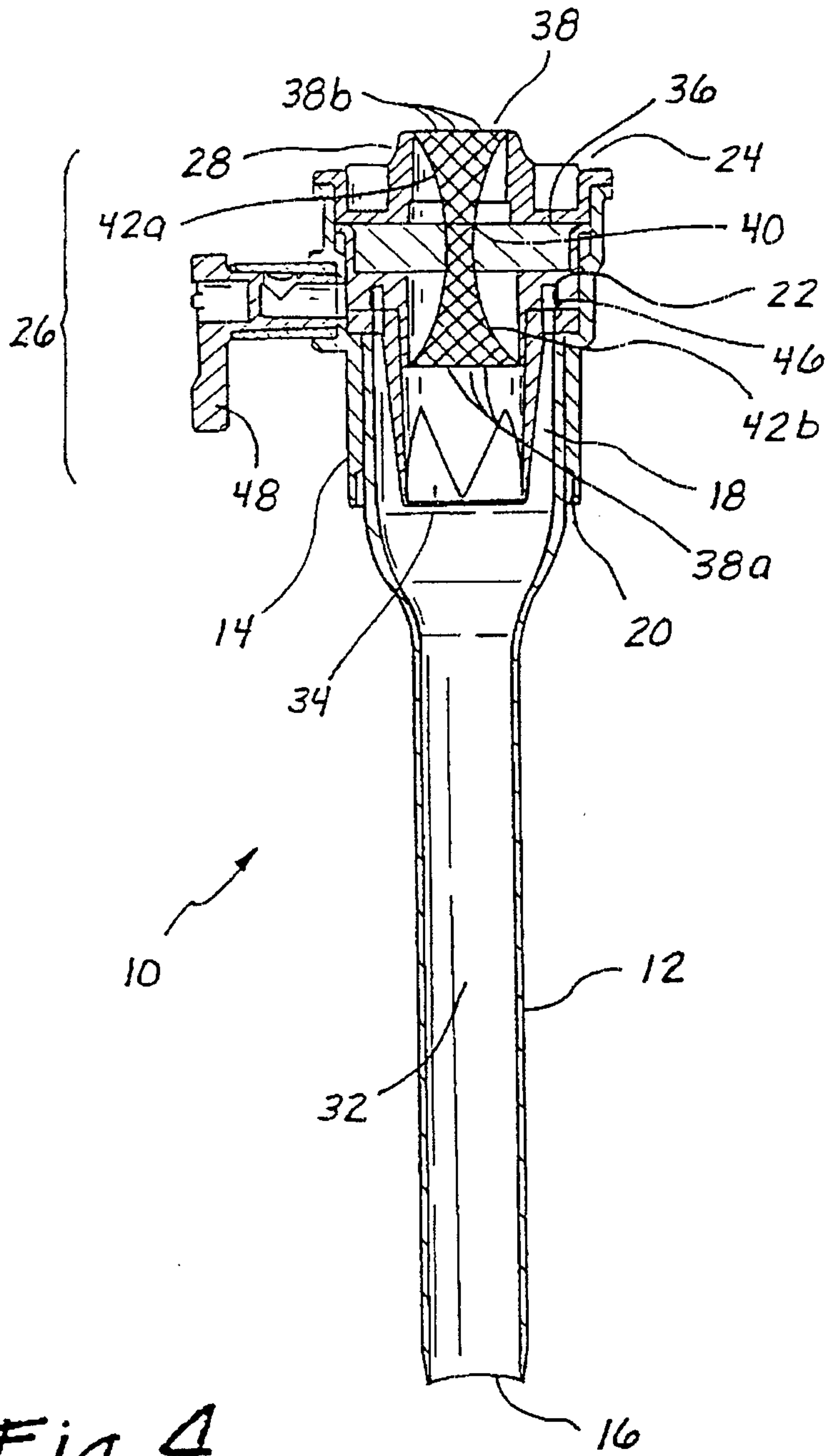


Fig. 4

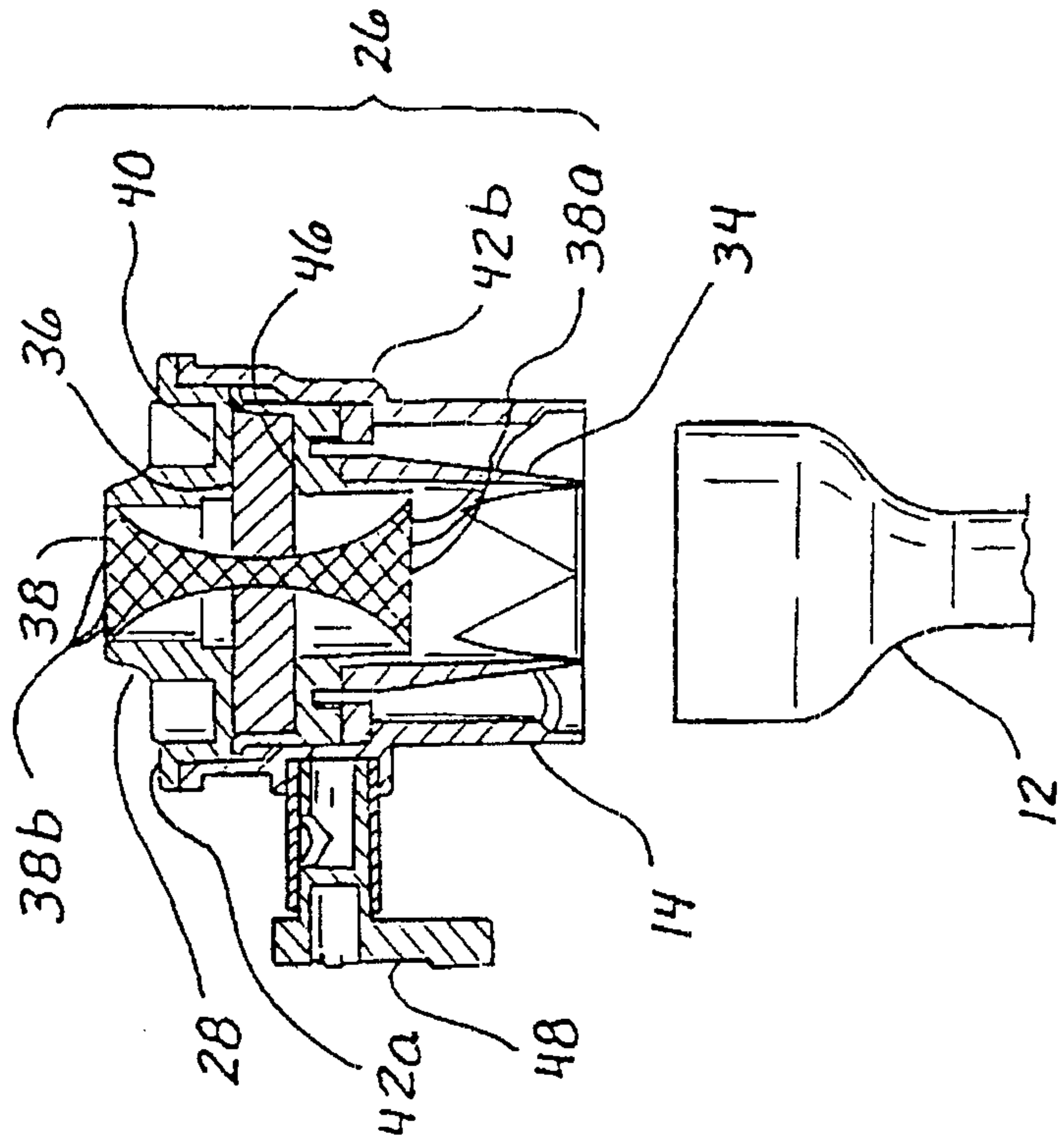
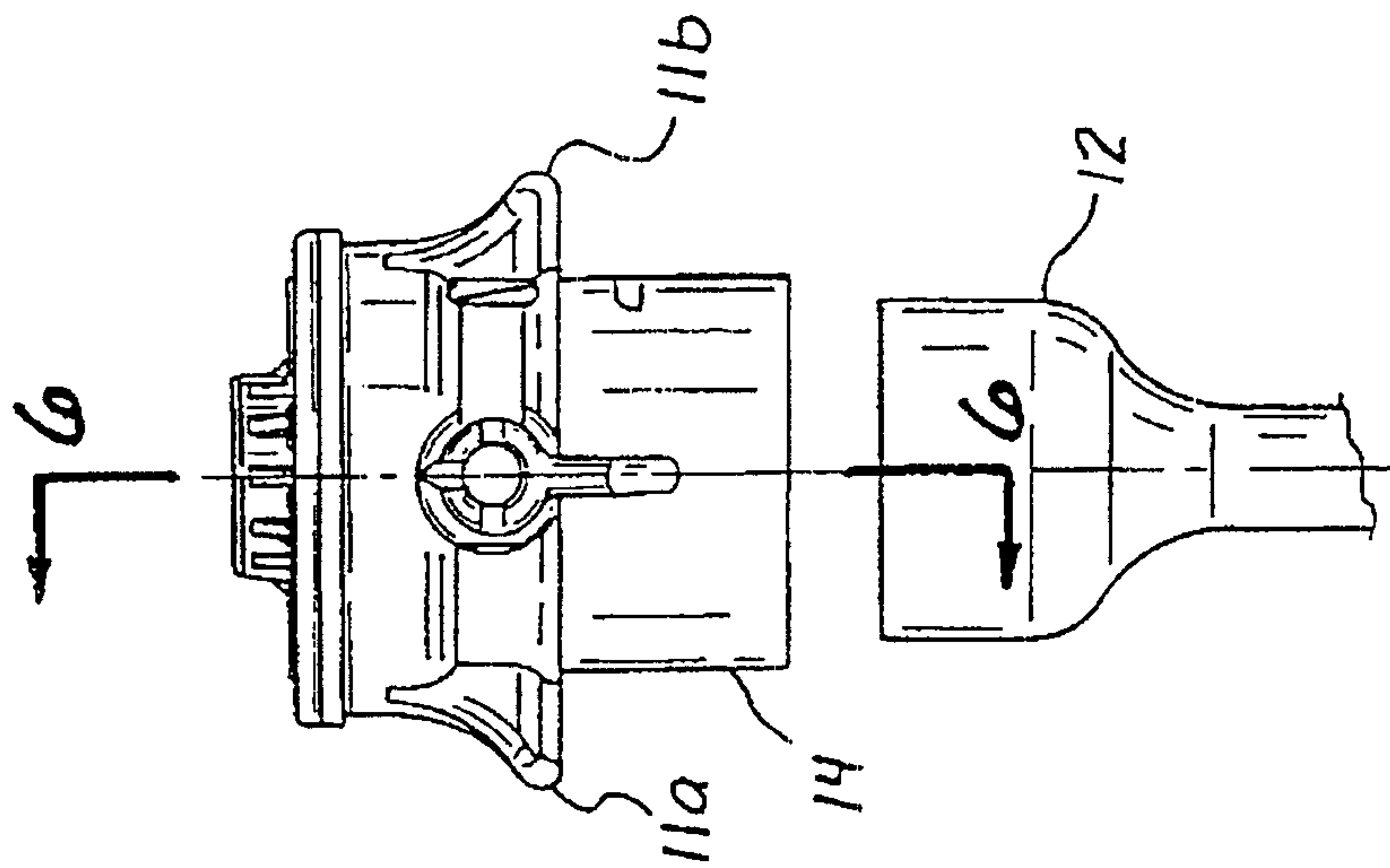


Fig. 5

Fig. 6

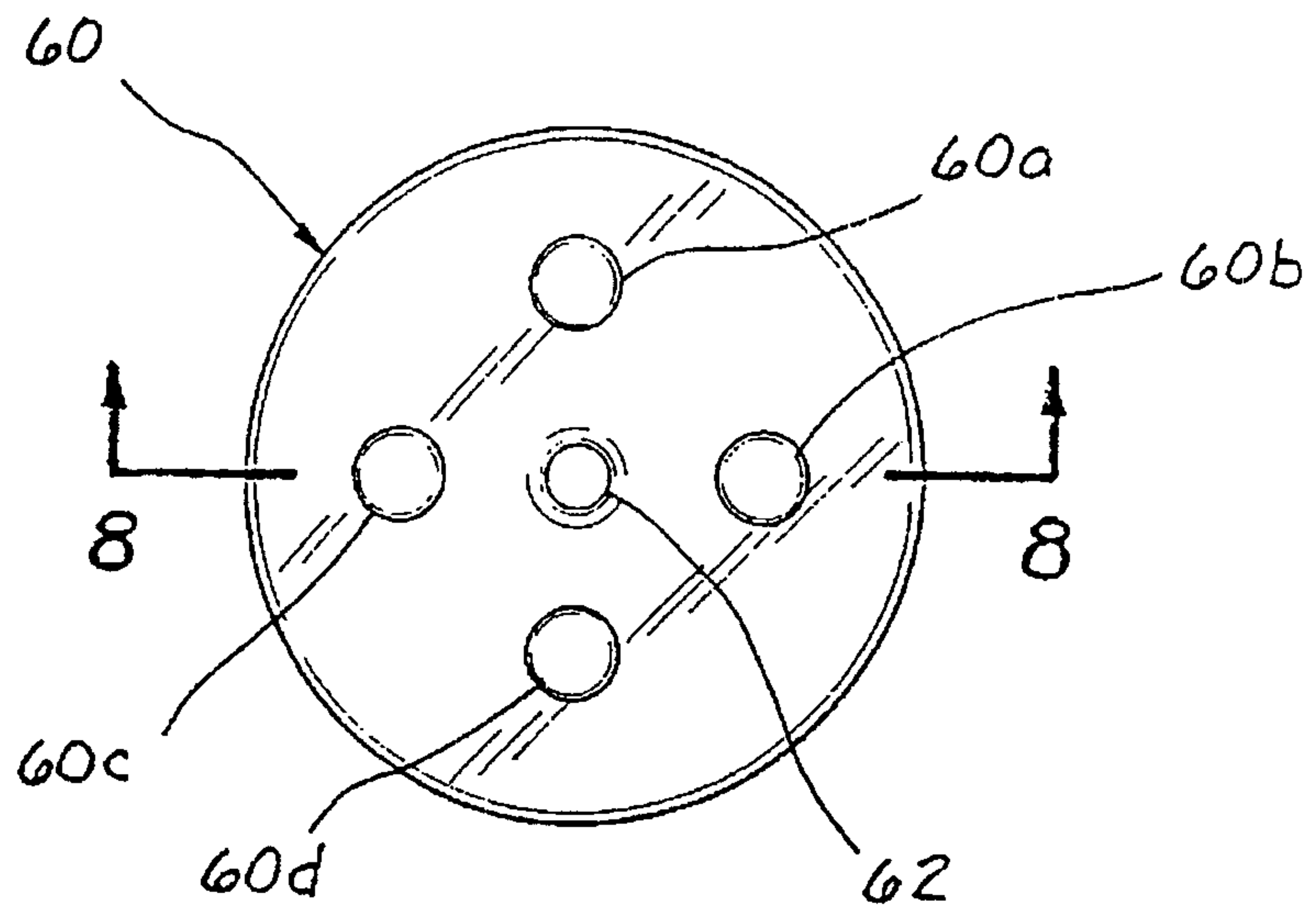


Fig. 7

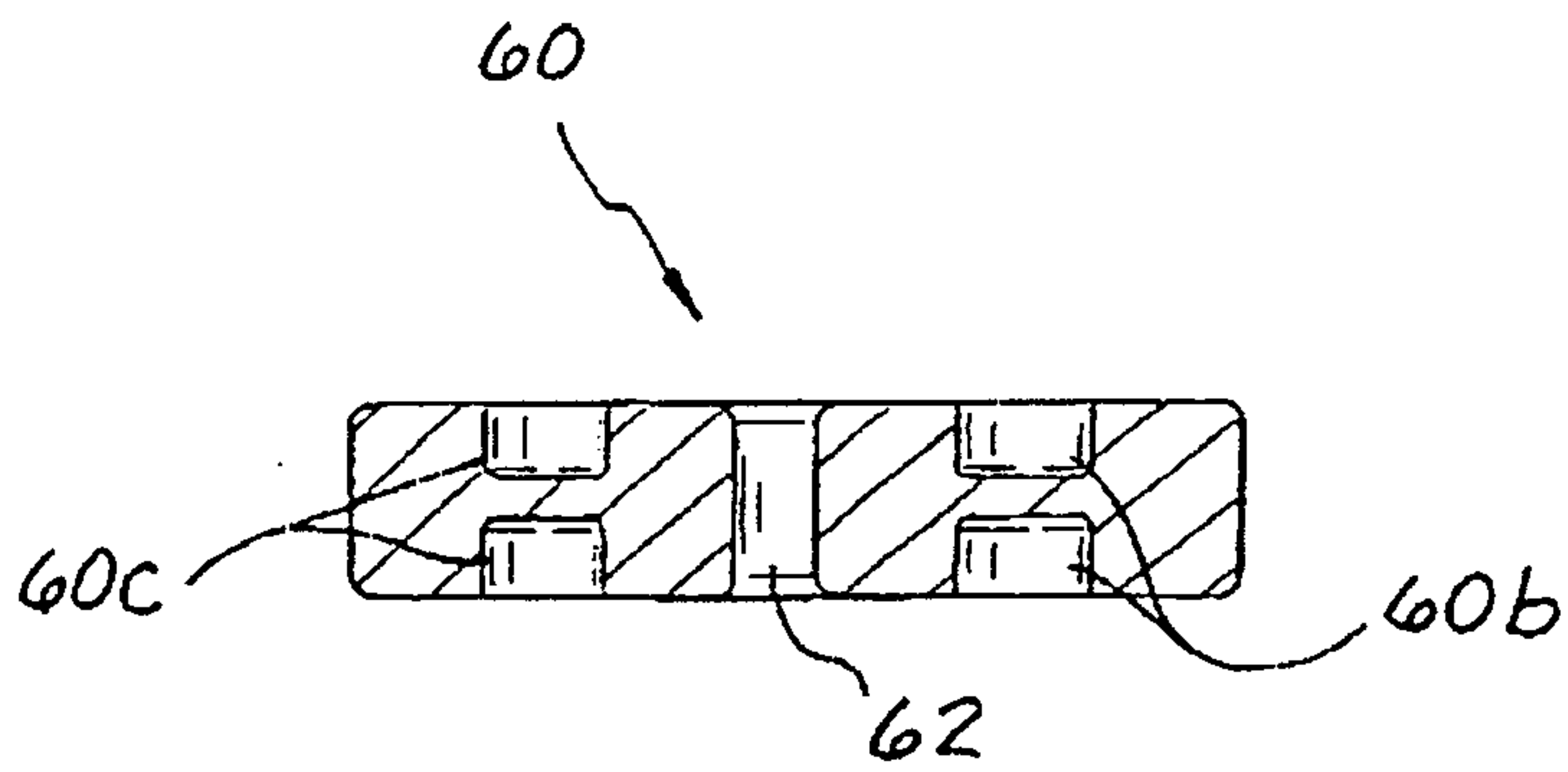


Fig. 8

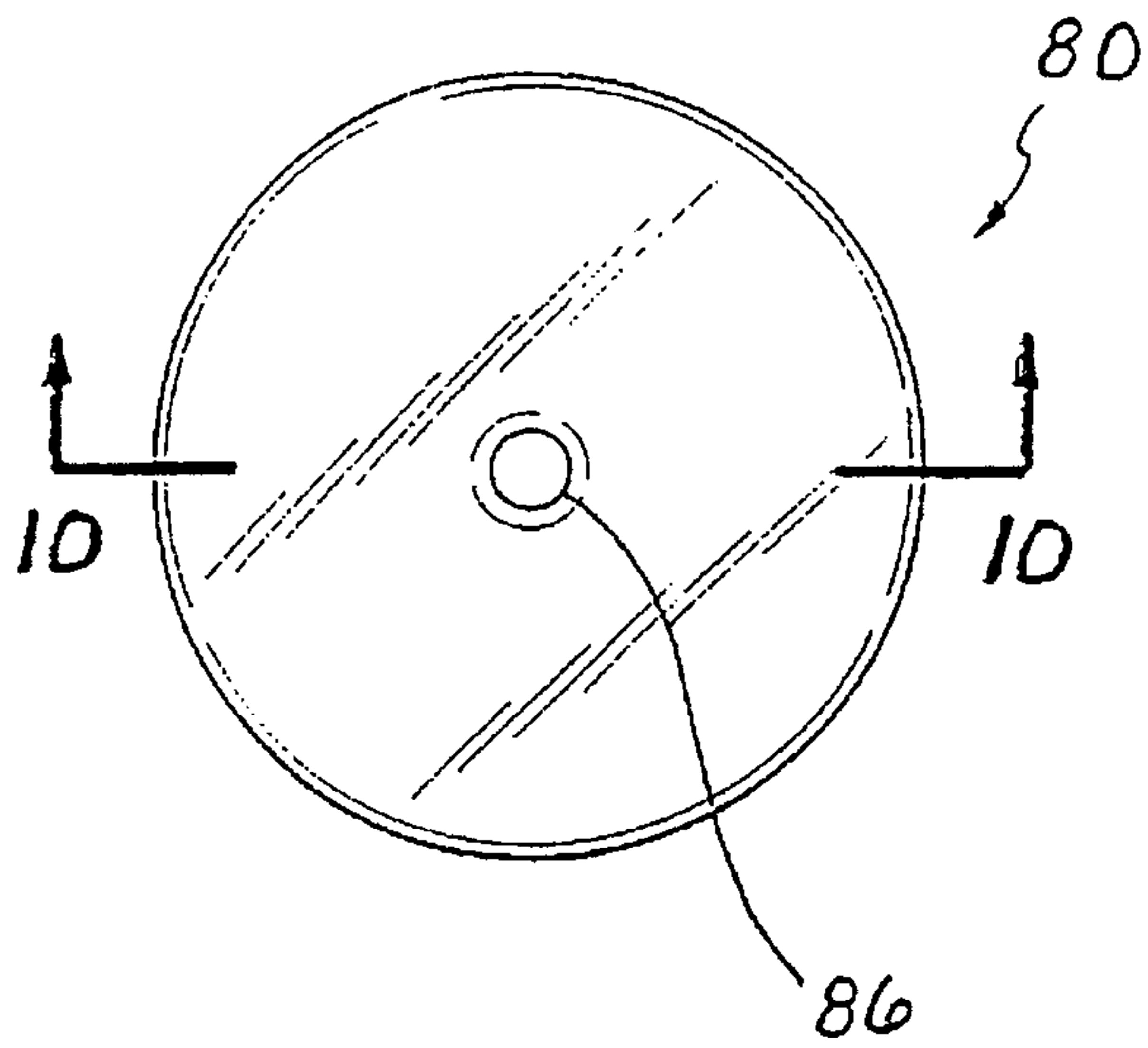


Fig. 9

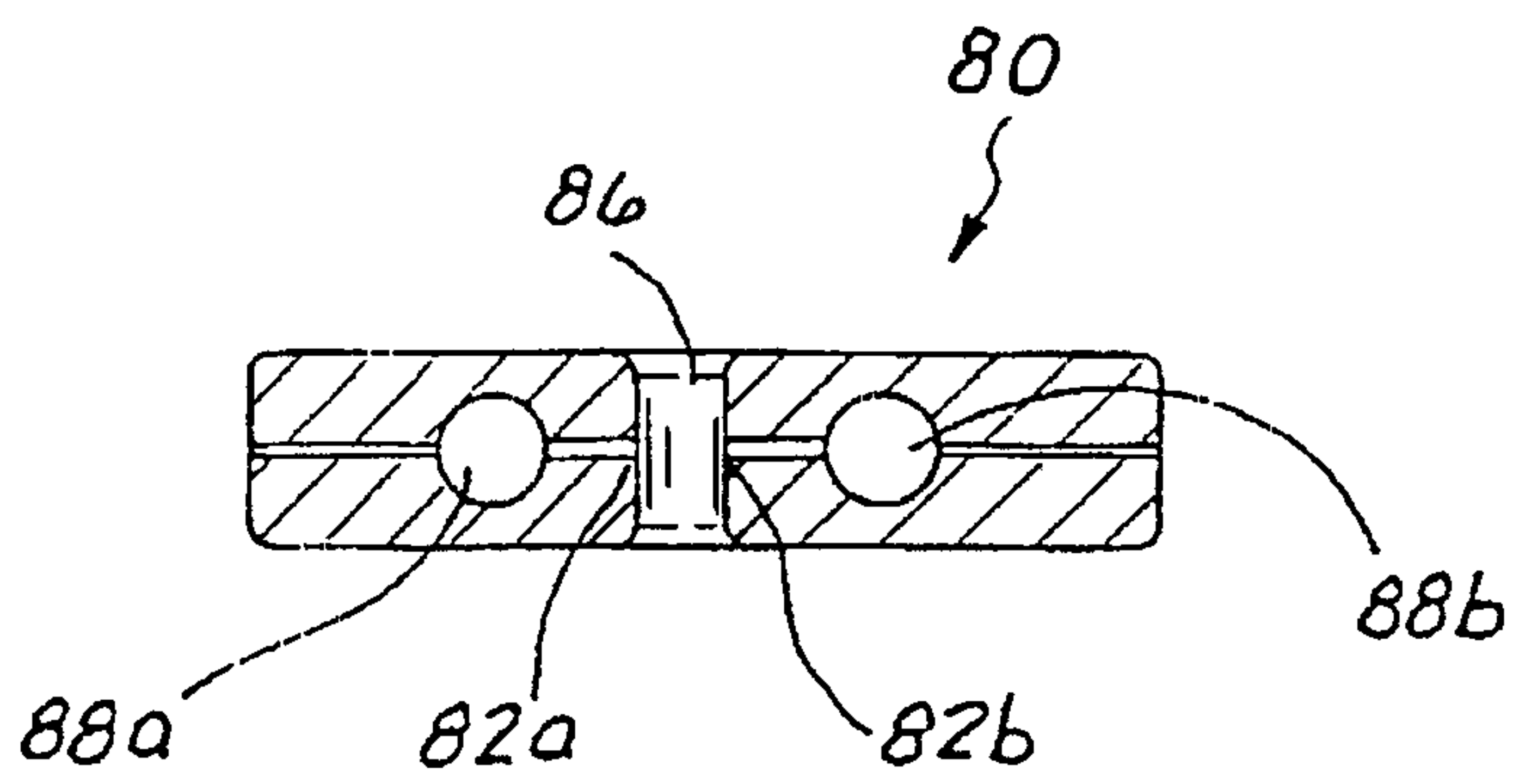


Fig. 10

