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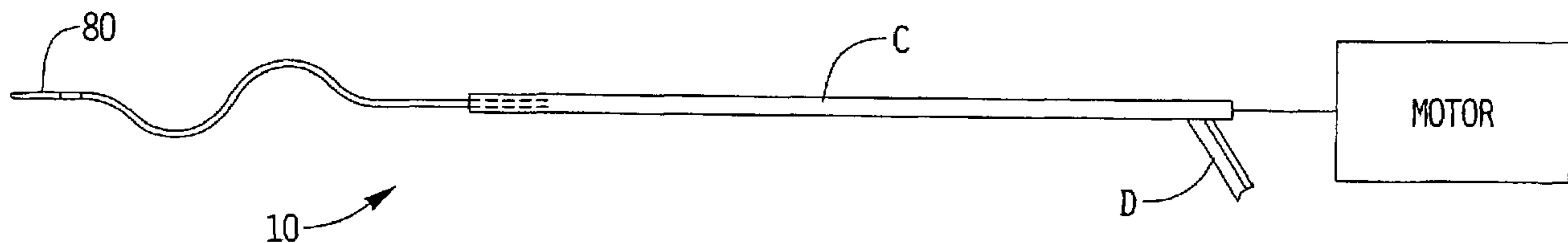
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(54) Title: ROTATIONAL THROMBECTOMY WIRE



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A rotatable thrombectomy wire for breaking up thrombus or other obstructive material comprising an inner core composed of a flexible material and an outer wire surrounding at least a portion of the inner core. The outer wire has a sinuous shaped portion at a distal region. The inner core limits the compressibility of the outer wire. The outer wire is operatively connectable at a proximal end to a motor for rotating the wire to macerate thrombus.

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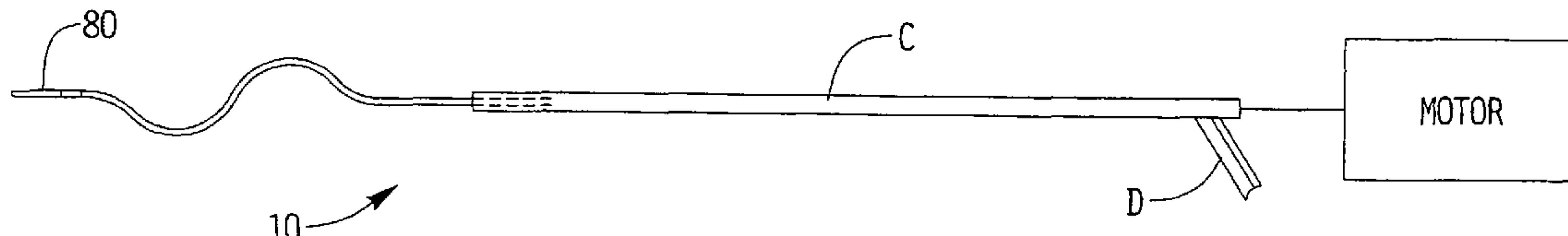
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(54) Title: ROTATIONAL THROMBECTOMY WIRE



(57) Abstract: A rotatable thrombectomy wire for breaking up thrombus or other obstructive material comprising an inner core composed of a flexible material and an outer wire surrounding at least a portion of the inner core. The outer wire has a sinuous shaped portion at a distal region. The inner core limits the compressibility of the outer wire. The outer wire is operatively connectable at a proximal end to a motor for rotating the wire to macerate thrombus.

ROTATIONAL THROMBECTOMY WIRE

BACKGROUND

This application claims priority from provisional application serial number 60/628,623, filed November 17, 2004, the entire contents of which are incorporated herein by reference.

Technical Field

This application relates to a rotational thrombectomy wire for clearing thrombus from native vessels.

Background of Related Art

In one method of hemodialysis, dialysis grafts, typically of PTFE, are implanted under the patient's skin, e.g. the patient's forearm, and sutured at one end to the vein for outflow and at the other end to the artery for inflow. The graft functions as a shunt creating high blood flow from the artery to the vein and enables access to the patient's blood without having to directly puncture the vein. (Repeated puncture of the vein could eventually damage the vein and cause blood clots, resulting in vein failure.) One needle is inserted into the graft to withdraw blood from the patient for transport to a dialysis machine (kidney machine); the other needle is inserted into the graft to return the filtered blood from the dialysis machine to the patient. In the dialysis machine, toxins and other waste products diffuse through a semi-permeable membrane into a dialysis fluid closely matching the chemical composition of the blood. The filtered blood, i.e. with the waste products removed, is then returned to the patient's body.

Over a period of time, thrombus or clots may form in the graft. Thrombus or clots may also form in the vessel. One approach to break up these clots and other obstructions in the graft and vessel is the injection of thrombolytic agents. The disadvantages of these agents are they are expensive, require lengthier hospital procedures and create risks of drug toxicity and bleeding complications as the clots are broken.

U.S. Patent No. 5,766,191 provides another approach to breaking up clots and obstructions via a mechanical thrombectomy device. The patent discloses a basket having six memory wires expandable to press against the inner lumen to conform to the

size and shape of the lumen. This device could be traumatic if used in the vessel, could denude endothelium, create vessel spasms and the basket and drive shaft could fracture.

U.S. Patent No. 6,090,118 discloses a mechanical thrombectomy device for breaking up clots. The single thrombectomy wire is rotated to create a standing wave to break-up or macerate thrombus. U.S. Patent Publication No. 2002/0173812 discloses another example of a rotational thrombectomy wire for breaking up clots. The thrombectomy wire has a sinuous shape at its distal end and is contained within a sheath in a substantially straight non-deployed position. When the sheath is retracted, the distal portion of the wire is exposed to enable the wire to return to its non-linear sinuous configuration. The wire is composed of stainless steel. Actuation of the motor causes rotational movement of the wire, creating a wave pattern, to macerate thrombus. The device of the '812 patent publication is effective in atraumatically and effectively breaking up blood clots in the graft and is currently being marketed by Datascope, Inc. as the Pro-Lumen* thrombectomy catheter. In the marketed device, the wire is a bifilar wire, composed of two stainless steel wires wound side by side with a metal tip and an elastomeric tip at the distalmost end.

Although the sinuous wire of the '812 publication is effective in proper clinical use to macerate thrombus in dialysis grafts, it is not suited for use in native vessels. The device is indicated for use in grafts, and if improperly used the wire can kink or knot, and perhaps even break. The wire can also bend, making it difficult to withdraw after use, and can lose its shape. Additionally, the wire would be abrasive to the vessel and the vessel could get caught in the interstices of the wire. It could also cause vessels spasms which can cause the vessel to squeeze down on the wire which could break the wire. Similar problems would occur with the use of the device of the '118 patent in native vessels.

The need therefore exists for a rotational thrombectomy wire which can be used to clear clots or other obstructions from the native vessels. Such wire could advantageously be used not only in native vessels adjacent dialysis grafts but for deep vein thrombosis and pulmonary embolisms.

SUMMARY

The present invention advantageously provides a rotational thrombectomy wire for breaking up thrombus or other obstructive material in a lumen of a native vessel.

The present invention provides a rotational thrombectomy wire comprising an inner core composed of a flexible material and a multifilar outer wire surrounding at least a portion of the inner core. The outer wire includes at least first and second metal wires wound side by side and having a sinuous shaped portion at a distal region. The inner core at a distal portion has a sinuous shaped portion within the sinuous portion of the outer wire. The inner core limits the compressibility of the multifilar wire. The multifilar wire is operatively connectable at a proximal end to a motor for rotating the wire to macerate thrombus within the vessel.

In a preferred embodiment, the inner core is composed of nylon material. In another embodiment, the inner core is composed of shape memory material wherein the inner core assumes its sinuous shape in the memorized configuration. In another embodiment, the core comprises at least two twisted wires of stainless steel.

The thrombectomy wire preferably further includes a polymeric material surrounding at least a distal portion of the multifilar wire. In a preferred embodiment, the polymeric material comprises a shrink wrap material attached to the multifilar wire. In another embodiment, the polymeric material is a coating over the multifilar wire.

The thrombectomy wire preferably comprises a flexible and blunt tip positioned at a distal end.

The inner core can have in one embodiment an enlarged distal end to form a connection portion and a metal tip secured to a distal end of the multifilar wire has a recess to receive the enlarged end of the inner core to frictionally engage the inner core.

In one embodiment, the first and second metal wires are wound together such that the coils of the first wire occupy the space between adjacent turns of the second wire and the coils of the multifilar outer wire have an inner diameter approximately equal to an outer diameter of the inner core.

The present invention also provides a rotatable thrombectomy wire for breaking up thrombus or other obstructive material in a lumen of a vessel comprising a multifilar outer wire including at least two metal wires wound side by side and operatively

connectable at a proximal end to a motor for rotating the wire to macerate thrombus. The multifilar wire has a sinuous shaped portion at a distal region. A polymeric material surrounds at least a region of the sinuous portion of the multifilar outer wire to block the interstices of the multifilar wire.

In a preferred embodiment, the polymeric material comprises a shrink wrap material. In another embodiment, the polymeric material is a coating over the bifilar wire.

The present invention also provides a thrombectomy apparatus for breaking up thrombus or other obstructive material comprising a handle, a sheath, a battery, a motor powered by the battery, and a sinuous thrombectomy wire having at least one wire wound to form a coil and an inner core composed of a material to limit the compressibility of the coil. The coil has a sinuous portion and surrounds at least a distal region of the inner core. The inner core has a sinuous portion within the sinuous portion of the coil. The sinuous portion of the inner core and first and second wires are movable from a straighter configuration within the sheath for delivery to a sinuous configuration when exposed from the sheath.

In a preferred embodiment, a polymeric material surrounds at least a distal portion of the coil to cover the interstices of the coil. In one embodiment, the core is composed of a shape memory material wherein the memorized position of the core has a sinuous configuration. In another embodiment, the core is composed of Nylon. In another embodiment, the core is composed of at least two twisted wires of stainless steel.

The present invention also provides a method for breaking up thrombus or other obstructive material in a native vessel comprising:

providing a thrombectomy wire having an inner core composed of a flexible material and at least one outer wire surrounding at least a portion of the inner core, the outer wire has a sinuous shaped portion at a distal region and the inner core has a sinuous shaped portion within the sinuous portion of the outer wire, and a polymeric material surrounding at least a distal portion of the at least one outer wire to block the interstices of the at least one outer wire;

delivering the wire to the lumen of the native vessel such that the sinuous shaped portions of the inner core and bifilar outer wire are in a more linear configuration within a sheath;

exposing the sinuous portion of the inner core and the at least one outer wire; and actuating a motor operatively connected to the thrombectomy wire so the sinuous portion of the at least one outer wire contacts the inner wall of the native vessel to macerate thrombus in the vessel.

BRIEF DESCRIPTION OF THE DRAWINGS

Preferred embodiment(s) of the present disclosure are described herein with reference to the drawings wherein:

Figure 1 is a side view in partial cross-section of a first embodiment of a thrombectomy wire of the present invention shown inside a catheter sleeve for delivery;

Figure 2 is a schematic view illustrating motorized rotation of the wire and a port for fluid delivery;

Figure 3 is a schematic side elevational view of the sinuous portion of the thrombectomy wire to depict a first embodiment of the inner core positioned therein;

Figure 4 is an enlarged cross-sectional view of the distalmost region of the rotational thrombectomy wire of Figure 3;

Figure 5 is schematic side elevational view of the sinuous portion of the thrombectomy wire to depict a second embodiment of the inner core positioned therein; and

Figure 6 is an enlarged side view of the distalmost region of the rotational wire of Figure 5;

Figure 7 is a schematic side elevational view of the sinuous portion of the thrombectomy wire to depict a third embodiment of the inner core positioned therein; and

Figure 8 is a cross-sectional view taken along line 8-8 of Figure 7.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

Referring now in detail to the drawings where like reference numerals identify similar or like components throughout the several views, Figures 3 and 4 illustrate a first embodiment of the thrombectomy wire of the present invention. The thrombectomy wire, designated generally by reference numeral 10, includes a core 20, a bifilar wire

(coil) 30, and shrink wrap 50. The bifilar wire 30 is formed by two stainless steel wires 32, 34, wound together. As shown they are wound side by side so the cross-sectional area or diameter "a" of the wire fills the space between adjacent turns of the other wire. For example, turns 32a and 32b are filled by respective turns 34a, 34b as shown. Preferably the bifilar wire 30 has a length of about 30 inches and a diameter of about .030 inches to about .040 inches and more preferably about .035 inches. When used in deeper native vessels, e.g. deep veins of the legs or pulmonary circuit, the wire 30 can have a length of about 52 inches. Other dimensions are also contemplated.

The distal region 16 of the bifilar wire 30 is formed into a sinuous or s-shape to contact the vessel wall as the wire rotates.

Although in the preferred illustrated and described embodiments, the outer wire is a multifilar wire in the form of a bifilar wire (two wires), a different number of wires could be wound to form the outer wire component of the thrombectomy wire of the present invention. In yet another embodiment the outer wire can comprise a single wound wire.

The bifilar wire 30 is preferably cold formed into an over-formed s-shape. The bifilar wire is heated, for example at about 670 degrees Fahrenheit, which removes residual stresses and changes the shape of the "s" so it warps back to its desired shape. This stress relief process makes the wire more dimensionally stable.

A tip 80, preferably composed of rubber, Pebax, or other elastomeric materials, is mounted at the distalmost tip of the wire 10 to provide the wire 10 with an atraumatic distal tip to prevent damage to the vessel wall during manipulation and rotation of the wire. A metal tip 60 is attached by laser welding or other methods to the distal end of the bifilar wire 30. The metal tip 60 has an enlarged dumbbell shaped head 62 to facilitate attachment to tip 80. The flexible tip 80 is attached by injection molding over the machined tip. Other attachment methods are also contemplated.

With continued reference to Figure 4, a core 20 is positioned within the bifilar wire 30 and preferably has an outer diameter E substantially equal to the inner diameter D of the coil. The core at a distal portion has a sinuous shaped portion within the sinuous shaped portion of the outer wire 30, corresponding to and formed by the sinuous shape of outer wire 30. In one embodiment, the core extends the entire length of the bifilar wire

30 and this is shown in the schematic drawing of Figure 3. The core 20 can alternatively have a length of about 4-5 inches so it extends through the distal linear portion and sinuous portion of the wire 30. That is, in such embodiment, the core extends through the portion of the wire that is exposed from the sheath and used to macerate thrombus. It is also contemplated that the core can extend within a shorter or longer length of the bifilar wire.

The core 20 is composed of a flexible material which will limit the compressibility of the wire 30 during use. The core in the embodiment of Figure 3 is composed of Nylon, and preferably a drawn Nylon monofilament. Other possible materials include, for example, Teflon, polypropylene, PET, and fluorocarbon. The Nylon provides a non-compressible material to limit the compressibility of the wire 30 during use. That is, as noted above, the Nylon core preferably has a diameter E to fill the inside of the coil 30, e.g. a diameter of about .008 inches to about .013 inches, and preferably about .012 inches. (Other dimensions are also contemplated.) This enables the coil (bifilar wire) 30 to compress only to that diameter. By limiting compressibility it strengthens the wire as it reduces its degree of elongation if it is under torque. It also prevents bending or knotting of the wire which could otherwise occur in native vessels. It increases the torsional strength of the wire and also strengthens the wire to accommodate spasms occurring in the vessel. An enlarged distal head, such as ball tip (not shown), can be provided on the core 20 to fit in a recess of machined tip 60. As an alternative, core 20 can be attached by adhesive at the tip, welded, crimped, soldered or can alternatively be free floating.

The shrink wrap material 50 covers a portion of the bifilar wire 30 proximal of the flexible tip 80 to block the interstices of the coil and provide a less abrasive surface. As shown in Figure 4, the distal end of the shrink wrap abuts the proximal end of the tip 60. The shrink wrap can be made of PET, Teflon, Pebax, polyurethane or other polymeric materials. The material extends over the exposed portion of the wire 30 (preferably for about 3 inches to about 4 inches) and helps to prevent the native vessel from being caught in the coil and reduces vessel spasms. Alternatively, instead of shrink wrap, a coating can be applied to the coil formed by the bifilar wire to cover the interstices

Figures 5 and 6 illustrate an alternate embodiment of the thrombectomy wire of the present invention, designated generally by reference numeral 100. Wire 100 is identical to wire 10 of Figure 1, except for the inner core 120. It is identical in that it has a bifilar wire 130, a shrink wrap 170, an elastomeric tip 180 and metal, e.g. stainless steel, tip 160.

In this embodiment, the core 120 is composed of a shape memory material, preferably Nitinol (a nickel titanium alloy), which has a memorized configuration of a sinuous or s-shape substantially corresponding to the s-shape of the bifilar wire 130. In the softer martensitic state within the sheath, core 120 is in a substantially linear configuration. This state is used for delivering the wire to the surgical site. When the wire is exposed to warmer body temperature, the core 120 transforms to its austenitic state, assuming the s-shaped memorized configuration. Cold saline is delivered through the catheter during delivery to maintain the core 120 in this martensitic state; the warming occurs by exposure to body temperature to transform the core 120 to the memorized state. Such memorized s-shape helps maintain the s-shape of the bifilar wire 130 during use. Cold saline can also be delivered to the core 120 at the end of the procedure to facilitate withdrawal.

The Nitinol core 120, like the Nylon core 20, is not compressible so it will also limit the compressibility of the bifilar wire 130. The Nitinol core 120 also will increase the stiffness of the wire 100, thereby reducing the chance of knotting and kinking and increase the strength of the wire to accommodate any spasms in the vessel. Its shape memory helps hold the amplitude of the bifilar wire 130 during use to maintain its force against the clot for maceration upon rotation. It preferably extends about 4-5 inches so it extends through the distal linear portion and sinuous portion of the wire 130, terminating at end 122. Alternately it can extend a shorter or longer length within the wire 130, or even the entire length as shown in the schematic view of Figure 5. It preferably has an outer diameter of about .008 inches to about .013 inches, and more preferably about .012 inches, corresponding to the inner diameter of the coil. Other dimensions are also contemplated.

In another embodiment, a stainless steel braid, cable, or strand of wires twisted together provides the inner core member to limit compressibility of the coil (bifilar wire)

and provide increased stiffness, strength and other advantages of the core enumerated above. This is shown in the embodiment of Figures 7 and 8 where wire 200 has inner core 220 of seven twisted stainless steel wires. A different number of twisted wires is also contemplated. The other elements of the wire 200, e.g., outer bifilar wire 230, metal tip 260, tip 280 shrink wrap 250, etc., are the same as in wires 10 and 100 described herein.

The rotational thrombectomy wires 10, 100 and 200 of the present invention can be used with various thrombectomy catheters to macerate thrombus within the vessel. The rotational thrombectomy wire 10 (or wire 100 or 200) is contained within a flexible sheath or sleeve C of a catheter as shown in Figure 1. Relative movement of the wire and sheath C will enable the wire 10 to be exposed to assume the curved (sinuous) configuration described below to enable removal of obstructions, such as blood clots, from the lumen of the vessel.

A motor powered by a battery is contained within a housing to macerate and liquefy the thrombus into small particles within the vessel lumen. This is shown schematically in Figure 2. Wire 10 (or 100 or 200) is operatively connected to the motor. Operative connection encompasses direct connection or connection via interposing components to enable rotation when the motor is actuated. The curved regions of the wire 10 or (100 or 200) are compressed so the wire (including the distal region 16, 116 or 216, respectively) is in a substantially straight or linear non-deployed configuration when in the sheath C. This covering of the wire 10 (or 100 or 200) by sheath C facilitates insertion through an introducer sheath and manipulation within the vessel. When the flexible sheath C is retracted, the wire is exposed to enable the wire to return to its non-linear substantially sinuous configuration for rotation about its longitudinal axis within the lumen of the vessel.

Fluids, such as imaging dye can be injected through the port D into the lumen of the sheath C in the space between wire 10 (or 100 or 200) and the inner wall of the sheath C, and exiting the distal opening to flow into the vessel. This imaging dye provides an indication that fluid flow has resumed in the vessel. The lumen of the sheath can also receive cold saline to cool the Nitinol core 120 as described above.

The rotational thrombectomy wires 10, 100 and 200 of the present invention can also be used with the thrombectomy catheters having one or more balloons such as the balloon described in the '812 publication. The wires 10, 100 and 200 can further be used with other thrombectomy catheters.

While the above description contains many specifics, those specifics should not be construed as limitations on the scope of the disclosure, but merely as exemplifications of preferred embodiments thereof. Those skilled in the art will envision many other possible variations that are within the scope and spirit of the disclosure as defined by the claims appended hereto.

WHAT IS CLAIMED IS:

1. A rotatable thrombectomy wire for breaking up thrombus or other obstructive material, the wire comprising:
 - an inner core composed of a flexible material; and
 - a multifilar outer wire surrounding at least a portion of the inner core, the multifilar outer wire including at least first and second metal wires wound side by side and having a sinuous shaped portion at a distal region, the inner core having a sinuous shaped portion within the sinuous shaped portion of the multifilar outer wire, the inner core limiting the compressibility of the multifilar wire, the multifilar wire operatively connectable at a proximal end to a motor for rotating the wire to macerate thrombus.
2. The thrombectomy wire of claim 1, wherein the inner core is composed of nylon material.
3. The thrombectomy wire of claim 1, wherein the inner core is composed of shape memory material, the inner core having a memorized configuration, the inner core assuming its sinuous shape in the memorized configuration.
4. The thrombectomy wire of claim 1, wherein the inner core is composed of at least two twisted wires of stainless steel.
5. The thrombectomy wire of claim 1, further comprising a polymeric material surrounding at least a distal portion of the multifilar wire.
6. The thrombectomy wire of claim 5, wherein the polymeric material is a coating over the multifilar wire.
7. The thrombectomy wire of claim 5, wherein the polymeric material comprises a shrink wrap material attached to the multifilar wire.

8. The thrombectomy wire of claim 5, wherein the inner core is composed of nylon material.
9. The thrombectomy wire of claim 5, wherein the inner core is composed of at least two twisted wires of stainless steel.
10. The thrombectomy wire of claim 1, wherein the first and second metal wires are wound together such that the coils of the first wire occupy the space between adjacent turns of the second wire.
11. The thrombectomy wire of claim 5, wherein the first and second wires are wound together such that the coils of the first wire occupy the space between adjacent turns of the second wire, and further including a flexible and blunt tip positioned at a distal end of the wire.
12. The thrombectomy apparatus of claim 1, wherein the outer wire forms coils with essentially no spaces between adjacent coils and the coils of the outer wire have an inner diameter approximately equal to an outer diameter of the inner core.
13. A rotatable thrombectomy wire for breaking up thrombus or other obstructive material in a lumen of a vessel, the wire comprising:
 - a multifilar outer wire including at least first and second metal wires wound side by side and operatively connectable at a proximal end to a motor for rotating the wire to macerate thrombus, the multifilar wire having a sinuous shaped portion at a distal region; and
 - a polymeric material surrounding at least a distal region of the sinuous of the first and second wires to block the interstices of the multifilar wire.
14. The thrombectomy wire of claim 14, wherein the polymeric material comprises a shrink wrap material attached to the multifilar wire.

15. A thrombectomy apparatus for breaking up thrombus or other obstructive material comprising a handle, a sheath, a battery, a motor powered by the battery, and a sinuous thrombectomy wire having at least one wire wound to form a coil and an inner core composed of a material to limit the compressibility of the coil, the coil having a sinuous portion and surrounding at least a distal region of the inner core, the inner core having a sinuous portion within the sinuous portion of the coil, the sinuous portion of the inner core and the coil movable from a straighter configuration within the sheath for delivery to a sinuous configuration when exposed from the sheath.

16. The thrombectomy apparatus of claim 15, further comprising a polymeric material surrounding at least a distal portion of the coil to cover the interstices of the coil.

17. The thrombectomy apparatus of claim 16, wherein the core is composed of nylon material.

18. The thrombectomy apparatus of claim 16, wherein the core is composed of at least two twisted wires of stainless steel.

19. A method for breaking up thrombus or other obstructive material in a native vessel comprising:

providing a thrombectomy wire having an inner core composed of a flexible material, at least one outer wire surrounding at least a portion of the inner core wherein the outer wire has a sinuous shaped portion at a distal region and the inner core has a sinuous portion within the sinuous portion of the outer wire, and a polymeric material surrounding at least a distal portion of the at least one outer wire to block the interstices of the at least one outer wire;

delivering the wire to the lumen of the native vessel such that the sinuous shaped portions of the inner core and outer wire are in a more linear configuration within a sheath;

exposing the sinuous portion of the inner core and the at least one outer wire; and

actuating a motor operatively connected to the thrombectomy wire so the sinuous portion of the at least one outer wire contacts the inner wall of the native vessel to macerate thrombus in the vessel.

20. The method of claim 19, further comprising the step of retracting the thrombectomy wire during rotation.

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FIG. 1

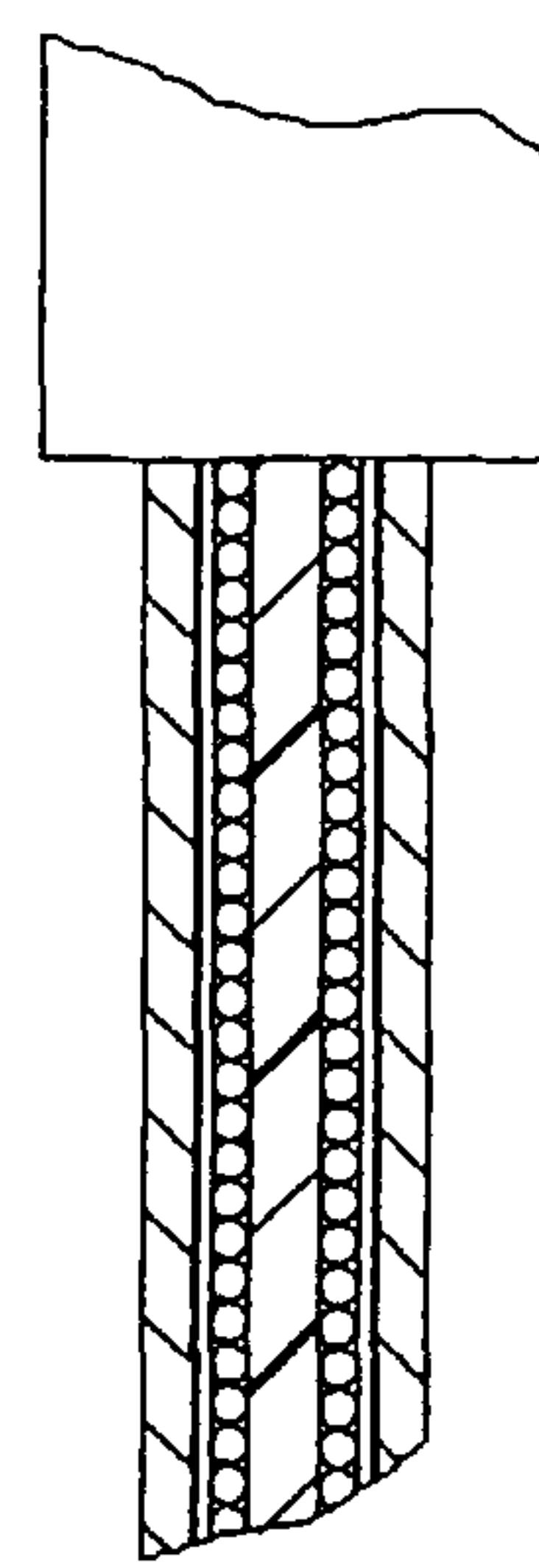
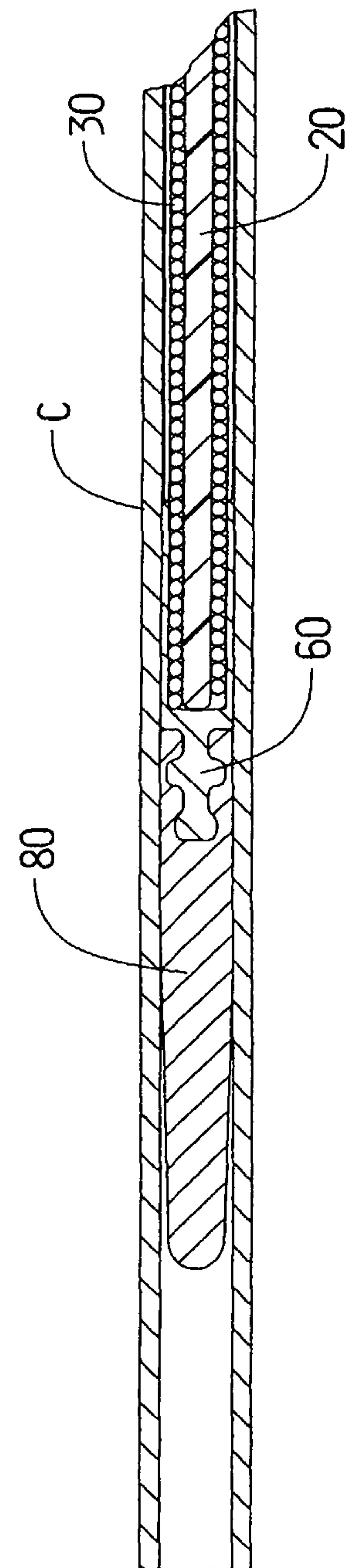
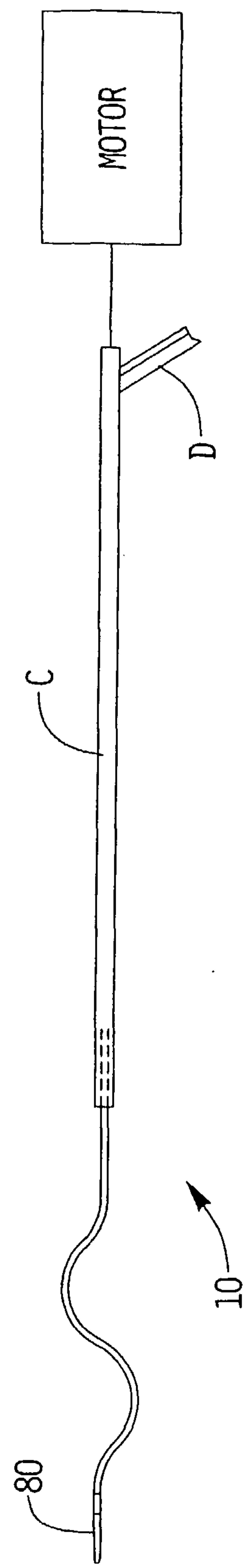
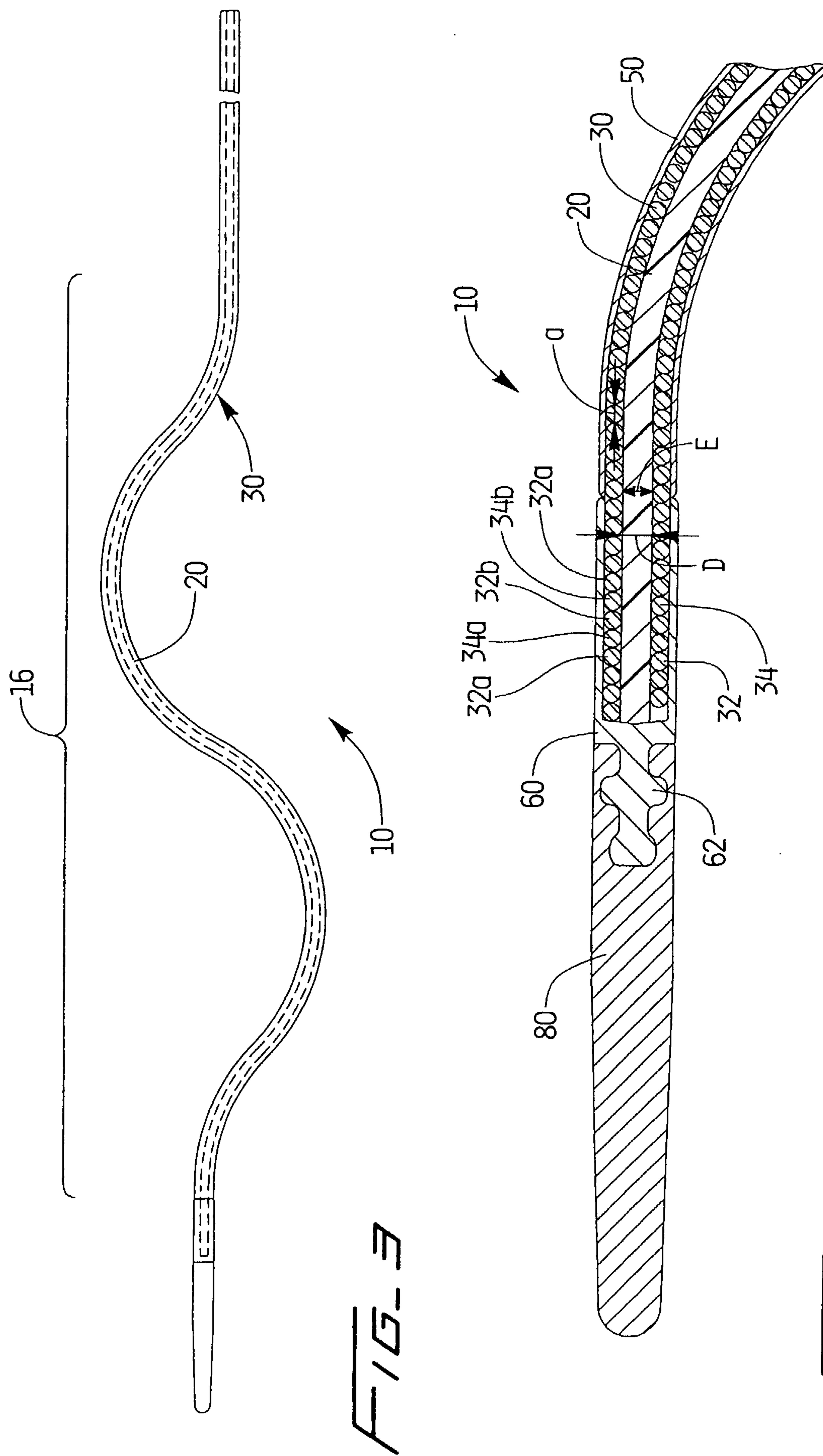


FIG. 2



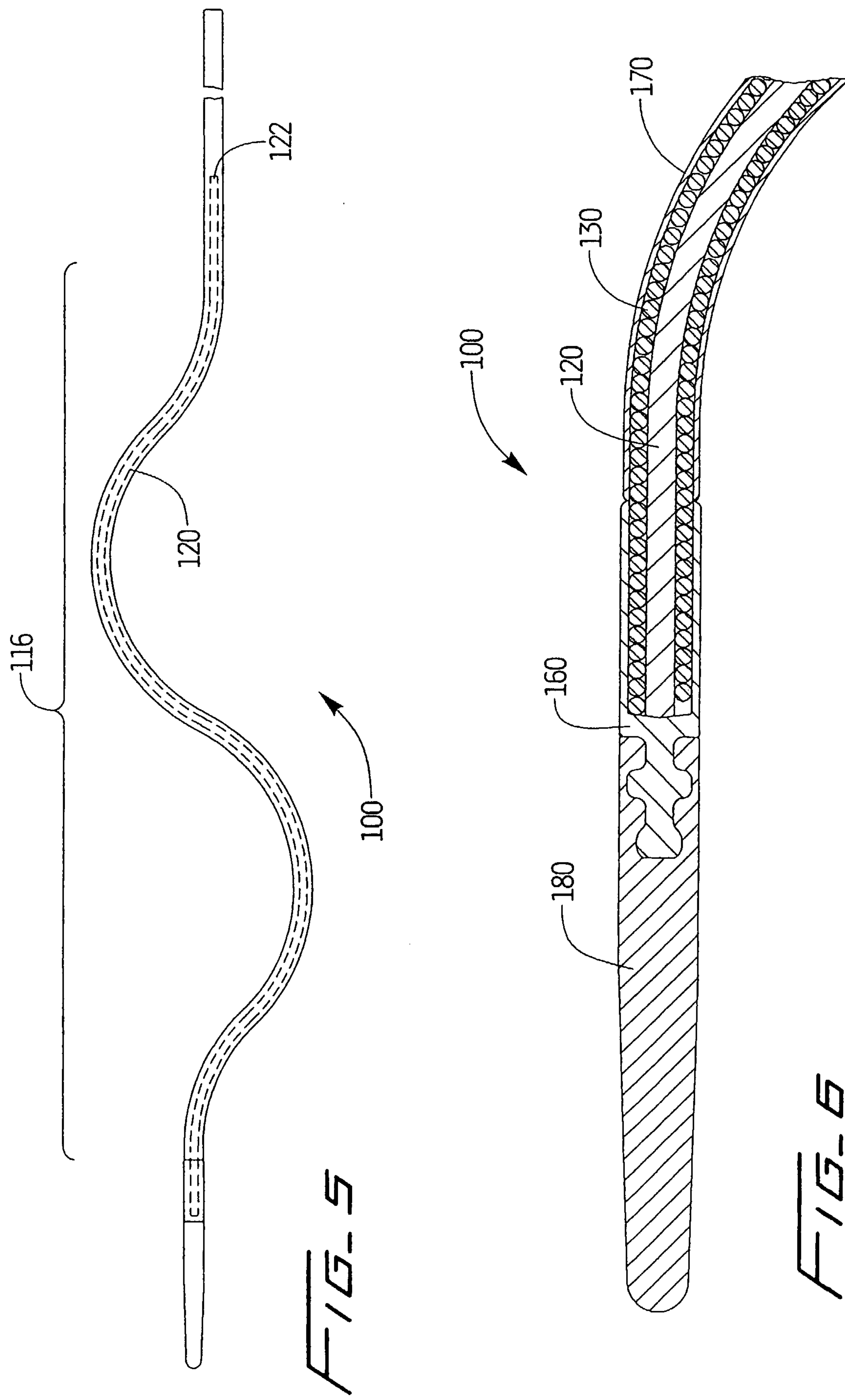
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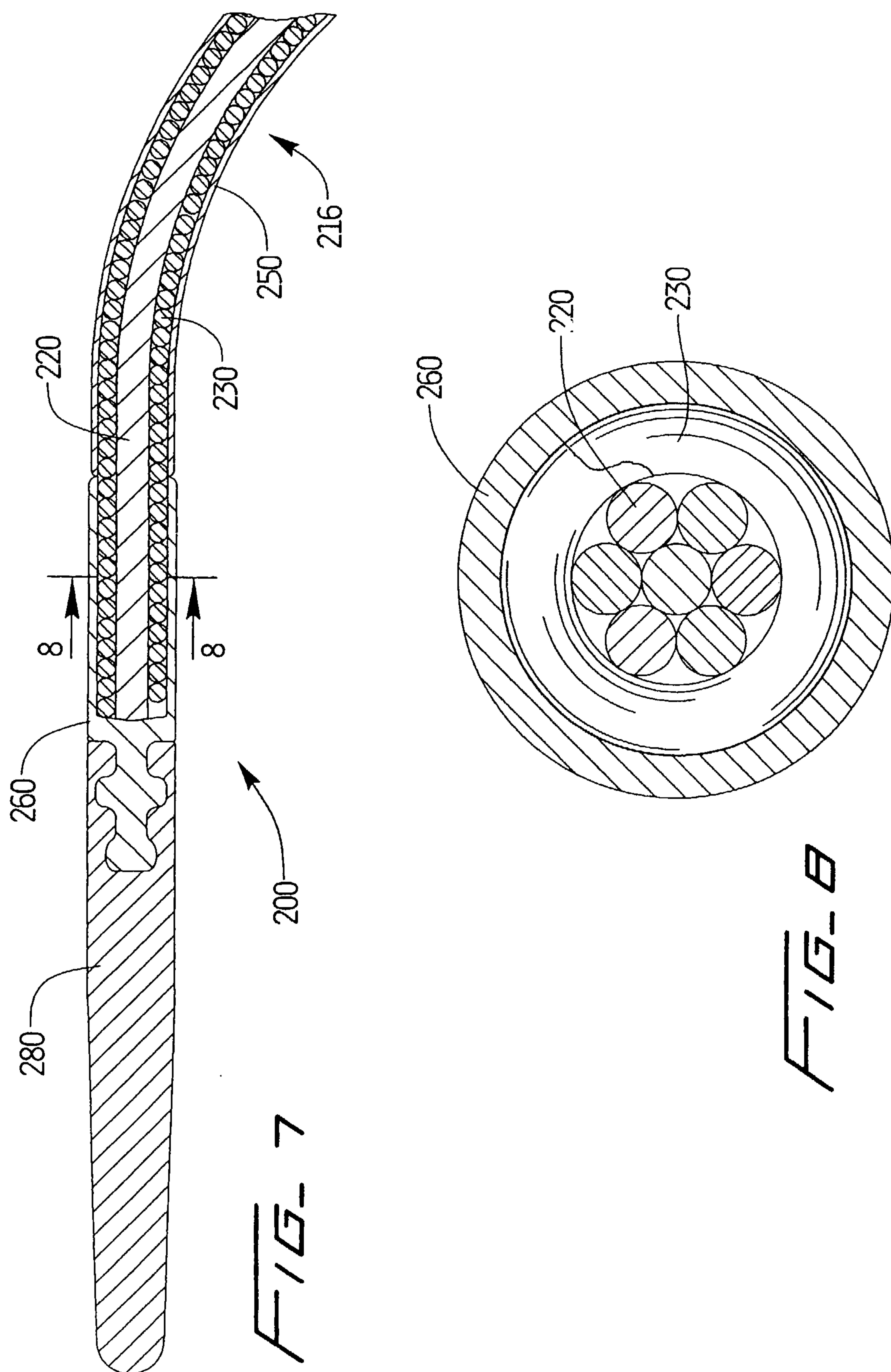
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